EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

11 July 2013
EMA/402820/2013
Patient Health Protection

Initial Notices for Parallel Distribution –2012

Please note that the highlighted products are Veterinary medicines

Product name	Parallel distributor	Date of notification	Strength	Pharmaceutical dosage form	Pack size	EU-number	Member State(s) of Origin	Member State(s) of Destination	Date of Notice letter
Abilify	2 Care 4	21/03/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/04/2012
Abilify	2 Care 4	21/03/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012
Abilify	2 Care 4	31/07/2012	30 mg	Tablet	28 x 1	EU/1/04/276/017	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	31/08/2012
Abilify	2 Care 4	31/07/2012	30 mg	Tablet	56 x 1	EU/1/04/276/019	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	31/08/2012

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Abilify	ABACUS MEDICINE A/S	17/05/2012	15 mg	Tablet	56 x 1	EU/1/04/276/014	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/06/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	10 mg	Tablet	14 x 1	EU/1/04/276/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	10 mg	Tablet	28 x 1	EU/1/04/276/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	10 mg	Tablet	49 x 1	EU/1/04/276/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	15 mg	Tablet	14 x 1	EU/1/04/276/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012

Abilify	ABACUS MEDICINE A/S	31/08/2012	15 mg	Tablet	28 x 1	EU/1/04/276/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	15 mg	Tablet	49 x 1	EU/1/04/276/013	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	15 mg	Tablet	98 x 1	EU/1/04/276/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	30 mg	Tablet	14 x 1	EU/1/04/276/016	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	30 mg	Tablet	28 x 1	EU/1/04/276/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	30 mg	Tablet	49 x 1	EU/1/04/276/018	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012

Abilify	ABACUS MEDICINE A/S	31/08/2012	30 mg	Tablet	98 x 1	EU/1/04/276/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	5 mg	Tablet	49 x 1	EU/1/04/276/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	31/08/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Abilify	ABACUS MEDICINE A/S	12/11/2012	15 mg	Orodispersible tablet	28 x 1	EU/1/04/276/028	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/11/2012

Abilify	AMIMED DIRECT LTD	25/04/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	26/04/2012
Abilify	axicorp Pharma B.V.	25/07/2012	10 mg	Tablet	56 x 1	EU/1/04/276/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/08/2012
Abilify	axicorp Pharma B.V.	25/07/2012	15 mg	Tablet	56 x 1	EU/1/04/276/014	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/08/2012
Abilify	axicorp Pharma B.V.	25/07/2012	30 mg	Tablet	56 x 1	EU/1/04/276/019	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/08/2012
Abilify	axicorp Pharma B.V.	25/07/2012	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/08/2012
Abilify	Axicorp Pharma GmbH	24/05/2012	10 mg	Orodispersible tablet	49 x 1	EU/1/04/276/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Abilify	Axicorp Pharma GmbH	24/05/2012	15 mg	Orodispersible tablet	49 x 1	EU/1/04/276/029	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Abilify	Axicorp Pharma GmbH	25/09/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/10/2012
Abilify	BB Farma s.r.l.	16/11/2012	30 mg	Tablet	98 x 1	EU/1/04/276/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012

Abilify	BB Farma s.r.l.	16/11/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012
Abilify	BB Farma s.r.l.	16/11/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012
Abilify	BB Farma s.r.l.	16/11/2012	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012
Abilify	BB Farma s.r.l.	16/11/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012
Abilify	Beragena Arzneimittel GmbH	29/11/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	11/12/2012
Abilify	Cambridge Major Laboratories Europe B.V	30/03/2012	5 mg	Tablet	49 x 1	EU/1/04/276/003	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012
Abilify	Cambridge Major Laboratories Europe B.V	30/03/2012	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012
Abilify	Cambridge Major Laboratories Europe B.V	30/03/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012

Abilify	Cambridge Major Laboratories Europe B.V	30/03/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012
Abilify	CD Pharma Limited	23/01/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Abilify	CD Pharma Limited	24/01/2012	10 mg	Tablet	49 x 1	EU/1/04/276/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Abilify	CD Pharma Limited	24/01/2012	5 mg	Tablet	49 x 1	EU/1/04/276/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Abilify	CD Pharma Limited	24/01/2012	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Abilify	CD Pharma Limited	26/01/2012	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Abilify	CD Pharma Limited	27/01/2012	30 mg	Tablet	49 x 1	EU/1/04/276/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/02/2012
Abilify	CD Pharma Limited	27/01/2012	30 mg	Tablet	98 x 1	EU/1/04/276/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/02/2012
Abilify	CD Pharma Limited	30/01/2012	15 mg	Tablet	49 x 1	EU/1/04/276/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/02/2012

Abilify	CD Pharma Limited	30/01/2012	15 mg	Tablet	98 x 1	EU/1/04/276/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/02/2012
Abilify	COPHARMA ApS	18/01/2012	10 mg	Orodispersible tablet	28 x 1	EU/1/04/276/025	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/02/2012
Abilify	COPHARMA ApS	18/01/2012	15 mg	Orodispersible tablet	28 x 1	EU/1/04/276/028	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/02/2012
Abilify	Cross Pharma AB	07/06/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Abilify	Cross Pharma AB	07/06/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Abilify	Cross Pharma AB	07/06/2012	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Abilify	Docpharm GmbH&CoKGaA	03/04/2012	30 mg	Tablet	28 x 1	EU/1/04/276/017	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/04/2012
Abilify	Docpharm GmbH&CoKGaA	04/04/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/05/2012
Abilify	Docpharm GmbH&CoKGaA	04/04/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/05/2012
Abilify	Docpharm GmbH&CoKGaA	08/05/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/05/2012

Abilify	Emra-Med	13/11/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/11/2012
Abilify	HAEMATO PHARM AG	23/10/2012	15 mg	Tablet	28 x 1	EU/1/04/276/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/10/2012
Abilify	Interport Ltd	30/10/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands	United Kingdom	06/11/2012
Abilify	kohlpharma GmbH	11/06/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/06/2012
Abilify	Kosei Pharma UK Limited	09/01/2012	10 mg	Tablet	28 x 1	EU/1/04/276/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	16/02/2012
Abilify	Medartuum AB	09/02/2012	10 mg	Orodispersible tablet	28 x 1	EU/1/04/276/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	05/03/2012
Abilify	O.P.D. Laboratories Ltd	18/09/2009	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	04/01/2012
Abilify	Orifarm AB	25/01/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/02/2012
Abilify	Paranova Danmark AS	26/04/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/05/2012

Abilify	Paranova Läkemedel AB	16/12/2011	15 mg	Orodispersible tablet	28 x 1	EU/1/04/276/028	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/01/2012
Abilify	Paranova Läkemedel AB	30/01/2012	30 mg	Tablet	56 x 1	EU/1/04/276/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	01/02/2012
Abilify	Pharma Westen GmbH	04/10/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/10/2012
Abilify	PHARMACHIM AB	24/10/2012	1 mg/ml	Oral solution	1 bottle + 1 cup + 1 calibrated dropper	EU/1/04/276/034	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Abilify	Profind Wholesale Ltd.	03/07/2012	10 mg	Tablet	28 x 1	EU/1/04/276/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/08/2012
Abilify	Profind Wholesale Ltd.	03/07/2012	15 mg	Tablet	28 x 1	EU/1/04/276/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/08/2012
Abilify	REMEDIX GMBH	16/11/2012	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/11/2012
Abilify	REMEDIX GMBH	16/11/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/11/2012

Abilify	s.c. HC PHARMA STORE s.r.l.	09/02/2012	15 mg	Tablet	98 x 1	EU/1/04/276/015	Romania, Spain	Germany	02/03/2012
Abilify	s.c. HC PHARMA STORE s.r.l.	09/02/2012	15 mg	Tablet	49 x 1	EU/1/04/276/013	Romania, Spain	Germany	02/03/2012
Abilify	Veron Pharma Vertriebs GmbH	02/03/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/03/2012
Abilify	Veron Pharma Vertriebs GmbH	02/03/2012	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/03/2012
Abilify	Veron Pharma Vertriebs GmbH	02/03/2012	5 mg	Tablet	98 x 1	EU/1/04/276/005	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/03/2012
Abilify	Veron Pharma Vertriebs GmbH	03/04/2012	30 mg	Tablet	14 x 1	EU/1/04/276/016	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	17/04/2012
Abilify	Veron Pharma Vertriebs GmbH	03/04/2012	30 mg	Tablet	28 x 1	EU/1/04/276/017	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	17/04/2012
Abilify	Veron Pharma Vertriebs GmbH	03/04/2012	30 mg	Tablet	98 x 1	EU/1/04/276/020	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	17/04/2012
Abilify	Veron Pharma Vertriebs GmbH	09/11/2012	15 mg	Orodispersible tablet	49 x 1	EU/1/04/276/029	Austria, Belgium, Finland, France, Greece, Italy, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Abilify	WAYMADE PLC	22/06/2012	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/07/2012
Aclasta	Cross Pharma AB	29/05/2012	5 mg/100 ml	Solution for infusion	1 bottle	EU/1/05/308/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	01/06/2012
Actos	Doncaster Pharmaceuticals Group Ltd	20/09/2012	45 mg	Tablet	28	EU/1/00/150/012	United Kingdom	Ireland, Malta	27/09/2012
Actos	kohlpharma GmbH	14/11/2012	15 mg	Tablet	112	EU/1/00/150/025	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012

Actos	kohlpharma GmbH	14/11/2012	45 mg	Tablet	112	EU/1/00/150/029	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Actrapid	BB Farma s.r.l.	07/06/2012	100 IU/ml	Solution for injection	5 cartridges	EU/1/02/230/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Actrapid	BB Farma s.r.l.	07/06/2012	100 IU/ml	Solution for injection	10 cartridges	EU/1/02/230/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Actrapid	Eureco-Pharma B.V.	14/05/2012	100 IU/ml	Solution for injection	10 cartridges	EU/1/02/230/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	29/05/2012
Adrovance	Euro Registratie Collectief BV	28/12/2011	70 mg / 5600 IU	Tablet	4	EU/1/06/364/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	01/02/2012
Advagraf	CC Pharma	20/09/2012	5 mg	Prolonged-release capsule, hard	100 x 1	EU/1/07/387/026	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
Advagraf	CC Pharma	11/10/2012	5 mg	Prolonged-release capsule, hard	50 x 1	EU/1/07/387/025	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/10/2012
Advagraf	CC Pharma	27/11/2012	3 mg	Prolonged-release capsule, hard	50 x 1	EU/1/07/387/022	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/12/2012

Advagraf	Cross Pharma AB	31/01/2012	0.5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/04/2012
Advagraf	Cross Pharma AB	31/01/2012	1 mg	Prolonged-release capsule, hard	50	EU/1/07/387/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/04/2012
Advagraf	Cross Pharma AB	31/01/2012	5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/04/2012
Advagraf	Cross Pharma AB	12/06/2012	3 mg	Prolonged-release capsule, hard	50	EU/1/07/387/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/07/2012
Advagraf	Dr Fisher Farma BV	23/02/2012	5 mg	Prolonged-release capsule, hard	100 x 1	EU/1/07/387/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/03/2012
Advagraf	Emra-Med	06/12/2012	3 mg	Prolonged-release capsule, hard	50	EU/1/07/387/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Advagraf	Emra-Med	06/12/2012	3 mg	Prolonged-release capsule, hard	100	EU/1/07/387/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012

Advagraf	Euro Registratie Collectief BV	03/10/2012	5 mg	Prolonged-release capsule, hard	50 x 1	EU/1/07/387/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/10/2012
Advagraf	Medartuum AB	19/12/2011	3 mg	Prolonged-release capsule, hard	50	EU/1/07/387/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	19/01/2012
Advagraf	Medartuum AB	19/04/2012	1 mg	Prolonged-release capsule, hard	60 x 1	EU/1/07/387/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/05/2012
Advagraf	Medcor Pharma-ceuticals BV	03/04/2012	3 mg	Prolonged-release capsule, hard	100 x 1	EU/1/07/387/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	11/05/2012
Advagraf	Medcor Pharma-ceuticals BV	01/05/2012	0.5 mg	Prolonged-release capsule, hard	30	EU/1/07/387/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/11/2012
Advagraf	Medcor Pharma-ceuticals BV	01/05/2012	1 mg	Prolonged-release capsule, hard	30	EU/1/07/387/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/11/2012
Advagraf	Orifarm AB	13/06/2012	5 mg	Prolonged-release capsule, hard	50 x 1	EU/1/07/387/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	02/07/2012
Advagraf	Orifarm AS	06/09/2012	3 mg	Prolonged-release capsule, hard	30	EU/1/07/387/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/10/2012

Advagraf	Orifarm AS	06/09/2012	5 mg	Prolonged-release capsule, hard	30	EU/1/07/387/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/10/2012
Advagraf	Paranova Läkemedel AB	24/05/2012	0.5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	28/06/2012
Advagraf	Paranova Oy	28/02/2012	5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	26/03/2012
Advagraf	Paranova Oy	25/06/2012	1 mg	Prolonged-release capsule, hard	50	EU/1/07/387/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	27/07/2012
Advagraf	Pharma Westen GmbH	13/01/2012	1 mg	Prolonged-release capsule, hard	100 x 1	EU/1/07/387/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Advagraf	REMEDIX GMBH	30/04/2012	0.5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	13/06/2012
Advagraf	REMEDIX GMBH	30/04/2012	5 mg	Prolonged-release capsule, hard	50	EU/1/07/387/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	13/06/2012
Advocate Spot-on solution for Extra-Large Dogs	CC Pharma	21/05/2012	10 % imidacloprid & 2.5 % moxidectrin	Spot-on Solution	4 ml 3 pipettes	EU/2/03/039/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Lithuania, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012

Advocate Spot-on solution for Large Dogs	CC Pharma	21/05/2012	10 % imidacloprid & 2.5 % moxidectin	Spot-on solution	2.5 ml- 3 pipettes	EU/2/03/039/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Advocate Spot-on solution for Medium Dogs	CC Pharma	21/05/2012	10 % imidacloprid & 2.5 % moxidectin	Spot-on solution	1 ml- 3 pipettes	EU/2/03/039/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Advocate Spot-on solution for Small Dogs	CC Pharma	10/04/2012	10 % imidacloprid & 2.5 % moxidectin	Spot-on solution	3 pipettes	EU/2/03/039/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/04/2012
Aerius	CC Pharma	16/02/2012	5 mg	Film Coated Tablet	20	EU/1/00/160/009	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/03/2012
Aerius	CC Pharma	16/02/2012	5 mg	Film-coated tablet	50	EU/1/00/160/012	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/03/2012
Aerius	CC Pharma	16/02/2012	5 mg	Film-coated tablet	100	EU/1/00/160/013	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/03/2012
Aerius	Empower Pharma s.r.o.	19/10/2012	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/063	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	14/11/2012
Aerius	Empower Pharma s.r.o.	19/10/2012	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/065	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	14/11/2012
Aerius	Eureco-Pharma B.V.	21/12/2011	2.5 mg	Orodispersible tablet	30	EU/1/00/160/044	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	22/10/2012

Aerius	Eureco-Pharma B.V.	21/12/2011	2.5 mg	Orodispersible tablet	90	EU/1/00/160/047	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	22/10/2012
Aerius	Eureco-Pharma B.V.	21/12/2011	2.5 mg	Orodispersible tablet	60	EU/1/00/160/046	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	22/10/2012
Aerius	hvd medical GmbH	06/03/2012	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/066	Belgium	Germany	05/04/2012
Aerius	InPharm Spzoo	30/11/2011	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/065	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	19/01/2012
Aerius	InPharm Spzoo	07/02/2012	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/066	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	05/03/2012
Aerius	Medartuum AB	13/02/2012	5 mg	Film-coated tablet	90	EU/1/00/160/036	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/02/2012
Aerius	Medcor Pharma-ceuticals BV	03/01/2012	5 mg	Orodispersible tablet	30	EU/1/00/160/056	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	06/02/2012

Aerius	S.C. PICARA TRADING SRL	08/07/2012	5 mg	Film-coated tablet	20	EU/1/00/160/009	Poland	Romania	25/07/2012
Aerius	S.C. PICARA TRADING SRL	08/07/2012	5 mg	Film-coated tablet	30	EU/1/00/160/011	Austria, Czech Republic, Hungary, Ireland, Poland, United Kingdom	Romania	25/07/2012
Afinitor	Dr Fisher Farma BV	23/10/2012	5 mg	Tablet	30	EU/1/09/538/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/11/2012
Aldara	2 Care 4	06/12/2011	5% w/w	Cream	12 sachets	EU/1/98/080/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Sweden	07/02/2012
Aldara	CC Pharma	22/02/2012	5% w/w	Cream	12 sachets	EU/1/98/080/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/03/2012
Aldara	Ginova Ltd	27/09/2012	5% w/w	Cream	12 sachets	EU/1/98/080/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	30/10/2012
Alimta	Pharma Westen GmbH	11/10/2012	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/10/2012
Aloxi	Max Pharma Group sro	31/10/2011	250 µg	Solution for injection	1 vial	EU/1/04/306/001	Germany	Czech Republic	13/02/2012
Aloxi	Medartuum AB	09/11/2012	250 µg	Solution for injection	1 vial	EU/1/04/306/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	20/11/2012
Apidra	Dr Fisher Farma BV	21/12/2011	100 Units/ml	Solution for injection	5 cartridges	EU/1/04/285/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/01/2012
Apidra	Medcor Pharma-ceuticals BV	28/06/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/04/285/032	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/09/2012

Apidra	Medcor Pharma-ceuticals BV	06/09/2012	100 Units/ml	Solution for injection	5 cartridges	EU/1/04/285/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, United Kingdom	The Netherlands	19/11/2012
Apidra	Pharma Westen GmbH	01/08/2012	100 Units/ml	Solution for injection	10 cartridges	EU/1/04/285/012	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/08/2012
Apidra	PHARMACHIM AB	18/01/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/04/285/032	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Aprovel	BB Farma s.r.l.	18/06/2012	150 mg	Film-coated tablet	98	EU/1/97/046/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Aprovel	BB Farma s.r.l.	18/06/2012	150 mg	Film-coated tablet	56	EU/1/97/046/023	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Aprovel	BB Farma s.r.l.	18/06/2012	150 mg	Film-coated tablet	28	EU/1/97/046/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Aprovel	BB Farma s.r.l.	19/06/2012	300 mg	Film-coated tablet	98	EU/1/97/046/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012

Aprovel	BB Farma s.r.l.	19/06/2012	300 mg	Film-coated tablet	28	EU/1/97/046/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Aprovel	BB Farma s.r.l.	19/06/2012	300 mg	Film-coated tablet	56	EU/1/97/046/028	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Aprovel	Imed Healthcare Ltd	24/09/2012	150 mg	Film-coated tablet	28	EU/1/97/046/022	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	08/10/2012
Aprovel	Imed Healthcare Ltd	24/09/2012	300 mg	Film-coated tablet	28	EU/1/97/046/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	08/10/2012
Aprovel	kohlpharma GmbH	11/05/2012	150 mg	Tablet	98	EU/1/97/046/006	Greece	Germany	30/05/2012
Aprovel	kohlpharma GmbH	17/07/2012	300 mg	Tablet	98	EU/1/97/046/009	Greece	Germany	03/08/2012
Aprovel	PI-Pharma N.V.	13/08/2012	150 mg	Film-coated tablet	28	EU/1/97/046/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	24/09/2012
Aprovel	PI-Pharma N.V.	13/08/2012	300 mg	Film-coated tablet	28	EU/1/97/046/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	03/10/2012

Aprovel	PI-Pharma N.V.	14/11/2012	300 mg	Film-coated tablet	98	EU/1/97/046/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	06/12/2012
Aptivus	HAEMATO PHARM AG	09/07/2012	250 mg	Capsule, soft	120	EU/1/05/315/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/08/2012
Aranesp	Axicorp Pharma B.V.	13/12/2012	300 µg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/01/185/096	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/12/2012
Aranesp	axicorp Pharma GmbH	13/12/2012	80 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/089	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/12/2012
Aranesp	Cross Pharma AB	21/11/2012	30 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/081	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/12/2012
Aranesp	Cross Pharma AB	21/11/2012	40 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/083	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/12/2012
Aranesp	Euro Registratie Collectief BV	18/01/2012	30 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/081	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/02/2012
Aranesp	Euro Registratie Collectief BV	23/01/2012	40 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/083	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/02/2012
Aranesp	Euro Registratie Collectief BV	23/01/2012	50 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/085	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/02/2012

Aranesp	Euro Registratie Collectief BV	23/01/2012	60 µg	Solution for injection	4 pre-filled syringe with needle guard	EU/1/01/185/087	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/02/2012
Aranesp	Euro Registratie Collectief BV	23/01/2012	80 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/089	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/02/2012
Aranesp	Inopha GmbH	07/03/2012	300 µg	Solution for injection	1 pre-filled syringe	EU/1/01/185/021	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/03/2012
Aranesp	Inopha GmbH	07/03/2012	500 µg	Solution for injection	1 pre-filled syringe	EU/1/01/185/031	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/03/2012
Aranesp	Medartuum AB	09/03/2012	300 µg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/01/185/096	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012
Aranesp	Medartuum AB	23/03/2012	150 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/095	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012
Aranesp	Medartuum AB	23/03/2012	30 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/081	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012
Aranesp	Medartuum AB	23/03/2012	40 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/083	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012

Aranesp	Medcor Pharma-ceuticals BV	05/03/2012	150 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/095	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/04/2012
Aranesp	Paranova Läkemedel AB	20/06/2012	40 µg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/01/185/082	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Aranesp	Paranova Läkemedel AB	20/06/2012	40 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/083	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Aranesp	Paranova Läkemedel AB	21/06/2012	150 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/095	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Aranesp	Paranova Läkemedel AB	21/06/2012	150 µg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/01/185/094	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/07/2012
Aranesp	PHARMACHIM AB	04/10/2012	20 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/079	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012
Aranesp	PHARMACHIM AB	04/10/2012	30 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/081	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012

Aranesp	PHARMACHIM AB	04/10/2012	40 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/083	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012
Aranesp	PHARMACHIM AB	04/10/2012	50 µg	Solution for injection	4 pre-filled syringes with needle guard	EU/1/01/185/085	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012
Aranesp	PHARMACHIM AB	04/10/2012	60 µg	Solution for injection	4 pre-filled syringe with needle guard	EU/1/01/185/087	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012
Arava	Adripharma GmbH	31/08/2012	10 mg	Film-coated tablet	30	EU/1/99/118/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/11/2012
Arava	Adripharma GmbH	31/08/2012	20 mg	Film-coated tablet	30	EU/1/99/118/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/11/2012
Arava	Axicorp Pharma GmbH	13/03/2012	20 mg	Film-coated tablet	100	EU/1/99/118/008	Austria, Belgium, Bulgaria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Lithuania, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Arava	BB Farma s.r.l.	07/05/2012	10 mg	Film-coated tablet	30	EU/1/99/118/003	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Romania, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	16/05/2012

Arava	BB Farma s.r.l.	07/05/2012	20 mg	Film-coated tablet	30	EU/1/99/118/007	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Romania, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	23/05/2012
Arava	Cambridge Major Laboratories Europe B.V	29/02/2012	10 mg	Film-coated tablet	100	EU/1/99/118/004	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/09/2012
Arava	isopharm GmbH	06/01/2012	20 mg	Film-coated tablet	100	EU/1/99/118/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	17/01/2012
Arixtra	BR Pharma International Ltd	21/09/2012	2.5 mg/ 0.5 ml	Solution for injection	20 pre-filled syringes	EU/1/02/206/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, Malta, United Kingdom	10/10/2012
Arixtra	Delfarma SpZoo	30/01/2012	2.5 mg/ 0.5 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	20/02/2012
Arixtra	Euromedicines , S.L.	13/12/2011	2.5 mg/ 0.5 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	04/01/2012
Arixtra	EuroPharma DK ApS	27/12/2011	2.5 mg/ 0.5 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/01/2012
Arixtra	EuroPharma DK ApS	27/12/2011	2.5 mg/ 0.5 ml	Solution for injection	20 pre-filled syringes	EU/1/02/206/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/01/2012

Atripla	Cross Pharma AB	14/12/2012	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	20/12/2012
Atripla	Docpharm GmbH&CoKGaA	18/07/2012	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/08/2012
Atripla	Docpharm GmbH&CoKGaA	08/08/2012	--	Film-coated tablet	3 x 30	EU/1/07/430/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/08/2012
Atripla	eurorx Arzneimittel GmbH	17/11/2010	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/05/2012
Atripla	propharmed GmbH	16/07/2012	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/08/2012
Atripla	propharmed GmbH	17/07/2012	--	Film-coated tablet	3 x 30	EU/1/07/430/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/08/2012
Avamys	BB Farma s.r.l.	01/06/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	10/07/2012
Avamys	Crystal Pharma Ltd	22/10/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	19/11/2012

Avamys	Ethigen Limited	06/11/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	20/11/2012
Avamys	Euromedicines S.L.	25/05/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	05/07/2012
Avamys	Ginova Ltd	30/11/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	17/12/2012
Avamys	Interport Ltd	07/02/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands	United Kingdom	28/02/2012
Avamys	Kosei Pharma UK Limited	29/10/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	30/10/2012
Avamys	Mediwin Ltd	29/08/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	06/09/2012
Avamys	O.P.D. Laboratories Ltd	16/07/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	23/07/2012

Avamys	Paranova Läkemedel AB	19/10/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	26/10/2012
Avamys	Swingward Ltd TA Medihealth	18/10/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	05/11/2012
Avamys	WAYMADE PLC	13/07/2012	27.5 micrograms/ spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/08/2012
Avastin	Cambridge Major Laboratories Europe B.V	29/02/2012	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/002	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/09/2012
Avastin	eurorx Arzneimittel GmbH	17/11/2010	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/06/2012
Avastin	Inopha GmbH	13/04/2012	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	14/05/2012
Avastin	Medartuum AB	08/11/2012	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/11/2012
Avonex	ABACUS MEDICINE A/S	19/10/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes +4 needles	EU/1/97/033/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012

Avonex	ABACUS MEDICINE A/S	12/11/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/11/2012
Avonex	axicorp Pharma B.V.	11/09/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled pens	EU/1/97/033/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/11/2012
Avonex	axicorp Pharma B.V.	23/10/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled pens	EU/1/97/033/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/11/2012
Avonex	CC Pharma	19/12/2011	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled pens	EU/1/97/033/006	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Hungary, Ireland, Italy, Norway, Portugal, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/01/2012
Avonex	Docpharm GmbH&CoKGaA	13/08/2012	30 µg (6 million IU)	Powder and solvent for solution for injection	4 vials with Bioset device +4 pre-filled syringes	EU/1/97/033/002	Belgium	Germany	18/09/2012
Avonex	Dr Fisher Farma BV	30/03/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/06/2012
Avonex	Emra-Med	03/05/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled pens	EU/1/97/033/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/05/2012
Avonex	Emra-Med	03/05/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled pens	EU/1/97/033/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/05/2012

Avonex	eurorx Arzneimittel GmbH	17/11/2010	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/06/2012
Avonex	Hecht-Pharma GmbH	28/06/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Austria	Germany	29/08/2012
Avonex	kohlpharma GmbH	26/04/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled pens	EU/1/97/033/005	France, Sweden	Germany	11/05/2012
Avonex	Medicopharm AG	13/12/2011	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled pens	EU/1/97/033/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Avonex	Medicopharm AG	03/07/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled pens	EU/1/97/033/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/10/2012
Avonex	MTK Pharma GmbH	16/12/2011	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Czech Republic, Finland, France, Greece, Sweden	Germany	19/01/2012
Avonex	Orifarm AB	14/03/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled pens	EU/1/97/033/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/06/2012
Avonex	Pharma Westen GmbH	01/10/2012	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled pens	EU/1/97/033/006	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
Axura	axicorp Pharma GmbH	12/03/2012	10 mg	Film-coated tablet	98	EU/1/02/218/014	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/03/2012
Axura	B&S Healthcare	03/09/2012	10 mg	Film-coated tablet	56	EU/1/02/218/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	10/09/2012
Axura	Elpis Ltd.	28/06/2012	10 mg	Film-coated tablet	56	EU/1/02/218/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Latvia	20/08/2012

Axura	Emra-Med	31/07/2012	10 mg	Film-coated tablet	98	EU/1/02/218/014	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/10/2012
Axura	PHARMACHIM AB	03/05/2012	10 mg	Film-coated tablet	98	EU/1/02/218/014	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/05/2012
Axura	PHARMACHIM AB	03/05/2012	10 mg	Film-coated tablet	56	EU/1/02/218/008	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/05/2012
Axura	PHARMACHIM AB	03/05/2012	10 mg	Film-coated tablet	28	EU/1/02/218/007	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	22/05/2012
Azarga	ABACUS MEDICINE A/S	10/12/2012	10 mg/ml / 5 mg/ml	Eye drops, suspension	3 bottles	EU/1/08/482/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/12/2012
Azarga	Copharma ApS	05/07/2012	10 mg/ml / 5 mg/ml	Eye drops, suspension	3 bottles	EU/1/08/482/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012
Azarga	Copharma ApS	05/07/2012	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012
Azarga	Cross Pharma AB	09/10/2012	10 mg/ml / 5 mg/ml	Eye drops, suspension	3 bottles	EU/1/08/482/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	22/10/2012

Azarga	PCO Manufacturing	16/03/2012	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	02/04/2012
AZILECT	Abacus Medicine A/S	25/11/2011	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012
AZILECT	Abacus Medicine A/S	25/11/2011	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012
AZILECT	B2B Medical GmbH	24/11/2011	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012
AZILECT	B2B Medical GmbH	04/01/2012	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/01/2012
AZILECT	Copharma ApS	20/08/2012	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/08/2012
AZILECT	Copharma ApS	20/08/2012	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/08/2012
AZILECT	Copharma ApS	20/08/2012	1 mg	Tablet	10	EU/1/04/304/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/08/2012
AZILECT	Orifarm AB	01/10/2012	1 mg	Tablet	28	EU/1/04/304/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	11/10/2012

AZILECT	ORIFARM AB	03/12/2012	1 mg	Tablet	112	EU/1/04/304/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	06/12/2012
AZILECT	Paranova Läkemedel AB	10/02/2012	1 mg	Tablet	28	EU/1/04/304/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/03/2012
AZILECT	Paranova Läkemedel AB	10/02/2012	1 mg	Tablet	112	EU/1/04/304/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/03/2012
Azopt	Copharma ApS	19/10/2012	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/10/2012
Azopt	Copharma ApS	19/10/2012	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/10/2012
Azopt	Pharmachim AB	05/10/2012	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Azopt	REMEDIX GmbH	14/03/2012	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	18/04/2012

Baraclude	Paranova Danmark AS	26/07/2012	0.5 mg	Film-coated tablet	30 x 1	EU/1/06/343/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
Baraclude	PHARMACHIM AB	16/03/2012	0.5 mg	Film-coated tablet	30 x 1	EU/1/06/343/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	02/05/2012
Benlysta	CC Pharma	03/12/2012	120 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/11/700/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Benlysta	CC PHARMA GMBH	03/12/2012	400 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/11/700/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Benlysta	Kohlpharma GmbH	13/04/2012	120 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/11/700/001	The Netherlands	Germany	26/04/2012
Benlysta	kohlpharma GmbH	13/04/2012	400 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/11/700/002	The Netherlands	Germany	26/04/2012
Benlysta	Pharma Westen GmbH	16/01/2012	400 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/11/700/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/01/2012
Betaferon	ADL Pharma GmbH	14/05/2012	0.25 mg/ml	Powder and solvent for solution for injection	14 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/09/2012
Betaferon	Adripharma GmbH	17/12/2012	0.25 mg/ml	Powder and solvent for solution for injection	14 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/12/2012

Betaferon	Adripharma GmbH	17/12/2012	0.25 mg/ml	Powder and solvent for solution for injection	3 x 14 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/12/2012
Betaferon	Cross Pharma AB	05/10/2012	0.25 mg/ml	Powder and solvent for solution for injection	15 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/10/2012
Betaferon	Docpharm GmbH& CoKGaA	22/02/2012	0.25 mg/ml	Powder and solvent for solution for injection	14 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Betaferon	Docpharm GmbH& CoKGaA	22/02/2012	0.25 mg/ml	Powder and solvent for solution for injection	3 x 15 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Betaferon	Docpharm GmbH& CoKGaA	23/02/2012	0.25 mg/ml	Powder and solvent for solution for injection	3 x 14 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012

Betaferon	HAEMATO PHARM AG	27/09/2012	0.25 mg/ml	Powder and solvent for solution for injection	3 x 15 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/10/2012
Betaferon	Paranova Läkemedel AB	18/01/2012	0.25 mg/ml	Powder and solvent for solution for injection	15 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/03/2012
Betaferon	remedix Gmbh	22/06/2012	0.25 mg/ml	Powder and solvent for solution for injection	15 x (1 vial + 1 pre-filled syringe + 1 vial adapter with needle + 2 alcohol wipes)	EU/1/95/003/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/08/2012
Binocrit	Eureco-Pharma B.V.	27/06/2012	40000 IU/1 ml	Solution for injection in pre-filled syringe	1 pre-filled syringe	EU/1/07/410/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	11/07/2012
Bondronat	Emra-Med	29/02/2012	6 mg/6 ml	Concentrate for solution for infusion	1 vial	EU/1/96/012/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/04/2012

Bondronat	Emra-Med	29/02/2012	6 mg/6 ml	Concentrate for solution for infusion	5 vials	EU/1/96/012/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/04/2012
Bonviva	Combiphar Europe BV	23/02/2012	150 mg	Film-coated tablet	1	EU/1/03/265/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	10/04/2012
Bonviva	Orifarm OY	03/01/2012	150 mg	Film-coated tablet	3	EU/1/03/265/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	16/01/2012
Bonviva	Paranova Oy	08/03/2012	150 mg	Film-coated tablet	3	EU/1/03/265/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	12/04/2012
Bonviva	REMEDIX GMBH	02/07/2012	3 mg/3ml	Solution for injection	1 pre-filled syringe	EU/1/03/265/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	13/07/2012
Bridion	Delfarma SpZoo	28/05/2012	100 mg/ml	Solution for injection	10 vials	EU/1/08/466/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	20/06/2012
Brilique	Euro Registratie Collectief BV	27/02/2012	90 mg	Film-coated tablet	56	EU/1/10/655/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/04/2012

Brilique	HAEMATO PHARM AG	11/07/2012	90 mg	Film-coated tablet	56	EU/1/10/655/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Brilique	HAEMATO PHARM AG	11/07/2012	90 mg	Film-coated tablet	100 x 1	EU/1/10/655/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Bydureon	Orifarm AS	15/02/2012	2 mg	Powder and solvent for prolonged release suspension for injection	4 x 1 single-dose kit (1 vial + 1 syringe)	EU/1/11/696/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/03/2012
BYETTA	ABACUS MEDICINE A/S	16/07/2012	10 micrograms	Solution for injection	1 pre-filled pen	EU/1/06/362/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	31/07/2012
BYETTA	Emra-Med	22/10/2012	5 micrograms	Solution for injection	1 pre-filled pen	EU/1/06/362/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/10/2012
BYETTA	PCO Manufacturing	15/02/2012	10 micrograms	Solution for injection	1 pre-filled pen	EU/1/06/362/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	15/03/2012
Caelyx	Aaston Healthcare GmbH	16/10/2012	2 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/001	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/10/2012

Cancidas	ABACUS MEDICINE A/S	16/11/2012	50 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/11/2012
Cancidas	ABACUS MEDICINE A/S	16/11/2012	70 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/11/2012
Cancidas	Dr Fisher Farma BV	29/05/2012	70 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	25/06/2012
Cancidas	Dr Fisher Farma BV	20/06/2012	50 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	11/07/2012
Cancidas	Medartuum AB	30/03/2012	50 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/05/2012
Cancidas	Medartuum AB	30/03/2012	70 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/05/2012
CellCept	Adripharma GmbH	23/08/2012	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/10/2012

CellCept	Adripharma GmbH	23/08/2012	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/11/2012
CellCept	Axicorp Pharma B.V.	25/05/2012	1 g/5 ml	Powder for oral suspension	1 bottle	EU/1/96/005/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/06/2012
CellCept	Clear Pharmacy	15/12/2011	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/01/2012
CellCept	Cross Pharma AB	02/03/2012	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/03/2012
CellCept	EU-Pharma BV	09/10/2012	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	18/10/2012
CellCept	EuroPharma DK ApS	04/07/2012	1 g/5 ml	Powder for oral suspension	1 bottle	EU/1/96/005/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/07/2012
CellCept	Mediwin Ltd	24/02/2012	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/03/2012
CellCept	Mediwin Ltd	24/02/2012	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/03/2012

CellCept	O.P.D. Laboratories Ltd	17/10/2012	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	20/11/2012
CellCept	O.P.D. Laboratories Ltd	04/12/2012	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	12/12/2012
Cetrotide	ABACUS MEDICINE A/S	12/04/2012	0.25 mg	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 2 injection needles + 2 alcohol swabs	EU/1/99/100/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/04/2012
Cetrotide	Emra-Med	22/11/2012	0.25 mg	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 2 injection needles + 2 alcohol swabs	EU/1/99/100/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	30/11/2012
Cetrotide	Orifarm AS	12/01/2012	0.25 mg	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 2 injection needles + 2 alcohol swabs	EU/1/99/100/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/02/2012
Champix	Euro Registratie Collectief BV	27/02/2012	1.0 mg	Film-coated tablet	56	EU/1/06/360/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	28/03/2012
Champix	Euromedicines S.L.	01/03/2012	0.5 mg and 1.0 mg	Film-coated tablet	11 x 0.5 mg + 14 x 1 mg	EU/1/06/360/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	22/06/2012

Champix	Euromedicines S.L.	01/03/2012	0.5 mg and 1.0 mg	Film-coated tablet	11 x 0.5 mg + 14 x 1 mg plus 28 x 1mg	EU/1/06/360/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	22/06/2012
Champix	Euromedicines S.L.	01/03/2012	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	22/06/2012
Champix	Euromedicines S.L.	01/03/2012	1.0 mg	Film-coated tablet	56	EU/1/06/360/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	22/06/2012
Champix	Euromedicines S.L.	01/03/2012	1.0 mg	Film-coated tablet	112	EU/1/06/360/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	22/06/2012
Champix	kohlpharma GmbH	26/04/2012	0.5 mg and 1.0 mg	Film-coated tablet	11 x 0.5 mg + 14 x 1 mg plus 28 x 1mg	EU/1/06/360/012	Czech Republic, Poland	Germany	25/07/2012
Champix	Orifarm AB	12/04/2012	1.0 mg	Film-coated tablet	56	EU/1/06/360/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/05/2012
Champix	Orifarm AS	02/07/2012	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012

Champix	STRATHCLYDE PHARMACEUTICALS LTD	29/12/2011	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/01/2012
Champix	STRATHCLYDE PHARMACEUTICALS LTD	29/12/2011	1.0 mg	Film-coated tablet	56	EU/1/06/360/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/01/2012
Cialis	2 Care 4	27/04/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/05/2012
Cialis	4pharma S.r.l.	25/11/2011	20 mg	Film-coated tablet	4	EU/1/02/237/003	Bulgaria, Romania	Italy	11/01/2012
Cialis	ABACUS MEDICINE A/S	02/05/2012	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/05/2012
Cialis	ABACUS MEDICINE A/S	10/10/2012	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012
Cialis	axicorp Pharma GmbH	13/02/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/03/2012
Cialis	CC Pharma	19/12/2011	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/01/2012

Cialis	Combiphar Europe BV	21/02/2012	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/03/2012
Cialis	Copharma ApS	29/05/2012	5 mg	Film-coated tablet	14	EU/1/02/237/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/06/2012
Cialis	Copharma ApS	29/05/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/06/2012
Cialis	Cross Pharma AB	05/11/2012	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cialis	Cross Pharma AB	05/11/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cialis	Cross Pharma AB	05/11/2012	20 mg	Film-coated tablet	8	EU/1/02/237/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cialis	Cross Pharma AB	05/11/2012	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012

Cialis	Cross Pharma AB	05/11/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cialis	Dineras Czech Republic, s.r.o.	05/10/2012	20 mg	Film-coated tablet	2	EU/1/02/237/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	23/11/2012
Cialis	Dineras Czech Republic, s.r.o.	05/10/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	23/11/2012
Cialis	Dineras Czech Republic, s.r.o.	05/10/2012	20 mg	Film-coated tablet	8	EU/1/02/237/004	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	23/11/2012
Cialis	Eureco-Pharma B.V.	26/09/2012	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	12/10/2012
Cialis	Haemato Pharm AG	08/02/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	21/02/2012
Cialis	HAEMATO PHARM AG	23/03/2012	20 mg	Film-coated tablet	8	EU/1/02/237/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	29/03/2012
Cialis	Lex ano UAB	09/12/2011	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Lithuania	09/01/2012

Cialis	Medartuum AB	18/09/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Cialis	Medartuum AB	28/11/2012	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Cialis	Mediwin Ltd	21/03/2012	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	France	30/03/2012
Cialis	Mediwin Ltd	21/03/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	France	30/03/2012
Cialis	NewNeopharm BV	08/08/2012	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/08/2012
Cialis	NewNeopharm BV	08/08/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/08/2012
Cialis	Orifarm AB	28/02/2012	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	14/03/2012
Cialis	Paranova Danmark AS	27/04/2012	2.5 mg	Film-coated tablet	28	EU/1/02/237/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/06/2012

Cialis	Paranova Läkemedel AB	13/11/2012	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/12/2012
Cialis	Pharmazet Pharmaceuticals s.r.o.	18/06/2012	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	24/07/2012
Cimzia	Cross Pharma AB	03/09/2012	200 mg	Solution for injection	2 pre-filled syringes + 2 alcohol swabs	EU/1/09/544/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	11/09/2012
Cimzia	PHARMACHIM AB	15/12/2011	200 mg	Solution for injection	2 pre-filled syringes + 2 alcohol swabs	EU/1/09/544/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Latvia, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	10/01/2012
Circadin	Interport Ltd	13/12/2012	2 mg	Prolonged-release tablet	21	EU/1/07/392/001	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands	United Kingdom	21/12/2012
Circadin	Waymade PLC	17/07/2012	2 mg	Prolonged-release tablet	30	EU/1/07/392/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	17/10/2012
Clopidogrel Teva Pharma	Doncaster Pharmaceuticals Group Ltd	30/05/2012	75 mg	Film-coated tablet	28 x 1 in a calendar pack	EU/1/09/561/010	United Kingdom	Ireland, Malta	13/07/2012
CoAprovel	ADL Pharma GmbH	22/05/2012	300 mg/ 12.50 mg	Film-coated tablet	56	EU/1/98/086/018	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012
CoAprovel	ADL Pharma GmbH	22/05/2012	300 mg/ 12.50 mg	Film-coated tablet	98	EU/1/98/086/020	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012

CoAprovel	BB Farma s.r.l.	07/06/2012	300 mg/ 25 mg	Film-coated tablet	98	EU/1/98/086/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/06/2012
CoAprovel	BB Farma s.r.l.	02/10/2012	150 mg/ 12.50 mg	Film-coated tablet	98	EU/1/98/086/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
CoAprovel	PI-Pharma N.V.	13/08/2012	300 mg/ 12.50 mg	Film-coated tablet	28	EU/1/98/086/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	03/10/2012
CoAprovel	PI-Pharma N.V.	13/08/2012	300 mg/ 25 mg	Film-coated tablet	28	EU/1/98/086/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	03/10/2012
CoAprovel	PI-Pharma N.V.	14/11/2012	300 mg/ 12.50 mg	Film-coated tablet	98	EU/1/98/086/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	06/12/2012
CoAprovel	PI-Pharma N.V.	14/11/2012	300 mg/ 25 mg	Film-coated tablet	98	EU/1/98/086/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	06/12/2012

Combivir	HAEMATO PHARM AG	14/06/2012	150 mg/ 300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands	Ireland, United Kingdom	10/07/2012
Combivir	Medcor Pharma-ceuticals BV	02/05/2012	150 mg/ 300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/06/2012
Combivir	PHARMACHIM AB	26/01/2012	150 mg/ 300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, France, Germany, Greece, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	02/02/2012
Combivir	Tisida GmbH	07/06/2012	150 mg/ 300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/07/2012
Competact	CC Pharma	14/03/2012	15 mg/ 850 mg	Film-coated tablet	112	EU/1/06/354/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/03/2012
Competact	CC Pharma	14/03/2012	15 mg/ 850 mg	Film-coated tablet	196 (2 x 98)	EU/1/06/354/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Competact	Emra-Med	12/04/2012	15 mg/ 850 mg	Film-coated tablet	112	EU/1/06/354/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Competact	O.P.D. Laboratories Ltd	29/04/2009	15 mg/ 850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/02/2012
Competact	Paranova Läkemedel AB	27/03/2012	15 mg/ 850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/05/2012
Competact	PCO Manufacturing	24/02/2012	15 mg / 850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	11/04/2012

Comtess	Cross Pharma AB	13/01/2012	200 mg	Film-coated tablet	100	EU/1/98/082/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/01/2012
Conbriza	CC Pharma	09/07/2012	20 mg	Film-coated tablet	28	EU/1/09/511/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Conbriza	CC Pharma	09/07/2012	20 mg	Film-coated tablet	84	EU/1/09/511/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Cymbalta	Cambridge Major Laboratories Europe B.V	05/03/2012	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Cymbalta	Cambridge Major Laboratories Europe B.V	05/03/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Cymbalta	Cambridge Major Laboratories Europe B.V	05/03/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Cymbalta	Cambridge Major Laboratories Europe B.V	05/03/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Cymbalta	CD Pharma Limited	07/02/2012	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/02/2012
Cymbalta	CD Pharma Limited	06/03/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/03/2012

Cymbalta	Dr Fisher Farma BV	17/02/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	30/03/2012
Cymbalta	Dr Fisher Farma BV	20/02/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	30/03/2012
Cymbalta	Euro Registratie Collectief BV	04/10/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/10/2012
Cymbalta	Medartuum AB	23/11/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/11/2012
Cymbalta	MPT Pharma	10/09/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	25/09/2012
Cymbalta	MPT Pharma	10/09/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	25/09/2012
Cymbalta	MPT Pharma	14/11/2012	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/296/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	28/11/2012

Cymbalta	Orifarm AB	06/11/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cymbalta	Orifarm AB	06/11/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
Cymbalta	Tisida GmbH	20/06/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/06/2012
Cymbalta	Tisida GmbH	20/06/2012	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/07/2012
Cymbalta	Tisida GmbH	21/06/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/07/2012
Cymbalta	Tisida GmbH	21/06/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/07/2012
Dasselta	BR Pharma International Ltd	17/08/2012	5 mg	Film-coated tablet	50	EU/1/11/739/005	Austria	Germany	05/10/2012
Dasselta	BR Pharma International Ltd	17/08/2012	5 mg	Film-coated tablet	100	EU/1/11/739/007	Austria	Germany	05/10/2012
Dasselta	Emra-Med	25/06/2012	5 mg	Film-coated tablet	10	EU/1/11/739/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Dasselta	Emra-Med	25/06/2012	5 mg	Film-coated tablet	100	EU/1/11/739/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012

Dasselta	Emra-Med	25/06/2012	5 mg	Film-coated tablet	30	EU/1/11/739/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Daxas	Beragena Arzneimittel GmbH	05/07/2012	500 µg	Film-coated tablet	30	EU/1/10/636/002	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	24/07/2012
Daxas	Emra-Med	21/09/2012	500 µg	Film-coated tablet	30	EU/1/10/636/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/10/2012
Daxas	Emra-Med	21/09/2012	500 µg	Film-coated tablet	90	EU/1/10/636/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/10/2012
Daxas	kohlpharma GmbH	16/11/2012	500 µg	Film-coated tablet	30	EU/1/10/636/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/11/2012
Daxas	kohlpharma GmbH	29/11/2012	500 µg	Film-coated tablet	90	EU/1/10/636/003	Slovak Republic	Germany	10/12/2012
Daxas	Paranova Danmark AS	16/11/2012	500 µg	Film-coated tablet	30	EU/1/10/636/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/11/2012
Daxas	Paranova Danmark AS	16/11/2012	500 µg	Film-coated tablet	90	EU/1/10/636/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/11/2012

Daxas	Paranova Läkemedel AB	29/10/2012	500 µg	Film-coated tablet	30	EU/1/10/636/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Daxas	Paranova Läkemedel AB	29/10/2012	500 µg	Film-coated tablet	90	EU/1/10/636/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Daxas	Pharma Gerke GmbH	28/11/2012	500 µg	Film-coated tablet	90	EU/1/10/636/003	Austria, Belgium, Bulgaria, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Desloratadine ratiopharm	2 Care 4	11/10/2012	5 mg	Film-coated tablet	90	EU/1/11/746/010	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/10/2012
Desloratadine ratiopharm	2 Care 4	11/10/2012	5 mg	Film-coated tablet	30	EU/1/11/746/006	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/10/2012
Draxxin	Bioptivet Tierarzneimittel GmbH& CO	03/09/2012	100 mg/ml	Solution for injection	100 ml	EU/2/03/041/003	Romania	Germany	05/10/2012
DuoPlavin	Axicorp Pharma GmbH	13/02/2012	75 mg/100 mg	Film-coated tablet	28	EU/1/10/619/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/02/2012
DuoPlavin	B2B Medical GmbH	25/04/2012	75 mg/100 mg	Film-coated tablet	28	EU/1/10/619/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/08/2012
DuoPlavin	B2B Medical GmbH	25/04/2012	75 mg/100 mg	Film-coated tablet	100 x 1	EU/1/10/619/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/08/2012
DuoPlavin	Beragena Arzneimittel GmbH	08/03/2012	75 mg/100 mg	Film-coated tablet	100 x 1	EU/1/10/619/014	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	25/06/2012

DuoPlavin	Beragena Arzneimittel GmbH	08/03/2012	75 mg/100 mg	Film-coated tablet	28	EU/1/10/619/009	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/03/2012
DuoPlavin	CC Pharma	14/03/2012	75 mg/100 mg	Film-coated tablet	100 x 1	EU/1/10/619/014	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/11/2012
DuoPlavin	Emra-Med	23/01/2012	75 mg/100 mg	Film-coated tablet	28	EU/1/10/619/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/05/2012
DuoPlavin	Milinda GmbH Milinda GmbH & Co. KG& Co. KG	30/11/2012	75 mg/100 mg	Film-coated tablet	28	EU/1/10/619/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Germany	21/12/2012
DuoTrav	Amimed Direct Ltd	01/08/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	14/08/2012
DuoTrav	axicorp Pharma GmbH	24/02/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/338/002	Austria, Belgium, Bulgaria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/03/2012
DuoTrav	Interport Ltd	01/11/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, The Netherlands	United Kingdom	12/11/2012
DuoTrav	Interport Ltd	01/11/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/338/002	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, The Netherlands	United Kingdom	13/11/2012
DuoTrav	O.P.D. Laboratories Ltd	17/10/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	30/10/2012
DuoTrav	Paranova Oy	03/04/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/338/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	11/05/2012

DuoTrav	PHARMACHIM AB	11/10/2012	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Poland, Portugal, Spain, The Netherlands, United Kingdom	Sweden	05/11/2012
Ebixa	ABACUS MEDICINE A/S	17/05/2012	10 mg	Film-coated tablet	112	EU/1/02/219/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/05/2012
Ebixa	ABACUS MEDICINE A/S	12/07/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/07/2012
Ebixa	axicorp Pharma GmbH	26/01/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/04/2012
Ebixa	CC Pharma	28/12/2011	10 mg	Film-coated tablet	42	EU/1/02/219/017	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Ebixa	CC Pharma	28/12/2011	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Ebixa	Copharma ApS	25/05/2012	10 mg	Film-coated tablet	30	EU/1/02/219/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/05/2012
Ebixa	Copharma ApS	25/05/2012	10 mg	Film-coated tablet	50	EU/1/02/219/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/05/2012
Ebixa	Copharma ApS	25/05/2012	10 mg	Film-coated tablet	100	EU/1/02/219/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/05/2012

Ebixa	Copharma ApS	16/11/2012	10 mg	Film-coated tablet	28	EU/1/02/219/007	Austria, Belgium, Bulgaria, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/11/2012
Ebixa	Copharma ApS	16/11/2012	10 mg	Film-coated tablet	56	EU/1/02/219/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/11/2012
Ebixa	Copharma ApS	16/11/2012	10 mg	Film-coated tablet	112	EU/1/02/219/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/11/2012
Ebixa	Copharma ApS	16/11/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/11/2012
Ebixa	Copharma ApS	16/11/2012	20 mg	Film-coated tablet	28	EU/1/02/219/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/11/2012
Ebixa	Copharma ApS	16/11/2012	20 mg	Film-coated tablet	98	EU/1/02/219/031	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/11/2012

Ebixa	Cross Pharma AB	06/09/2012	20 mg	Film-coated tablet	28	EU/1/02/219/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	14/09/2012
Ebixa	Cross Pharma AB	06/09/2012	20 mg	Film-coated tablet	98	EU/1/02/219/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	14/09/2012
Ebixa	Emra-Med	07/06/2012	10 mg	Film-coated tablet	42	EU/1/02/219/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012
Ebixa	Emra-Med	07/06/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012
Ebixa	EuroPharma DK ApS	23/12/2011	5 mg/pump actuation	Oral solution	1 bottle of 50 ml	EU/1/02/219/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/02/2012
Ebixa	Impexeco S.A.	17/09/2012	10 mg	Film-coated tablet	56	EU/1/02/219/008	Austria, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium	25/10/2012
Ebixa	Mediwin Ltd	06/12/2012	10 mg	Film-coated tablet	28	EU/1/02/219/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	13/12/2012
Ebixa	Orifarm AB	24/01/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	01/02/2012

Ebixa	Orifarm AS	30/01/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/03/2012
Ebixa	Paranova Danmark AS	22/03/2012	5 mg/pump actuation	Oral solution	1 bottle	EU/1/02/219/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/05/2012
Ebixa	Paranova Läkemedel AB	25/10/2012	5 mg/pump actuation	Oral solution	1 bottle	EU/1/02/219/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/11/2012
Ebixa	Paranova Läkemedel AB	26/10/2012	20 mg	Film-coated tablet blister (alu/PP) in carton	98	EU/1/02/219/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	29/10/2012
Ebixa	Pharma Westen GmbH	15/03/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/03/2012
Ebixa	Profind Wholesale Ltd.	01/02/2012	10 mg	Film-coated tablet	56	EU/1/02/219/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	02/03/2012
Ebixa	Profind Wholesale Ltd.	24/04/2012	10 mg	Film-coated tablet	28	EU/1/02/219/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/05/2012
Ebixa	S.C. PICARA TRADING SRL	04/05/2012	10 mg	Film-coated tablet	56	EU/1/02/219/008	Austria, Czech Republic, Hungary, Ireland, Italy, Poland, Slovenia, United Kingdom	Romania	28/06/2012

Ebixa	STRATHCLYDE PHARMACEUTICALS LTD	31/03/2012	10 mg	Film-coated tablet	42	EU/1/02/219/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/05/2012
Ebixa	STRATHCLYDE PHARMACEUTICALS LTD	31/03/2012	10 mg	Film-coated tablet	98	EU/1/02/219/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/05/2012
Ebixa	STRATHCLYDE PHARMACEUTICALS LTD	04/07/2012	10 mg	Film-coated tablet	56	EU/1/02/219/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/07/2012
Ebixa	STRATHCLYDE PHARMACEUTICALS LTD	04/09/2012	10 mg	Film-coated tablet	28	EU/1/02/219/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	13/09/2012
Edurant	HAEMATO PHARM AG	08/02/2012	25 mg	Film-coated tablet	30	EU/1/11/736/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/02/2012
Efient	Doncaster Pharmaceuticals Group Ltd	12/12/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	20/12/2012
Efient	EU-Pharma BV	04/01/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	31/01/2012
Efient	EU-Pharma BV	04/01/2012	5 mg	Film-coated tablet	28	EU/1/08/503/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	30/01/2012
Efient	Ginova Ltd	19/06/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	09/07/2012

Efient	Interport Ltd	10/10/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, The Netherlands	United Kingdom	30/10/2012
Efient	Medartuum AB	22/10/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/11/2012
Efient	Mediwin Ltd	27/03/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/03/2012
Efient	O.P.D. Laboratories Ltd	29/03/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	26/04/2012
Efient	O.P.D. Laboratories Ltd	16/07/2012	5 mg	Film-coated tablet	28	EU/1/08/503/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	20/07/2012
Efient	PCO Manufacturing	24/02/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	09/03/2012
Efient	Polyfarma Ltd	05/11/2012	5 mg	Film-coated tablet	28	EU/1/08/503/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	26/11/2012
Efient	STRATHCLYDE PHARMACEUTICALS LTD	04/09/2012	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	13/09/2012
Efient	WAYMADE PLC	10/09/2012	10 mg	Film-coated tablet	30	EU/1/08/503/016	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/09/2012
Elaprase	Orifarm AS	15/02/2012	2 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/06/365/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/03/2012

ellaOne	2 Care 4	20/03/2012	30 mg	Tablet	1	EU/1/09/522/001	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/04/2012
ellaOne	2 Care 4	30/03/2012	30 mg	Tablet	1	EU/1/09/522/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/04/2012
Emselex	2 Care 4	01/03/2012	7.5 mg	Prolonged-release tablet	28	EU/1/04/294/017	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/03/2012
Emselex	2 Care 4	01/03/2012	7.5 mg	Prolonged-release tablet	98	EU/1/04/294/020	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	08/03/2012
Emselex	Copharma ApS	05/07/2012	7.5 mg	Prolonged-release tablet	98	EU/1/04/294/020	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012
Emselex	Copharma ApS	05/07/2012	7.5 mg	Prolonged-release tablet	28	EU/1/04/294/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012
Emselex	Copharma ApS	18/07/2012	15 mg	Prolonged-release tablet	98	EU/1/04/294/026	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
Emselex	Copharma ApS	18/07/2012	15 mg	Prolonged-release tablet	28	EU/1/04/294/023	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
Enbrel	ABACUS MEDICINE A/S	04/09/2012	50 mg	Solution for injection in pre-filled pen	12 pre-filled pens + 12 alcohol swabs	EU/1/99/126/021	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/09/2012

Enbrel	ABACUS MEDICINE A/S	05/09/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/09/2012
Enbrel	Axicorp Pharma B.V.	20/01/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/05/2012
Enbrel	Axicorp Pharma B.V.	13/06/2012	50 mg	Solution for injection in pre-filled pen	12 pre-filled pens + 12 alcohol swabs	EU/1/99/126/021	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/07/2012
Enbrel	Beragena Arzneimittel GmbH	14/06/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	28/06/2012
Enbrel	Beragena Arzneimittel GmbH	14/06/2012	50 mg	Solution for injection in pre-filled pen	12 pre-filled pens + 12 alcohol swabs	EU/1/99/126/021	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	28/06/2012
Enbrel	Delphi Pharmaceuticals BV	17/10/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	22/11/2012
Enbrel	Emra-Med	19/12/2011	50 mg	Solution for injection in pre-filled pen	12 pre-filled pens + 24 alcohol swabs	EU/1/99/126/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/01/2012
Enbrel	Inopha GmbH	29/04/2011	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/11/2012
Enbrel	kohlpharma GmbH	05/03/2012	25 mg	Powder and solvent for solution for injection	24 vials + 24 pre-filled syringes + 24 needles + 24 vial adaptors + 48 alcohol swabs	EU/1/99/126/005	Norway	Germany	21/05/2012

Enbrel	Medartuum AB	02/01/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 8 alcohol swabs	EU/1/99/126/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	01/02/2012
Enbrel	Paranova Läkemedel AB	05/10/2012	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 4 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/10/2012
Enbrel	PHARMACHIM AB	07/04/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/04/2012
Enbrel	Propharmed GmbH	13/07/2012	25 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 4 alcohol swabs	EU/1/99/126/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Enbrel	Propharmed GmbH	13/07/2012	25 mg	Solution for injection in pre-filled syringe	8 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/014	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Enbrel	Propharmed GmbH	13/07/2012	25 mg	Solution for injection in pre-filled syringe	24 pre-filled syringes + 24 alcohol swabs	EU/1/99/126/015	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Enbrel	propharmed GmbH	31/07/2012	50 mg	Solution for injection in pre-filled pen	12 pre-filled pens + 12 alcohol swabs	EU/1/99/126/021	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/08/2012
Enbrel	propharmed GmbH	31/07/2012	50 mg	Solution for injection in pre-filled pen	4 pre-filled pens + 4 alcohol swabs	EU/1/99/126/020	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/08/2012
Enbrel	propharmed GmbH	31/07/2012	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 4 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/08/2012
Enbrel	propharmed GmbH	31/07/2012	50 mg	Solution for injection in pre-filled syringe	12 pre-filled syringes + 12 alcohol swabs	EU/1/99/126/018	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/08/2012

Epivir	Haemato Pharm AG	03/01/2012	150 mg	Film-coated tablet	60	EU/1/96/015/001	Austria, Belgium, Cyprus, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/02/2012
Epivir	HAEMATO PHARM AG	20/06/2012	150 mg	Film-coated tablet	60	EU/1/96/015/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	09/07/2012
Esbriet	HAEMATO PHARM AG	21/05/2012	267 mg	Capsule, hard	252 (4 x 63)	EU/1/11/667/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/06/2012
Eucreas	Eureco-Pharma B.V.	24/04/2012	50 mg / 1000 mg	Film-coated tablet	60	EU/1/07/425/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	25/04/2012
Eucreas	Eureco-Pharma B.V.	24/04/2012	50 mg / 850 mg	Film-coated tablet	60	EU/1/07/425/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	25/04/2012
Eucreas	Orifarm AS	17/07/2012	50 mg / 850 mg	Film-coated tablet	180	EU/1/07/425/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/08/2012
Eviplera	CC Pharma	09/11/2012	- -	Film-coated tablet	30	EU/1/11/737/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Eviplera	CC Pharma	09/11/2012	- -	Film-coated tablet	3 x 30	EU/1/11/737/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012

Eviplera	HAEMATO PHARM AG	13/04/2012	- -	Film-coated tablet	30	EU/1/11/737/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/04/2012
Eviplera	Orifarm AS	30/07/2012	- -	Film-coated tablet	30	EU/1/11/737/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/09/2012
Eviplera	Pharma Westen GmbH	13/07/2012	- -	Film-coated tablet	30	EU/1/11/737/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/08/2012
Evista	ABACUS MEDICINE A/S	09/08/2012	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/08/2012
Evista	ABACUS MEDICINE A/S	22/08/2012	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Evista	BB Farma s.r.l.	21/08/2012	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Romania, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	10/09/2012
Evista	BB Farma s.r.l.	21/08/2012	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Romania, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	10/09/2012
Evista	Docpharm GmbH & CoKGaA	18/06/2012	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/07/2012

Evista	Docpharm GmbH & CoKGaA	18/06/2012	60 mg	Film-coated tablet	14	EU/1/98/073/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/07/2012
Evista	Docpharm GmbH & CoKGaA	10/07/2012	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Evista	Galmed a.s.	16/11/2011	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	18/01/2012
Evista	O.P.D. Laboratories Ltd	16/04/2012	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	15/05/2012
Evista	PHARMA LAB	04/10/2012	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	25/10/2012
Evista	PHARMA LAB	04/10/2012	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	25/10/2012
Evoltra	Orifarm AB	30/07/2012	1 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/06/334/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012
Evoltra	Orifarm AB	30/07/2012	1 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/06/334/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012
EVRA	PCO Manufacturing	08/06/2012	- -	Transdermal patch	9	EU/1/02/223/002	United Kingdom	Ireland	05/07/2012

Exelon	2 Care 4	08/06/2012	9.5 mg / 24 hours	Transdermal patch	60 sachets (30 x 2 sachets)	EU/1/98/066/025	Austria, Belgium, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/07/2012
Exelon	ABACUS MEDICINE A/S	28/12/2011	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/01/2012
Exelon	ABACUS MEDICINE A/S	28/12/2011	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/01/2012
Exelon	ABACUS MEDICINE A/S	03/07/2012	9.5 mg / 24 hours	Transdermal patch	60 sachets (30 x 2 sachets)	EU/1/98/066/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/07/2012
Exelon	Amimed Direct Ltd	02/07/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	13/07/2012
Exelon	BB Farma s.r.l.	23/11/2012	1.5 mg	Capsule, hard	56	EU/1/98/066/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	06/12/2012
Exelon	Copharma ApS	25/09/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/10/2012

Exelon	Cross Pharma AB	05/12/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/12/2012
Exelon	Cross Pharma AB	05/12/2012	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/12/2012
Exelon	Cross Pharma AB	05/12/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/12/2012
Exelon	Cross Pharma AB	05/12/2012	9.5 mg / 24 hours	Transdermal patch	90 sachets (30 x 3 sachets)	EU/1/98/066/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/12/2012
Exelon	Emra-Med	12/07/2012	4.6 mg / 24 hours	Transdermal patch	60 (30 x 2) sachets	EU/1/98/066/021	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Exelon	Emra-Med	12/07/2012	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012

Exelon	Ethigen Limited	23/08/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	15/10/2012
Exelon	Ethigen Limited	23/08/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	15/10/2012
Exelon	Forfarm	22/06/2011	1.5 mg	Capsule, hard	28	EU/1/98/066/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	12/11/2012
Exelon	Interport Ltd	27/11/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands	United Kingdom	05/12/2012
Exelon	Interport Ltd	27/11/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands	United Kingdom	05/12/2012
Exelon	kohlpharma GmbH	23/07/2012	4.5 mg	Capsule, hard	56	EU/1/98/066/008	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/07/2012
Exelon	kohlpharma GmbH	23/07/2012	4.5 mg	Capsule, hard	112	EU/1/98/066/009	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/07/2012
Exelon	Medartuum AB	19/10/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012

Exelon	Medartuum AB	19/10/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Exelon	Medartuum AB	24/10/2012	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Exelon	Medartuum AB	24/10/2012	9.5 mg / 24 hours	Transdermal patch	90 sachets (30 x 3 sachets)	EU/1/98/066/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
Exelon	Mediwin Ltd	26/09/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	05/10/2012
Exelon	MPT Pharma	15/08/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	03/09/2012
Exelon	MPT Pharma	15/08/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	03/09/2012
Exelon	O.P.D. Laboratories Ltd	12/08/2010	4.6 mg / 24 hours	Transdermal patch	20 sachets	EU/1/98/066/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/02/2012

Exelon	O.P.D. Laboratories Ltd	12/08/2010	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/02/2012
Exelon	Orifarm AB	01/06/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	25/06/2012
Exelon	Orifarm AB	01/06/2012	9.5 mg / 24 hours	Transdermal patch	90 sachets (30 x 3 sachets)	EU/1/98/066/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	25/06/2012
Exelon	Paranova Läkemedel AB	30/05/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/06/2012
Exelon	Paranova Läkemedel AB	30/05/2012	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/06/2012
Exelon	PHARMACHIM AB	27/11/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	06/12/2012
Exelon	PHARMACHIM AB	27/11/2012	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012
Exelon	PHARMACHIM AB	27/11/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	06/12/2012

Exelon	PHARMACHIM AB	27/11/2012	9.5 mg / 24 hours	Transdermal patch	90 sachets (30 x 3 sachets)	EU/1/98/066/026	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012
Exelon	Polyfarma Ltd	21/09/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	01/10/2012
Exelon	Polyfarma Ltd	21/09/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	01/10/2012
Exelon	STRATHCLYDE PHARMACEUTICALS LTD	27/08/2012	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/08/2012
Exelon	STRATHCLYDE PHARMACEUTICALS LTD	27/08/2012	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/08/2012
Exforge	Clear Pharma Limited	04/07/2012	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/07/2012
Exforge	Clear Pharma Limited	04/07/2012	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/07/2012

Exforge	Docpharm GmbH & CoKGaA	23/04/2012	5 mg/80 mg	Film-coated tablet	28	EU/1/06/370/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/05/2012
Exforge	Docpharm GmbH & CoKGaA	23/04/2012	5 mg/80 mg	Film-coated tablet	98	EU/1/06/370/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/05/2012
Exforge	Dr Fisher Farma BV	24/07/2012	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/10/2012
Exforge	Dr Fisher Farma BV	24/07/2012	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/10/2012
Exforge	Imed Healthcare Ltd	01/08/2012	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	14/08/2012
Exforge	Imed Healthcare Ltd	01/08/2012	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	14/08/2012
Exforge	Polyfarma Ltd	20/09/2012	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	United Kingdom	Ireland	09/10/2012
Exforge	Swingward Ltd TA Medihealth	29/02/2012	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	20/03/2012

Exforge HCT	Dr Fisher Farma BV	18/07/2012	10 mg/ 160 mg/ 25 mg	Film-coated tablet	28	EU/1/09/569/038	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	08/10/2012
Exforge HCT	Dr Fisher Farma BV	18/07/2012	5 mg/ 160 mg/ 25 mg	Film-coated tablet	28	EU/1/09/569/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	08/10/2012
Exforge HCT	Emra-Med	30/10/2012	10 mg/ 160 mg/ 12.5 mg	Film-coated tablet	28	EU/1/09/569/014	Austria, Belgium, Liechtenstein, Luxembourg, The Netherlands	Germany	05/11/2012
Exforge HCT	Emra-Med	30/10/2012	10 mg/ 160 mg/ 25 mg	Film-coated tablet	28	EU/1/09/569/038	Austria, Belgium, Liechtenstein, Luxembourg, The Netherlands	Germany	05/11/2012
Exforge HCT	Emra-Med	30/10/2012	5 mg / 160 mg / 12.5 mg	Film-coated tablet	28	EU/1/09/569/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	Germany	05/11/2012
Exforge HCT	Emra-Med	30/10/2012	5 mg/ 160 mg/ 25 mg	Film-coated tablet	28	EU/1/09/569/026	Austria, Belgium, Liechtenstein, Luxembourg, The Netherlands	Germany	05/11/2012
Exforge HCT	Eureco-Pharma B.V.	27/04/2012	10 mg/ 160 mg/ 25 mg	Film-coated tablet	14	EU/1/09/569/037	Austria, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	07/05/2012
Exforge HCT	Eureco-Pharma B.V.	27/04/2012	5 mg/ 160 mg/ 12.5 mg	Film-coated tablet	14	EU/1/09/569/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	07/05/2012
Exforge HCT	PCO Manufacturing	16/02/2012	10 mg/ 160 mg/ 12.5 mg	Film-coated tablet	30	EU/1/09/569/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	19/03/2012

Exforge HCT	PCO Manufacturing	16/02/2012	5 mg/ 160 mg/ 12.5 mg	Film-coated tablet	30	EU/1/09/569/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	19/03/2012
Exjade	Clear Pharmacy	19/12/2011	250 mg	Dispersible tablet	28	EU/1/06/356/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	06/01/2012
Exjade	Clear Pharmacy	19/12/2011	500 mg	Dispersible tablet	28	EU/1/06/356/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	06/01/2012
Exjade	Orifarm AB	17/07/2012	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/08/2012
Exjade	Orifarm AS	18/07/2012	500 mg	Dispersible tablet	28	EU/1/06/356/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/09/2012
Exjade	Orifarm AS	28/09/2012	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/09/2012
Extavia	Axicorp Pharma B.V.	13/07/2012	250 µg / ml	Powder and solvent for solution for injection	3 x 15 vials + 3 x 15 pre-filled syringes	EU/1/08/454/005	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Lithuania, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/09/2012
Extavia	kohlpharma GmbH	01/02/2012	250 µg / ml	Powder and solvent for solution for injection	3 x 15 vials + 3 x 15 pre-filled syringes	EU/1/08/454/005	Greece, Poland	Germany	01/03/2012
Extavia	Pharma Westen GmbH	08/11/2012	250 µg / ml	Powder and solvent for solution for injection	3 x 15 vials + 3 x 15 pre-filled syringes (luer cone)	EU/1/08/454/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/11/2012

Fampyra	HAEMATO PHARM AG	30/08/2012	10 mg	Prolonged-release tablet	56 (4 bottles of 14)	EU/1/11/699/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Fampyra	HAEMATO PHARM AG	30/08/2012	10 mg	Prolonged-release tablet	28 (2 bottles of 14)	EU/1/11/699/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Faslodex	Beragena Arzneimittel GmbH	22/10/2012	250 mg/5 ml	Solution for injection	1 pre-filled syringe + 1 safety needle	EU/1/03/269/001	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Germany	12/11/2012
Faslodex	Emra-Med	22/11/2012	250 mg/5 ml	Solution for injection	2 pre-filled syringes + 2 safety needles	EU/1/03/269/002	Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	30/11/2012
Faslodex	Eureco-Pharma B.V.	05/10/2012	250 mg/5 ml	Solution for injection	1 pre-filled syringe + 1 safety needle	EU/1/03/269/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	19/10/2012
Faslodex	Inopha GmbH	30/03/2012	250 mg/5 ml	Solution for injection	2 pre-filled syringes + 2 safety needles	EU/1/03/269/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	25/04/2012
Faslodex	kohlpharma GmbH	17/07/2012	250 mg/5 ml	Solution for injection	2 pre-filled syringes + 2 safety needles	EU/1/03/269/002	France	Germany	07/08/2012

Fasturtec	HAEMATO PHARM AG	23/03/2012	1.5 mg/ml	Powder and solvent for concentrate for solution for infusion	3 vials + 3 ampoules	EU/1/00/170/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/04/2012
Fasturtec	HAEMATO PHARM AG	23/03/2012	1.5 mg/ml	Powder and solvent for concentrate for solution for infusion	1 vial + 1 ampoule	EU/1/00/170/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/04/2012
Fendrix	Euro Registratie Collectief BV	03/10/2012	20 µg/0.5 ml	Suspension for injection	1 pre-filled syringe + 1 needle	EU/1/04/299/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/10/2012
Firazyr	Paranova Danmark AS	08/11/2012	30 mg	Solution for injection	1 pre-filled syringe + 1 needle	EU/1/08/461/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/11/2012
Firazyr	Paranova Läkemedel AB	08/11/2012	30 mg	Solution for injection	1 pre-filled syringe + 1 needle	EU/1/08/461/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/11/2012
Firmagon	Euro Registratie Collectief BV	15/02/2012	120 mg	Powder and solvent for solution for injection	2 vials + 2 vials + 2 syringes + 4 needles	EU/1/08/504/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/03/2012
Firmagon	Orifarm AS	17/07/2012	80 mg	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 1 plunger rod + 1 vial adapter + 1 needle	EU/1/08/504/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/08/2012
Forsteo	ABACUS MEDICINE A/S	18/04/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/04/2012

Forsteo	ABACUS MEDICINE A/S	22/08/2012	20 µg/80 µl	Solution for injection	3 pre-filled pens	EU/1/03/247/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/09/2012
Forsteo	ABACUS MEDICINE A/S	22/08/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/09/2012
Forsteo	Copharma ApS	31/05/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/07/2012
Forsteo	Copharma ApS	31/05/2012	20 µg/80 µl	Solution for injection	3 pre-filled pens	EU/1/03/247/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/07/2012
Forsteo	Cross Pharma AB	13/11/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/11/2012
Forsteo	EuroPharma DK ApS	14/02/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/03/2012
Forsteo	HAEMATO PHARM AG	17/07/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/08/2012

Forsteo	Medartuum AB	04/04/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	22/05/2012
Forsteo	NewNeopharm BV	22/06/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/07/2012
Forsteo	Paranova Danmark AS	01/05/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Forsteo	Paranova Läkemedel AB	02/10/2012	20 µg/80 µl	Solution for injection	1 pre-filled pen	EU/1/03/247/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/10/2012
FOSAVANCE	B&S Healthcare	17/11/2011	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	16/02/2012
FOSAVANCE	Imed Healthcare Ltd	16/07/2012	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/07/2012
FOSAVANCE	Mediwin Ltd	12/03/2012	70 mg / 2800 IU	Tablet	4	EU/1/05/310/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	12/04/2012
FOSAVANCE	Profind Wholesale Ltd.	22/08/2012	70 mg / 2800 IU	Tablet	4	EU/1/05/310/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	04/12/2012

Fycompa	Emra-Med	23/10/2012	2 mg	Film-coated tablet	7	EU/1/12/776/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	23/10/2012	4 mg	Film-coated tablet	7	EU/1/12/776/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/11/2012
Fycompa	Emra-Med	23/10/2012	4 mg	Film-coated tablet	28	EU/1/12/776/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	23/10/2012	4 mg	Film-coated tablet	84	EU/1/12/776/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	23/10/2012	6 mg	Film-coated tablet	28	EU/1/12/776/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	23/10/2012	6 mg	Film-coated tablet	84	EU/1/12/776/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	24/10/2012	10 mg	Film-coated tablet	28	EU/1/12/776/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	24/10/2012	10 mg	Film-coated tablet	84	EU/1/12/776/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	24/10/2012	12 mg	Film-coated tablet	28	EU/1/12/776/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012

Fycompa	Emra-Med	24/10/2012	12 mg	Film-coated tablet	84	EU/1/12/776/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	24/10/2012	8 mg	Film-coated tablet	28	EU/1/12/776/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Fycompa	Emra-Med	24/10/2012	8 mg	Film-coated tablet	84	EU/1/12/776/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Galvus	ABACUS MEDICINE A/S	02/11/2012	50 mg	Tablet	112	EU/1/07/414/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/11/2012
Galvus	EuroPharma DK ApS	19/12/2011	50 mg	Tablet	28	EU/1/07/414/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/01/2012
Galvus	EuroPharma DK ApS	19/12/2011	50 mg	Tablet	112	EU/1/07/414/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/01/2012
Galvus	Orifarm AS	20/01/2012	50 mg	Tablet	90	EU/1/07/414/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/03/2012
Ganfort	axicorp Pharma GmbH	20/03/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/04/2012

Ganfort	axicorp Pharma GmbH	20/03/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/04/2012
Ganfort	Ethigen Limited	27/08/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	08/10/2012
Ganfort	Euro Registratie Collectief BV	26/09/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/10/2012
Ganfort	Ginova Ltd	20/11/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	10/12/2012
Ganfort	G-Pharma Ltd	08/03/2012	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/03/2012
Ganfort	Pharma Lab	09/09/2011	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	07/03/2012
Ganfort	Polyfarma Ltd	07/12/2012	300 µg/ml / 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	14/12/2012
Gilenya	CC Pharma	17/12/2012	0.5 mg	Capsule, hard	28	EU/1/11/677/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/12/2012
Glivec	Cambridge Major Laboratories Europe B.V	24/02/2012	400 mg	Film-coated tablet	30	EU/1/01/198/010	Austria, Belgium, Czech Republic, Finland, France, Greece, Ireland, Italy, Lithuania, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/09/2012

Glivec	Cambridge Major Laboratories Europe B.V	24/02/2012	400 mg	Film-coated tablet	90	EU/1/01/198/013	Austria, Belgium, Czech Republic, Finland, France, Greece, Ireland, Italy, Lithuania, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/09/2012
Glivec	Cancernova GmbH onkologische Arzneimittel	07/08/2012	100 mg	Film-coated tablet	60	EU/1/01/198/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/08/2012
Glivec	Cancernova GmbH onkologische Arzneimittel	07/08/2012	400 mg	Film-coated tablet	30	EU/1/01/198/010	Greece	Germany	24/08/2012
Glivec	Dr Fisher Farma BV	20/06/2012	100 mg	Capsule, hard	120	EU/1/01/198/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/09/2012
Glivec	Emra-Med	22/11/2012	100 mg	Film-coated tablet	180	EU/1/01/198/012	Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria	28/11/2012
Glivec	Medartuum AB	31/01/2012	100 mg	Film-coated tablet	60	EU/1/01/198/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/02/2012
Glivec	Medartuum AB	31/01/2012	400 mg	Film-coated tablet	30	EU/1/01/198/010	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/02/2012
Glivec	Orifarm OY	18/07/2012	100 mg	Film-coated tablet	120	EU/1/01/198/011	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	14/08/2012

Glivec	Paranova Läkemedel AB	25/07/2012	100 mg	Film-coated tablet	60	EU/1/01/198/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012
Glivec	Propharmed GmbH	11/06/2012	400 mg	Film-coated tablet	30	EU/1/01/198/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/07/2012
Glivec	Propharmed GmbH	11/06/2012	400 mg	Film-coated tablet	90	EU/1/01/198/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/07/2012
GONAL-f	ABACUS MEDICINE A/S	23/04/2012	300 IU/ 0.5 ml (22 micrograms/ 0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 8 needles	EU/1/95/001/033	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
GONAL-f	ABACUS MEDICINE A/S	23/04/2012	450 IU/ 0.75 ml(33 micrograms/ 0.75 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 12 needles	EU/1/95/001/034	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
GONAL-f	axicorp Pharma GmbH	11/05/2012	300 IU/ 0.5 ml (22 micrograms/ 0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 8 needles	EU/1/95/001/033	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/06/2012
GONAL-f	Copharma ApS	31/05/2012	300 IU/ 0.5 ml (22 micrograms/ 0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 8 needles	EU/1/95/001/033	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/06/2012

GONAL-f	HAEMATO PHARM AG	15/02/2012	300 IU/ 0.5 ml (22 micrograms/ 0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 8 needles	EU/1/95/001/033	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/03/2012
GONAL-f	HAEMATO PHARM AG	15/02/2012	450 IU/ 0.75 ml (33 micrograms/ 0.75 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 7 needles	EU/1/95/001/034	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/04/2012
GONAL-f	HAEMATO PHARM AG	15/02/2012	900 IU/ 1.5 ml (66 micrograms/ 1.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 20 needles	EU/1/95/001/035	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/03/2012
GONAL-f	PHARMACHIM AB	21/05/2012	900 IU/ 1.5 ml (66 micrograms/ 1.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 20 needles	EU/1/95/001/035	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/06/2012
HBVAXPRO	hvd medical GmbH	16/04/2012	40 µg/ml	Suspension for injection	1 vial	EU/1/01/183/015	United Kingdom	Germany	23/04/2012
HBVAXPRO	Pharma Gerke GmbH	26/11/2012	40 µg/ml	Suspension for injection	1 vial	EU/1/01/183/015	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/12/2012
HBVAXPRO	PHARMACHIM AB	30/03/2012	10 µg/ml	Suspension for injection	1 pre-filled syringe with 2 separate needles	EU/1/01/183/028	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/04/2012
HBVAXPRO	PHARMACHIM AB	30/03/2012	5 µg/0.5 ml	Suspension for injection	1 pre-filled syringe with 2 separate needles	EU/1/01/183/024	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/04/2012
Hepsera	axicorp Pharma B.V.	10/12/2012	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/12/2012

Hepsera	axicorp Pharma B.V.	10/12/2012	10 mg	Tablet	3 x 30	EU/1/03/251/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/12/2012
Hepsera	Omnia Läkemedel AB	06/12/2011	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	20/01/2012
Herceptin	Cambridge Major Laboratories Europe B.V	12/03/2012	150 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/00/145/001	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/09/2012
Herceptin	Propharmed GmbH	11/06/2012	150 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/00/145/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/07/2012
Hirobriz Breezhaler	CC Pharma	28/03/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/594/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/04/2012
Hirobriz Breezhaler	CC Pharma	28/03/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/594/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/04/2012
Humalog	Eureco-Pharma B.V.	04/05/2012	100 U/ml	Solution for injection	5 cartridges	EU/1/96/007/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	24/05/2012
Humalog	EuroPharma DK ApS	11/09/2012	100 U/ml	Solution for injection	5 pens	EU/1/96/007/031	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/09/2012
Humalog	Ginova Ltd	18/09/2012	100 U/ml	Suspension for injection	5 cartridges	EU/1/96/007/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	16/10/2012

Humalog	MPT Pharma	31/07/2012	100 U/ml	Suspension for injection	5 cartridges	EU/1/96/007/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	21/08/2012
Humalog	MPT Pharma	31/07/2012	100 U/ml	Suspension for injection	5 cartridges	EU/1/96/007/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	21/08/2012
Humalog	MPT Pharma	31/07/2012	100 U/ml	Solution for injection	5 pens	EU/1/96/007/031	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	21/08/2012
Humalog	MPT Pharma	31/07/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/033	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	21/08/2012
Humalog	MPT Pharma	31/07/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/035	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	20/08/2012
Humalog	MPT Pharma	10/09/2012	100 U/ml	Solution for injection	5 cartridges	EU/1/96/007/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	20/09/2012
Humalog	O.P.D. Laboratories Ltd	07/09/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/035	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	10/10/2012

Humalog	O.P.D. Laboratories Ltd	07/09/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/033	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	03/10/2012
Humalog KwikPen	Ginova Ltd	28/06/2012	100 U/ml	Solution for injection	5 pens	EU/1/96/007/031	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	17/07/2012
Humalog Mix25 KwikPen	Ginova Ltd	28/06/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/033	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	17/07/2012
Humalog Mix50 KwikPen	Ginova Ltd	28/06/2012	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/035	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	17/07/2012
Humira	cambridge Major Laboratories Europe B.V	12/03/2012	40 mg	Solution for injection	6 pre-filled pens + 6 alcohol pads in a blister	EU/1/03/256/010	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/09/2012
Humira	Cambridge Major Laboratories Europe B.V	12/03/2012	40 mg	Solution for injection	2 pre-filled pens + 2 alcohol pads in a blister	EU/1/03/256/008	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/09/2012
Humira	kohlpharma GmbH	29/10/2012	40 mg/0.8 ml	Solution for injection	2 carton pack. Each carton contains: 1 vial + 1 syringe + 1 sterile adapter + 2 alcohol pads	EU/1/03/256/001	Austria, France	Germany	03/12/2012
Humira	propharmed GmbH	20/07/2012	40 mg	Solution for injection	2 pre-filled syringes + 2 alcohol pads	EU/1/03/256/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012
Humira	propharmed GmbH	20/07/2012	40 mg	Solution for injection	6 pre-filled syringes + 6 alcohol pads	EU/1/03/256/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012

Humira	propharmed GmbH	20/07/2012	40 mg	Solution for injection	2 pre-filled pens + 2 alcohol pads in a blister	EU/1/03/256/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012
Humira	propharmed GmbH	20/07/2012	40 mg	Solution for injection	6 pre-filled pens + 6 alcohol pads in a blister	EU/1/03/256/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012
Humira	Veron Pharma Vertriebs GmbH	27/09/2012	40 mg	Solution for injection	6 pre-filled syringes + 6 alcohol pads	EU/1/03/256/005	Austria	Germany	23/10/2012
Hycamtin	ABACUS MEDICINE A/S	08/08/2012	4 mg	Powder for concentrate for solution for infusion	1 vial (17 ml)	EU/1/96/027/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/08/2012
Hycamtin	Paranova Oy	31/08/2012	1 mg	Capsule, hard	10	EU/1/96/027/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	17/09/2012
INCIVO	Haemato Pharm AG	12/04/2012	375 mg	Film-coated tablet	168 (4 x 42)	EU/1/11/720/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/06/2012
INCIVO	Orifarm AB	21/08/2012	375 mg	Film-coated tablet	1 x 42	EU/1/11/720/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012
INCIVO	Orifarm AB	21/08/2012	375 mg	Film-coated tablet	168 (4 x 42)	EU/1/11/720/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012
INCIVO	Orifarm AS	08/08/2012	375 mg	Film-coated tablet	1 x 42	EU/1/11/720/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/08/2012

Infanrix hexa	Emra-Med	28/06/2012	- -	Powder and suspension for suspension for injection	10 vials + 10 pre-filled syringes + 20 needles	EU/1/00/152/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/07/2012
Infanrix hexa	Emra-Med	05/07/2012	- -	Powder and suspension for suspension for injection	20 vials + 20 pre-filled syringes + 40 needles	EU/1/00/152/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Infanrix hexa	Emra-Med	05/07/2012	- -	Powder and suspension for suspension for injection	50 vials + 50 pre-filled syringes + 100 needles	EU/1/00/152/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Inovelon	Amimed Direct Ltd	01/11/2012	100 mg	Film-coated tablet	10	EU/1/06/378/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	07/11/2012
Inovelon	Orifarm AB	12/01/2012	400 mg	Film-coated tablet	60	EU/1/06/378/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/02/2012
Inovelon	Paranova Danmark AS	04/06/2012	400 mg	Film-coated tablet	100	EU/1/06/378/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/07/2012
Instanyl	Dr Fisher Farma BV	21/11/2012	100 micrograms / dose	Nasal spray, solution	1 bottle (20 doses)	EU/1/09/531/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/12/2012
Instanyl	Dr Fisher Farma BV	21/11/2012	200 micrograms / dose	Nasal spray, solution	1 bottle (20 doses)	EU/1/09/531/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/12/2012

Instanyl	Dr Fisher Farma BV	21/11/2012	50 micrograms / dose	Nasal spray, solution	1 bottle (20 doses)	EU/1/09/531/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/12/2012
Insulatard	Dr Fisher Farma BV	09/03/2012	100 IU/ml	Suspension for injection	5 cartridges	EU/1/02/233/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	27/03/2012
Insulatard	Emra-Med	14/05/2012	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/02/233/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012
Insulatard	Emra-Med	14/05/2012	100 IU/ml	Suspension for injection	10 pre-filled pens	EU/1/02/233/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012
Insulatard	O.P.D. Laboratories Ltd	17/10/2012	100 IU/ml	Suspension for injection	5 cartridges	EU/1/02/233/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	23/10/2012
Insulatard Penfill	Eureco-Pharma B.V.	15/05/2012	100 IU/ml	Suspension for injection	10 cartridges	EU/1/02/233/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	28/06/2012
Insuman	2 Care 4	02/08/2012	100 IU/ml	Solution for injection	5 pre-filled pens	EU/1/97/030/142	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012
Insuman	axicorp Pharma GmbH	10/12/2012	100 IU/ml	Solution for injection	5 pre-filled pens	EU/1/97/030/142	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/12/2012
Insuman	CC Pharma	19/10/2012	100 IU/ml	Solution for injection	10 pre-filled pens	EU/1/97/030/145	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/10/2012

Insuman	CC Pharma	19/10/2012	100 IU/ml	Solution for injection	5 pre-filled pens	EU/1/97/030/142	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/10/2012
Insuman	Emra-Med	10/05/2012	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/97/030/148	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/06/2012
Insuman	Emra-Med	10/05/2012	100 IU/ml	Suspension for injection	10 pre-filled pens	EU/1/97/030/151	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/06/2012
Insuman	Paranova Danmark AS	14/11/2012	100 IU/ml	Solution for injection	5 pre-filled pens	EU/1/97/030/142	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/11/2012
Insuman	Pharma Westen GmbH	21/12/2011	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/064	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Insuman	Pharma Westen GmbH	21/12/2011	100 IU/ml	Solution for injection	10 pre-filled pens	EU/1/97/030/145	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012
Insuman Basal	axicorp Pharma GmbH	22/06/2012	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/058	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Insuman Basal	kohlpharma GmbH	26/04/2012	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/97/030/148	Portugal	Germany	12/06/2012
Insuman Basal	kohlpharma GmbH	26/04/2012	100 IU/ml	Suspension for injection	10 pre-filled pens	EU/1/97/030/151	Portugal	Germany	12/06/2012

Insuman Basal	Pharma Westen GmbH	22/11/2011	100 IU/ml	SoloStar, Suspension for injection	10 pre-filled pens	EU/1/97/030/151	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/01/2012
Insuman Comb 25	Emra-Med	16/02/2012	100 IU/ml	SoloStar, Suspension for injection	5 pre-filled pens	EU/1/97/030/160	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012
Insuman Comb 25	Emra-Med	16/02/2012	100 IU/ml	SoloStar, Suspension for injection	10 pre-filled pens	EU/1/97/030/163	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012
Insuman Comb 25	Emra-Med	01/03/2012	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/062	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012
Insuman comb 25	kohlpharma GmbH	30/04/2012	100 IU/ml	Solostar, suspension for injection	5 pre-filled pens	EU/1/97/030/160	Portugal	Germany	05/06/2012
Insuman comb 25	kohlpharma GmbH	30/04/2012	100 IU/ml	Solostar, suspension for injection	10 pre-filled pens	EU/1/97/030/163	Portugal	Germany	05/06/2012
Insuman Comb 50	Emra-Med	13/12/2011	100 IU/ml	OptiSet, suspension for injection	5 pens	EU/1/97/030/083	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/01/2012
Insuman Comb 50	Emra-Med	13/12/2011	100 IU/ml	OptiSet, suspension for injection	10 pens	EU/1/97/030/084	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/01/2012

Insuman Comb 50	Emra-Med	01/03/2012	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/064	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012
Insuman Rapid	kohlpharma GmbH	19/01/2012	100 IU/ml	SoloStar, Solution for injection	5 pre-filled pens	EU/1/97/030/142	Italy	Germany	07/03/2012
Insuman Rapid	kohlpharma GmbH	25/01/2012	100 IU/ml	SoloStar, Solution for injection	10 pre-filled pens	EU/1/97/030/145	Italy	Germany	07/03/2012
Insuman Rapid (Solostar)	Beragena Arzneimittel GmbH	29/11/2012	100 IU/ml	Solution for injection	5 pre-filled pens	EU/1/97/030/142	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	11/12/2012
Insuman Rapid (Solostar)	Beragena Arzneimittel GmbH	29/11/2012	100 IU/ml	Solution for injection	10 pre-filled pens	EU/1/97/030/145	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	11/12/2012
Insuman Rapid Solostar	axicorp Pharma GmbH	24/08/2012	100 IU/ml	Solution for injection	10 pre-filled pens, SoloStar	EU/1/97/030/145	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012
Integrilin	Delfarma SpZoo	31/01/2012	2 mg/ml	Solution for injection	1 vial	EU/1/99/109/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	01/02/2012
INTELENCE	Dr Fisher Farma BV	20/01/2012	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/02/2012
INTELENCE	Medcor Pharmaceuticals BV	02/05/2012	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	10/09/2012
INTELENCE	Medicopharm AG	31/07/2012	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/08/2012

INTELENCE	Paranova Danmark AS	19/01/2012	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/01/2012
INTELENCE	Pharma Westen GmbH	24/07/2012	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/08/2012
IntronA	ACA Müller ADAG Pharma AG	04/12/2012	18 million IU	Solution for injection	2 pens + 24 injection sets	EU/1/99/127/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/12/2012
IntronA	CC Pharma	20/09/2012	18 million IU	Solution for injection or infusion	2 vials	EU/1/99/127/025	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
IntronA	Orifarm AS	04/10/2012	30 million IU	Solution for injection	1 pen + 12 injection sets	EU/1/99/127/034	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/12/2012
Invanz	Paranova Läkemedel AB	09/10/2012	1 g	Powder for concentrate for solution for infusion	1 vial	EU/1/02/216/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Invanz	Paranova Läkemedel AB	09/10/2012	1 g	Powder for concentrate for solution for infusion	10 vials	EU/1/02/216/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Invanz	PHARMACHIM AB	09/03/2012	1 g	Powder for concentrate for solution for infusion	1 vial	EU/1/02/216/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	02/04/2012
Invega	B&S Healthcare	18/01/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	21/02/2012

Invega	Copharma ApS	18/07/2012	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/09/2012
Invega	Copharma ApS	25/09/2012	3 mg	Prolonged-release tablet	28	EU/1/07/395/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/10/2012
Invega	Copharma ApS	25/09/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/10/2012
Invega	Dr Fisher Farma BV	05/01/2012	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/03/2012
Invega	Dr Fisher Farma BV	23/02/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	07/03/2012
Invega	Medartuum AB	30/11/2012	3 mg	Prolonged-release tablet	28	EU/1/07/395/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
Invega	Medartuum AB	30/11/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
Invega	Paranova Danmark AS	13/03/2012	3 mg	Prolonged-release tablet	28	EU/1/07/395/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/04/2012

Invega	Paranova Danmark AS	13/03/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/04/2012
Invega	Paranova Danmark AS	13/03/2012	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/04/2012
Invega	Polyfarma Ltd	21/09/2012	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	08/10/2012
Invega	Polyfarma Ltd	21/09/2012	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	08/10/2012
Invirase	Haemato Pharm AG	22/06/2012	500 mg	Film-coated tablet	1 bottle	EU/1/96/026/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	Ireland, United Kingdom	04/07/2012
Invirase	Medcor Pharma-ceuticals BV	06/07/2012	500 mg	Film-coated tablet	1 bottle	EU/1/96/026/002	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, United Kingdom	The Netherlands	28/08/2012
Invirase	REMEDIX GmbH	14/03/2012	500 mg	Film-coated tablet	1 bottle	EU/1/96/026/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	02/05/2012
Iressa	Cancernova GmbH onkologische Arzneimittel	07/08/2012	250 mg	Film-coated tablet	30	EU/1/09/526/002	Lithuania	Germany	16/08/2012

Ixiaro	Orifarm AB	18/07/2012	6 µg	Suspension for injection	1 pre-filled syringe	EU/1/08/501/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/08/2012
Ixiaro	Orifarm AS	13/01/2012	6 µg	Suspension for injection	1 pre-filled syringe	EU/1/08/501/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/02/2012
Ixiaro	Orifarm OY	15/02/2012	6 µg	Suspension for injection	1 pre-filled syringe	EU/1/08/501/002	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	02/03/2012
Ixiaro	Paranova Danmark AS	30/01/2012	6 µg	Suspension for injection	1 pre-filled syringe	EU/1/08/501/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/02/2012
Janumet	2 Care 4	29/11/2011	50 mg / 850 mg	Film-coated tablet	56	EU/1/08/455/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/01/2012
Janumet	2 Care 4	29/11/2011	50 mg / 850 mg	Film-coated tablet	196	EU/1/08/455/006	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/01/2012
Janumet	B&S Healthcare	29/10/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	31/10/2012
Janumet	COPHARMA ApS	10/02/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/03/2012
Janumet	Cross Pharma AB	04/09/2012	50 mg / 1000 mg	Film-coated tablet	196	EU/1/08/455/013	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/09/2012

Janumet	Doncaster Pharmaceuticals Group Ltd	09/12/2011	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	27/02/2012
Janumet	Interport Ltd	26/11/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands	United Kingdom	30/11/2012
Janumet	Medartuum AB	06/11/2012	50 mg / 850 mg	Film-coated tablet	196 (2 x 98)	EU/1/08/455/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/11/2012
Janumet	O.P.D. Laboratories Ltd	02/04/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	10/05/2012
Janumet	Orifarm AS	27/07/2012	50 mg / 850 mg	Film-coated tablet	196	EU/1/08/455/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/08/2012
Janumet	Orifarm AS	30/08/2012	50 mg / 850 mg	Film-coated tablet	56	EU/1/08/455/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/09/2012
Janumet	Paranova Läkemedel AB	21/02/2012	50 mg / 850 mg	Film-coated tablet	56	EU/1/08/455/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	27/03/2012
Janumet	Paranova Läkemedel AB	21/02/2012	50 mg / 850 mg	Film-coated tablet	196 (2 x 98)	EU/1/08/455/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	27/03/2012

Janumet	PCO Manufacturing	24/05/2012	50 mg / 1000 mg	Film-coated tablet	28	EU/1/08/455/009	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Romania, Spain, Sweden, The Netherlands	Ireland, United Kingdom	29/05/2012
Janumet	PCO Manufacturing	24/05/2012	50 mg / 850 mg	Film-coated tablet	28	EU/1/08/455/002	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Romania, Spain, Sweden, The Netherlands	Ireland, United Kingdom	29/05/2012
Janumet	Swingward Ltd TA Medihealth	23/08/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	04/10/2012
Janumet	WAYMADE PLC	22/06/2012	50 mg / 1000 mg	Film-coated tablet	56	EU/1/08/455/010	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	02/07/2012
Januvia	COPHARMA ApS	18/01/2012	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/02/2012
Januvia	Interport Ltd	26/11/2012	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands	United Kingdom	30/11/2012
Januvia	Medcor Pharma-ceuticals BV	29/11/2012	25 mg	Film-coated tablet	28	EU/1/07/383/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/12/2012
Januvia	Mediwin Ltd	31/10/2012	100 mg	Film-coated tablet	28	EU/1/07/383/014	France	Ireland, Romania, United Kingdom	08/11/2012
Januvia	Polyfarma Ltd	07/06/2012	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	21/06/2012

Javlor	HAEMATO PHARM AG	10/07/2012	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/09/550/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Javlor	HAEMATO PHARM AG	10/07/2012	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/09/550/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Jevtana	HAEMATO PHARM AG	09/02/2012	60 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/11/676/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Jevtana	Orifarm AB	11/12/2012	60 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/11/676/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Kaletra	Docpharm GmbH & CoKGaA	25/06/2012	200 mg / 50 mg	Film-coated tablet	120 (10 x 12)	EU/1/01/172/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/09/2012
Kaletra	Docpharm GmbH & CoKGaA	10/10/2012	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/10/2012
Kaletra	Orifarm OY	15/02/2012	200 mg / 50 mg	Film-coated tablet	120 (10 x 12)	EU/1/01/172/008	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	09/03/2012
Karvea	B&S Healthcare	15/05/2012	75 mg	Film-coated tablet	28	EU/1/97/049/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	08/06/2012

Karvea	Doncaster Pharma-ceuticals Group Ltd	07/03/2012	150 mg	Film-coated tablet	28	EU/1/97/049/022	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	10/04/2012
Karvea	Doncaster Pharma-ceuticals Group Ltd	07/03/2012	300 mg	Film-coated tablet	28	EU/1/97/049/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	10/04/2012
Karvea	Doncaster Pharma-ceuticals Group Ltd	07/03/2012	75 mg	Film-coated tablet	28	EU/1/97/049/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	10/04/2012
Karvea	Emra-Med	03/05/2012	150 mg	Film-coated tablet	98	EU/1/97/049/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Karvea	Emra-Med	03/05/2012	150 mg	Film-coated tablet	56	EU/1/97/049/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Karvea	Emra-Med	03/05/2012	150 mg	Film-coated tablet	28	EU/1/97/049/022	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Karvea	kohlpharma GmbH	14/05/2012	300 mg	Film-coated tablet	98	EU/1/97/049/030	Spain	Germany	08/06/2012
Karvea	kohlpharma GmbH	14/06/2012	150 mg	Film-coated tablet	98	EU/1/97/049/025	Spain	Germany	05/07/2012
Karvea	kohlpharma GmbH	14/06/2012	75 mg	Film-coated tablet	98	EU/1/97/049/020	Spain	Germany	05/07/2012
Karvezide	Cambridge Major Laboratories Europe B.V	21/06/2012	150 mg/ 12.50 mg	Tablet	98	EU/1/98/085/003	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/10/2012
Karvezide	Doncaster Pharmaceuticals Group Ltd	07/03/2012	150 mg/ 12.50 mg	Film-coated tablet	28	EU/1/98/085/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	12/04/2012

Karvezide	kohlpharma GmbH	10/01/2012	150 mg/ 12.50 mg	Film-coated tablet	98	EU/1/98/085/015	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012
Karvezide	kohlpharma GmbH	10/01/2012	300 mg/ 12.50 mg	Film-coated tablet	98	EU/1/98/085/020	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Keppra	2 Care 4	14/12/2011	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, The Netherlands, United Kingdom	Sweden	09/03/2012
Keppra	Adripharma GmbH	20/08/2012	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/10/2012
Keppra	Adripharma GmbH	23/08/2012	1000 mg	Film-coated tablet	50	EU/1/00/146/023	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/11/2012
Keppra	Amimed Direct Ltd	08/08/2012	100 mg/ml	Oral solution	1 bottle + 1 oral syringe 10 ml	EU/1/00/146/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	22/08/2012
Keppra	BB Farma s.r.l.	23/11/2011	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012
Keppra	BB Farma s.r.l.	23/11/2011	500 mg	Film-coated tablet	100	EU/1/00/146/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012

Keppra	BB Farma s.r.l.	23/11/2011	500 mg	Film-coated tablet	200	EU/1/00/146/013	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/01/2012
Keppra	BB Farma s.r.l.	16/12/2011	1000 mg	Film-coated tablet	50	EU/1/00/146/023	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/02/2012
Keppra	BB Farma s.r.l.	16/12/2011	1000 mg	Film-coated tablet	100	EU/1/00/146/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/02/2012
Keppra	BB Farma s.r.l.	16/12/2011	1000 mg	Film-coated tablet	200	EU/1/00/146/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/02/2012
Keppra	Cross Pharma AB	05/06/2012	250 mg	Film-coated tablet	50	EU/1/00/146/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/07/2012
Keppra	Cross Pharma AB	05/06/2012	250 mg	Film-coated tablet	100	EU/1/00/146/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/07/2012

Keppra	CROSS PHARMA AB	15/06/2012	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/06/2012
Keppra	Dr Fisher Farma BV	15/11/2012	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	26/11/2012
Keppra	Imed Healthcare Ltd	05/06/2012	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/06/2012
Keppra	Kosei Pharma UK Limited	22/11/2012	1000 mg	Film-coated tablet	30	EU/1/00/146/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	26/11/2012
Keppra	Kosei Pharma UK Limited	22/11/2012	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	26/11/2012
Keppra	Medartuum AB	18/07/2012	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	06/09/2012

Keppra	Medartuum AB	08/11/2012	100 mg/ml	Concentrate for solution for infusion	10 vials	EU/1/00/146/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/11/2012
Keppra	Mediwin Ltd	29/06/2012	1000 mg	Film-coated tablet	60	EU/1/00/146/024	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	11/07/2012
Keppra	Mediwin Ltd	29/06/2012	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	11/07/2012
Keppra	MPT Pharma	18/01/2012	100 mg/ml	Oral solution	1 bottle + 1 oral syringe 10 ml	EU/1/00/146/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	30/01/2012
Keppra	MPT Pharma	16/08/2012	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	11/09/2012
Keppra	MPT Pharma	16/08/2012	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	11/09/2012
Keppra	MPT Pharma	16/08/2012	500 mg	Film-coated tablet	100	EU/1/00/146/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	11/09/2012

Keppra	Orifarm AS	03/07/2012	100 mg/ml	Concentrate for solution for infusion	10 vials	EU/1/00/146/030	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/07/2012
Keppra	Paranova Danmark AS	25/07/2012	100 mg/ml	Concentrate for solution for infusion	10 vials	EU/1/00/146/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/09/2012
Keppra	Paranova Läkemedel AB	02/12/2011	1000 mg	Film-coated tablet	200	EU/1/00/146/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/01/2012
Keppra	Paranova Läkemedel AB	02/12/2011	1000 mg	Film-coated tablet	100	EU/1/00/146/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/01/2012
Keppra	Paranova Läkemedel AB	31/05/2012	100 mg/ml	Oral solution	1 bottle + 1 oral syringe 10 ml	EU/1/00/146/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/06/2012
Keppra	Pharmachim AB	20/12/2011	750 mg	Film-coated tablet	50	EU/1/00/146/016	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/01/2012
Keppra	Pharmachim AB	20/12/2011	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Bulgaria, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/01/2012

Keppra	STRATHCLYDE PHARMACEUTICALS LTD	20/11/2012	250 mg	Film-coated tablet	50	EU/1/00/146/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/11/2012
Keppra	STRATHCLYDE PHARMACEUTICALS LTD	20/11/2012	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/11/2012
Kineret	Dr Fisher Farma BV	17/09/2012	100 mg	Solution for injection	7 pre-filled syringes	EU/1/02/203/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	24/10/2012
Kiovig	CC Pharma	23/03/2012	100 mg/ml	Solution for infusion	1 vial of 10 g/ 100 ml	EU/1/05/329/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2012
Kiovig	CC Pharma	23/03/2012	100 mg/ml	Solution for infusion	1 vial of 20 g/ 200 ml	EU/1/05/329/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2012
Kiovig	CC Pharma	30/04/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Kiovig	CC Pharma	30/04/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Kiovig	kohlpharma GmbH	01/08/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012
Kiovig	kohlpharma GmbH	01/08/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/08/2012

Kiovig	kohlpharma GmbH	01/08/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012
Kiovig	Medcor Pharmaceuticals BV	08/03/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/05/329/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/03/2012
Kivexa	ABACUS MEDICINE A/S	12/11/2012	- -	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Kivexa	ABACUS MEDICINE A/S	16/11/2012	- -	Film-coated tablet	3 x 30	EU/1/04/298/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/11/2012
Kivexa	ABACUS MEDICINE A/S	29/11/2012	- -	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012
Kivexa	REMEDIX GmbH	14/03/2012	- -	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	02/05/2012
Kivexa	Veron Pharma Vertriebs GmbH	30/08/2012	- -	Film-coated tablet	3 x 30	EU/1/04/298/003	Austria	Germany	14/09/2012
KOGENATE Bayer	Omnia Läkemedel AB	12/04/2012	1000 IU	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 1 vial adapter	EU/1/00/143/009	Austria, Belgium, Bulgaria, Czech Republic, France, Germany, Greece, Ireland, Italy, Lithuania, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	25/04/2012

KOGENATE Bayer	Omnia Läkemedel AB	12/04/2012	2000 IU	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 1 vial adapter**	EU/1/00/143/011	Austria, Belgium, Bulgaria, Czech Republic, France, Germany, Greece, Ireland, Italy, Lithuania, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	25/04/2012
Lantus	ABACUS MEDICINE A/S	13/08/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/09/2012
Lantus	Empower Pharma s.r.o.	19/10/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Czech Republic	19/11/2012
Lantus	EU-Pharma BV	28/02/2012	100 Units/ml	Solution for injection in cartridge	5 cartridges	EU/1/00/134/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/03/2012
Lantus	Paranova Läkemedel AB	21/06/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/07/2012
Lantus	Paranova Läkemedel AB	08/11/2012	100 Units/ml	Solution for injection	5 cartridges	EU/1/00/134/025	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/11/2012
Lantus	PCO Manufacturing	15/02/2012	100 Units/ml	Solution for injection in cartridge	5 cartridges	EU/1/00/134/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	09/03/2012
Lantus	WAYMADE PLC	10/08/2012	100 Units/ml	Solution for injection	5 pre-filled pens (SoloStar)	EU/1/00/134/033	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/08/2012

Lantus cartridge	Medartuum AB	23/10/2012	100 Units/ml	Solution for injection in cartridge	5 cartridges	EU/1/00/134/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/11/2012
Lantus Opticlick	PHARMACHIM AB	30/08/2012	100 Units/ml	Solution for injection	5 cartridges for OptiClik	EU/1/00/134/025	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012
Lantus SoloStar	ABACUS MEDICINE A/S	12/12/2011	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/01/2012
Lantus SoloStar	EU-Pharma BV	20/02/2012	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/03/2012
Leflunomide ratiopharm	isopharm GmbH	14/05/2012	20 mg	Film-coated tablet	100	EU/1/10/654/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/07/2012
Levemir	ABACUS MEDICINE A/S	12/10/2012	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/04/278/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/10/2012
Levemir	EU-Pharma BV	14/06/2012	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/04/278/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	14/08/2012
Levemir Flexpen	Paranova Läkemedel AB	05/07/2012	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/04/278/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012

Levetiracetam Actavis	Ebb Medical AB	15/11/2012	250 mg	Film-coated tablet	100	EU/1/11/713/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levetiracetam Actavis	Ebb Medical AB	15/11/2012	250 mg	Film-coated tablet	100	EU/1/11/713/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levetiracetam Actavis	Ebb Medical AB	15/11/2012	500 mg	Film-coated tablet	100	EU/1/11/713/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levetiracetam Actavis	Ebb Medical AB	15/11/2012	500 mg	Film-coated tablet	200	EU/1/11/713/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levetiracetam Actavis	Ebb Medical AB	15/11/2012	500 mg	Film-coated tablet	100	EU/1/11/713/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levetiracetam Actavis	Ebb Medical AB	15/11/2012	500 mg	Film-coated tablet	200	EU/1/11/713/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/12/2012
Levitra	4pharma S.r.l.	14/05/2012	20 mg	Film-coated tablet	4	EU/1/03/248/010	Bulgaria, Romania	Italy	12/06/2012
Levitra	Elpis Ltd.	28/08/2012	20 mg	Film-coated tablet	4	EU/1/03/248/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Latvia	04/09/2012

Levitra	Impexeco S.A.	24/09/2012	20 mg	Film-coated tablet	4	EU/1/03/248/010	Austria, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium	15/10/2012
Levitra	Impexeco S.A.	24/09/2012	20 mg	Film-coated tablet	12	EU/1/03/248/012	Austria, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium	15/10/2012
Levitra	Medartuum AB	13/09/2012	20 mg	Film-coated tablet	12	EU/1/03/248/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	25/09/2012
Litak	ABACUS MEDICINE A/S	08/08/2012	2 mg/ml	Solution for injection	5 vials	EU/1/04/275/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012
Lumigan	Doncaster Pharmaceuticals Group Ltd	13/01/2012	0.1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/205/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	02/03/2012
Lumigan	O.P.D. Laboratories Ltd	19/10/2012	0.3 mg/ml	Eye drops, solution	1 bottle	EU/1/02/205/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	30/10/2012
Lumigan	Paranova Oy	31/05/2012	0.3 mg/ml	Eye drops, solution	3 bottles	EU/1/02/205/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	03/07/2012
Lyrica	axicorp Pharma GmbH	02/11/2012	50 mg	Capsule, hard	21	EU/1/04/279/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/11/2012

Lyrica	BR Pharma International Ltd	14/09/2012	300 mg	Capsule, hard	56	EU/1/04/279/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, Malta, United Kingdom	08/10/2012
Lyrica	Chemilines Ltd	10/04/2012	100 mg	Capsule, hard	84	EU/1/04/279/015	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands	Ireland, United Kingdom	30/05/2012
Lyrica	Crystal Pharma Ltd	25/05/2012	100 mg	Capsule, hard	84	EU/1/04/279/015	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	22/08/2012
Lyrica	Crystal Pharma Ltd	08/06/2012	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	22/08/2012
Lyrica	Crystal Pharma Ltd	07/09/2012	50 mg	Capsule, hard	84	EU/1/04/279/009	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	05/10/2012
Lyrica	Doncaster Pharmaceuticals Group Ltd	29/09/2011	50 mg	Capsule, hard	56	EU/1/04/279/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	04/01/2012
Lyrica	Emra-Med	24/08/2012	100 mg	Capsule, hard	100	EU/1/04/279/039	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Lyrica	hvd medical GmbH	06/03/2012	50 mg	Capsule, hard	21	EU/1/04/279/007	Austria	Germany	05/04/2012
Lyrica	hvd medical GmbH	05/09/2012	75 mg	Capsule, hard	100 x 1	EU/1/04/279/013	Norway	Germany	08/10/2012

Lyrica	hvd medical GmbH	08/11/2012	75 mg	Capsule, hard	100	EU/1/04/279/038	Austria, Belgium, Finland, Greece, Italy, Slovak Republic, The Netherlands	Germany	16/11/2012
Lyrica	Kohlpharma GmbH	31/07/2008	25 mg	Capsule, hard	100	EU/1/04/279/036	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Lyrica	Kohlpharma GmbH	23/11/2010	100 mg	Capsule, hard	100	EU/1/04/279/039	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2012
Lyrica	Kohlpharma GmbH	23/11/2010	150 mg	Capsule, hard	100	EU/1/04/279/040	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Lyrica	kohlpharma GmbH	27/01/2012	225 mg	Capsule, hard	100	EU/1/04/279/042	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/04/2012
Lyrica	Kosei Pharma UK Limited	23/03/2012	200 mg	Capsule, hard	21	EU/1/04/279/020	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	07/05/2012
Lyrica	Kosei Pharma UK Limited	16/04/2012	50 mg	Capsule, hard	56	EU/1/04/279/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	07/05/2012
Lyrica	Medartuum AB	03/07/2012	225 mg	Capsule, hard	56	EU/1/04/279/034	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/07/2012
Lyrica	Medcor Pharmaceuticals BV	19/10/2012	75 mg	Capsule, hard	112 (2 x 56)	EU/1/04/279/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/12/2012

Lyrica	Mediwin Ltd	13/01/2012	50 mg	Capsule, hard	21	EU/1/04/279/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Latvia, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	16/02/2012
Lyrica	Orifarm AB	17/07/2012	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	07/09/2012
Lyrica	Paranova Oy	21/05/2012	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	27/06/2012
Lyrica	PCO Manufacturing	15/02/2012	100 mg	Capsule, hard	21	EU/1/04/279/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	09/03/2012
Lyrica	PCO Manufacturing	07/03/2012	150 mg	Capsule, hard	14	EU/1/04/279/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	11/04/2012
Lyrica	Pharma Gerke GmbH	07/06/2012	300 mg	Capsule, hard	56	EU/1/04/279/024	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2012
Lyrica	Tisida GmbH	02/07/2012	75 mg	Capsule, hard	100	EU/1/04/279/038	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012
Lyrica	Tisida GmbH	02/07/2012	75 mg	Capsule, hard	56	EU/1/04/279/012	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012
Lyrica	Tisida GmbH	04/07/2012	150 mg	Capsule, hard	100	EU/1/04/279/040	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012

Lyrica	Tisida GmbH	04/07/2012	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012
Lyrica	Tisida GmbH	04/07/2012	300 mg	Capsule, hard	56	EU/1/04/279/024	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012
Lyrica	Tisida GmbH	04/07/2012	300 mg	Capsule, hard	100	EU/1/04/279/043	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/07/2012
Lyrica	WAYMADE PLC	15/05/2012	300 mg	Capsule, hard	56	EU/1/04/279/024	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	29/06/2012
MabCampath	Delfarma SpZoo	30/01/2012	30 mg/ml	Concentrate for solution for infusion	3 vials	EU/1/01/193/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	27/04/2012
MabCampath	Inopha GmbH	16/01/2012	30 mg/ml	Concentrate for solution for infusion	3 vials	EU/1/01/193/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	13/02/2012
Mabthera	ABACUS MEDICINE A/S	04/09/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/09/2012

Mabthera	axicorp Pharma B.V.	24/10/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/10/2012
Mabthera	Cambridge Major Laboratories Europe B.V	05/03/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/09/2012
Mabthera	Cancernova GmbH onkologische Arzneimittel	28/03/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Greece	Germany	10/04/2012
Mabthera	Cancernova GmbH onkologische Arzneimittel	28/03/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Greece	Germany	09/04/2012
Mabthera	Haemato Pharm AG	19/07/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
Mabthera	Haemato Pharm AG	19/07/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
Mabthera	Medartuum AB	18/07/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/09/2012
Mabthera	Medartuum AB	04/10/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012
Mabthera	MedicoPharm AG	24/02/2010	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012

Mabthera	Orifarm AB	30/04/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/06/2012
Mabthera	Orifarm AB	30/04/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/06/2012
Mabthera	Paranova Läkemedel AB	24/07/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012
Mabthera	Paranova Läkemedel AB	15/10/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	25/10/2012
Mabthera	propharmed GmbH	11/07/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/08/2012
Mabthera	propharmed GmbH	11/07/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/08/2012
Mabthera	Veron Pharma Vertriebs GmbH	30/03/2012	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	18/04/2012
Mabthera	Veron Pharma Vertriebs GmbH	30/03/2012	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	18/04/2012
Metacam	2 Care 4	23/10/2012	0.5 mg/ml	Oral suspension	1 x 15 ml bottle	EU/2/97/004/026	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	05/11/2012

Metacam	CC Pharma	09/01/2012	0.5 mg/ml	Oral suspension	1 bottle	EU/2/97/004/034	Belgium, Norway, United Kingdom	Germany	14/02/2012
Metacam	CC Pharma	06/02/2012	0.5 mg/ml	Oral suspension	1 Polyethylene bottle	EU/2/97/004/026	Belgium, Norway, United Kingdom	Germany	14/02/2012
Metacam	CC Pharma	27/03/2012	1 mg	Chewable	84	EU/2/97/004/044	Norway	Germany	02/04/2012
Metacam	CC Pharma	27/03/2012	2.5 mg	Chewable	84	EU/2/97/004/047	Norway	Germany	02/04/2012
Metacam	CC Pharma	28/03/2012	5 mg/ml	Solution for injection	20 ml	EU/2/97/004/011	Belgium, Norway	Germany	02/04/2012
Metacam	CC Pharma	10/04/2012	2 mg/ml	Solution for injection	10 ml	EU/2/97/004/039	Belgium, Norway	Germany	23/04/2012
Metacam	CC Pharma	11/04/2012	1.5 mg/ml	Oral suspension	10 ml	EU/2/97/004/003	Belgium, Norway, United Kingdom	Germany	24/04/2012
Metacam	CC Pharma	11/04/2012	1.5 mg/ml	Oral suspension	32 ml	EU/2/97/004/004	Belgium, Norway, Slovak Republic, United Kingdom	Germany	24/04/2012
Metacam	CC Pharma	11/04/2012	1.5 mg/ml	Oral suspension	100 ml	EU/2/97/004/005	Belgium, United Kingdom	Germany	24/04/2012
Metacam	CC Pharma	11/04/2012	1.5 mg/ml	Oral suspension	180 ml	EU/2/97/004/029	Belgium, Norway, United Kingdom	Germany	24/04/2012
Metacam	CC Pharma	17/04/2012	15 mg/ml	Oral suspension	100 ml	EU/2/97/004/009	Belgium, Norway	Germany	26/04/2012
Metacam	CC Pharma	17/04/2012	15 mg/ml	Oral suspension	250 ml	EU/2/97/004/030	Norway, United Kingdom	Germany	26/04/2012
Metacam	CC Pharma	21/05/2012	20 mg/ml	Solution for injection	50 ml	EU/2/97/0 04/007	Belgium, Norway	Germany	05/06/2012
Metacam	CC Pharma	21/05/2012	20 mg/ml	Solution for injection	100 ml	EU/2/97/0 04/008	Belgium	Germany	05/06/2012

Micardis	ADL Pharma GmbH	30/11/2012	80 mg	Tablet	56	EU/1/98/090/007	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/12/2012
Micardis	ADL Pharma GmbH	30/11/2012	80 mg	Tablet	98	EU/1/98/090/008	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/12/2012
Micardis	EU-Pharma BV	05/03/2012	80 mg	Tablet	14	EU/1/98/090/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	19/04/2012
Micardis	EU-Pharma BV	06/12/2012	80 mg	Tablet	28	EU/1/98/090/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/12/2012
Micardis	Kosei Pharma UK Limited	01/05/2012	80 mg	Tablet	28	EU/1/98/090/006	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	16/05/2012
Micardis	MPT Pharma	10/04/2012	40 mg	Tablet	28	EU/1/98/090/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	23/04/2012
Micardis	MPT Pharma	10/04/2012	80 mg	Tablet	28	EU/1/98/090/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	23/04/2012
MicardisPlus	ADL Pharma GmbH	07/03/2012	80 mg/ 12.5 mg	Tablet	56	EU/1/02/213/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/04/2012

MicardisPlus	ADL Pharma GmbH	07/03/2012	80 mg/ 12.5 mg	Tablet	98	EU/1/02/213/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/04/2012
MicardisPlus	ADL Pharma GmbH	15/05/2012	80 mg/ 25 mg	Tablet	56	EU/1/02/213/021	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
MicardisPlus	ADL Pharma GmbH	15/05/2012	80 mg/ 25 mg	Tablet	98	EU/1/02/213/023	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
MicardisPlus	axicorp Pharma GmbH	15/10/2012	40 mg/ 12.5 mg	Tablet	98	EU/1/02/213/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/10/2012
MicardisPlus	B2B Medical GmbH	23/10/2012	80 mg/ 25 mg	Tablet	98	EU/1/02/213/023	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/11/2012
MicardisPlus	Clear Pharmacy	15/12/2011	80 mg/ 25 mg	Tablet	28	EU/1/02/213/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	17/01/2012
MicardisPlus	EU-Pharma BV	10/01/2012	80 mg/ 12.5 mg	Tablet	14	EU/1/02/213/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/01/2012
MicardisPlus	Kosei Pharma UK Limited	31/01/2012	80 mg/ 12.5 mg	Tablet	28	EU/1/02/213/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	14/03/2012
MicardisPlus	Mediwin Ltd	30/04/2012	80 mg/ 12.5 mg	Tablet	30 x 1	EU/1/02/213/015	France	United Kingdom	22/05/2012
Mimpara	ABACUS MEDICINE A/S	27/12/2011	30 mg	Film-coated tablet	28	EU/1/04/292/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/01/2012

Mimpara	Orifarm AB	30/07/2012	60 mg	Film-coated tablet	28	EU/1/04/292/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	20/08/2012
Mirapexin	2 Care 4	13/03/2012	0.26 mg	Prolonged-release tablet	10	EU/1/97/051/013	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/04/2012
Mirapexin	2 Care 4	13/03/2012	0.26 mg	Prolonged-release tablet	30	EU/1/97/051/014	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/04/2012
Mirapexin	2 Care 4	13/03/2012	0.52 mg	Prolonged-release tablet	10	EU/1/97/051/016	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/04/2012
Mirapexin	AMIMED DIRECT LTD	25/04/2012	0.26 mg	Prolonged-release tablet	10	EU/1/97/051/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	18/05/2012
Mirapexin	AMIMED DIRECT LTD	25/04/2012	0.52 mg	Prolonged-release tablet	10	EU/1/97/051/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	18/05/2012
Mirapexin	Chemilines Ltd	20/04/2012	0.52 mg	Prolonged-release tablet	30	EU/1/97/051/017	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands	United Kingdom	18/05/2012
Mirapexin	Cross Pharma AB	24/01/2012	2.1 mg	Prolonged-release tablet	100	EU/1/97/051/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/02/2012

Mirapexin	Cross Pharma AB	02/05/2012	0.26 mg	Prolonged-release tablet	100	EU/1/97/051/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	22/05/2012
Mirapexin	Cross Pharma AB	02/05/2012	0.52 mg	Prolonged-release tablet	100	EU/1/97/051/018	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	22/05/2012
Mirapexin	Cross Pharma AB	02/05/2012	1.05 mg	Prolonged-release tablet	100	EU/1/97/051/021	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	22/05/2012
Mirapexin	Docpharm GmbH & CoKGaA	14/12/2011	0.26 mg	Prolonged-release tablet	10	EU/1/97/051/013	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Mirapexin	Docpharm GmbH & CoKGaA	14/12/2011	0.52 mg	Prolonged-release tablet	30	EU/1/97/051/017	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Mirapexin	Docpharm GmbH & CoKGaA	14/12/2011	0.52 mg	Prolonged-release tablet	100	EU/1/97/051/018	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Mirapexin	EuroPharma DK ApS	07/06/2012	0.26 mg	Prolonged-release tablet	10	EU/1/97/051/013	Austria, Belgium, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/06/2012
Mirapexin	EuroPharma DK ApS	07/06/2012	0.26 mg	Prolonged-release tablet	30	EU/1/97/051/014	Austria, Belgium, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/06/2012

Mirapexin	EuroPharma DK ApS	07/06/2012	0.26 mg	Prolonged-release tablet	100	EU/1/97/051/015	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/06/2012
Mirapexin	Ginova Ltd	11/04/2012	3.15 mg	Prolonged-release tablet	30	EU/1/97/051/026	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	15/05/2012
Mirapexin	Mediwin Ltd	10/12/2012	0.52 mg	Prolonged-release tablet	30	EU/1/97/051/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	18/12/2012
Mirapexin	O.P.D. Laboratories Ltd	02/07/2010	0.26 mg	Prolonged-release tablet	10	EU/1/97/051/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	01/02/2012
Mirapexin	O.P.D. Laboratories Ltd	02/07/2010	0.52 mg	Prolonged-release tablet	10	EU/1/97/051/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	01/02/2012
Mirapexin	O.P.D. Laboratories Ltd	02/07/2010	1.05 mg	Prolonged-release tablet	30	EU/1/97/051/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	01/02/2012
Mirapexin	O.P.D. Laboratories Ltd	02/07/2010	2.1 mg	Prolonged-release tablet	30	EU/1/97/051/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	31/01/2012
Mirapexin	O.P.D. Laboratories Ltd	02/07/2010	3.15 mg	Prolonged-release tablet	30	EU/1/97/051/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	01/02/2012
Mirapexin	O.P.D. Laboratories Ltd	10/08/2012	0.7 mg	Tablet	30	EU/1/97/051/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	23/08/2012
Mirapexin	Orifarm AB	24/01/2012	0.26 mg	Prolonged-release tablet	100	EU/1/97/051/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	01/02/2012

Mirapexin	PCO Manufacturing	28/05/2012	1.05 mg	Prolonged-release tablet	30	EU/1/97/051/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	07/06/2012
Mirapexin	PCO Manufacturing	29/05/2012	2.1 mg	Prolonged-release tablet	30	EU/1/97/051/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	07/06/2012
Mirapexin	PCO Manufacturing	05/11/2012	0.18 mg	Tablet	100	EU/1/97/051/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	09/11/2012
Mirapexin	PCO Manufacturing	05/11/2012	0.7 mg	Tablet	100	EU/1/97/051/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	09/11/2012
Mircera	CC Pharma	21/11/2012	30 µg/0.3 ml	Solution for injection	3 x 1 pre-filled syringe	EU/1/07/400/022	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/11/2012
Mircera	Euro Registratie Collectief BV	06/03/2012	100 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/03/2012
Mircera	Euro Registratie Collectief BV	07/03/2012	50 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/03/2012
Mircera	kohlpharma GmbH	10/01/2012	360 µg/0.6 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/021	Greece	Germany	24/01/2012
Mircera	kohlpharma GmbH	08/06/2012	120 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/020	Greece, Spain	Germany	12/07/2012
Mircera	Paranova Läkemedel AB	28/12/2011	100 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/01/2012

Mircera	Paranova Läkemedel AB	28/12/2011	50 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/01/2012
Mircera	Paranova Läkemedel AB	28/12/2011	75 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/01/2012
Mircera	Paranova Oy	15/03/2012	75 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	24/04/2012
Mixtard	Emra-Med	17/04/2012	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/02/231/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/06/2012
M-M-RVAXPRO	CC Pharma	12/12/2012	- -	Powder and solvent for suspension for injection in pre-filled syringe	1 vial + 1 pre-filled syringe + 2 needles	EU/1/06/337/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/12/2012
M-M-RVAXPRO	hvd medical GmbH	16/03/2012	- -	Powder and solvent for suspension for injection in pre-filled syringe	1 vial + 1 pre-filled syringe + 2 needles	EU/1/06/337/011	France	Germany	30/03/2012
Mozobil	Orifarm AS	13/08/2012	20 mg/ml	Solution for injection	1 vial	EU/1/09/537/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012

MULTAQ	ABACUS MEDICINE A/S	02/08/2012	400 mg	Film-coated tablet	100 x 1	EU/1/09/591/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/08/2012
MULTAQ	Amimed Direct Ltd	05/07/2012	400 mg	Film-coated tablet	60	EU/1/09/591/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	16/07/2012
MULTAQ	Ginova Ltd	19/06/2012	400 mg	Film-coated tablet	20	EU/1/09/591/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/07/2012
MULTAQ	Ginova Ltd	19/06/2012	400 mg	Film-coated tablet	60	EU/1/09/591/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/07/2012
MULTAQ	Medartuum AB	03/10/2012	400 mg	Film-coated tablet	60	EU/1/09/591/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012
MULTAQ	O.P.D. Laboratories Ltd	16/07/2012	400 mg	Film-coated tablet	60	EU/1/09/591/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	23/07/2012
MULTAQ	Orifarm AB	15/02/2012	400 mg	Film-coated tablet	100 x 1	EU/1/09/591/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/06/2012
MULTAQ	PHARMACHIM AB	14/12/2012	400 mg	Film-coated tablet	20	EU/1/09/591/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/12/2012
MULTAQ	PHARMACHIM AB	14/12/2012	400 mg	Film-coated tablet	100 x 1	EU/1/09/591/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/12/2012
Mycamine	Orifarm AS	07/09/2012	100 mg	Powder for solution for infusion	1 vial	EU/1/08/448/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/10/2012

Mycamine	Orifarm AS	07/09/2012	50 mg	Powder for solution for infusion	1 vial	EU/1/08/448/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/10/2012
Myocet	ABACUS MEDICINE A/S	24/07/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	2 sets of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/08/2012
Myocet	CC Pharma	28/12/2011	50 mg	Powder and pre-admixtures for concentrate for liposomal dispersion for infusion	One set of one vial for each of the three components	EU/1/00/141/002	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/01/2012
Myocet	HAEMATO PHARM AG	03/05/2012	50 mg	Powder and pre-admixtures for concentrate for liposomal dispersion for infusion	One set of one vial for each of the three components	EU/1/00/141/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2012
Myocet	Inopha GmbH	14/08/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	1 set of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/08/2012
Myocet	Orifarm AB	27/09/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	2 sets of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	19/11/2012
Myocet	Orifarm AS	27/09/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	2 sets of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/11/2012

Myocet	Paranova Danmark AS	20/07/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	2 sets of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/08/2012
Myocet	Shipping Dynamics Scandinavia AB	06/07/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	2 sets of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	03/09/2012
Myocet	Tisida GmbH	25/06/2012	50 mg	Powder, dispersion and solvent for concentrate for dispersion for infusion	1 set of 3 vials (1 powder + 1 liposomes + 1 buffer)	EU/1/00/141/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/07/2012
Naxcel	Dopharma B.V	21/12/2011	100 mg/ml	Suspension for injection	50 ml	EU/2/05/053/002	Lithuania, Romania, Spain	Belgium, France, Germany, The Netherlands	06/03/2012
NeoRecormon	ABACUS MEDICINE A/S	23/12/2011	30000 IU	Solution for injection	4 pre-filled syringes	EU/1/97/031/046	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/01/2012
NeoRecormon	Cross Pharma AB	12/10/2012	10000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/036	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012
NeoRecormon	Cross Pharma AB	12/10/2012	4000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/042	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	30/10/2012

NeoRecormon	Cross Pharma AB	12/10/2012	6000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/044	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/10/2012
NeoRecormon	Omnia Läkemedel AB	22/03/2012	10000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/036	Austria, Belgium, Bulgaria, Finland, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	14/05/2012
NeoRecormon	Omnia Läkemedel AB	22/03/2012	4000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/042	Austria, Belgium, Bulgaria, Finland, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	11/05/2012
NeoRecormon	Omnia Läkemedel AB	22/03/2012	5000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/034	Austria, Belgium, Bulgaria, Finland, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	11/05/2012
NeoRecormon	Omnia Läkemedel AB	22/03/2012	6000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/044	Austria, Belgium, Bulgaria, Finland, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	11/05/2012
NeoRecormon	Orifarm AB	24/09/2012	3000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/10/2012
NeoRecormon	Paranova Läkemedel AB	27/08/2012	6000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/044	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/09/2012
NeoRecormon	Paranova Läkemedel AB	29/08/2012	2000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/09/2012

NeoRecormon	Paranova Läkemedel AB	29/08/2012	3000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/032	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/09/2012
NeoRecormon	Paranova Läkemedel AB	29/08/2012	4000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/042	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	18/09/2012
NeoRecormon	Paranova Läkemedel AB	21/09/2012	30000 IU	Solution for injection	4 pre-filled syringes	EU/1/97/031/046	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	08/10/2012
NeoRecormon	Paranova Oy	21/06/2012	10000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/036	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	10/07/2012
NeoRecormon	Paranova Oy	21/06/2012	30000 IU	Solution for injection	4 pre-filled syringes	EU/1/97/031/046	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	10/07/2012
NeoRecormon	Paranova Oy	21/06/2012	4000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/042	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	10/07/2012

NeoRecormon	Paranova Oy	21/06/2012	5000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/034	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	10/07/2012
NeoRecormon	Paranova Oy	21/06/2012	6000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/044	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	10/07/2012
Neulasta	Aaston Healthcare GmbH	24/11/2011	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012
Neulasta	ABACUS MEDICINE A/S	24/08/2012	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/09/2012
Neulasta	Adripharma GmbH	27/08/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/10/2012
Neulasta	Axicorp Pharma B.V.	28/09/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Hungary, Italy, Latvia, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/10/2012
Neulasta	Cross Pharma AB	20/08/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012

Neulasta	HAEMATO PHARM AG	18/06/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	Ireland, United Kingdom	11/07/2012
Neulasta	Inopha GmbH	07/03/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	14/03/2012
Neulasta	Northern Medical Group ApS.	26/01/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	11/07/2012
Neulasta	Orifarm OY	29/04/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	14/05/2012
Neulasta	REMEDIX GMBH	14/05/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/06/2012
Neulasta	Tisida GmbH	11/06/2012	6 mg	Solution for injection	1 pre-filled syringe with needle guard	EU/1/02/227/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Neupro	Amimed Direct Ltd	05/11/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	13/11/2012

Neupro	Amimed Direct Ltd	03/12/2012	2 mg/24h	Transdermal patch	28	EU/1/05/331/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	17/12/2012
Neupro	axicorp Pharma GmbH	24/02/2012	2 mg/24h	Transdermal patch	28	EU/1/05/331/002	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/03/2012
Neupro	axicorp Pharma GmbH	12/04/2012	2 mg/24h	Transdermal patch	7	EU/1/05/331/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Neupro	axicorp Pharma GmbH	12/04/2012	2 mg/24h	Transdermal patch	56	EU/1/05/331/016	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Neupro	axicorp Pharma GmbH	12/04/2012	3 mg/24h	Transdermal patch	28	EU/1/05/331/049	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Neupro	axicorp Pharma GmbH	24/04/2012	3 mg/24h	Transdermal patch	56	EU/1/05/331/051	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Neupro	axicorp Pharma GmbH	14/06/2012	4 mg/24h	Transdermal patch	84	EU/1/05/331/024	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/07/2012
Neupro	axicorp Pharma GmbH	21/08/2012	2 mg/24h	Transdermal patch	84	EU/1/05/331/018	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Neupro	axicorp Pharma GmbH	22/08/2012	6 mg/24h	Transdermal patch	84	EU/1/05/331/030	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Neupro	axicorp Pharma GmbH	22/08/2012	8 mg/24h	Transdermal patch	84	EU/1/05/331/036	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012

Neupro	Dr Fisher Farma BV	21/11/2012	2 mg/24h	Transdermal patch	7	EU/1/05/331/001	Austria, Belgium, Bulgaria, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	12/12/2012
Neupro	Eureco-Pharma B.V.	30/05/2012	2 mg/24h	Transdermal patch	7	EU/1/05/331/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	07/06/2012
Neupro	Ginova Ltd	21/09/2012	2 mg/24h	Transdermal patch	28	EU/1/05/331/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	14/11/2012
Neupro	Ginova Ltd	21/09/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	14/11/2012
Neupro	Ginova Ltd	21/09/2012	6 mg/24h	Transdermal patch	28	EU/1/05/331/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	14/11/2012
Neupro	Ginova Ltd	21/09/2012	8 mg/24h	Transdermal patch	28	EU/1/05/331/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	14/11/2012
Neupro	HAEMATO PHARM AG	24/04/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/05/2012

Neupro	HAEMATO PHARM AG	24/04/2012	6 mg/24h	Transdermal patch	28	EU/1/05/331/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/05/2012
Neupro	HAEMATO PHARM AG	24/04/2012	8 mg/24h	Transdermal patch	28	EU/1/05/331/011	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/05/2012
Neupro	hvd medical GmbH	06/03/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Czech Republic, Spain	Germany	10/05/2012
Neupro	MPT Pharma	29/06/2012	2 mg/24h	Transdermal patch	28	EU/1/05/331/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012
Neupro	MPT Pharma	29/06/2012	4 mg/24h	Transdermal patch	7	EU/1/05/331/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012
Neupro	MPT Pharma	29/06/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012
Neupro	MPT Pharma	29/06/2012	6 mg/24h	Transdermal patch	7	EU/1/05/331/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012

Neupro	MPT Pharma	29/06/2012	6 mg/24h	Transdermal patch	28	EU/1/05/331/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012
Neupro	MPT Pharma	29/06/2012	8 mg/24h	Transdermal patch	28	EU/1/05/331/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/07/2012
Neupro	Pharma Westen GmbH	04/10/2011	4 mg/24h	Transdermal patch	56	EU/1/05/331/022	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/01/2012
Neupro	REMEDIX GMBH	17/05/2012	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	11/06/2012
Neupro	REMEDIX GMBH	17/05/2012	6 mg/24h	Transdermal patch	28	EU/1/05/331/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	12/06/2012
Neupro	REMEDIX GMBH	17/05/2012	8 mg/24h	Transdermal patch	28	EU/1/05/331/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	12/06/2012
Neupro	S.C.A.C Ltd	13/12/2011	2 mg/24h	Transdermal patch	28	EU/1/05/331/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/01/2012

Neupro	S.C.A.C Ltd	13/12/2011	4 mg/24h	Transdermal patch	28	EU/1/05/331/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/01/2012
Neupro	S.C.A.C Ltd	13/12/2011	6 mg/24h	Transdermal patch	28	EU/1/05/331/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/01/2012
Neupro	S.C.A.C Ltd	13/12/2011	8 mg/24h	Transdermal patch	28 patches	EU/1/05/331/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	19/01/2012
Neupro	WAYMADE PLC	16/12/2011	4 mg/24h	Transdermal patch	7	EU/1/05/331/004	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/01/2012
Neupro	WAYMADE PLC	16/12/2011	6 mg/24h	Transdermal patch	7	EU/1/05/331/007	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/01/2012
Neupro	WAYMADE PLC	16/12/2011	8 mg/24h	Transdermal patch	7	EU/1/05/331/010	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	09/01/2012
Nevanac	2 Care 4	05/09/2012	1 mg/ml	Eye drops, suspension	1 bottle	EU/1/07/433/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	26/09/2012
Nevanac	2 Care 4	05/09/2012	1 mg/ml	Eye drops, suspension	1 bottle	EU/1/07/433/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/09/2012
Nevanac	Copharma ApS	26/09/2012	1 mg/ml	Eye drops, suspension	1 bottle	EU/1/07/433/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/10/2012

Nexavar	Beragena Arzneimittel GmbH	12/07/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	25/07/2012
Nexavar	Cambridge Major Laboratories Europe B.V	24/02/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Bulgaria, Czech Republic, Finland, France, Greece, Ireland, Italy, Lithuania, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Nexavar	Emra-Med	16/11/2011	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	12/01/2012
Nexavar	MTK Pharma GmbH	29/10/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Czech Republic, Poland	Germany	09/11/2012
Nexavar	Paranova Läkemedel AB	05/01/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/01/2012
Nexavar	Paranova Oy	20/03/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	21/05/2012
Nexavar	PHARMACHIM AB	31/01/2012	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/02/2012
Norvir	axicorp Pharma GmbH	26/07/2012	100 mg	Film-coated tablet	30	EU/1/96/016/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/08/2012

Norvir	Cross Pharma AB	20/06/2012	100 mg	Film-coated tablet	90 (3 x 30)	EU/1/96/016/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/06/2012
Norvir	Emra-Med	12/12/2011	100 mg	Film-coated tablet	30	EU/1/96/016/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/01/2012
Norvir	Orifarm AB	16/07/2012	100 mg	Film-coated tablet	90 (3 x 30)	EU/1/96/016/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/07/2012
Norvir	Orifarm AS	16/07/2012	100 mg	Film-coated tablet	90 (3 x 30)	EU/1/96/016/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/07/2012
Norvir	Orifarm AS	16/07/2012	100 mg	Film-coated tablet	30	EU/1/96/016/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/07/2012
Norvir	Paranova Läkemedel AB	25/10/2012	100 mg	Film-coated tablet	90 (3 x 30)	EU/1/96/016/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/11/2012
NovoMix	Eureco-Pharma B.V.	23/12/2011	100 U/ml	Suspension for injection	5 cartridges	EU/1/00/142/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	17/02/2012

NovoMix	Eureco-Pharma B.V.	11/05/2012	100 U/ml	Suspension for injection	10 cartridges	EU/1/00/142/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	05/06/2012
NovoMix	kohlpharma GmbH	01/02/2012	100 U/ml	Suspension for injection	5 pre-filled pens	EU/1/00/142/009	France, Italy, Spain	Germany	21/02/2012
NovoMix	kohlpharma GmbH	01/02/2012	100 U/ml	Suspension for injection	10 pre-filled pens	EU/1/00/142/010	France, Italy, Spain	Germany	27/02/2012
NovoMix	kohlpharma GmbH	02/02/2012	100 U/ml	Suspension for injection	5 cartridges	EU/1/00/142/004	Austria, Poland	Germany	27/02/2012
NovoMix	kohlpharma GmbH	02/02/2012	100 U/ml	Suspension for injection	10 cartridges	EU/1/00/142/005	Austria, Poland	Germany	27/02/2012
NovoMix	PHARMACHIM AB	05/12/2012	100 U/ml	Suspension for injection	5 pre-filled pens	EU/1/00/142/009	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
NovoMix 30 Penfill	EU-Pharma BV	10/01/2012	100 U/ml	Suspension for injection	5 cartridges	EU/1/00/142/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/01/2012
NovoNorm	2 Care 4	13/04/2012	0.5 mg	Tablet	90	EU/1/98/076/005	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/04/2012
NovoNorm	2 Care 4	13/04/2012	1 mg	Tablet	90	EU/1/98/076/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/04/2012
NovoNorm	2 Care 4	13/04/2012	2 mg	Tablet	90	EU/1/98/076/019	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/04/2012

NovoNorm	BB Farma s.r.l.	30/04/2012	0.5 mg	Tablet	90	EU/1/98/076/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	24/05/2012
NovoNorm	BB Farma s.r.l.	30/04/2012	0.5 mg	Tablet	270	EU/1/98/076/023	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	24/05/2012
NovoNorm	BB Farma s.r.l.	30/04/2012	1 mg	Tablet	90	EU/1/98/076/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	24/05/2012
NovoNorm	BB Farma s.r.l.	30/04/2012	1 mg	Tablet	270	EU/1/98/076/024	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	24/05/2012
NovoNorm	BB Farma s.r.l.	09/05/2012	0.5 mg	Tablet	30	EU/1/98/076/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/06/2012
NovoNorm	BB Farma s.r.l.	17/05/2012	0.5 mg	Tablet	90	EU/1/98/076/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	07/06/2012

NovoNorm	BB Farma s.r.l.	06/06/2012	1 mg	Tablet	120	EU/1/98/076/013	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/07/2012
NovoNorm	BB Farma s.r.l.	06/06/2012	1 mg	Tablet	30	EU/1/98/076/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/07/2012
NovoNorm	BB Farma s.r.l.	13/06/2012	2 mg	Tablet	30	EU/1/98/076/018	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/07/2012
NovoNorm	BB Farma s.r.l.	13/06/2012	2 mg	Tablet	120	EU/1/98/076/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/07/2012
NovoNorm	BB Farma s.r.l.	15/06/2012	1 mg	Tablet	90	EU/1/98/076/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	03/07/2012
NovoNorm	Euro Registratie Collectief BV	01/06/2012	2 mg	Tablet	90	EU/1/98/076/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	03/07/2012
NovoNorm	Mediwin Ltd	10/09/2012	0.5 mg	Tablet	90	EU/1/98/076/005	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/09/2012

NovoNorm	Mediwin Ltd	10/09/2012	1 mg	Tablet	90	EU/1/98/076/012	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/09/2012
NovoRapid	Eureco-Pharma B.V.	11/05/2012	100 U/ml	Solution for injection	10 cartridges	EU/1/99/119/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	28/06/2012
Noxafil	Medartuum AB	10/04/2012	40 mg/ml	Oral suspension	1 bottle	EU/1/05/320/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/04/2012
Noxafil	Paranova Läkemedel AB	13/08/2012	40 mg/ml	Oral suspension	1 bottle	EU/1/05/320/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/08/2012
Noxafil	PHARMACHIM AB	26/09/2012	40 mg/ml	Oral suspension	1 bottle	EU/1/05/320/001	Austria, Belgium, Bulgaria, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	08/10/2012
Nplate	Medartuum AB	20/06/2012	500 µg	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 1 vial adapter + 1 needle + 1 syringe + 4 alcohol pads	EU/1/08/497/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	16/07/2012
Nulojix	CC Pharma	21/02/2012	250 mg	Powder for concentrate for solution for infusion	1 vial + 1 syringe	EU/1/11/694/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012

Nulojix	Emra-Med	03/08/2012	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/11/694/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/08/2012
Nulojix	Emra-Med	06/08/2012	250 mg	Powder for concentrate for solution for infusion	1 vial + 1 syringe	EU/1/11/694/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/08/2012
NutropinAq	Medartuum AB	09/10/2012	10 mg/2 ml	Solution for injection	3 cartridges	EU/1/00/164/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/10/2012
NutropinAq	Pharma Lab	15/02/2012	10 mg/2 ml	Solution for injection	1 cartridge	EU/1/00/164/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	23/05/2012
Omnitrope	CC Pharma	22/11/2012	10 mg/1.5ml	Solution for injection in cartridge	1 cartridge	EU/1/06/332/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/11/2012
Omnitrope	COPHARMA ApS	26/04/2012	10 mg/1.5ml	Solution for injection in cartridge	1 cartridge	EU/1/06/332/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	02/05/2012
Omnitrope	Emra-Med	02/12/2011	5 mg/1.5ml	Solution for injection in cartridge	10 cartridges	EU/1/06/332/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/02/2012
Omnitrope	Orifarm AB	07/09/2012	10 mg/1.5ml	Solution for injection in cartridge	1 cartridge	EU/1/06/332/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/11/2012
Onbrez Breezhaler	2 Care 4	09/12/2011	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Sweden	21/02/2012

Onbrez Breezhaler	2 Care 4	09/12/2011	150 µg	Inhalation powder, hard capsule	3 x (30+1 inhaler)	EU/1/09/593/004	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Sweden	22/02/2012
Onbrez Breezhaler	2 Care 4	09/12/2011	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Sweden	21/02/2012
Onbrez Breezhaler	2 Care 4	09/12/2011	300 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/009	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Sweden	22/02/2012
Onbrez Breezhaler	2 Care 4	24/05/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/05/2012
Onbrez Breezhaler	2 Care 4	24/05/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/05/2012
Onbrez Breezhaler	2 Care 4	25/05/2012	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/05/2012
Onbrez Breezhaler	2 Care 4	25/05/2012	300 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/009	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/05/2012
Onbrez Breezhaler	ABACUS MEDICINE A/S	02/05/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/05/2012
Onbrez Breezhaler	Beragena Arzneimittel GmbH	28/11/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	10/12/2012
Onbrez Breezhaler	Beragena Arzneimittel GmbH	28/11/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	10/12/2012
Onbrez Breezhaler	Chemilines Ltd	08/12/2011	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	11/01/2012

Onbrez Breezhaler	Cross Pharma AB	16/05/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/05/2012
Onbrez Breezhaler	Cross Pharma AB	16/05/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/05/2012
Onbrez Breezhaler	Cross Pharma AB	16/05/2012	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/05/2012
Onbrez Breezhaler	Cross Pharma AB	16/05/2012	300 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/05/2012
Onbrez Breezhaler	Doncaster Pharmaceuticals Group Ltd	12/12/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	14/12/2012
Onbrez Breezhaler	Ginova Ltd	12/11/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/12/2012
Onbrez Breezhaler	Ginova Ltd	12/11/2012	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/12/2012

Onbrez Breezhaler	Medartuum AB	29/12/2011	300 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Onbrez Breezhaler	Medartuum AB	03/01/2012	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Onbrez Breezhaler	Orifarm AS	18/07/2012	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/09/2012
Onbrez Breezhaler	PCO Manufacturing	16/02/2012	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	19/03/2012
Onbrez Breezhaler	PCO Manufacturing	16/02/2012	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	19/03/2012
Onbrez Breezhaler	Pharmachim AB	02/12/2011	150 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/004	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
Onbrez Breezhaler	Pharmachim AB	02/12/2011	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Czech Republic, Denmark, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
Onbrez Breezhaler	Pharmachim AB	02/12/2011	300 µg	Inhalation powder, hard capsule	3 x (30 + 1 inhaler)	EU/1/09/593/009	Austria, Belgium, Czech Republic, Denmark, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012

Onbrez Breezhaler	Pharmachim AB	02/12/2011	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Czech Republic, Denmark, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
Onbrez Breezhaler	STRATHCLYDE PHARMACEUTICALS LTD	14/12/2011	150 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	05/01/2012
Onbrez Breezhaler	STRATHCLYDE PHARMACEUTICALS LTD	15/12/2011	300 µg	Inhalation powder, hard capsule	30 (3 x 10) + 1 inhaler	EU/1/09/593/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	05/01/2012
Onglyza	ABACUS MEDICINE A/S	13/08/2012	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012
Onglyza	CC Pharma	15/09/2010	5 mg	Film-coated tablet	28	EU/1/09/545/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/04/2012
Onglyza	CC Pharma	15/09/2010	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/04/2012
Onglyza	Medartuum AB	08/10/2012	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	29/10/2012
Onglyza	Mevita HandelsGmbH	22/12/2011	5 mg	Film-coated tablet	28	EU/1/09/545/006	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	30/01/2012
Onglyza	Mevita HandelsGmbH	22/12/2011	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	30/01/2012
Onglyza	O.P.D. Laboratories Ltd	30/06/2010	5 mg	Film-coated tablet	28	EU/1/09/545/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/02/2012

Onglyza	Paranova Läkemedel AB	02/04/2012	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	11/05/2012
Onglyza	PHARMACHIM AB	10/12/2012	5 mg	Film-coated tablet	28	EU/1/09/545/006	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	17/12/2012
Onglyza	PHARMACHIM AB	10/12/2012	5 mg	Film-coated tablet	98	EU/1/09/545/008	Austria, Belgium, Czech Republic, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	17/12/2012
Opatanol	Copharma ApS	19/06/2012	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Opatanol	Copharma ApS	19/06/2012	1 mg/ml	Eye drops, solution	3 bottles	EU/1/02/217/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Opatanol	Medartuum AB	29/11/2011	1 mg/ml	Eye drops, solution	3 bottles	EU/1/02/217/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	04/01/2012
Optruma	Orifarm AB	04/10/2012	60 mg	Film-coated tablet	28	EU/1/98/074/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	16/10/2012
Optruma	PHARMA LAB	03/10/2012	60 mg	Film-coated tablet	28	EU/1/98/074/002	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	24/10/2012

Optruma	PHARMA LAB	03/10/2012	60 mg	Film-coated tablet	84	EU/1/98/074/003	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	24/10/2012
Orencia	Paranova Läkemedel AB	16/03/2012	250 mg	Powder for concentrate for solution for infusion	1 vial + 1 syringe	EU/1/07/389/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/04/2012
Orfadin	Euro Registratie Collectief BV	10/01/2012	10 mg	Capsule, hard	60	EU/1/04/303/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/01/2012
Orgalutran	ABACUS MEDICINE A/S	13/06/2012	0.25 mg/ 0.5 ml	Solution for injection	5 pre-filled syringes	EU/1/00/130/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/07/2012
Orgalutran	axicorp Pharma GmbH	10/12/2012	0.25 mg/ 0.5 ml	Solution for injection	5 pre-filled syringes	EU/1/00/130/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/12/2012
Orgalutran	HAEMATO PHARM AG	14/03/2012	0.25 mg/ 0.5 ml	Solution for injection	1 pre-filled syringe	EU/1/00/130/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/03/2012
Orgalutran	HAEMATO PHARM AG	14/03/2012	0.25 mg/ 0.5 ml	Solution for injection	5 pre-filled syringes	EU/1/00/130/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/03/2012
Orgalutran	Medcor Pharmaceuticals BV	29/10/2012	0.25 mg/ 0.5 ml	Solution for injection	1 pre-filled syringe	EU/1/00/130/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	30/10/2012

Orgalutran	Paranova Oy	03/08/2012	0.25 mg/ 0.5 ml	Solution for injection	5 pre-filled syringes	EU/1/00/130/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	29/08/2012
Ovitrelle	Eureco-Pharma B.V.	12/09/2012	250 micrograms	Solution for injection	1 pre-filled pen + 2 needles	EU/1/00/165/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	25/10/2012
Ovitrelle	kohlpharma GmbH	22/12/2011	250 micrograms/ 0.5 ml	Solution for injection	1 pre-filled syringe	EU/1/00/165/007	Lithuania	Germany	16/02/2012
Ovitrelle	kohlpharma GmbH	02/07/2012	250 micrograms	Solution for injection	1 pre-filled pen + 2 needles	EU/1/00/165/008	Lithuania	Germany	02/08/2012
Ovitrelle	PHARMACHIM AB	09/11/2012	250 micrograms	Solution for injection	1 pre-filled pen + 1 needle	EU/1/00/165/008	Austria, Belgium, Bulgaria, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	26/11/2012
PANTECTA Control	CC Pharma	18/04/2012	227mg	Chewable tablets	180	EU/2/04/045/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/04/2012
Pegasys	ABACUS MEDICINE A/S	07/03/2012	135 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/03/2012
Pegasys	CC Pharma	18/06/2012	135 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/012	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/07/2012
Pegasys	CC Pharma	18/06/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/07/2012

Pegasys	CC Pharma	18/06/2012	180 µg	Solution for injection	12 pre-filled pens	EU/1/02/221/016	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/07/2012
Pegasys	CC Pharma	08/10/2012	135 µg	Solution for injection	1 pre-filled pen	EU/1/02/221/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/10/2012
Pegasys	CC Pharma	08/10/2012	180 µg	Solution for injection	1 pre-filled pen	EU/1/02/221/014	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/10/2012
Pegasys	Emra-Med	13/11/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/11/2012
Pegasys	kohlpharma GmbH	02/10/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
Pegasys	Medartuum AB	02/11/2012	135 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	03/12/2012
Pegasys	Medartuum AB	02/11/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	03/12/2012
Pegasys	Orifarm AB	28/02/2012	135 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	02/05/2012

Pegasys	Orifarm AB	08/03/2012	135 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	02/05/2012
Pegasys	Orifarm AB	08/03/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	02/05/2012
Pegasys	Paranova Läkemedel AB	07/11/2012	135 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/11/2012
Pegasys	Pharma Westen GmbH	05/03/2012	180 µg	Solution for injection	4 pre-filled pens	EU/1/02/221/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/05/2012
PegIntron	ABACUS MEDICINE A/S	09/11/2012	100 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/040	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/11/2012
PegIntron	Emra-Med	26/01/2012	100 micrograms	Powder and solvent for solution for injection in pre-filled pen	12 pens + 12 injection needles + 24 cleansing swabs	EU/1/00/131/042	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/03/2012
PegIntron	Orifarm AS	15/08/2012	50 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/032	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/08/2012

Plavix	B2B Medical GmbH	25/04/2012	75 mg	Film-coated tablet	100	EU/1/98/069/004a	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/06/2012
Plavix	B2B Medical GmbH	25/04/2012	75 mg	Film-coated tablet	28	EU/1/98/069/001a	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/06/2012
Plavix	hvd medical GmbH	15/03/2012	75 mg	Film-coated tablet	100	EU/1/98/069/004a	Austria, Poland, The Netherlands	Germany	11/06/2012
Plavix	S.C. Farmastore Prest S.R.L.	08/10/2012	75 mg	Film-coated tablet	28	EU/1/98/069/001a	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, France, Germany, Greece, Hungary, Italy, Lithuania, Poland, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Romania	14/11/2012
Pradaxa	2 Care 4	14/12/2011	110 mg	Capsule, hard	60 x 1	EU/1/08/442/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	25/04/2012
Pradaxa	2 Care 4	14/12/2011	75 mg	Capsule, hard	60 x 1	EU/1/08/442/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	25/04/2012
Pradaxa	Cross Pharma AB	21/09/2012	150 mg	Capsule, hard	60 x 1 (unite dose)	EU/1/08/442/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012
Pradaxa	Cross Pharma AB	21/09/2012	150 mg	Capsule, hard	180 (3 x 60 x 1) (unite dose) (multipack)	EU/1/08/442/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/10/2012

Pradaxa	Paranova Läkemedel AB	24/01/2012	110 mg	Capsule, hard	30 x 1	EU/1/08/442/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/03/2012
Pradaxa	Paranova Läkemedel AB	24/01/2012	110 mg	Capsule, hard	60 x 1	EU/1/08/442/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	23/03/2012
Pradaxa	PCO Manufacturing	25/06/2012	110 mg	Capsule, hard	30 x 1 (unite dose)	EU/1/08/442/006	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Romania, Spain, Sweden, The Netherlands	Ireland, United Kingdom	10/07/2012
Pradaxa	PHARMACHIM AB	15/11/2012	110 mg	Capsule, hard	60 x 1 (unite dose)	EU/1/08/442/007	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/12/2012
Pradaxa	PHARMACHIM AB	15/11/2012	150 mg	Capsule, hard	60 x 1 (unite dose)	EU/1/08/442/011	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/12/2012
Pradaxa	PHARMACHIM AB	15/11/2012	75 mg	Capsule, hard	60 x 1 (unite dose)	EU/1/08/442/003	Austria, Belgium, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/12/2012
Preotact	ABACUS MEDICINE A/S	13/04/2012	100 µg	Powder and solvent for solution for injection	6 cartridges	EU/1/06/339/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/07/2012

Preotact	ABACUS MEDICINE A/S	18/04/2012	100 µg	Powder and solvent for solution for injection	2 cartridges	EU/1/06/339/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/07/2012
Prevenar 13	Copharma ApS	26/11/2012	- -	Suspension for injection	1 pre-filled syringe with separate needle	EU/1/09/590/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/12/2012
Prevenar 13	Copharma ApS	26/11/2012	- -	Suspension for injection	10 pre-filled syringes with separate needle	EU/1/09/590/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/12/2012
Previcox	CC Pharma	16/02/2012	57 mg	Chewable tablets	30	EU/2/04/045/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/03/2012
Previcox	CC Pharma	18/04/2012	227 mg	Chewable tablets	10	EU/2/04/045/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/04/2012
Previcox	CC Pharma	18/04/2012	227 mg	Chewable tablets	30	EU/2/04/045/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/04/2012
Prezista	Pharma Westen GmbH	10/05/2012	600 mg	Film-coated tablet	60	EU/1/06/380/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/05/2012
Prezista	Tisida GmbH	28/05/2012	400 mg	Film-coated tablet	60	EU/1/06/380/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012

Prezista	Tisida GmbH	28/05/2012	600 mg	Film-coated tablet	60	EU/1/06/380/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012
Pritor	CC Pharma	15/12/2011	80 mg	Tablet	98	EU/1/98/089/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/01/2012
Pritor	CC Pharma	31/05/2012	40 mg	Tablet	98	EU/1/98/089/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/07/2012
Pritor	kohlpharma GmbH	26/03/2012	80 mg	Tablet	98	EU/1/98/089/009	France	Germany	06/08/2012
Pritor	kohlpharma GmbH	11/05/2012	40 mg	Tablet	98	EU/1/98/089/004	France	Germany	06/08/2012
Pritor	WAYMADE PLC	15/10/2012	40 mg	Tablet	30	EU/1/98/089/021	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	25/10/2012
Pritor	WAYMADE PLC	15/10/2012	80 mg	Tablet	30	EU/1/98/089/022	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	25/10/2012
PritorPlus	Amimed Direct Ltd	30/11/2012	40 mg/ 12.5 mg	Tablet	28	EU/1/02/215/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	17/12/2012
PritorPlus	AMIMED DIRECT LTD	03/12/2012	80 mg/ 25mg	Tablet	28	EU/1/02/215/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	17/12/2012
PritorPlus	CC Pharma	06/12/2012	40 mg / 12.5 mg	Tablet	98	EU/1/02/215/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012

PritorPlus	CC Pharma	06/12/2012	80 mg / 25 mg	Tablet	98	EU/1/02/215/021	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
PritorPlus	CC Pharma	06/12/2012	80 mg/ 12.5 mg	Tablet	98	EU/1/02/215/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
PritorPlus	PHARMACHIM AB	25/04/2012	80 mg/ 12.5 mg	Tablet	28	EU/1/02/215/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/05/2012
PritorPlus	PHARMACHIM AB	25/04/2012	80 mg/ 12.5 mg	Tablet	98	EU/1/02/215/010	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/05/2012
Privigen	Axicorp Pharma B.V.	27/12/2011	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/003	Austria, Greece, Italy	Germany	30/01/2012
Privigen	axicorp Pharma B.V.	26/06/2012	100 mg/ml	Solution for infusion	3 vials	EU/1/08/446/005	Austria	Germany	27/07/2012
Privigen	axicorp Pharma B.V.	26/06/2012	100 mg/ml	Solution for infusion	3 vials	EU/1/08/446/006	Austria	Germany	27/07/2012
Privigen	Emra-Med	30/01/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Privigen	HAEMATO PHARM AG	23/03/2012	100 mg/ml	Solution for infusion	3 vials	EU/1/08/446/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/04/2012
Privigen	HAEMATO PHARM AG	23/03/2012	100 mg/ml	Solution for infusion	3 vials	EU/1/08/446/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/04/2012
Privigen	kohlpharma GmbH	27/06/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/002	Austria	Germany	10/07/2012

Privigen	kohlpharma GmbH	27/06/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/003	Austria	Germany	10/07/2012
Privigen	Pharma Westen GmbH	07/02/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/02/2012
Privigen	Pharma Westen GmbH	02/04/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/05/2012
Privigen	Tisida GmbH	08/06/2012	100 mg/ml	Solution for infusion	1 vial	EU/1/08/446/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/06/2012
Procoralan	COPHARMA ApS	26/04/2012	5 mg	Film-coated tablet	56	EU/1/05/316/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/04/2012
Procoralan	COPHARMA ApS	26/04/2012	7.5 mg	Film-coated tablet	56	EU/1/05/316/010	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/04/2012
Procoralan	COPHARMA ApS	30/04/2012	5 mg	Film-coated tablet	112	EU/1/05/316/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/05/2012
Procoralan	COPHARMA ApS	30/04/2012	7.5 mg	Film-coated tablet	112	EU/1/05/316/014	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/05/2012
Procoralan	Doncaster Pharmaceuticals Group Ltd	13/12/2012	7.5 mg	Film-coated tablet	56	EU/1/05/316/010	United Kingdom	Ireland, Malta	19/12/2012
Procoralan	Impexeco S.A.	11/07/2012	5 mg	Film-coated tablet	56	EU/1/05/316/003	Austria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Belgium	28/08/2012

Procoralan	Impexeco S.A.	12/07/2012	7.5 mg	Film-coated tablet	56	EU/1/05/316/010	Austria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Belgium	28/08/2012
Procoralan	PCO Manufacturing	27/02/2012	7.5 mg	Film-coated tablet	56	EU/1/05/316/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	04/04/2012
Procoralan	Pharma Lab	20/04/2012	5 mg	Film-coated tablet	56	EU/1/05/316/003	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	25/05/2012
Profender	CC Pharma	23/04/2012	150 mg praziquantel &30 mg emodepside	Modified-release tablets	2	EU/2/05/054/028	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/04/2012
Profender	CC Pharma	23/04/2012	150 mg praziquantel &30 mg emodepside	Modified-release tablets	24	EU/2/05/054/030	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/05/2012
Profender	CC Pharma	16/05/2012	15 mg praziquantel &3 mg emodepside	Modified-release tablets	24	EU/2/05/054/021	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/05/2012
Profender	CC Pharma	16/05/2012	50 mg praziquantel &10 mg emodepside	Modified-release tablets	24	EU/2/05/054/026	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/05/2012
Profender	CC Pharma	27/03/2012	50 mg praziquantel &10 mg emodepside	Modified-release tablets	2	EU/2/05/054/023	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/04/2012
Prolia	2 Care 4	12/07/2012	60 mg/ml	Solution for injection	1 pre-filled syringe with automatic needle guard (blistered)	EU/1/10/618/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/07/2012

Prolia	Cross Pharma AB	27/06/2012	60 mg/ml	Solution for injection	1 pre-filled syringe with automatic needle guard (blistered)	EU/1/10/618/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	11/07/2012
Prolia	kohlpharma GmbH	14/02/2012	60 mg/ml	Solution for injection	1 pre-filled syringe with automatic needle guard (blistered)	EU/1/10/618/003	Austria	Germany	17/02/2012
Prolia	Orifarm AB	30/07/2012	60 mg/ml	Solution for injection	1 pre-filled syringe with automatic needle guard (blistered)	EU/1/10/618/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/09/2012
Prolia	Orifarm AS	24/01/2012	60 mg/ml	Solution for injection	1 pre-filled syringe with automatic needle guard (blistered)	EU/1/10/618/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/03/2012
Protelos	Chemilines Ltd	22/11/2012	2 g	Granules for oral suspension	28 sachets	EU/1/04/288/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	03/12/2012
Protelos	Interport Ltd	06/12/2012	2 g	Granules for oral suspension	28 sachets	EU/1/04/288/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands	United Kingdom	17/12/2012
Protelos	Orifarm AB	30/07/2012	2 g	Granules for oral suspension	84 sachets	EU/1/04/288/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	07/09/2012
Protelos	Polyfarma Ltd	06/11/2012	2 g	Granules for oral suspension	28 sachets	EU/1/04/288/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/11/2012
Protopic	axicorp Pharma GmbH	14/03/2012	0.1%	Ointment	1 tube	EU/1/02/201/004	Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/04/2012

Protopic	B&S Healthcare	03/04/2012	0.03%	Ointment	1 tube	EU/1/02/201/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/05/2012
Protopic	B&S Healthcare	03/04/2012	0.1%	Ointment	1 tube	EU/1/02/201/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/05/2012
Protopic	Cross Pharma AB	13/02/2012	0.1%	Ointment	1 tube	EU/1/02/201/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012
Protopic	Cross Pharma AB	13/02/2012	0.1%	Ointment	1 tube	EU/1/02/201/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012
Protopic	Emra-Med	05/07/2012	0.1%	Ointment	1 tube	EU/1/02/201/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012
Protopic	EuroPharma DK ApS	04/04/2012	0.1%	Ointment	1 tube	EU/1/02/201/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	08/06/2012
Protopic	EuroPharma DK ApS	04/04/2012	0.1%	Ointment	1 tube	EU/1/02/201/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	08/06/2012
Puregon	2 Care 4	20/09/2012	900 IU/ 1.08 ml	Solution for injection	1 cartridge + 3 packs with 3 pen needles	EU/1/96/008/041	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/10/2012

Puregon	Axicorp Pharma B.V.	30/11/2012	600 IU/ 0.72 ml	Solution for injection	1 cartridge + 2 packs with 3 pen needles	EU/1/96/008/039	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/12/2012
Puregon	Eureco-Pharma B.V.	30/05/2012	300 IU/ 0.36 ml	Solution for injection	1 cartridge + 2 packs with 3 pen needles	EU/1/96/008/038	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	06/06/2012
Puregon	Eureco-Pharma B.V.	11/09/2012	900 IU/ 1.08 ml	Solution for injection	1 cartridge + 3 packs with 3 pen needles	EU/1/96/008/041	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	04/10/2012
Qutenza	HAEMATO PHARM AG	24/07/2012	179 mg	Cutaneous patch	1 patch + 1 cleansing gel	EU/1/09/524/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Rapamune	Axicorp Pharma B.V.	13/06/2012	1 mg/ml	Oral solution	1 bottle + 30 syringes + 1 syringe adapter + 1 carrying case	EU/1/01/171/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/06/2012
Rapamune	axicorp Pharma B.V.	05/12/2012	0.5 mg	Coated tablet	10 x 10	EU/1/01/171/014	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Rapamune	Beragena Arzneimittel GmbH	08/03/2012	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	11/04/2012
Rapamune	hvd medical GmbH	06/03/2012	1 mg	Coated tablet	100	EU/1/01/171/008	Italy, Spain, United Kingdom	Germany	05/04/2012

Rapamune	Orifarm AS	28/06/2012	1 mg/ml	Oral solution	1 bottle + 30 syringes + 1 syringe adapter + 1 carrying case	EU/1/01/171/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/07/2012
Rapamune	Paranova Danmark A/S	06/01/2012	2 mg	Coated tablet	100	EU/1/01/171/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/01/2012
Rapamune	Paranova Läkemedel AB	19/01/2012	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/02/2012
Rapamune	remedix Gmbh	23/11/2012	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/12/2012
Rasilez	ABACUS MEDICINE A/S	03/08/2012	150 mg	Film-coated tablet	98	EU/1/07/405/028	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012
Rasilez	ABACUS MEDICINE A/S	03/08/2012	300 mg	Film-coated tablet	98	EU/1/07/405/038	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012
Rasilez	ABACUS MEDICINE A/S	15/08/2012	150 mg	Film-coated tablet	28	EU/1/07/405/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012

Rasilez	Amimed Direct Ltd	02/10/2012	150 mg	Film-coated tablet	28	EU/1/07/405/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	31/10/2012
Rasilez	Amimed Direct Ltd	30/10/2012	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	13/11/2012
Rasilez	CC Pharma	22/02/2012	150 mg	Film-coated tablet	28	EU/1/07/405/022	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/03/2012
Rasilez	CC Pharma	22/02/2012	150 mg	Film-coated tablet	56	EU/1/07/405/025	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/03/2012
Rasilez	CC Pharma	22/02/2012	150 mg	Film-coated tablet	98	EU/1/07/405/028	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/03/2012
Rasilez	Clear Pharmacy	30/03/2012	150 mg	Film-coated tablet	28	EU/1/07/405/022	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	17/04/2012
Rasilez	Clear Pharmacy	30/03/2012	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	17/04/2012
Rasilez	CROSS HEALTHCARE LTD	28/03/2012	150 mg	Film-coated tablet	28	EU/1/07/405/022	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	14/05/2012

Rasilez	CROSS HEALTHCARE LTD	28/03/2012	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	14/05/2012
Rasilez	Eureco-Pharma B.V.	20/10/2011	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	The Netherlands	03/01/2012
Rasilez	Ginova Ltd	08/11/2012	150 mg	Film-coated tablet	30	EU/1/07/405/023	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/12/2012
Rasilez	Ginova Ltd	08/11/2012	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/12/2012
Rasilez	Kohlpharma GmbH	11/10/2012	300 mg	Film-coated tablet	28	EU/1/07/405/032	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/10/2012
Rasilez	Kohlpharma GmbH	11/10/2012	300 mg	Film-coated tablet	98	EU/1/07/405/038	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/10/2012
Rasilez HCT	Cambridge Major Laboratories Europe B.V	30/08/2012	150 mg / 12.5 mg	Film-coated tablet	98	EU/1/08/491/018	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/09/2012
Rasilez HCT	Cambridge Major Laboratories Europe B.V	03/09/2012	300 mg / 12.5 mg	Film-coated tablet	98	EU/1/08/491/058	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/09/2012
Rebif	axicorp Pharma B.V.	22/06/2012	44 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/009	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/07/2012
Rebif	axicorp Pharma B.V.	20/08/2012	22 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/008	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012

Rebif	axicorp Pharma B.V.	06/09/2012	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/09/2012
Rebif	cambridge Major Laboratories Europe B.V	12/03/2012	44 µg/0.5 ml	Solution for injection	12 cartridges	EU/1/98/063/019	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/08/2012
Rebif	cambridge Major Laboratories Europe B.V	12/03/2012	44 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/009	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/09/2012
Rebif	cambridge Major Laboratories Europe B.V	14/03/2012	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/09/2012
Rebif	cambridge Major Laboratories Europe B.V	14/03/2012	22 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/008	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/09/2012
Rebif	cambridge Major Laboratories Europe B.V	14/03/2012	22 µg/0.5 ml	Solution for injection	12 cartridges	EU/1/98/063/018	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/08/2012
Rebif	Cambridge Major Laboratories Europe B.V	14/03/2012	44 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/006	Austria, Belgium, Czech Republic, Finland, France, Greece, Ireland, Italy, Latvia, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/09/2012
Rebif	CC Pharma	18/10/2012	8.8 µg (2.4 m IU) + 22 µg (6 m IU)	Solution for injection	Initiation pack: 6 pre-filled pens of 8.8 mcg + 6 pre-filled pens of 22 mcg	EU/1/98/063/017	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/10/2012
Rebif	Cross Pharma AB	12/11/2012	44 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/11/2012

Rebif	Emra-Med	03/05/2012	22 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Rebif	Emra-Med	03/05/2012	44 µg/0.5 ml	Solution for injection	12 cartridges	EU/1/98/063/019	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Rebif	Emra-Med	03/05/2012	44 µg/0.5 ml	Solution for injection	4 cartridges	EU/1/98/063/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Rebif	kohlpharma GmbH	15/02/2012	44 µg/0.5 ml	Solution for injection	12 cartridges	EU/1/98/063/019	France	Germany	13/03/2012
Rebif	Orifarm AB	04/10/2012	22 µg	Solution for injection	12 pre-filled pens	EU/1/98/063/013	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/10/2012
Rebif	Orifarm AB	04/10/2012	44 µg	Solution for injection	12 pre-filled pens	EU/1/98/063/016	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/10/2012
Remicade	ABACUS MEDICINE A/S	05/11/2012	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/99/116/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/11/2012

Remicade	Eureco-Pharma B.V.	04/12/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	12/12/2012
Remicade	eurorx Arzneimittel GmbH	17/11/2010	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/07/2012
Remicade	Medartuum AB	19/06/2012	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/99/116/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/07/2012
Remicade	Medartuum AB	19/06/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/07/2012
Remicade	Orifarm ABOrifarm AB	23/04/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/06/2012
Remicade	Orifarm OY	23/04/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	16/05/2012
Remicade	PHARMACHIM AB	20/03/2012	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/99/116/001	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	02/05/2012
Remicade	PHARMACHIM AB	28/06/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/07/2012

Remicade	Veron Pharma Vertriebs GmbH	20/02/2012	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	22/03/2012
Remicade	Veron Pharma Vertriebs GmbH	20/02/2012	100 mg	Powder for concentrate for solution for infusion	4 vials	EU/1/99/116/004	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	22/03/2012
Remicade	Veron Pharma Vertriebs GmbH	20/02/2012	100 mg	Powder for concentrate for solution for infusion	5 vials	EU/1/99/116/005	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	22/03/2012
Renagel	ADL Pharma GmbH	13/03/2012	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	16/04/2012
Renvela	Euro Registratie Collectief BV	09/02/2012	2.4 g	Powder for oral suspension	60 sachets	EU/1/09/521/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	02/03/2012
Renvela	HAEMATO PHARM AG	13/04/2012	800 mg	Film-coated tablet	180 (without outer carton)	EU/1/09/521/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2012
Renvela	Medartuum AB	15/12/2011	800 mg	Film-coated tablet	180 (without outer carton)	EU/1/09/521/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/02/2012
Renvela	Orifarm AB	31/07/2012	800 mg	Film-coated tablet	180 (without outer carton)	EU/1/09/521/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/08/2012
Renvela	Paranova Läkemedel AB	28/03/2012	800 mg	Film-coated tablet	180 (without outer carton)	EU/1/09/521/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	27/06/2012

Renvela	Paranova Läkemedel AB	19/07/2012	2.4 g	Powder for oral suspension	90 sachets	EU/1/09/521/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/09/2012
Renvela	Paranova Oy	07/03/2012	800 mg	Film-coated tablet	180 (without outer carton)	EU/1/09/521/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	28/03/2012
Replagal	Orifarm AB	16/07/2012	1 mg/ml	Concentrate for solution for infusion	10 vials	EU/1/01/189/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	19/07/2012
Replagal	Orifarm AB	16/07/2012	1 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/01/189/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	19/07/2012
Replagal	Orifarm AB	16/07/2012	1 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/01/189/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	19/07/2012
Replagal	Orifarm AS	14/03/2012	1 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/01/189/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	23/04/2012
Revatio	cambridge Major Laboratories Europe B.V	12/03/2012	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/09/2012
Revatio	Combiphar Europe BV	21/02/2012	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/04/2012

Revatio	Paranova Oy	02/05/2012	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	19/06/2012
Revatio	REMEDIX GmbH	13/03/2012	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	02/05/2012
Revlimid	Euro Registratie Collectief BV	16/03/2012	10 mg	Capsule, hard	21	EU/1/07/391/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/04/2012
Revlimid	Euro Registratie Collectief BV	16/03/2012	15 mg	Capsule, hard	21	EU/1/07/391/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/04/2012
Revlimid	Euro Registratie Collectief BV	16/03/2012	25 mg	Capsule, hard	21	EU/1/07/391/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/04/2012
Revlimid	Orifarm AS	13/01/2012	25 mg	Capsule, hard	21	EU/1/07/391/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	08/03/2012
Reyataz	ABACUS MEDICINE A/S	04/09/2012	200 mg	Capsule, hard	60	EU/1/03/267/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/09/2012
Reyataz	ABACUS MEDICINE A/S	04/09/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/09/2012

Reyataz	ABACUS MEDICINE A/S	11/10/2012	300 mg	Capsule, hard	3 x 30	EU/1/03/267/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/10/2012
Reyataz	ADL Pharma GmbH	23/02/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	07/03/2012
Reyataz	B2B Medical GmbH	09/10/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
Reyataz	B2B Medical GmbH	09/10/2012	300 mg	Capsule, hard	3 x 30	EU/1/03/267/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Germany	23/10/2012
Reyataz	BB Farma s.r.l.	16/12/2011	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/01/2012
Reyataz	BB Farma s.r.l.	16/12/2011	300 mg	Capsule, hard	3 x 30	EU/1/03/267/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/01/2012
Reyataz	BR Pharma International Ltd	24/01/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/02/2012
Reyataz	BR Pharma International Ltd	31/01/2012	300 mg	Capsule, hard	3 x 30	EU/1/03/267/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/02/2012

Reyataz	Cross Pharma AB	30/11/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012
Reyataz	Emra-Med	17/09/2012	200 mg	Capsule, hard	60	EU/1/03/267/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/09/2012
Reyataz	Euro Registratie Collectief BV	13/03/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/03/2012
Reyataz	Haemato Pharm AG	05/01/2012	200 mg	Capsule, hard	60	EU/1/03/267/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/01/2012
Reyataz	Haemato Pharm AG	05/01/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/01/2012
Reyataz	HAEMATO PHARM AG	06/01/2012	150 mg	Capsule, hard	60	EU/1/03/267/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/01/2012
Reyataz	Inopha GmbH	07/09/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
Reyataz	Orifarm AB	18/07/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/09/2012

Reyataz	Paranova Läkemedel AB	07/08/2012	150 mg	Capsule, hard	60	EU/1/03/267/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/08/2012
Reyataz	s.c. HC PHARMA STORE s.r.l.	15/05/2012	150 mg	Capsule, hard	60	EU/1/03/267/004	Romania	Germany	05/06/2012
Reyataz	Tisida GmbH	08/06/2012	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/07/2012
Reyataz	Tisida GmbH	25/09/2012	300 mg	Capsule, hard	3 x 30	EU/1/03/267/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/10/2012
Reyataz	Veron Pharma Vertriebs GmbH	23/11/2011	150 mg	Capsule, hard	60	EU/1/03/267/004	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	04/01/2012
Reyataz	Veron Pharma Vertriebs GmbH	23/11/2011	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	04/01/2012
Rilutek	Paranova Oy	29/10/2012	50 mg	Film-coated tablet	56	EU/1/96/010/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	08/11/2012
Rilutek	STRATHCLYDE PHARMACEUTICALS LTD	24/08/2012	50 mg	Film-coated tablet	56	EU/1/96/010/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	11/09/2012

RoActemra	ABACUS MEDICINE A/S	18/01/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
RoActemra	ABACUS MEDICINE A/S	18/01/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
RoActemra	ABACUS MEDICINE A/S	18/01/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	23/01/2012
RoActemra	axicorp Pharma GmbH	09/05/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, Denmark, Estonia, Finland, France, Greece, Ireland, Italy, Latvia, Lithuania, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/06/2012
RoActemra	Beragena Arzneimittel GmbH	20/06/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	16/07/2012
RoActemra	Beragena Arzneimittel GmbH	20/06/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/003	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	16/07/2012
RoActemra	Beragena Arzneimittel GmbH	20/06/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/001	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	16/07/2012
RoActemra	Inopha GmbH	01/08/2012	20 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/08/492/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/08/2012
RoActemra	Inopha GmbH	01/08/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/08/2012

RoActemra	kohlpharma GmbH	26/04/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/001	Greece	Germany	14/05/2012
RoActemra	kohlpharma GmbH	26/04/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/003	Greece	Germany	14/05/2012
RoActemra	kohlpharma GmbH	26/04/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Greece	Germany	14/05/2012
RoActemra	kohlpharma GmbH	02/07/2012	20 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/08/492/002	Greece	Germany	12/07/2012
RoActemra	Orifarm AB	27/07/2012	20 mg/ml	Concentrate for solution for infusion	4 vials	EU/1/08/492/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/08/2012
RoActemra	Orifarm AB	27/07/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/08/2012
RoActemra	Orifarm AB	27/07/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/08/2012
RoActemra	Orifarm AB	27/07/2012	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/08/2012
Sifrol	2 Care 4	16/12/2011	0.52 mg	Prolonged-release tablet	100	EU/1/97/050/018	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/02/2012
Sifrol	2 Care 4	16/12/2011	1.05 mg	Prolonged-release tablet	100	EU/1/97/050/021	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/02/2012
Sifrol	2 Care 4	16/12/2011	2.1 mg	Prolonged-release tablet	100	EU/1/97/050/024	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/02/2012

Sifrol	Axicorp Pharma B.V.	20/12/2011	2.1 mg	Prolonged-release tablet	100	EU/1/97/050/024	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/01/2012
Sifrol	Axicorp Pharma B.V.	20/12/2011	3.15 mg	Prolonged-release tablet	100	EU/1/97/050/027	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/01/2012
Sifrol	Axicorp Pharma B.V.	25/05/2012	1.05 mg	Prolonged-release tablet	100	EU/1/97/050/021	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/05/2012
Sifrol	BB Farma s.r.l.	08/06/2012	1.05 mg	Prolonged-release tablet	30	EU/1/97/050/020	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Sifrol	BB Farma s.r.l.	08/06/2012	1.05 mg	Prolonged-release tablet	100	EU/1/97/050/021	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Sifrol	BB Farma s.r.l.	11/06/2012	3.15 mg	Prolonged-release tablet	30	EU/1/97/050/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Sifrol	BB Farma s.r.l.	11/06/2012	3.15 mg	Prolonged-release tablet	100	EU/1/97/050/027	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/07/2012
Sifrol	Beragena Arzneimittel GmbH	19/06/2012	1.05 mg	Prolonged-release tablet	30	EU/1/97/050/020	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	09/07/2012
Sifrol	Beragena Arzneimittel GmbH	19/06/2012	1.05 mg	Prolonged-release tablet	100	EU/1/97/050/021	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	06/07/2012

Sifrol	Beragena Arzneimittel GmbH	09/07/2012	2.1 mg	Prolonged-release tablet	100	EU/1/97/050/024	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	27/07/2012
Sifrol	Beragena Arzneimittel GmbH	09/07/2012	2.1 mg	Prolonged-release tablet	30	EU/1/97/050/023	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	27/07/2012
Sifrol	Beragena Arzneimittel GmbH	09/07/2012	3.15 mg	Prolonged-release tablet	30	EU/1/97/050/026	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	31/07/2012
Sifrol	Beragena Arzneimittel GmbH	09/07/2012	3.15 mg	Prolonged-release tablet	100	EU/1/97/050/027	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	27/07/2012
Sifrol	Beragena Arzneimittel GmbH	26/07/2012	0.52 mg	Prolonged-release tablet	30	EU/1/97/050/017	Austria	Germany	02/08/2012
Sifrol	Beragena Arzneimittel GmbH	26/07/2012	0.52 mg	Prolonged-release tablet	100	EU/1/97/050/018	Austria, Belgium, Finland, France, Greece, Italy, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/08/2012
Sifrol	hvd medical GmbH	16/03/2012	1.05 mg	Prolonged-release tablet	30	EU/1/97/050/020	Austria, The Netherlands	Germany	27/03/2012
Sifrol	MTK Pharma GmbH	26/04/2012	2.62 mg	Prolonged-release tablet	30	EU/1/97/050/032	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/05/2012
Sifrol	MTK Pharma GmbH	26/04/2012	2.62 mg	Prolonged-release tablet	100	EU/1/97/050/033	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Silgard	Paranova Danmark AS	26/09/2012	--	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/358/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/10/2012
Silodyx	BCN Farma SL	20/06/2012	8 mg	Capsule, hard	30	EU/1/09/607/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	18/07/2012

Silodyx	Euro Registratie Collectief BV	21/02/2012	4 mg	Capsule, hard	30	EU/1/09/607/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/03/2012
Silodyx	Euro Registratie Collectief BV	21/02/2012	8 mg	Capsule, hard	30	EU/1/09/607/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/03/2012
Silodyx	Medcor Pharmaceuticals BV	05/01/2012	8 mg	Capsule, hard	30	EU/1/09/607/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/01/2012
Silodyx	Medcor Pharmaceuticals BV	15/03/2012	4 mg	Capsule, hard	30	EU/1/09/607/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/03/2012
Simponi	Axicorp Pharma B.V.	08/12/2011	50 mg	Solution for injection in pre-filled syringe	1 pre-filled syringe	EU/1/09/546/003	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Lithuania, Luxembourg, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/01/2012
Simponi	CC Pharma	26/09/2012	50 mg	Solution for injection in pre-filled syringe	3 (3 x 1) pre-filled syringes	EU/1/09/546/004	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/10/2012
Somavert	HAEMATO PHARM AG	15/12/2011	20 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
Somavert	HAEMATO PHARM AG	10/01/2012	10 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012

Somavert	HAEMATO PHARM AG	10/01/2012	15 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
SonoVue	Orifarm AS	29/02/2012	8 µl/ml	Powder and solvent for dispersion for injection	1 vial + 1 pre-filled syringe + 1 'MiniSpike' rec. device	EU/1/01/177/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/03/2012
Sprycel	axicorp Pharma B.V.	11/10/2012	100 mg	Film-coated tablet	30 x 1 (unit dose)	EU/1/06/363/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
Sprycel	axicorp Pharma B.V.	11/10/2012	50 mg	Film-coated tablet	60 x 1 (unit dose)	EU/1/06/363/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
Sprycel	Dr Fisher Farma BV	08/05/2012	100 mg	Film-coated tablet	30 x 1 (unit dose)	EU/1/06/363/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/05/2012
Sprycel	Emra-Med	31/07/2012	100 mg	Film-coated tablet	30 x 1 (unit dose)	EU/1/06/363/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	22/08/2012
Sprycel	HAEMATO PHARM AG	12/07/2012	100 mg	Film-coated tablet	30	EU/1/06/363/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/07/2012
Sprycel	Omnia Läkemedel AB	10/04/2012	100 mg	Film-coated tablet	30 x 1 (unit dose)	EU/1/06/363/011	Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/04/2012
Sprycel	Orifarm AB	30/07/2012	50 mg	Film-coated tablet	60 x 1 (unit dose)	EU/1/06/363/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	07/09/2012

Stalevo	2 Care 4	13/08/2012	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/007	Austria, Belgium, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Stalevo	ADL Pharma GmbH	09/02/2012	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/03/2012
Stalevo	axicorp Pharma GmbH	13/02/2012	150 mg / 37.5 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/011	Italy, Portugal, United Kingdom	Germany	01/03/2012
Stalevo	BR Pharma International Ltd	26/01/2012	175 mg / 43.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/036	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/03/2012
Stalevo	BR Pharma International Ltd	16/05/2012	200 mg / 50 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2012
Stalevo	CC Pharma	06/08/2012	125 mg / 31.25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/031	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/08/2012
Stalevo	CC Pharma	06/08/2012	200 mg / 50 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/021	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/08/2012
Stalevo	CC Pharma	06/08/2012	75 mg / 18.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/026	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/08/2012
Stalevo	CC Pharma	09/10/2012	200 mg / 50 mg / 200 mg	Film-coated tablet	30	EU/1/03/260/020	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/10/2012
Stalevo	CD Pharma Limited	14/02/2012	75 mg / 18.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/02/2012

Stalevo	CD Pharma Limited	15/02/2012	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/02/2012
Stalevo	CD Pharma Limited	15/02/2012	125 mg / 31.25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/02/2012
Stalevo	Docpharm GmbH&CoKGaA	14/11/2011	125 mg / 31.25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/031	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/01/2012
Stalevo	Docpharm GmbH&CoKGaA	14/11/2011	75 mg / 18.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/01/2012
Stalevo	Doncaster Pharmaceuticals Group Ltd	25/09/2012	75 mg / 18.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/026	United Kingdom	Ireland, Malta	08/10/2012
Stalevo	EurimPharm Arzneimittel GmbH	23/11/2012	175 mg / 43.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/036	Finland	Germany	21/12/2012
Stalevo	kohlpharma GmbH	16/04/2012	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/036	United Kingdom	Germany	03/08/2012
Stalevo	Medcor Pharmaceuticals BV	21/09/2012	125 mg / 31.25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/10/2012
Stalevo	PCO Manufacturing	16/02/2012	125 mg / 31.25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/031	United Kingdom	Ireland	19/03/2012
Stalevo	PCO Manufacturing	16/02/2012	200 mg / 50 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/021	United Kingdom	Ireland	19/03/2012
Stalevo	PCO Manufacturing	16/02/2012	75 mg / 18.75 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/026	United Kingdom	Ireland	19/03/2012

Stalevo	REMEDIX GmbH	13/04/2012	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	27/04/2012
Stalevo	WAYMADE PLC	31/07/2012	150 mg / 37.5 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	07/08/2012
Stocrin	NewNeopharm BV	10/01/2012	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/01/2012
Stocrin	NewNeopharm BV	16/10/2012	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	26/11/2012
Stocrin	NewNeopharm BV	27/11/2012	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	06/12/2012
Stocrin	PHARMACHIM AB	10/12/2012	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	21/12/2012
Stronghold	CC Pharma	16/02/2012	15 mg	Spot-on solution	0.25 ml (60mg/ml) Unit dose tubes (polypropylene)	EU/2/99/014/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/03/2012
Stronghold	CC Pharma	18/04/2012	30 mg	Spot-on solution	3 tubes	EU/2/99/014/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/05/2012

Stronghold	CC Pharma	31/05/2012	30 mg	Spot-on solution	6 x (0.25 ml (60mg/ml) Unit dose tubes (polypropylene)	EU/2/99/014/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Stronghold	CC Pharma	31/05/2012	45 mg	Spot-on solution	3 x (0.75 ml (60mg/ml) Unit dose tubes (polypropylene)	EU/2/99/014/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Stronghold	CC Pharma	31/05/2012	45 mg	Spot-on solution	6 x (0.75 ml (60mg/ml)) Unit dose tubes (polypropylene)	EU/2/99/014/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/06/2012
Stronghold	CC Pharma	19/06/2012	60 mg	Spot-on solution	3 x 0.5 ml (120mg/ml) Unit dose tubes (polypropylene)	EU/2/99/014/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/07/2012
Stronghold	CC Pharma	19/06/2012	60 mg	Spot-on solution	6 x 0.5 ml (120 mg/ml)Unit dose tubes (polypropylene)	EU/2/99/014/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/07/2012
Stronghold	CC Pharma	07/08/2012	120 mg	Spot-on solution	Unit dose tubes (polypropylene) 1.0ml (120mg/ml)	EU/2/99/014/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012
Stronghold	CC Pharma	07/08/2012	120 mg	Spot-on solution	Unit dose tubes (polypropylene) 1.0 ml (120mg/ml)	EU/2/99/014/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012

Stronghold	CC Pharma	11/10/2012	240 mg	Unit dose tubes (polypropylene) 2.0 ml(120mg/ml)	3 tubes	EU/2/99/014/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
Stronghold	CC Pharma	11/10/2012	240 mg	Unit dose tubes (polypropylene) 2.0 ml (120mg/ml)	6 tubes	EU/2/99/014/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/10/2012
Suboxone	Chemilines Ltd	15/08/2007	2 mg/0.5 mg	Sublingual tablet	28	EU/1/06/359/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	20/03/2012
Suboxone	Chemilines Ltd	15/08/2007	8 mg/2 mg	Sublingual tablet	28	EU/1/06/359/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	20/03/2012
Suboxone	Chemilines Ltd	10/10/2012	2 mg/0.5 mg	Sublingual tablet	7	EU/1/06/359/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	25/10/2012
Suboxone	Chemilines Ltd	10/10/2012	8 mg/2 mg	Sublingual tablet	7	EU/1/06/359/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	25/10/2012
Suboxone	Cross Healthcare Ltd	20/09/2012	2 mg/0.5 mg	Sublingual tablet	28	EU/1/06/359/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	28/09/2012
Suboxone	Cross Healthcare Ltd	20/09/2012	8 mg/2 mg	Sublingual tablet	28	EU/1/06/359/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	28/09/2012
Suboxone	MPT Pharma	12/11/2012	2 mg/0.5 mg	Sublingual tablet	7	EU/1/06/359/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	27/11/2012

Suboxone	MPT Pharma	12/11/2012	2 mg/0.5 mg	Sublingual tablet	28	EU/1/06/359/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	27/11/2012
Suprelorin	2 Care 4	09/11/2012	4.7 mg	Implant	2 implants preloaded in implanters + 1 actuator	EU/2/07/072/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	26/11/2012
Sustiva	axicorp Pharma B.V.	06/12/2012	600 mg	Film-coated tablet	30	EU/1/99/110/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/12/2012
Sustiva	CC Pharma	30/03/2012	200 mg	Capsule, hard	90	EU/1/99/110/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/04/2012
Sustiva	Emra-Med	14/05/2012	600 mg	Film-coated tablet	30	EU/1/99/110/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Sustiva	Emra-Med	24/07/2012	600 mg	Film-coated tablet	9 (3 x 30 x 1)	EU/1/99/110/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2012
Sutent	Cancernova GmbH onkologische Arzneimittel	23/05/2012	25 mg	Capsule, hard	30	EU/1/06/347/002	Greece	Germany	28/05/2012
Sutent	Cancernova GmbH onkologische Arzneimittel	23/05/2012	50 mg	Capsule, hard	30	EU/1/06/347/003	Greece	Germany	28/05/2012
Sutent	Cancernova GmbH onkologische Arzneimittel	26/11/2012	25 mg	Capsule, hard	28	EU/1/06/347/005	United Kingdom	Germany	07/12/2012

Sutent	Cancernova GmbH onkologische Arzneimittel	26/11/2012	50 mg	Capsule, hard	28	EU/1/06/347/006	United Kingdom	Germany	07/12/2012
Sutent	Dr Fisher Farma BV	22/06/2012	12.5 mg	Capsule, hard	30	EU/1/06/347/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/08/2012
Sutent	Dr Fisher Farma BV	22/06/2012	25 mg	Capsule, hard	30	EU/1/06/347/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/08/2012
Sutent	Dr Fisher Farma BV	22/06/2012	50 mg	Capsule, hard	30	EU/1/06/347/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/08/2012
Sutent	Omnia Läkemedel AB	16/02/2012	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Spain, The Netherlands, United Kingdom	Sweden	29/03/2012
Sutent	Orifarm AB	09/07/2012	25 mg	Capsule, hard	30	EU/1/06/347/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	17/07/2012
Sutent	Orifarm AB	06/08/2012	12.5 mg	Capsule, hard	30	EU/1/06/347/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/08/2012
Sutent	Orifarm AB	06/08/2012	50 mg	Capsule, hard	30	EU/1/06/347/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/08/2012
Sycrest	Axicorp Pharma GmbH	24/05/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Bulgaria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Lithuania, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012

Sycrest	Axicorp Pharma GmbH	24/05/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Bulgaria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Lithuania, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/06/2012
Sycrest	B2B Medical GmbH	29/08/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/10/2012
Sycrest	B2B Medical GmbH	29/08/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/10/2012
Sycrest	Beragena Arzneimittel GmbH	29/11/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	13/12/2012
Sycrest	Beragena Arzneimittel GmbH	29/11/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	13/12/2012
Sycrest	BR Pharma International Ltd	22/02/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Sycrest	BR Pharma International Ltd	22/02/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Sycrest	CC Pharma	11/05/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012
Sycrest	CC Pharma	11/05/2012	10 mg	Sublingual tablet	20	EU/1/10/640/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012
Sycrest	CC Pharma	11/05/2012	5 mg	Sublingual tablet	20	EU/1/10/640/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012

Sycrest	CC Pharma	11/05/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/05/2012
Sycrest	Emra-Med	03/05/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Sycrest	Emra-Med	03/05/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Sycrest	HAEMATO PHARM AG	13/04/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/04/2012
Sycrest	HAEMATO PHARM AG	13/04/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/04/2012
Sycrest	kohlpharma GmbH	14/03/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Italy	Germany	16/04/2012
Sycrest	kohlpharma GmbH	14/03/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Italy	Germany	16/04/2012
Sycrest	MPT Pharma	16/04/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	10/05/2012

Sycrest	NewNeopharm BV	20/12/2011	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/01/2012
Sycrest	NewNeopharm BV	21/12/2011	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/01/2012
Sycrest	Pharma Westen GmbH	06/02/2012	10 mg	Sublingual tablet	60	EU/1/10/640/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012
Sycrest	Pharma Westen GmbH	06/02/2012	5 mg	Sublingual tablet	60	EU/1/10/640/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012
Synagis	Delfarma SpZoo	25/04/2012	100 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/99/117/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	03/05/2012
Synflorix	PHARMACHIM AB	25/03/2012	- -	Suspension for injection	1 pre-filled syringe + 1 needle	EU/1/09/508/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/05/2012
TachoSil	Delfarma SpZoo	25/11/2011	- -	Medicated sponge	1 sponge of 3.0 x 2.5 cm	EU/1/04/277/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	04/01/2012
TachoSil	Medartuum AB	07/11/2012	2 IU/cm² / 5.5 mg/cm²	Medicated sponge	1 sponge of 9.5 x 4.8 cm	EU/1/04/277/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/11/2012

TachoSil	Paranova Läkemedel AB	14/09/2012	- -	Medicated sponge	1 sponge of 9.5 x 4.8 cm	EU/1/04/277/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	27/09/2012
Tarceva	ABACUS MEDICINE A/S	27/12/2011	100 mg	Film-coated tablet	30	EU/1/05/311/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/01/2012
Tarceva	ABACUS MEDICINE A/S	27/12/2011	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/01/2012
Tarceva	Cancernova GmbH onkologische Arzneimittel	05/03/2012	100 mg	Film-coated tablet	30	EU/1/05/311/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/03/2012
Tarceva	Cancernova GmbH onkologische Arzneimittel	05/03/2012	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/03/2012
Tarceva	eurorx Arzneimittel GmbH	17/11/2010	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/06/2012
Tarceva	Orifarm AB	14/12/2011	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/02/2012

Targretin	ABACUS MEDICINE A/S	09/08/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/08/2012
Targretin	Emra-Med	11/04/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/04/2012
Targretin	kohlpharma GmbH	11/01/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Belgium, Greece, The Netherlands	Germany	01/03/2012
Targretin	Orifarm AS	17/07/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/09/2012
Targretin	Paranova Läkemedel AB	19/06/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	05/07/2012
Targretin	Pharma Westen GmbH	19/01/2012	75 mg	Capsule, soft	100	EU/1/01/178/001	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
Tasigna	CC Pharma	09/07/2012	200 mg	Capsule, hard	112 (4x28) (multipack)	EU/1/07/422/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/08/2012
Tasigna	Emra-Med	15/10/2012	200 mg	Capsule, hard	112 (4x28) (multipack)	EU/1/07/422/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/10/2012
Tasigna	Orifarm AB	27/09/2012	200 mg	Capsule, hard	112 (4x28) (multipack)	EU/1/07/422/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	26/10/2012

Tasigna	Orifarm AS	27/09/2012	200 mg	Capsule, hard	112 (4x28) (multipack)	EU/1/07/422/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/10/2012
Tasmar	kohlpharma GmbH	27/04/2012	100 mg	Film-coated tablet	200 (2 X 100) multipack	EU/1/97/044/009	Italy	Germany	25/06/2012
Temodal	Docpharm GmbH & CoKGaA	15/02/2012	100 mg	Capsule, hard	5	EU/1/98/096/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/02/2012
Temodal	Docpharm GmbH & CoKGaA	15/02/2012	100 mg	Capsule, hard	20	EU/1/98/096/016	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/04/2012
Thyrogen	ABACUS MEDICINE A/S	31/07/2012	0.9 mg	Powder for solution for injection	2 vials	EU/1/99/122/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/08/2012
TOBI Podhaler	ABACUS MEDICINE A/S	20/07/2012	28 mg	Inhalation powder, hard capsule	224 (4x 56) + 5 inhalers (monthly multipack)	EU/1/10/652/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/08/2012
TOBI Podhaler	kohlpharma GmbH	02/07/2012	28 mg	Inhalation powder, hard capsule	224 (4x 56) + 5 inhalers (monthly multipack)	EU/1/10/652/ 002	Austria	Germany	13/07/2012
TOBI Podhaler	Paranova Danmark AS	29/06/2012	28 mg	Inhalation powder, hard capsule	224 (4x 56) + 5 inhalers (monthly multipack)	EU/1/10/652/ 002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/07/2012

Toviaz	ABACUS MEDICINE A/S	02/08/2012	4 mg	Prolonged-release tablet	28	EU/1/07/386/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/08/2012
Toviaz	ABACUS MEDICINE A/S	02/08/2012	4 mg	Prolonged-release tablet	84	EU/1/07/386/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/08/2012
Toviaz	ABACUS MEDICINE A/S	02/08/2012	8 mg	Prolonged-release tablet	28	EU/1/07/386/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/08/2012
Toviaz	ABACUS MEDICINE A/S	02/08/2012	8 mg	Prolonged-release tablet	84	EU/1/07/386/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/08/2012
Toviaz	B&S Healthcare	12/09/2012	4 mg	Prolonged-release tablet	28	EU/1/07/386/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	16/10/2012
Toviaz	BB Farma s.r.l.	31/07/2012	8 mg	Prolonged-release tablet	14	EU/1/07/386/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	16/08/2012

Toviaz	BB Farma s.r.l.	31/07/2012	8 mg	Prolonged-release tablet	28	EU/1/07/386/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	16/08/2012
Toviaz	Copharma ApS	31/05/2012	4 mg	Prolonged-release tablet	28	EU/1/07/386/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/06/2012
Toviaz	Copharma ApS	31/05/2012	4 mg	Prolonged-release tablet	84	EU/1/07/386/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/06/2012
Toviaz	Euromedicines S.L.	13/12/2011	8 mg	Prolonged-release tablet	28	EU/1/07/386/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	03/01/2012
Toviaz	WAYMADE PLC	18/09/2012	8 mg	Prolonged-release tablet	28	EU/1/07/386/008	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	12/11/2012
Tracleer	Medartuum AB	06/09/2012	125 mg	Film-coated tablet	56	EU/1/02/220/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	01/10/2012
Tracleer	Medartuum AB	18/10/2012	62.5 mg	Film-coated tablet	56	EU/1/02/220/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	26/10/2012
Tracleer	Omnia Läkemedel AB	28/03/2012	125 mg	Film-coated tablet	56	EU/1/02/220/004	Bulgaria, France, Germany, Italy, Poland, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	25/04/2012

Trajenta	Medcor Pharmaceuticals BV	29/11/2012	5 mg	Film-coated tablet	30	EU/1/11/707/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/12/2012
Travatan	ABACUS MEDICINE A/S	14/08/2012	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/09/2012
Travatan	ABACUS MEDICINE A/S	14/08/2012	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/09/2012
Travatan	Copharma ApS	25/09/2012	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/10/2012
Travatan	Copharma ApS	25/09/2012	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/10/2012
Travatan	EuroPharma DK ApS	11/09/2012	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/09/2012
Travatan	EuroPharma DK ApS	11/09/2012	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Cyprus, Czech Republic, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/09/2012
Travatan	O.P.D. Laboratories Ltd	19/10/2012	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands	United Kingdom	08/11/2012

Travatan	Paranova Oy	22/03/2012	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	02/05/2012
Travatan	PHARMACHIM AB	11/10/2012	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Poland, Portugal, Spain, The Netherlands, United Kingdom	Sweden	29/10/2012
Tredaptive	kohlpharma GmbH	01/08/2012	1000 mg / 20 mg	Modified-release tablet	28	EU/1/08/459/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/08/2012
Tredaptive	Swingward Ltd TA Medihealth	21/05/2012	1000 mg / 20 mg	Modified-release tablet	28	EU/1/08/459/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	12/06/2012
Trizivir	BB Farma s.r.l.	11/07/2012	- -	Film-coated tablet	60	EU/1/00/156/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/07/2012
Trizivir	Dr Fisher Farma BV	27/03/2012	- -	Film-coated tablet	60	EU/1/00/156/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, United Kingdom	The Netherlands	12/06/2012
Trizivir	Euro Registratie Collectief BV	12/01/2012	- -	Film-coated tablet	60	EU/1/00/156/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/02/2012
Trizivir	Medicopharm AG	12/09/2012	- -	Film-coated tablet	60	EU/1/00/156/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	11/10/2012
Trobalt	Emra-Med	01/06/2012	100 mg	Film-coated tablet	168	EU/1/11/681/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012

Trobalt	Emra-Med	01/06/2012	100 mg	Film-coated tablet	84	EU/1/11/681/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012
Trobalt	Emra-Med	01/06/2012	200 mg	Film-coated tablet	84	EU/1/11/681/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012
Trobalt	Emra-Med	01/06/2012	400 mg	Film-coated tablet	84	EU/1/11/681/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012
Trobalt	Emra-Med	01/06/2012	50 mg/100mg	Film-coated tablet	Initiation pack: 21 tablets 50 mg + 42 tablets 100 mg	EU/1/11/681/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012
Trobalt	kohlpharma GmbH	03/08/2012	50 mg	Film-coated tablet	168	EU/1/11/681/003	Austria	Germany	15/08/2012
Trobalt	kohlpharma GmbH	04/09/2012	100 mg	Film-coated tablet	168	EU/1/11/681/006	Austria	Germany	04/10/2012
Trobalt	kohlpharma GmbH	04/09/2012	200 mg	Film-coated tablet	168 (2 x 84) (multipack)	EU/1/11/681/008	Austria	Germany	04/10/2012
Trobalt	kohlpharma GmbH	04/09/2012	300 mg	Film-coated tablet	168 (2 x 84) (multipack)	EU/1/11/681/010	Austria	Germany	04/10/2012
Trobalt	Orifarm AS	12/01/2012	400 mg	Film-coated tablet	84	EU/1/11/681/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/02/2012
Twinrix Adult	ABACUS MEDICINE A/S	27/06/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	11/07/2012

Twinrix Adult	ABACUS MEDICINE A/S	14/08/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	28/08/2012
Twinrix Adult	ADL Pharma GmbH	14/11/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Latvia, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/11/2012
Twinrix Adult	EuroPharma DK ApS	06/09/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/10/2012
Twinrix Adult	Paranova Danmark AS	02/04/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/05/2012
Twinrix Adult	PHARMACHIM AB	29/02/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	10/04/2012
Twinrix Adult	PI-Pharma N.V.	19/11/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Belgium, Luxembourg	30/11/2012
Twinrix Adult	Tisida GmbH	11/06/2012	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/07/2012
Twinrix Paediatric	CC Pharma	09/07/2012	- -	Suspension for injection	10 pre-filled syringes + 10 separate needles	EU/1/97/029/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/07/2012

Twinrix Paediatric	PHARMACHIM AB	22/03/2012	- -	Suspension for injection	1 pre-filled syringe + 1 separate needle	EU/1/97/029/006	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	11/04/2012
Twynsta	CC Pharma	14/11/2012	40 mg/10 mg	Tablet	98	EU/1/10/648/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Twynsta	CC Pharma	14/11/2012	40 mg/5 mg	Tablet	98	EU/1/10/648/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Twynsta	CC Pharma	14/11/2012	80 mg/10 mg	Tablet	98	EU/1/10/648/027	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
Twynsta	CC Pharma	14/11/2012	80 mg/5 mg	Tablet	98	EU/1/10/648/020	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/11/2012
TWYNSTA	Doncaster Pharmaceuticals Group Ltd	17/04/2012	80 mg/10 mg	Tablet	28	EU/1/10/648/023	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Italy, Liechtenstein, Luxembourg, Portugal, Slovak Republic, Spain, Sweden, The Netherlands	Ireland, Malta, United Kingdom	21/08/2012
TWYNSTA	Doncaster Pharmaceuticals Group Ltd	17/04/2012	80 mg/5 mg	Tablet	28	EU/1/10/648/016	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Italy, Liechtenstein, Luxembourg, Portugal, Slovak Republic, Spain, Sweden, The Netherlands	Ireland, Malta, United Kingdom	21/08/2012
TWYNSTA	Emra-Med	11/04/2012	80 mg/5 mg	Tablet	98	EU/1/10/648/020	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/05/2012
Twynsta	Emra-Med	12/07/2012	80 mg/10 mg	Tablet	56	EU/1/10/648/025	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/08/2012

Twynsta	Emra-Med	12/07/2012	80 mg/10 mg	Tablet	98	EU/1/10/648/027	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/08/2012
TWYNSTA	kohlpharma GmbH	13/04/2012	80 mg/10 mg	Tablet	56	EU/1/10/648/025	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/04/2012
TWYNSTA	kohlpharma GmbH	13/04/2012	80 mg/5 mg	Tablet	56	EU/1/10/648/018	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/04/2012
Tysabri	ABACUS MEDICINE A/S	12/01/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/01/2012
Tysabri	Cambridge Major Laboratories Europe B.V	05/03/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Tysabri	Orifarm AS	30/07/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/08/2012
Tysabri	Paranova Danmark AS	01/03/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/03/2012
Tysabri	Paranova Läkemedel AB	13/03/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	17/04/2012
Tysabri	PHARMACHIM AB	08/06/2012	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	15/06/2012

Tyverb	Haemato Pharm AG	17/12/2012	250 mg	Film-coated tablet	70	EU/1/07/440/004	Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Austria	21/12/2012
Tyverb	kohlpharma GmbH	27/02/2012	250 mg	Film-coated tablet	70	EU/1/07/440/001	Czech Republic, Greece	Germany	01/03/2012
Tyverb	kohlpharma GmbH	27/02/2012	250 mg	Film-coated tablet	70	EU/1/07/440/004	Czech Republic, Greece	Germany	14/03/2012
Tyverb	Medartuum AB	22/11/2012	250 mg	Film-coated tablet	70	EU/1/07/440/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	26/11/2012
Tyverb	Orifarm AS	09/08/2012	250 mg	Film-coated tablet	70	EU/1/07/440/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/09/2012
Tyverb	Orifarm AS	14/08/2012	250 mg	Film-coated tablet	140	EU/1/07/440/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/09/2012
Tyverb	Paranova Oy	02/05/2012	250 mg	Film-coated tablet	70	EU/1/07/440/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	14/05/2012
Urorec	CC Pharma	04/06/2012	8 mg	Capsule, hard	30	EU/1/09/608/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012

Urorec	kohlpharma GmbH	27/02/2012	4 mg	Capsule, hard	50	EU/1/09/608/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/03/2012
Urorec	Mediwin Ltd	20/07/2012	4 mg	Capsule, hard	30	EU/1/09/608/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	30/08/2012
Urorec	Mediwin Ltd	20/07/2012	8 mg	Capsule, hard	30	EU/1/09/608/011	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	30/08/2012
Urorec	PCO Manufacturing	16/02/2012	8 mg	Capsule, hard	30	EU/1/09/608/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	09/03/2012
Valdoxan	axicorp Pharma GmbH	20/12/2011	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/01/2012
Valdoxan	Beragena Arzneimittel GmbH	20/06/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	19/07/2012
Valdoxan	Beragena Arzneimittel GmbH	20/06/2012	25 mg	Film-coated tablet	98	EU/1/08/499/007	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	19/07/2012
Valdoxan	CC Pharma	14/12/2011	25 mg	Film-coated tablet	98	EU/1/08/499/007	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Valdoxan	CC Pharma	14/12/2011	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/01/2012
Valdoxan	Clear Pharmacy	19/12/2011	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	23/01/2012
Valdoxan	Ebb Medical AB	29/11/2012	25 mg	Film-coated tablet	84	EU/1/08/499/006	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012

Valdoxan	Ebb Medical AB	29/11/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	10/12/2012
Valdoxan	HAEMATO PHARM AG Haemato Pharm AG	13/04/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/05/2012
Valdoxan	Medartuum AB	11/09/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/09/2012
Valdoxan	Medartuum AB	11/09/2012	25 mg	Film-coated tablet	84	EU/1/08/499/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/09/2012
Valdoxan	O.P.D. Laboratories Ltd	16/07/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	26/07/2012
Valdoxan	Paranova Läkemedel AB	12/12/2012	25 mg	Film-coated tablet	84	EU/1/08/499/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/12/2012
Valdoxan	PHARMACHIM AB	29/02/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012
Valdoxan	PHARMACHIM AB	29/02/2012	25 mg	Film-coated tablet	84	EU/1/08/499/006	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012
Valdoxan	PharmaCoDane ApS	10/09/2012	25 mg	Film-coated tablet	28	EU/1/08/499/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/10/2012

Valdoxan	PharmaCoDane ApS	10/10/2012	25 mg	Film-coated tablet	84	EU/1/08/499/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/10/2012
Valdoxan	REMEDIX GMBH	07/08/2012	25 mg	Film-coated tablet	98	EU/1/08/499/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	11/10/2012
VANTAVO	Dr Fisher Farma BV	07/02/2012	70 mg/2800 IU	Tablet	4	EU/1/09/572/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/03/2012
VELCADE	Haemato Pharm AG	08/12/2011	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Cyprus, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Denmark	10/01/2012
VELCADE	Medartuum AB	02/07/2012	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/07/2012
VELCADE	Paranova Läkemedel AB	18/07/2012	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	07/08/2012
VELCADE	Tisida GmbH	20/03/2012	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/04/2012
Vfend	2 Care 4	02/12/2011	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Denmark	30/01/2012

Vfend	2 Care 4	15/12/2011	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/01/2012
Vfend	2 Care 4	04/04/2012	50 mg	Tablet	56	EU/1/02/212/008	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Romania, Spain, The Netherlands, United Kingdom	Sweden	07/05/2012
Vfend	ABACUS MEDICINE A/S	31/08/2012	50 mg	Tablet	30	EU/1/02/212/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/09/2012
Vfend	axicorp Pharma B.V.	13/06/2012	50 mg	Tablet	30	EU/1/02/212/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/06/2012
Vfend	axicorp Pharma B.V.	26/06/2012	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/07/2012
Vfend	Delfarma SpZoo	30/01/2012	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	11/04/2012
Vfend	Orifarm AB	01/10/2012	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/10/2012
Vfend	Orifarm AS	27/07/2012	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/08/2012

Vfend	Paranova Danmark AS	23/07/2012	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/08/2012
Vfend	Paranova Danmark AS	27/07/2012	50 mg	Tablet	28	EU/1/02/212/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/08/2012
Vfend	PCO Manufacturing	31/08/2011	50 mg	Tablet	28	EU/1/02/212/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/01/2012
Viagra	ABACUS MEDICINE A/S	22/12/2011	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/01/2012
Viagra	ABACUS MEDICINE A/S	22/12/2011	25 mg	Film-coated tablet	12	EU/1/98/077/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/01/2012
Viagra	ABACUS MEDICINE A/S	22/12/2011	50 mg	Film-coated tablet	12	EU/1/98/077/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	19/01/2012
Viagra	ABACUS MEDICINE A/S	31/08/2012	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	13/09/2012

Viagra	ABACUS MEDICINE A/S	28/09/2012	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Viagra	Combiphar Europe BV	21/02/2012	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	02/04/2012
Viagra	Combiphar Europe BV	21/02/2012	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	02/04/2012
Viagra	Docpharm GmbH& CoKGaA	12/09/2012	100 mg	Film-coated tablet	8	EU/1/98/077/011	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/09/2012
Viagra	EU-Pharma BV	23/10/2012	100 mg	Film-coated tablet	8	EU/1/98/077/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/11/2012
Viagra	NewNeopharm BV	17/07/2012	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/08/2012
Viagra	NewNeopharm BV	18/07/2012	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/08/2012
Viagra	NewNeopharm BV	22/10/2012	25 mg	Film-coated tablet	4	EU/1/98/077/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/10/2012

Viagra	Orifarm AS	12/01/2012	50 mg	Film-coated tablet	8	EU/1/98/077/018	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	08/02/2012
Viagra	Orifarm AS	27/08/2012	50 mg	Film-coated tablet	8	EU/1/98/077/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/09/2012
Viagra	PCO Manufacturing	23/02/2012	25 mg	Film-coated tablet	4	EU/1/98/077/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	02/04/2012
Viagra	Pharma Westen GmbH	18/01/2012	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
Viagra	Pharma Westen GmbH	12/04/2012	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/06/2012
Viagra	Pharma Westen GmbH	12/04/2012	50 mg	Film-coated tablet	12	EU/1/98/077/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/06/2012
Viagra	PHARMACHIM AB	24/01/2012	25 mg	Film-coated tablet	4	EU/1/98/077/002	Austria, Belgium, France, Germany, Greece, Italy, Malta, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Viagra	PHARMACHIM AB	24/01/2012	25 mg	Film-coated tablet	12	EU/1/98/077/004	Austria, Belgium, France, Germany, Greece, Italy, Malta, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Viagra	PHARMACHIM AB	24/01/2012	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, France, Germany, Greece, Italy, Malta, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012
Viagra	PHARMACHIM AB	24/01/2012	50 mg	Film-coated tablet	12	EU/1/98/077/008	Austria, Belgium, France, Germany, Greece, Italy, Malta, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom	Sweden	31/01/2012

Victoza	ABACUS MEDICINE A/S	05/01/2012	6 mg/ml	Solution for injection in pre-filled pen	3 pre-filled pens	EU/1/09/529/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/01/2012
Victoza	EU-Pharma BV	22/08/2012	6 mg/ml	Solution for injection in pre-filled pen	2 pre-filled pens	EU/1/09/529/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	10/09/2012
Victoza	Euro Registratie Collectief BV	12/01/2012	6 mg/ml	Solution for injection in pre-filled pen	2 pre-filled pens	EU/1/09/529/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/02/2012
Victrelis	Orifarm AB	12/10/2012	200 mg	Capsule, hard	336 (4 packs of 84)	EU/1/11/704/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	03/12/2012
Victrelis	Orifarm AS	13/08/2012	200 mg	Capsule, hard	84	EU/1/11/704/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	17/08/2012
Vimpat	ABACUS MEDICINE A/S	29/06/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/07/2012
Vimpat	ABACUS MEDICINE A/S	29/06/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/07/2012
Vimpat	AMIMED DIRECT LTD	08/03/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/03/2012

Vimpat	Amimed Direct Ltd	09/03/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/03/2012
Vimpat	Amimed Direct Ltd	31/07/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/08/2012
Vimpat	Amimed Direct Ltd	31/07/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/08/2012
Vimpat	Amimed Direct Ltd	31/07/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/08/2012
Vimpat	axicorp Pharma GmbH	20/12/2011	150 mg	Film-coated tablet	168	EU/1/08/470/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/01/2012
Vimpat	axicorp Pharma GmbH	27/12/2011	200 mg	Film-coated tablet	168	EU/1/08/470/012	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/01/2012
Vimpat	axicorp Pharma GmbH	22/08/2012	50 mg	Film-coated tablet	168	EU/1/08/470/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012
Vimpat	Beragena Arzneimittel GmbH	29/11/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/12/2012
Vimpat	Beragena Arzneimittel GmbH	29/11/2012	150 mg	Film-coated tablet	168	EU/1/08/470/009	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/12/2012
Vimpat	Beragena Arzneimittel GmbH	29/11/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/12/2012
Vimpat	Beragena Arzneimittel GmbH	29/11/2012	200 mg	Film-coated tablet	168	EU/1/08/470/012	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	21/12/2012

Vimpat	CC Pharma	19/10/2012	150 mg	Film-coated tablet	168	EU/1/08/470/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/10/2012
Vimpat	CC Pharma	19/10/2012	50 mg	Film-coated tablet	168	EU/1/08/470/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/10/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	150 mg	Film-coated tablet	14	EU/1/08/470/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	50 mg	Film-coated tablet	56	EU/1/08/470/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2012
Vimpat	Docpharm GmbH&CoKGaA	17/07/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2012
Vimpat	Docpharm GmbH&CoKGaA	25/07/2012	100 mg	Film-coated tablet	168	EU/1/08/470/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/08/2012
Vimpat	Docpharm GmbH&CoKGaA	25/07/2012	150 mg	Film-coated tablet	168	EU/1/08/470/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/08/2012
Vimpat	Docpharm GmbH&CoKGaA	25/07/2012	50 mg	Film-coated tablet	168	EU/1/08/470/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/08/2012

Vimpat	Emra-Med	08/11/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/12/2012
Vimpat	Ginova Ltd	06/03/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	11/04/2012
Vimpat	Ginova Ltd	19/04/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	27/04/2012
Vimpat	Interport Ltd	27/11/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands	United Kingdom	13/12/2012
Vimpat	Interport Ltd	27/11/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands	United Kingdom	13/12/2012
Vimpat	Interport Ltd	27/11/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands	United Kingdom	13/12/2012
Vimpat	kohlpharma GmbH	27/02/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Italy	Germany	26/03/2012
Vimpat	Kosei Pharma UK Limited	08/06/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/06/2012
Vimpat	Kosei Pharma UK Limited	08/06/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/06/2012

Vimpat	Kosei Pharma UK Limited	08/06/2012	150 mg	Film-coated tablet	14	EU/1/08/470/007	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/06/2012
Vimpat	Kosei Pharma UK Limited	08/06/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/06/2012
Vimpat	Kosei Pharma UK Limited	08/06/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	29/06/2012
Vimpat	Mediwin Ltd	24/07/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	10/08/2012
Vimpat	Mediwin Ltd	18/10/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	23/10/2012
Vimpat	Mediwin Ltd	04/12/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	12/12/2012
Vimpat	O.P.D. Laboratories Ltd	16/07/2012	150 mg	Film-coated tablet	14	EU/1/08/470/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	27/07/2012
Vimpat	PCO Manufacturing	05/03/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	12/03/2012
Vimpat	PCO Manufacturing	23/03/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Hungary, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	04/04/2012
Vimpat	Pharma Westen GmbH	15/12/2011	50 mg	Film-coated tablet	168	EU/1/08/470/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012

Vimpat	PHARMACHIM AB	28/11/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
Vimpat	PHARMACHIM AB	28/11/2012	50 mg	Film-coated tablet	56	EU/1/08/470/002	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
Vimpat	PHARMACHIM AB	28/11/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Estonia, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	13/12/2012
Vimpat	S.C.A.C Ltd	04/01/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	10/01/2012
Vimpat	S.C.A.C Ltd	04/01/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	16/02/2012
Vimpat	S.C.A.C Ltd	04/01/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	10/01/2012
Vimpat	S.C.A.C Ltd	01/06/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	15/06/2012
Vimpat	STRATHCLYDE PHARMACEUTICALS LTD	28/02/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	05/04/2012
Vimpat	STRATHCLYDE PHARMACEUTICALS LTD	28/02/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	05/04/2012

Vimpat	STRATHCLYDE PHARMACEUTICALS LTD	28/02/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	05/04/2012
Vimpat	Veron Pharma Vertriebs GmbH	08/05/2012	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	26/06/2012
Vimpat	Veron Pharma Vertriebs GmbH	08/05/2012	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	26/06/2012
Vimpat	Vital Supplies UK Limited	04/01/2012	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	19/01/2012
Vimpat	WAYMADE PLC	20/02/2012	50 mg	Film-coated tablet	56	EU/1/08/470/002	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	28/02/2012
Vimpat	WAYMADE PLC	25/07/2012	150 mg	Film-coated tablet	14	EU/1/08/470/007	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/07/2012
Vimpat	WAYMADE PLC	25/07/2012	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/07/2012
Vimpat	WAYMADE PLC	25/07/2012	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	31/07/2012
Viramune	ADL Pharma GmbH	26/04/2012	200 mg	Tablet	60	EU/1/97/055/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Viramune	Axicorp Pharma B.V.	23/05/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012
Viramune	Axicorp Pharma B.V.	23/05/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012

Viramune	CC Pharma	02/03/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/03/2012
Viramune	CC Pharma	14/03/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/03/2012
Viramune	Dr Fisher Farma BV	21/05/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/09/2012
Viramune	Emra-Med	15/03/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Viramune	Emra-Med	15/03/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Viramune	Emra-Med	02/11/2012	100 mg	Prolonged-release tablet	90	EU/1/97/055/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/11/2012
Viramune	EU-Pharma BV	29/05/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/09/2012
Viramune	Eureco-Pharma B.V.	25/09/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/10/2012
Viramune	Haemato Pharm AG	08/02/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012

Viramune	HAEMATO PHARM AG	07/06/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/07/2012
Viramune	kohlpharma GmbH	27/04/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Norway, The Netherlands	Germany	03/05/2012
Viramune	kohlpharma GmbH	27/04/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Norway, The Netherlands	Germany	03/05/2012
Viramune	Medicopharm AG	07/03/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/03/2012
Viramune	Medicopharm AG	20/09/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/10/2012
Viramune	Orifarm AS	01/10/2012	200 mg	Tablet	60	EU/1/97/055/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	12/10/2012
Viramune	PHARMA LAB	02/10/2012	200 mg	Tablet	60	EU/1/97/055/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	24/10/2012
Viramune	Pharma Westen GmbH	20/01/2012	400 mg	Prolonged-release tablet	30	EU/1/97/055/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
Viramune	Pharma Westen GmbH	16/04/2012	100 mg	Prolonged-release tablet	90	EU/1/97/055/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/06/2012
Viramune	Pharma Westen GmbH	11/09/2012	400 mg	Prolonged-release tablet	90	EU/1/97/055/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/10/2012
Viramune	PHARMACHIM AB	27/02/2012	200 mg	Tablet	60	EU/1/97/055/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012

Viramune	Veron Pharma Vertriebs GmbH	25/10/2011	200 mg	Tablet	60	EU/1/97/055/001	Austria	Germany	05/01/2012
Viramune	Veron Pharma Vertriebs GmbH	25/10/2011	200 mg	Tablet	120	EU/1/97/055/003	Austria	Germany	05/01/2012
Viread	Combiphar Europe BV	21/02/2012	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	02/04/2012
Viread	HAEMATO PHARM AG	22/12/2011	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/01/2012
Viread	Veron Pharma Vertriebs GmbH	12/04/2012	245 mg	Film-coated tablet	3 x 30	EU/1/01/200/002	Austria	Germany	23/04/2012
Vistide	ABACUS MEDICINE A/S	27/08/2012	75 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/97/037/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/09/2012
Visudyne	ABACUS MEDICINE A/S	31/07/2012	15 mg	Powder for solution for infusion	1 vial	EU/1/00/140/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	16/08/2012
Visudyne	Orifarm AS	23/02/2012	15 mg	Powder for solution for infusion	1 vial	EU/1/00/140/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/02/2012
Votrient	Beragena Arzneimittel GmbH	27/06/2012	400 mg	Film-coated tablet	60	EU/1/10/628/004	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	04/07/2012

Votrient	Cancernova GmbH onkologische Arzneimittel	23/04/2012	400 mg	Film-coated tablet	60	EU/1/10/628/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/04/2012
Votrient	CC Pharma	21/12/2011	200 mg	Film-coated tablet	90	EU/1/10/628/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012
Votrient	CC Pharma	02/02/2012	400 mg	Film-coated tablet	60	EU/1/10/628/004	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/02/2012
Votrient	cc Pharma	23/04/2012	200 mg	Film-coated tablet	30	EU/1/10/628/001	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2012
Votrient	cc Pharma	23/04/2012	400 mg	Film-coated tablet	30	EU/1/10/628/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/05/2012
Votrient	Emra-Med	08/02/2012	200 mg	Film-coated tablet	90	EU/1/10/628/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/03/2012
Votrient	Emra-Med	08/02/2012	400 mg	Film-coated tablet	60	EU/1/10/628/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/03/2012
Votrient	Haemato Pharm AG	13/04/2012	200 mg	Film-coated tablet	90	EU/1/10/628/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012

Votrient	HAEMATO PHARM AG	13/04/2012	400 mg	Film-coated tablet	30	EU/1/10/628/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/05/2012
Votrient	HAEMATO PHARM AG	26/10/2012	200 mg	Film-coated tablet	30	EU/1/10/628/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/11/2012
Votrient	MTK Pharma GmbH	21/02/2012	200 mg	Film-coated tablet	30	EU/1/10/628/001	Czech Republic, Greece	Germany	05/03/2012
Votrient	MTK Pharma GmbH	21/02/2012	200 mg	Film-coated tablet	90	EU/1/10/628/002	Czech Republic, Greece	Germany	05/03/2012
Votrient	MTK Pharma GmbH	21/02/2012	400 mg	Film-coated tablet	60	EU/1/10/628/004	Greece	Germany	05/03/2012
Votrient	Orifarm AB	17/02/2012	200 mg	Film-coated tablet	90	EU/1/10/628/002	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	15/03/2012
VPRIV	Orifarm AB	16/07/2012	400 U	Powder for solution for infusion	1 vial	EU/1/10/646/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	03/08/2012
VPRIV	Orifarm AS	02/07/2012	400 U	Powder for solution for infusion	1 vial	EU/1/10/646/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/07/2012
Xarelto	2 Care 4	15/12/2011	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/01/2012

Xarelto	2 Care 4	21/12/2011	10 mg	Film-coated tablet	100 x 1	EU/1/08/472/008	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/01/2012
Xarelto	2 Care 4	12/04/2012	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	25/04/2012
Xarelto	CC Pharma	21/03/2012	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/04/2012
Xarelto	CC Pharma	30/04/2012	15 mg	Film-coated tablet	28	EU/1/08/472/012	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/05/2012
Xarelto	CC Pharma	30/04/2012	15 mg	Film-coated tablet	98	EU/1/08/472/014	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/05/2012
Xarelto	CC Pharma	30/04/2012	15 mg	Film-coated tablet	42	EU/1/08/472/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/05/2012
Xarelto	CC Pharma	30/04/2012	20 mg	Film-coated tablet	28	EU/1/08/472/018	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2012
Xarelto	CC Pharma	30/04/2012	20 mg	Film-coated tablet	98	EU/1/08/472/019	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2012
Xarelto	CC Pharma	08/10/2012	10 mg	Film-coated tablet	5	EU/1/08/472/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/10/2012
Xarelto	COPHARMA ApS	08/02/2012	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/03/2012
Xarelto	COPHARMA ApS	08/02/2012	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/03/2012

Xarelto	HAEMATO PHARM AG	15/10/2012	15 mg	Film-coated tablet	100 x 1	EU/1/08/472/016	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/10/2012
Xarelto	HAEMATO-PHARM AG	15/05/2012	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/05/2012
Xarelto	Medcor Pharmaceuticals BV	19/10/2012	20 mg	Film-coated tablet	28	EU/1/08/472/018	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/10/2012
Xarelto	Mediwin Ltd	29/06/2012	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/07/2012
Xarelto	Mediwin Ltd	29/06/2012	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/07/2012
Xarelto	Paranova OY	11/04/2012	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	24/04/2012
Xarelto	Paranova Oy	11/04/2012	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	24/04/2012
Xeloda	D-Pharm, a.s.	12/06/2012	500 mg	Film-coated tablet	120	EU/1/00/163/002	Slovak Republic	Czech Republic	04/07/2012

Xeloda	Medartuum AB	02/10/2012	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Xeloda	Paranova Läkemedel AB	18/06/2012	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	20/06/2012
Xeloda	Paranova Oy	20/04/2012	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Finland	07/05/2012
Xenical	BB Farma s.r.l.	23/01/2012	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	07/03/2012
Xenical	Chemilines Ltd	30/03/2012	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Ireland, United Kingdom	11/06/2012
Xenical	Ethigen Limited	23/05/2012	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	25/06/2012
Xenical	Euromedicines, S.L.	15/06/2012	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Portugal	11/09/2012

Xenical	Euromedicines, S.L.	15/06/2012	120 mg	Capsule, hard	42	EU/1/98/071/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Portugal	11/09/2012
Xenical	Impexeco S.A.	31/08/2011	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Czech Republic, Denmark, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Lithuania, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Belgium	06/02/2012
Xenical	Kosei Pharma UK Limited	09/12/2011	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	05/01/2012
Xenical	NewNeopharm BV	09/11/2012	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/11/2012
Xeplion	2 Care 4	28/02/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/03/2012
Xeplion	2 Care 4	28/02/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	08/03/2012
Xeplion	ACA Müller ADAG Pharma AG	18/12/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/12/2012
Xeplion	Axicorp Pharma B.V.	11/07/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/08/2012
Xeplion	Axicorp Pharma B.V.	22/08/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012

Xeplion	Beragena Arzneimittel GmbH	28/06/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, France, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	09/07/2012
Xeplion	BR Pharma International Ltd	20/04/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/05/2012
Xeplion	CC Pharma	09/07/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/08/2012
Xeplion	CC Pharma	06/09/2012	50 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/09/2012
Xeplion	Cross Pharma AB	01/11/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/11/2012
Xeplion	Cross Pharma AB	01/11/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/11/2012
Xeplion	Cross Pharma AB	01/11/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	16/11/2012
Xeplion	Dr Fisher Farma BV	21/02/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/03/2012

Xeplion	Dr Fisher Farma BV	21/02/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/03/2012
Xeplion	Dr Fisher Farma BV	21/02/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	08/03/2012
Xeplion	Emra-Med	28/02/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/03/2012
Xeplion	Emra-Med	13/11/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/12/2012
Xeplion	Emra-Med	26/11/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/11/2012
Xeplion	HAEMATO PHARM AG	16/04/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012

Xeplion	HAEMATO PHARM AG	16/04/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/05/2012
Xeplion	kohlpharma GmbH	22/12/2011	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/01/2012
Xeplion	kohlpharma GmbH	22/12/2011	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/01/2012
Xeplion	kohlpharma GmbH	29/08/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/09/2012
Xeplion	Medartuum AB	17/09/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	01/10/2012
Xeplion	Medartuum AB	20/09/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	01/10/2012
Xeplion	Medcor Pharmaceuticals BV	02/05/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	11/06/2012
Xeplion	Medcor Pharmaceuticals BV	02/05/2012	50 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/002	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	11/06/2012

Xeplion	Medcor Pharmaceuticals BV	02/05/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	11/06/2012
Xeplion	Medcor Pharmaceuticals BV	29/05/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	11/06/2012
Xeplion	Orifarm AB	18/05/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	05/06/2012
Xeplion	Orifarm AB	21/06/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/07/2012
Xeplion	Orifarm AB	21/06/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	04/07/2012
Xeplion	Orifarm AS	18/05/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	05/06/2012
Xeplion	Orifarm AS	21/06/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/07/2012
Xeplion	Orifarm AS	21/06/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	04/07/2012
Xeplion	Orifarm AS	01/08/2012	50 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/09/2012

Xeplion	Paranova Danmark AS	30/01/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/02/2012
Xeplion	Paranova Danmark AS	20/03/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/05/2012
Xeplion	Paranova Danmark AS	17/07/2012	50 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	31/07/2012
Xeplion	Paranova Danmark AS	17/07/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	31/07/2012
Xeplion	Paranova Läkemedel AB	22/02/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	24/02/2012
Xeplion	Paranova Läkemedel AB	20/03/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	09/05/2012

Xeplion	Paranova Läkemedel AB	17/07/2012	50 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/07/2012
Xeplion	Paranova Läkemedel AB	17/07/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	31/07/2012
Xeplion	PCO Manufacturing	26/06/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/07/2012
Xeplion	PCO Manufacturing	26/06/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/07/2012
Xeplion	Pharma Westen GmbH	18/01/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/01/2012
Xeplion	Pharma Westen GmbH	18/01/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/01/2012
Xeplion	Pharma Westen GmbH	18/01/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/01/2012
Xeplion	Tisida GmbH	16/07/2012	100 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/08/2012

Xeplion	Tisida GmbH	16/07/2012	150 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/08/2012
Xeplion	Tisida GmbH	16/07/2012	75 mg	Suspension for injection in pre-filled syringe	1 pre-filled syringe + 2 needles	EU/1/11/672/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/08/2012
Xeristar	axicorp Pharma GmbH	23/03/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/03/2012
Xeristar	axicorp Pharma GmbH	23/03/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/03/2012
Xeristar	axicorp Pharma GmbH	11/05/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2012
Xeristar	B2B Medical GmbH	03/03/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	Germany	02/04/2012
Xeristar	B2B Medical GmbH	03/03/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/04/2012
Xeristar	Cambridge Major Laboratories Europe B.V	28/06/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/08/2012
Xeristar	Cambridge Major Laboratories Europe B.V	28/06/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/08/2012

Xeristar	Cambridge Major Laboratories Europe B.V	28/06/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/08/2012
Xeristar	CC Pharma	13/02/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/02/2012
Xeristar	CD Pharma Limited	01/02/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/02/2012
Xeristar	Docpharm GmbH&CoKGaA	12/12/2011	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/01/2012
Xeristar	Docpharm GmbH&CoKGaA	10/01/2012	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/01/2012
Xeristar	Docpharm GmbH&CoKGaA	10/01/2012	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/01/2012
Xeristar	Docpharm GmbH&CoKGaA	13/02/2012	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/02/2012
Xeristar	O.P.D. Laboratories Ltd	19/07/2010	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	29/02/2012
XGEVA	Dr Fisher Farma BV	01/10/2012	120 mg	Solution for injection	1 vial	EU/1/11/703/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/10/2012
Xolair	HAEMATO PHARM AG	12/07/2012	150 mg	Solution for injection	1 pre-filled syringe	EU/1/05/319/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/07/2012

Xolair	HAEMATO PHARM AG	12/07/2012	75 mg	Solution for injection	1 pre-filled syringe	EU/1/05/319/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/07/2012
Xolair	Medcor Pharmaceuticals BV	28/06/2012	150 mg	Solution for injection	1 pre-filled syringe	EU/1/05/319/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/08/2012
Xolair	Orifarm AB	20/12/2011	150 mg	Solution for injection	1 pre-filled syringe	EU/1/05/319/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/01/2012
Xolair	Paranova Läkemedel AB	21/08/2012	150 mg	Solution for injection	1 pre-filled syringe	EU/1/05/319/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	23/08/2012
Yellox	2 Care 4	20/11/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	28/11/2012
Yellox	B&S Healthcare	29/10/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	05/11/2012
Yellox	Dr Fisher Farma BV	24/10/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/11/2012
Yellox	Emra-Med	23/04/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2012
Yellox	EU-Pharma BV	04/10/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/12/2012

Yellox	Euro Registratie Collectief BV	03/07/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	10/07/2012
Yellox	Ginova Ltd	13/12/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	20/12/2012
Yellox	kohlpharma GmbH	14/06/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Portugal	Germany	06/07/2012
Yellox	Medcor Pharmaceuticals BV	06/03/2012	0.9 mg/ml	Eye drops, solution	1 bottle	EU/1/11/692/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/03/2012
Yentreve	CD Pharma Limited	07/02/2012	20 mg	Gastro-resistant capsule, hard	98	EU/1/04/280/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/02/2012
Yentreve	CD Pharma Limited	07/02/2012	40 mg	Gastro-resistant capsule, hard	98	EU/1/04/280/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/02/2012
Yervoy	CC Pharma	05/12/2012	5 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/11/698/ 001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Yervoy	CC Pharma	05/12/2012	5 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/11/698/ 002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/12/2012
Zarzio	Orifarm AB	17/07/2012	48 MU (96 MU/ml)	Solution for injection or infusion	1 pre-filled syringe with needle guard	EU/1/08/495/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012
Zarzio	Orifarm AB	18/07/2012	30 MU (60 MU/ml)	Solution for injection or infusion	5 pre-filled syringes with needle guard	EU/1/08/495/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	12/10/2012

Zavesca	Orifarm AB	12/10/2012	100 mg	Capsule, hard	84	EU/1/02/238/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	29/10/2012
Zebinix	2 Care 4	24/09/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	18/10/2012
Zebinix	axicorp Pharma GmbH	02/11/2012	800 mg	Tablet	90	EU/1/09/514/019	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/11/2012
Zebinix	CC Pharma	19/06/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/07/2012
Zebinix	Emra-Med	25/05/2012	800 mg	Tablet	90	EU/1/09/514/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2012
Zebinix	HAEMATO PHARM AG	10/07/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/08/2012
Zebinix	kohlpharma GmbH	17/07/2012	800 mg	Tablet	90	EU/1/09/514/019	France, Portugal	Germany	17/08/2012
Zebinix	kohlpharma GmbH	17/07/2012	800 mg	Tablet	30	EU/1/09/514/017	France, Portugal	Germany	17/08/2012
Zebinix	Orifarm AB	03/12/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/12/2012

Zebinix	Orifarm AS	24/01/2012	800 mg	Tablet	30	EU/1/09/514/013	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/03/2012
Zebinix	Orifarm AS	30/05/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	29/06/2012
Zebinix	Paranova Danmark AS	26/11/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/12/2012
Zebinix	Paranova Läkemedel AB	07/11/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, United Kingdom	Sweden	14/11/2012
Zebinix	Pharma Westen GmbH	31/05/2012	800 mg	Tablet	30	EU/1/09/514/017	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	29/06/2012
Zebinix	Pharma Westen GmbH	14/06/2012	800 mg	Tablet	90	EU/1/09/514/019	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/07/2012
Zeffix	Combiphar Europe BV	22/05/2009	100 mg	Film-coated tablet	28	EU/1/99/114/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/01/2012
Zelboraf	CC Pharma	13/12/2012	240 mg	Film-coated tablet	56	EU/1/12/751/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/12/2012
Zelboraf	HAEMATO PHARM AG	12/07/2012	240 mg	Film-coated tablet	56	EU/1/12/751/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/07/2012

Ziagen	axicorp Pharma GmbH	13/03/2012	300 mg	Film-coated tablet	60	EU/1/99/112/001	Austria, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/04/2012
Ziagen	Orifarm AS	12/01/2012	300 mg	Film-coated tablet	60	EU/1/99/112/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	19/03/2012
Ziagen	REMEDIX GmbH	14/03/2012	300 mg	Film-coated tablet	60	EU/1/99/112/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Austria, Germany	23/04/2012
Zonegran	Abacus Medicine A/S	18/11/2011	100 mg	Capsule, hard	98	EU/1/04/307/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/01/2012
Zonegran	ABACUS MEDICINE A/S	28/09/2012	100 mg	Capsule, hard	196	EU/1/04/307/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/10/2012
Zonegran	Copharma ApS	19/06/2012	100 mg	Capsule, hard	98	EU/1/04/307/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
Zonegran	Copharma ApS	19/06/2012	100 mg	Capsule, hard	28	EU/1/04/307/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
Zonegran	Copharma ApS	19/06/2012	100 mg	Capsule, hard	56	EU/1/04/307/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012

Zonegran	Copharma ApS	19/06/2012	25 mg	Capsule, hard	14	EU/1/04/307/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Zonegran	Copharma ApS	19/06/2012	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Zonegran	Copharma ApS	19/06/2012	25 mg	Capsule, hard	28	EU/1/04/307/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2012
Zonegran	Copharma ApS	19/06/2012	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
Zonegran	Copharma ApS	19/06/2012	50 mg	Capsule, hard	28	EU/1/04/307/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
Zonegran	Copharma ApS	19/06/2012	50 mg	Capsule, hard	14	EU/1/04/307/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/06/2012
Zonegran	Kosei Pharma UK Limited	06/02/2012	25 mg	Capsule, hard	14	EU/1/04/307/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/02/2012
Zonegran	Kosei Pharma UK Limited	06/02/2012	50 mg	Capsule, hard	28	EU/1/04/307/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	27/02/2012

Zonegran	Kosei Pharma UK Limited	17/04/2012	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	24/04/2012
Zonegran	Kosei Pharma UK Limited	17/05/2012	50 mg	Capsule, hard	14	EU/1/04/307/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	30/05/2012
Zonegran	Kosei Pharma UK Limited	26/06/2012	25 mg	Capsule, hard	28	EU/1/04/307/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	06/07/2012
Zonegran	MPT Pharma	15/11/2012	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/11/2012
Zonegran	O.P.D. Laboratories Ltd	21/09/2012	50 mg	Capsule, hard	14	EU/1/04/307/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	08/10/2012
Zonegran	STRATHCLYDE PHARMACEUTICALS LTD	07/12/2012	50 mg	Capsule, hard	14	EU/1/04/307/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	18/12/2012
Zutectra	Axicorp Pharma B.V.	12/03/2012	500 IU	Solution for injection	5 pre-filled syringes	EU/1/09/600/001	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/03/2012

Zutectra	Emra-Med	15/05/2012	500 IU	Solution for injection	5 pre-filled syringes	EU/1/09/600/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/07/2012
Zutectra	kohlpharma GmbH	09/08/2012	500 IU	Solution for injection	5 pre-filled syringes	EU/1/09/600/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/09/2012
Zutectra	Orifarm AB	18/07/2012	500 IU	Solution for injection	5 pre-filled syringes	EU/1/09/600/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	03/08/2012
Zutectra	Pharma Westen GmbH	01/02/2012	500 IU	Solution for injection	5 pre-filled syringes	EU/1/09/600/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/05/2012
Zypadhera	CC Pharma	11/05/2012	210 mg	Powder and solvent for prolonged-release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	01/06/2012
Zypadhera	Emra-Med	31/08/2012	210 mg	Powder and solvent for prolonged release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	11/09/2012
Zypadhera	Eureco-Pharma B.V.	12/09/2012	210 mg	Powder and solvent for prolonged release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	26/09/2012

Zypadhera	Eureco-Pharma B.V.	12/09/2012	300 mg	Powder and solvent for prolonged release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	26/09/2012
Zypadhera	Eureco-Pharma B.V.	12/09/2012	405 mg	Powder and solvent for prolonged release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	26/09/2012
Zypadhera	Medcor Pharmaceuticals BV	06/07/2012	210 mg	Powder and solvent for prolonged-release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	02/10/2012
Zypadhera	Medcor Pharmaceuticals BV	06/07/2012	405 mg	Powder and solvent for prolonged-release suspension for injection	1 vial + 1 vial + 1 syringe with pre-attached safety needle + 2 safety needles	EU/1/08/479/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	The Netherlands	02/10/2012
Zyprexa	Adripharma GmbH	10/10/2012	10 mg	Coated tablet	70	EU/1/96/022/032	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/10/2012

Zyprexa	Adripharma GmbH	10/10/2012	5 mg	Coated tablet	70	EU/1/96/022/030	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	30/10/2012
Zyprexa	BB Farma s.r.l.	23/05/2012	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	28/06/2012
Zyprexa	BB Farma s.r.l.	23/05/2012	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	28/06/2012
Zyprexa	Mediwin Ltd	15/12/2011	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Cyprus, Denmark, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	13/01/2012
Zyprexa	Mediwin Ltd	15/12/2011	7.5 mg	Coated tablet	28	EU/1/96/022/011	Austria, Belgium, Cyprus, Denmark, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	20/01/2012
Zyprexa	Mediwin Ltd	10/07/2012	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	France	24/07/2012
Zyprexa	PHARMA LAB	02/07/2012	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	17/07/2012
Zyprexa	PHARMA LAB	02/07/2012	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	France	17/07/2012

Zyprexa Velotab	BB Farma s.r.l.	24/05/2012	10 mg	Orodispersible tablet	28	EU/1/99/125/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	28/06/2012
Zyprexa Velotab	BB Farma s.r.l.	24/05/2012	5 mg	Orodispersible tablet	28	EU/1/99/125/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	France	28/06/2012
Zyprexa Velotab	BB Farma s.r.l.	20/06/2012	5 mg	Orodispersible tablet	28	EU/1/99/125/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Latvia, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Italy	04/07/2012
Zyprexa Velotab	Emra-Med	06/12/2011	15 mg	Orodispersible tablet	98	EU/1/99/125/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/01/2012
Zyprexa Velotab	Mediwin Ltd	16/12/2011	10 mg	Orodispersible tablet	28	EU/1/99/125/002	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	13/01/2012
Zyprexa Velotab	Mediwin Ltd	16/12/2011	5 mg	Orodispersible tablet	28	EU/1/99/125/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Ireland, Italy, Lithuania, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	13/01/2012
Zyprexa Velotab	PHARMA LAB	02/07/2012	10 mg	Orodispersible tablet	28	EU/1/99/125/002	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/07/2012
Zyprexa Velotab	PHARMA LAB	02/07/2012	5 mg	Orodispersible tablet	28	EU/1/99/125/001	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	France	19/07/2012

Zytiga	axicorp Pharma B.V.	17/08/2012	250 mg	Tablet	120	EU/1/11/714/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/09/2012
Zytiga	CC Pharma	16/02/2012	250 mg	Tablet	120	EU/1/11/714/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Norway, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/03/2012
Zytiga	Emra-Med	01/06/2012	250 mg	Tablet	120	EU/1/11/714/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/06/2012
Zytiga	kohlpharma GmbH	06/07/2012	250 mg	Tablet	120	EU/1/11/714/001	Austria, Norway, Sweden, The Netherlands	Germany	18/07/2012
Zytiga	Pharma Westen GmbH	03/02/2012	250 mg	Tablet	120	EU/1/11/714/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	10/02/2012