02 August 2011
EMA/742248/2011
Patient Health Protection

Initial Notices for Parallel Distribution –2009

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	15/06/2009	15 mg	Tablet	28 x 1	EU/1/04/276/012	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland, Romania, Finland, Iceland	Denmark	26/11/2009
Abilify	2 Care 4	15/06/2009	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Iceland, Finland	Denmark	26/11/2009
Abilify	2 Care 4	19/10/2009	10 mg	Tablet	56 x 1	EU/1/04/276/009	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/11/2009
Abilify	2 Care 4	19/10/2009	15 mg	Tablet	56 x 1	EU/1/04/276/014	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom	Denmark	26/11/2009
Abilify	2 Care 4	19/10/2009	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	26/11/2009

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Abilify	2 Care 4	20/10/2009	10 mg	Tablet	28 x 1	EU/1/04/276/007	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden, United Kingdom	Denmark	26/11/2009
Abilify	Abis-Pharma	14/08/2009	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	12/10/2009
Abilify	Abis-Pharma	14/08/2009	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	12/10/2009
Abilify	Abis-Pharma	14/08/2009	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	12/10/2009
Abilify	Abis-Pharma	14/08/2009	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	Germany	12/10/2009
Abilify	Abis-Pharma	14/08/2009	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	12/10/2009
Abilify	Abis-Pharma	14/08/2009	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	12/10/2009

Abilify	Emra-Med	11/09/2008	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/01/2009
Abilify	EuroPharma DK	23/03/2009	30 mg	Tablet	28 x 1	EU/1/04/276/017	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/04/2009
Abilify	EuroPharma DK	23/03/2009	30 mg	Tablet	56 x 1	EU/1/04/276/019	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	14/04/2009
Abilify	Lexon (UK) Ltd	05/08/2009	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, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	07/08/2009
Abilify	Manichem Limited	16/09/2009	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Cyprus, Iceland, Liechtenstein, Norway	United Kingdom, Ireland	21/09/2009
Abilify	Mevita HandelsGmbH	29/07/2008	5 mg	Tablet	49 x 1	EU/1/04/276/003	France, Italy, Spain, United Kingdom	Germany	29/01/2009
Abilify	Milinda Arzneimittel GmbH	28/08/2009	10 mg	Tablet	14 x 1	EU/1/04/276/006	France, Spain	Germany	25/11/2009
Abilify	Milinda Arzneimittel GmbH	28/08/2009	10 mg	Tablet	49 x 1	EU/1/04/276/008	France, Spain	Germany	25/11/2009
Abilify	Milinda Arzneimittel GmbH	28/08/2009	10 mg	Tablet	98 x 1	EU/1/04/276/010	France, Spain		25/11/2009
Abilify	Milinda Arzneimittel GmbH	28/08/2009	5 mg	Tablet	49 x 1	EU/1/04/276/003	France, Spain	Germany	25/11/2009
Abilify	Orifarm AB	03/07/2009	10 mg	Tablet	28 x 1	EU/1/04/276/007	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	05/08/2009

Abilify	Pharma Gerke GmbH	02/05/2008	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	29/06/2009
Abilify	Pharmacodane Aps	09/07/2009	5 mg	Tablet	14 x 1	EU/1/04/276/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	02/09/2009
Abilify	Pharmacodane Aps	09/07/2009	5 mg	Tablet	28 x 1	EU/1/04/276/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	02/09/2009
Abilify	Pharmacodane Aps	09/07/2009	5 mg	Tablet	56 x 1	EU/1/04/276/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	02/09/2009
Abilify	Swingward Ltd TA Medihealth	04/11/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	27/11/2009
Abilify	Swingward Ltd TA Medihealth	04/11/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	27/11/2009
Abilify	Veron Pharma	27/10/2008	10 mg	Tablet	14 x 1	EU/1/04/276/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	09/02/2009
Abilify	Veron Pharma	27/10/2008	10 mg	Tablet	49 x 1	EU/1/04/276/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	09/02/2009
Abilify	Veron Pharma	27/10/2008	10 mg	Tablet	98 x 1	EU/1/04/276/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	09/02/2009
Abilify	Veron Pharma	28/09/2009	5 mg	Tablet	49 x 1	EU/1/04/276/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	13/11/2009
Aclasta	Emra-Med	07/09/2009	5 mg/100 ml	Solution for infusion	1 bottle	EU/1/05/308/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/11/2009

Aclasta	Kohlpharma GmbH	18/05/2009	5 mg/100 ml	Solution for infusion	1 bottle	EU/1/05/308/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Aclasta	Paranova Danmark AS	26/08/2009	5 mg/100 ml	Solution for infusion	1 bottle	EU/1/05/308/001	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	26/11/2009
Aclasta	Pharma Westen GmbH	19/01/2009	5 mg/100 ml	Solution for infusion	1 bottle	EU/1/05/308/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/04/2009
Actos	Abis-Pharma	14/08/2009	30 mg	Tablet	28	EU/1/00/150/004	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	12/10/2009
Actos	Abis-Pharma	14/08/2009	30 mg	Tablet	98	EU/1/00/150/006	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	12/10/2009
Actos	Abis-Pharma	14/08/2009	45 mg	Tablet	28	EU/1/00/150/012	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	12/10/2009
Actos	Abis-Pharma	14/08/2009	45 mg	Tablet	98	EU/1/00/150/015	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	12/10/2009
Actos	Beragena Arzneimittel GmbH	10/02/2009	15mg	Tablets	28	EU/1/00/150/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	24/03/2009

Actos	Beragena Arzneimittel GmbH	10/02/2009	15mg	Tablets	98	EU/1/00/150/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	24/03/2009
Actos	BR Pharma International Ltd	28/05/2009	15 mg	Tablet	98	EU/1/00/150/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Malta	Germany	30/06/2009
Actos	BR Pharma International Ltd	17/08/2009	45 mg	Tablet	98	EU/1/00/150/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/09/2009
Actos	Docpharm GmbH&CoKGaA	13/03/2009	15 mg	Tablet	28	EU/1/00/150/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/03/2009
Actos	Docpharm GmbH&CoKGaA	13/03/2009	15 mg	Tablet	56	EU/1/00/150/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	25/03/2009
Actos	Docpharm GmbH&CoKGaA	13/03/2009	15 mg	Tablet	98	EU/1/00/150/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	25/03/2009
Actos	Docpharm GmbH&CoKGaA	16/03/2009	30 mg	Tablet	28	EU/1/00/150/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	25/03/2009
Actos	Docpharm GmbH&CoKGaA	16/03/2009	30 mg	Tablet	56	EU/1/00/150/010	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/03/2009
Actos	Docpharm GmbH&CoKGaA	16/03/2009	30 mg	Tablet	98	EU/1/00/150/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/03/2009
Actos	Emra-Med	24/08/2009	45 mg	Tablet	98	EU/1/00/150/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Actos	Ginova Ltd	26/08/2009	30 mg	Tablet	28	EU/1/00/150/004	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	14/10/2009

Actos	Kohlpharma GmbH	20/10/2009	45 mg	Tablet	98	EU/1/00/150/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/10/2009
Actos	Mediwin Ltd	25/08/2009	15 mg	Tablet	28	EU/1/00/150/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Italy	14/10/2009
Actos	Milinda Arzneimittel GmbH	21/08/2009	30 mg	Tablet	28	EU/1/00/150/004	France, Spain	Germany	26/11/2009
Actos	Milinda Arzneimittel GmbH	21/08/2009	30 mg	Tablet	98	EU/1/00/150/006	France, Spain	Germany	26/11/2009
Actos	Paranova OY	03/11/2009	15 mg	Tablet	98	EU/1/00/150/003	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Latvia	Finland	27/11/2009
Actos	Pharma Westen GmbH	02/03/2009	45 mg	Tablet	98	EU/1/00/150/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/05/2009
Actrapid	Euro Registratie Collectief BV	28/08/2009	100 IU/ml	Solution for injection	1 vial	EU/1/02/230/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/09/2009
Adrovance	Medcor Pharmaceuticals BV	16/03/2009	70 mg / 5600 IU	Tablet	4	EU/1/06/364/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/05/2009
Advagraf	BR Pharma International Ltd	12/08/2008	0.5 mg	Prolonged release capsules, hard	50	EU/1/07/387/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	01/04/2009
Advagraf	BR Pharma International Ltd	12/08/2008	1 mg	Prolonged release capsules, hard	50	EU/1/07/387/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	01/04/2009

Advagraf	BR Pharma International Ltd	12/08/2008	1 mg	Prolonged release capsules, hard	100	EU/1/07/387/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	01/04/2009
Advagraf	BR Pharma International Ltd	03/10/2008	0.5 mg	Prolonged release capsules, hard	100	EU/1/07/387/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	01/04/2009
Advagraf	BR Pharma International Ltd	04/11/2009	3 mg	Prolonged-release capsules, hard	50	EU/1/07/387/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/11/2009
Advagraf	BR Pharma International Ltd	11/11/2009	3 mg	Prolonged-release capsules, hard	100	EU/1/07/387/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/12/2009
Advagraf	Cross Healthcare Ltd	11/08/2009	0.5 mg	Prolonged-release capsules, hard	30	EU/1/07/387/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/10/2009
Advagraf	Cross Healthcare Ltd	13/08/2009	1 mg	Prolonged-release capsules, hard	30	EU/1/07/387/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/10/2009
Advagraf	Cross Healthcare Ltd	02/09/2009	5 mg	Prolonged-release capsules, hard	30	EU/1/07/387/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/10/2009
Advagraf	Cross Healthcare Ltd	06/10/2009	1 mg	Prolonged-release capsules, hard	60	EU/1/07/387/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	26/10/2009
Advagraf	Dr Fisher Farma BV	23/02/2009	1 mg	Prolonged release capsules, hard	60	EU/1/07/387/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/04/2009

Advagraf	Emra-Med	06/11/2008	0.5 mg	Prolonged release capsules, hard	100	EU/1/07/387/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/03/2009
Advagraf	Emra-Med	26/05/2009	5 mg	Prolonged release capsules, hard	100	EU/1/07/387/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	21/09/2009
Advagraf	Emra-Med	26/05/2009	5 mg	Prolonged release capsules, hard	50	EU/1/07/387/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	21/09/2009
Advagraf	Euro Registratie Collectief BV	27/01/2009	5 mg	Prolonged release capsules, hard	30	EU/1/07/387/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	17/02/2009
Advagraf	Euro Registratie Collectief BV	27/01/2009	5 mg	Prolonged release capsules, hard	50	EU/1/07/387/008	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	17/02/2009
Advagraf	Haemato Pharm AG	03/11/2008	1 mg	Prolonged-release capsules, hard	50	EU/1/07/387/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	26/01/2009
Advagraf	Haemato Pharm AG	03/11/2008	5 mg	Prolonged-release capsules, hard	50	EU/1/07/387/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	26/01/2009
Advagraf	kohlpharma GmbH	16/11/2009	3 mg	Prolonged-release capsules, hard	50	EU/1/07/387/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009

Advagraf	Medartuum AB	12/01/2009	0.5 mg	Prolonged-release capsules, hard	50	EU/1/07/387/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	21/01/2009
Advagraf	Medartuum AB	07/09/2009	1 mg	Prolonged-release capsules, hard	60	EU/1/07/387/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	22/09/2009
Advagraf	NewNeopharm BV	01/09/2009	1 mg	Prolonged-release capsules, hard	60	EU/1/07/387/005	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	08/09/2009
Advagraf	Pharma Westen GmbH	18/12/2008	5 mg	Prolonged release capsules, hard	50	EU/1/07/387/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/02/2009
Advagraf	Singad Pharma ApS	06/07/2009	1 mg	Prolonged-release capsules, hard	50	EU/1/07/387/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	14/09/2009
Advagraf	Singad Pharma ApS	06/07/2009	5 mg	Prolonged-release capsules, hard	50	EU/1/07/387/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	14/09/2009
Advagraf	Swingward Ltd TA Medihealth	08/02/2008	1 mg	Prolonged-release capsules, hard	30	EU/1/07/387/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, The Netherlands, Portugal, Spain, Sweden, Malta, Norway	United Kingdom	16/09/2009
Advagraf	Swingward Ltd TA Medihealth	08/02/2008	1 mg	Prolonged-release capsules, hard	60	EU/1/07/387/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	16/09/2009
Advagraf	Swingward Ltd TA Medihealth	08/02/2008	5 mg	Prolonged-release capsules, hard	30	EU/1/07/387/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	16/09/2009

Aerius	2 Care 4	20/05/2009	5 mg	Film-coated tablet	30	EU/1/00/160/011	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	24/06/2009
Aerius	2 Care 4	08/06/2009	5 mg	Film-coated tablet	100	EU/1/00/160/013	Austria, Belgium, Czech Republic, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden, Spain, United Kingdom	Denmark	22/06/2009
Aerius	Beragena Arzneimittel GmbH	06/07/2009	5 mg	Film-coated tablet	100	EU/1/00/160/013	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/11/2009
Aerius	Beragena Arzneimittel GmbH	06/07/2009	5 mg	Film-coated tablet	50	EU/1/00/160/012	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/11/2009
Aerius	Beragena Arzneimittel GmbH	06/07/2009	5 mg	Film-coated tablet	20	EU/1/00/160/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/11/2009
Aerius	Delfarma SpZoo	28/11/2008	5 mg	Film-coated tablet	30	EU/1/00/160/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland , Poland	12/02/2009
Aerius	Delfarma SpZoo	24/04/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/069	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	19/05/2009
Aerius	Dr Fisher Farma BV	04/02/2009	0.5mg/ml	Oral solution	150ml	EU/1/00/160/066	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	20/02/2009
Aerius	Emra-Med	14/05/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/066	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Slovak Republic, Slovenia	Germany	09/07/2009

Aerius	Emra-Med	07/09/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/069	Austria, Belgium, Cyprus, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Denmark	Germany	05/10/2009
Aerius	Eureco-Pharma B.V.	01/10/2008	5 mg	Orodispersible tablet	30	EU/1/00/160/056	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	09/02/2009
Aerius	Euro Registratie Collectief BV	02/07/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/066	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/07/2009
Aerius	Kohlpharma GmbH	19/01/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/160/063	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/01/2009
Aerius	O.P.D. Laboratories Ltd	05/06/2009	0.5 mg/ml	Oral solution	1 bottle - 120 ml	EU/1/00/160/065	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	23/06/2009
Aldara	EU-Pharma BV	13/03/2009	5% w/w	Cream	12 sachets	EU/1/98/080/001	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain, United Kingdom, Ireland	The Netherlands	08/04/2009
Aldara	Mediwin Ltd	02/09/2009	5% w/w	Cream	12 sachets	EU/1/98/080/001	Austria, Belgium, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	France	14/10/2009
Alimta	Aaston Healthcare GmbH	24/04/2009	500 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	26/08/2009
Alimta	CC Pharma	22/05/2009	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/06/2009
Alimta	Haemato Pharm AG	08/12/2008	100mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	Germany, Austria	03/03/2009

Alimta	Inopha GmbH	17/03/2009	500 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, The Netherlands, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	25/03/2009
Alimta	MedicoPharm AG	13/03/2009	500 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/04/290/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Norway	Germany	30/04/2009
alli	Doncaster Pharmaceuticals Group Ltd	11/05/2009	60 mg	Capsule, hard	42	EU/1/07/401/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	17/06/2009
alli	Doncaster Pharmaceuticals Group Ltd	12/05/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	17/06/2009
alli	Emra-Med	17/11/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
alli	Emra-Med	17/11/2009	60 mg	Capsule, hard	42	EU/1/07/401/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
alli	Euro Registratie Collectief BV	10/08/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	24/08/2009
alli	Ginova Ltd	07/10/2009	60 mg	Capsule, hard	42	EU/1/07/401/007	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	20/10/2009
alli	Medcor Pharmaceuticals BV	14/08/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	25/09/2009

alli	Orifarm AB	28/09/2009	60 mg	Capsule, hard	42	EU/1/07/401/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	22/10/2009
alli	Orifarm AB	28/09/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	22/10/2009
alli	Orifarm AS	02/10/2009	60 mg	Capsule, hard	84	EU/1/07/401/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	28/10/2009
Apidra	CC Pharma	15/10/2009	100 Units/ml	Solution for injection	5 vials	EU/1/04/285/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009
Apidra	Dr Fisher Farma BV	15/12/2008	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/04/285/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/02/2009
Apidra	Euro Registratie Collectief BV	27/01/2009	100U/ml	Solution for injection in a pre-filled pen	5x3ml	EU/1/04/285/032	Austria, Belgium, France, Germany, Greece, Italy, Ireland, Portugal, Spain, United Kingdom	The Netherlands	17/02/2009
Apidra	Pharma Gerke GmbH	27/02/2008	100 Units/ml	Solution for injection	3 cartridges	EU/1/04/285/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	28/05/2009
Apidra Solostar	Eureco-Pharma B.V.	12/01/2009	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/04/285/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	21/04/2009
Aprovel	2 Care 4	10/11/2008	300mg	Film-coated tablets	28		Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	26/01/2009
Aprovel	BCN Farma SL	08/10/2008	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, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	12/01/2009

Aprovel	Emra-Med	24/08/2009	300 mg	Film-coated tablet	56	EU/1/97/046/028	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Aprovel	Emra-Med	24/08/2009	300 mg	Film-coated tablet	98	EU/1/97/046/030	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Aprovel	Emra-Med	24/08/2009	300 mg	Film-coated tablet	28	EU/1/97/046/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Aprovel	O.P.D. Laboratories Ltd	17/06/2005	150 mg	Film-coated tablet	28	EU/1/97/046/022	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	12/08/2009
Aprovel	O.P.D. Laboratories Ltd	17/06/2005	300 mg	Film-coated tablet	28	EU/1/97/046/027	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	12/08/2009
Aprovel	O.P.D. Laboratories Ltd	17/06/2005	75 mg	Film-coated tablet	28	EU/1/97/046/017	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	12/08/2009
Aprovel	O.P.D. Laboratories Ltd	06/04/2009	75 mg	Film-coated tablet	90	EU/1/97/046/037	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	12/08/2009
Aprovel	Pharma Westen GmbH	09/03/2009	300 mg	Film-coated tablet	98	EU/1/97/046/030	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Slovak Republic	Germany	30/03/2009
Aprovel	Pharma Westen GmbH	04/12/2009	75 mg	Film-coated tablet	98	EU/1/97/046/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/12/2009
Aptivus	CC Pharma	16/01/2009	250 mg	Capsule, soft	120	EU/1/05/315/001	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	28/01/2009

Aranesp	Haemato Pharm AG	12/11/2008	150 µg	Solution for injection	4 pre-filled syringes	EU/1/01/185/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Germany, Greece, Hungary, Estonia, Lithuania, Latvia, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	21/01/2009
Aranesp	Haemato Pharm AG	21/08/2009	300 µg	Solution for injection	1 pre-filled syringe	EU/1/01/185/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Hungary, Germany, Greece, Estonia, Latvia, Lithuania, Liechtenstein, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	22/10/2009
Aranesp	Haemato Pharm AG	21/08/2009	500 µg	Solution for injection	1 pre-filled syringe	EU/1/01/185/031	Austria, Belgium, Cyprus, Denmark, Finland, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Hungary, Germany, France, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	22/10/2009
Aranesp	MedicoPharm AG	26/10/2009	500 µg	Solution for injection	1 pre-filled syringe	EU/1/01/185/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Arava	Delfarma SpZoo	16/01/2009	10 mg	Film-coated tablet	30	EU/1/99/118/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	23/02/2009
Arava	InPharm Spzoo	08/05/2009	20 mg	Film-coated tablet	30	EU/1/99/118/007	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	01/07/2009
Arava	PCO Manufacturing	07/05/2009	20 mg	Film-coated tablet	30	EU/1/99/118/007	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Spain, Sweden, The Netherlands, United Kingdom, Portugal, Italy	Ireland, United Kingdom	17/06/2009

Arava	PCO Manufacturing	09/06/2009	10 mg	Film-coated tablet	30	EU/1/99/118/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	Ireland, United Kingdom	12/11/2009
Arixtra	Abacus Medicine A/S	05/06/2009	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	29/06/2009
Arixtra	Abacus Medicine A/S	05/06/2009	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	29/06/2009
Arixtra	Abacus Medicine A/S	02/09/2009	1.5 mg/0.3 ml	Solution for injection	7 pre-filled syringes	EU/1/02/206/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/09/2009
Arixtra	Abacus Medicine A/S	02/09/2009	10 mg/0.8 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/09/2009
Arixtra	Abacus Medicine A/S	02/09/2009	5 mg/0.4 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/09/2009
Arixtra	Abacus Medicine A/S	02/09/2009	7.5 mg/0.6 ml	Solution for injection	10 pre-filled syringes	EU/1/02/206/014	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/09/2009
Arixtra	Dr Fisher Farma BV	05/02/2009	2.5 mg/0.5 ml	Solution for injection in a pre-filled syringe	10	EU/1/02/206/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/04/2009
Atriance	Haemato Pharm AG	02/10/2009	5 mg/ml	Solution for infusion	6 vials	EU/1/07/403/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/10/2009

Atripla	Aaston Healthcare GmbH	27/04/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	19/05/2009
Atripla	Abacus Medicine A/S	10/06/2009	600mg/200mg /245mg	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	30/06/2009
Atripla	Beragena Arzneimittel GmbH	15/06/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	08/07/2009
Atripla	Beragena Arzneimittel GmbH	15/06/2009	--	Film-coated tablet	3 x 30	EU/1/07/430/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	08/07/2009
Atripla	CC Pharma	30/12/2008	--	Film-coated tablet	30	EU/1/07/430/001	Norway, The Netherlands, United Kingdom	Germany	31/03/2009
Atripla	CC Pharma	30/12/2008	--	Film-coated tablet	3 x 30	EU/1/07/430/002	The Netherlands, United Kingdom, Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain	Germany	12/03/2009
Atripla	Dr Fisher Farma BV	15/12/2008	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	18/02/2009
Atripla	Euro Registratie Collectief BV	18/02/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	13/03/2009
Atripla	Haemato Pharm AG	02/03/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Germany, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	11/03/2009
Atripla	Haemato Pharm AG	29/06/2009	600mg/200mg /245mg	Film-coated tablet	3 x 30	EU/1/07/430/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/07/2009

Atripla	Kohlpharma GmbH	02/07/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/07/2009
Atripla	MedicoPharm AG	30/04/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/05/2009
Atripla	Pharma Gerke GmbH	08/01/2009	--	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	24/02/2009
Atripla	Pharma Gerke GmbH	08/01/2009	--	Film-coated tablet	3 x 30	EU/1/07/430/002	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	24/02/2009
Atripla	Pharma Westen GmbH	19/01/2009	600mg/200mg/245mg	Film-coated tablet	30	EU/1/07/430/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/05/2009
Avamys	2 Care 4	28/09/2009	27.5 micrograms/spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Denmark	08/10/2009
Avamys	B&S Healthcare Ltd	08/09/2009	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, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Ireland, United Kingdom	22/09/2009
Avamys	Euro Registratie Collectief BV	17/06/2009	27.5 micrograms/spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	The Netherlands	08/09/2009
Avamys	Medcor Pharmaceuticals BV	10/08/2009	27.5 micrograms/spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic	The Netherlands	28/08/2009
Avamys	Singad Pharma ApS	27/11/2009	27.5 micrograms/spray	Nasal spray, suspension	1 bottle	EU/1/07/434/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Poland, Czech Republic	Denmark	16/12/2009

Avandamet	2 Care 4	20/10/2009	4 mg/1000 mg	Film-coated tablet	56	EU/1/03/258/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/11/2009
Avandamet	BR Pharma International Ltd	16/01/2009	4 mg/1000 mg	Film-coated tablet	112	EU/1/03/258/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/09/2009
Avandamet	Chemilines Ltd	06/11/2009	2 mg/1000 mg	Film-coated tablet	168	EU/1/03/258/017	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	15/12/2009
Avandamet	Chemilines Ltd	06/11/2009	4 mg/1000 mg	Film-coated tablet	168	EU/1/03/258/018	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	15/12/2009
Avandamet	GenPlus Ltd	03/04/2009	2 mg/500 mg	Film-coated tablet	56	EU/1/03/258/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Cyprus, Iceland, Liechtenstein, Norway	United Kingdom, Ireland	15/04/2009
Avandamet	Ginova Ltd	24/12/2008	2/500mg	Film-coated tablets	56	EU/1/03/258/005	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	20/01/2009
Avandamet	Haemato Pharm AG	12/02/2009	2 mg/1000 mg	Film-coated tablet	112 (2x56)	EU/1/03/258/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Greece	Germany	17/03/2009
Avandamet	Haemato Pharm AG	12/02/2009	4 mg/1000 mg	Film-coated tablet	112 (2x56)	EU/1/03/258/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/03/2009
Avandamet	Medartuum AB	05/03/2009	4 mg/1000 mg	Film-coated tablet	112	EU/1/03/258/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Sweden	07/04/2009
Avandamet	Orifarm OY	08/05/2009	4 mg/1000 mg	Film-coated tablet	112	EU/1/03/258/014	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Finland	13/05/2009

Avandamet	Pharma Gerke GmbH	14/07/2008	2 mg/500 mg	Film-coated tablet	112	EU/1/03/258/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/03/2009
Avandamet	Pharmacodane Aps	27/11/2008	4/100mg	Film-coated tablets	56		Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	19/01/2009
Avandia	CC Pharma	15/10/2009	4 mg	Film-coated tablet	28	EU/1/00/137/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	01/12/2009
Avandia	CC Pharma	15/10/2009	4 mg	Film-coated tablet	112	EU/1/00/137/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	01/12/2009
Avandia	CC Pharma	15/10/2009	8 mg	Film-coated tablet	28	EU/1/00/137/011	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	01/12/2009
Avandia	CC Pharma	15/10/2009	8 mg	Film-coated tablet	112	EU/1/00/137/012	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	01/12/2009
Avandia	Crystal Pharma Ltd	01/07/2009	4 mg	Film-coated tablet	28	EU/1/00/137/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden	United Kingdom	28/08/2009
Avandia	Euro Registratie Collectief BV	14/08/2009	4 mg	Film-coated tablet	84	EU/1/00/137/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/09/2009
Avandia	G-Pharma Ltd	25/11/2009	8 mg	Film-coated tablet	84	EU/1/00/137/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, Romania	United Kingdom	10/12/2009
Avandia	Interport Ltd	02/09/2009	4 mg	Film-coated tablet	84	EU/1/00/137/014	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	15/09/2009
Avandia	Interport Ltd	02/09/2009	8 mg	Film-coated tablet	84	EU/1/00/137/015	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	15/09/2009

Avastin	Inopha GmbH	18/02/2009	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	14/04/2009
Avastin	Pharma Westen GmbH	28/05/2009	25 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/04/300/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	27/07/2009
Avonex	Aaston Healthcare GmbH	16/03/2009	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	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	24/06/2009
Avonex	Axicorp Pharma GmbH	06/07/2009	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled syringes + 12 needles	EU/1/97/033/004	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Lithuania, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/09/2009
Avonex	Beragena Arzneimittel GmbH	11/03/2008	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled syringes + 12 needles	EU/1/97/033/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	06/01/2009
Avonex	Haemato Pharm AG	14/08/2009	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled syringes + 12 needles	EU/1/97/033/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	16/10/2009
Avonex	Inopha GmbH	30/11/2009	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, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/12/2009
Avonex	Kohlpharma GmbH	14/08/2009	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled syringes + 12 needles	EU/1/97/033/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/09/2009
Avonex	Mevita HandelsGmbH	12/01/2009	30 µg/0.5 ml (6 million IU)	Solution for injection	4 pre-filled syringes + 4 needles	EU/1/97/033/003	Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009

Avonex	Pharma Gerke GmbH	26/02/2009	30 µg/0.5 ml (6 million IU)	Solution for injection	12 pre-filled syringes + 12 needles	EU/1/97/033/004	Austria, Belgium, Czech Republic, France, Greece, Italy, Norway, Portugal, Slovak Republic, Spain, The Netherlands, United Kingdom, Slovenia	Germany	12/08/2009
Azarga	Doncaster Pharmaceuticals Group Ltd	28/09/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	12/10/2009
Azarga	Eureco-Pharma B.V.	14/08/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/09/2009
Azarga	Euro Registratie Collectief BV	14/08/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/09/2009
Azarga	Interport Ltd	10/09/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	17/09/2009
Azarga	Kohlpharma GmbH	16/11/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
Azarga	Medartuum AB	03/11/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	3 bottles	EU/1/08/482/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	27/11/2009
Azarga	Medcor Pharmaceuticals BV	14/08/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/09/2009
Azarga	Pharmachim AB	10/08/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	29/09/2009
Azarga	Pharmachim AB	10/08/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	3 bottles	EU/1/08/482/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	29/09/2009
Azarga	Waymade PLC	23/09/2009	10 mg/ml / 5 mg/ml	Eye drops, suspension	1 bottle	EU/1/08/482/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, Malta, United Kingdom	United Kingdom, Malta, Ireland	05/10/2009

AZILECT	2 Care 4	13/05/2009	1 mg	Tablet	28	EU/1/04/304/003	Austria, Belgium, France, Germany, Greece, Hungary, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Denmark	28/05/2009
AZILECT	2 Care 4	12/06/2009	1 mg	Tablet	112	EU/1/04/304/006	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Denmark	26/06/2009
AZILECT	Axicorp Pharma GmbH	13/11/2009	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
AZILECT	Axicorp Pharma GmbH	13/11/2009	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
AZILECT	Beragena Arzneimittel GmbH	14/04/2009	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	12/06/2009
AZILECT	Beragena Arzneimittel GmbH	14/04/2009	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	22/06/2009
AZILECT	BR Pharma International Ltd	05/03/2009	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/04/2009
Azilect	CC Pharma	16/01/2009	1mg	Tablets	100	EU/1/04/304/005	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Czech Republic, Denmark, Finland, Sweden	Germany	28/01/2009
Azilect	CC Pharma	04/02/2009	1mg	Tablets	30	EU/1/04/304/004	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Czech Republic, Denmark, Finland, Sweden	Germany	20/02/2009
Azilect	Docpharm GmbH&CoKGaA	22/12/2008	1mg	Tablets	28		Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	28/01/2009
Azilect	Docpharm GmbH&CoKGaA	22/12/2008	1mg	Tablets	30		Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	28/01/2009

Azilect	Docpharm GmbH&CoKGaA	22/12/2008	1mg	Tablets	100		Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	28/01/2009
AZILECT	Emra-Med	26/05/2009	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Romania, Poland, Slovak Republic, Slovenia	Germany	30/06/2009
AZILECT	Emra-Med	26/05/2009	1 mg	Tablet	30	EU/1/04/304/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Germany	30/06/2009
Azilect	Paranova Danmark AS	06/05/2009	1 mg	Tablet	28	EU/1/04/304/003	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania	Denmark	26/06/2009
AZILECT	PCO Manufacturing	09/06/2009	1 mg	Tablet	28	EU/1/04/304/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	Ireland, United Kingdom	22/06/2009
AZILECT	Pharma Westen GmbH	26/11/2008	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	28/01/2009
Azilect	Pharma Westen GmbH	26/11/2008	1mg	Tablets	30		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	28/01/2009
AZILECT	Veron Pharma	16/03/2009	1 mg	Tablet	30	EU/1/04/304/004	Belgium, Austria, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	27/03/2009
AZILECT	Veron Pharma	16/03/2009	1 mg	Tablet	100	EU/1/04/304/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	27/03/2009
Azopt	2 Care 4	22/10/2009	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/12/2009

Azopt	2 Care 4	22/10/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/12/2009
Azopt	Abis-Pharma	14/08/2009	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	12/10/2009
Azopt	Abis-Pharma	14/08/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	12/10/2009
Azopt	Beragena Arzneimittel GmbH	04/09/2009	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	07/10/2009
Azopt	Beragena Arzneimittel GmbH	04/09/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	07/10/2009
Azopt	Emra-Med	04/06/2009	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Slovak Republic	Germany	03/09/2009
Azopt	Mediwin Ltd	22/05/2009	10 mg/ml	Eye drops, suspension	1 bottle	EU/1/00/129/001	Austria, Belgium, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	France	15/06/2009
Azopt	Milinda Arzneimittel GmbH	26/08/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Ireland, Portugal, United Kingdom, Italy	Germany	25/11/2009
Azopt	Orifarm AS	27/10/2009	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, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic, Czech Republic	Denmark	09/11/2009

Azopt	Paranova Danmark AS	17/06/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	12/10/2009
Azopt	Paranova Läkemedel AB	21/04/2009	10 mg/ml	Eye drops, suspension	3 bottles	EU/1/00/129/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	27/04/2009
Baraclude	CC Pharma	17/02/2009	0.5 mg	Film-coated tablet	30 x 1	EU/1/06/343/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic, Denmark, Finland, Sweden	Germany	14/05/2009
Baraclude	CC Pharma	17/02/2009	1 mg	Film-coated tablet	30 x 1	EU/1/06/343/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic	Germany	14/05/2009
Baraclude	CC Pharma	24/08/2009	0.5 mg	Film-coated tablet	90 x 1	EU/1/06/343/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	04/09/2009
Baraclude	CC Pharma	24/08/2009	1 mg	Film-coated tablet	90 x 1	EU/1/06/343/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	04/09/2009
Baraclude	Emra-Med	29/09/2008	0.5 mg	Film-coated tablet	90 x 1	EU/1/06/343/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	27/01/2009
Baraclude	Haemato Pharm AG	02/10/2009	0.5 mg	Film-coated tablet	30 x 1	EU/1/06/343/003	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	Germany, Austria	22/10/2009
Baraclude	Haemato Pharm AG	02/10/2009	1 mg	Film-coated tablet	30 x 1	EU/1/06/343/004	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	Germany, Austria	22/10/2009

Baraclude	Haemato Pharm AG	13/10/2009	0.5 mg	Film-coated tablet	90 x 1	EU/1/06/343/006	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	22/10/2009
Baraclude	Haemato Pharm AG	13/10/2009	1 mg	Film-coated tablet	90 x 1	EU/1/06/343/007	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	22/10/2009
Betaferon	Aaston Healthcare GmbH	23/03/2009	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, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	06/05/2009
Betaferon	Axicorp Pharma GmbH	30/01/2009	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, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Lithuania, Luxembourg, Norway, Portugal, Spain, Sweden, United Kingdom, The Netherlands, Bulgaria, Poland, Romania	Germany	06/04/2009
Betaferon	Beragena Arzneimittel GmbH	16/04/2009	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, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Poland, Bulgaria	Germany	04/06/2009
Betaferon	CC Pharma	12/05/2009	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	Bulgaria, Hungary, Poland, Sweden, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, Romania, Finland	Germany	22/07/2009

Betaferon	CC Pharma	27/11/2009	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, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Finland, Sweden, Lithuania, Slovak Republic, Romania, Slovenia, Poland	Germany	07/12/2009
Betaferon	CC Pharma	27/11/2009	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, France, Italy, Greece, The Netherlands, Norway, Portugal, Spain, United Kingdom, Finland, Sweden, Lithuania, Slovak Republic, Romania, Slovenia, Poland	Germany	07/12/2009
Betaferon	Emra-Med	21/10/2009	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	02/12/2009
Betaferon	Haemato Pharm AG	06/04/2009	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, Denmark, Finland, France, Greece, Iceland, Italy, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	11/06/2009
Betaferon	Kohlpharma GmbH	15/04/2009	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, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Germany	07/05/2009

Betaferon	Kohlpharma GmbH	21/09/2009	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, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom, Slovak Republic	Germany	26/11/2009
Betaferon	MedicoPharm AG	16/04/2009	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, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	11/08/2009
Betaferon	Pharma Gerke GmbH	20/08/2007	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, France, Greece, Italy, Lithuania, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania, Finland, Poland, Latvia	Germany	18/02/2009
Betaferon	Pharma Gerke GmbH	22/06/2009	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, Finland, France, Greece, Hungary, Italy, Lithuania, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania, Poland	Germany	08/12/2009
Betaferon	Pharma Gerke GmbH	26/11/2009	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, Finland, France, Greece, Hungary, Italy, Lithuania, The Netherlands, Norway, Poland, Portugal, Romania, Spain, United Kingdom, Slovak Republic, Czech Republic	Germany	21/12/2009

Betaferon	Pharma Gerke GmbH	26/11/2009	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, Finland, France, Greece, Hungary, Italy, Lithuania, The Netherlands, Norway, Poland, Portugal, Romania, Spain, United Kingdom	Germany	21/12/2009
Bondronat	Abacus Medicine A/S	22/05/2009	50 mg	Film-coated tablet	28	EU/1/96/012/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/06/2009
Bondronat	Abacus Medicine A/S	22/05/2009	50 mg	Film-coated tablet	84	EU/1/96/012/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/06/2009
Bondronat	Haemato Pharm AG	01/06/2009	6mg/6 ml	Concentrate for solution for infusion	1 vial	EU/1/96/012/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany, Austria	09/06/2009
Bondronat	Haemato Pharm AG	29/06/2009	6mg/6 ml	Concentrate for solution for infusion	5 vials	EU/1/96/012/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/07/2009
Bondronat	Kohlpharma GmbH	16/03/2009	50 mg	Film-coated tablet	28	EU/1/96/012/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/03/2009
Bondronat	Kohlpharma GmbH	16/03/2009	50 mg	Film-coated tablet	84	EU/1/96/012/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/03/2009
Bondronat	Kohlpharma GmbH	03/08/2009	6mg/6 ml	Concentrate for solution for infusion	1 vial	EU/1/96/012/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Germany	16/09/2009

Bondronat	Parallell Pharma AB	13/07/2009	50 mg	Film-coated tablet	28	EU/1/96/012/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	11/09/2009
Bondronat	Paranova Danmark AS	04/12/2009	50 mg	Film-coated tablet	84	EU/1/96/012/010	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/12/2009
Bondronat	Paranova Danmark AS	07/12/2009	50 mg	Film-coated tablet	28	EU/1/96/012/009	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/12/2009
Bondronat	Paranova Läkemedel AB	22/05/2009	50 mg	Film-coated tablet	84	EU/1/96/012/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	30/06/2009
Bondronat	Singad Pharma ApS	14/07/2009	50 mg	Film-coated tablet	28	EU/1/96/012/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	30/11/2009
Bondronat	Singad Pharma ApS	20/07/2009	50 mg	Film-coated tablet	84	EU/1/96/012/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/11/2009
Bonviva	Axicorp Pharma GmbH	26/01/2009	3 mg/3ml	Solution for injection	1 pre-filled syringe	EU/1/03/265/005	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic	Germany	16/02/2009
Bonviva	B&S Healthcare Ltd	01/04/2009	3 mg/3ml	Solution for injection	1 pre-filled syringe	EU/1/03/265/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Hungary, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	United Kingdom, Ireland	26/05/2009
Bonviva	Beragena Arzneimittel GmbH	28/07/2009	150 mg	Film-coated tablet	1	EU/1/03/265/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/11/2009
Bonviva	Beragena Arzneimittel GmbH	28/07/2009	150 mg	Film-coated tablet	3	EU/1/03/265/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/11/2009

Bonviva	Delfarma SpZoo	18/06/2009	3 mg/3ml	Solution for injection	1 pre-filled syringe	EU/1/03/265/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	13/08/2009
Bonviva	Europharma DK	26/08/2008	150mg	Film-coated tablets	1		Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	29/01/2009
Bonviva	Europharma DK	26/08/2008	150mg	Film-coated tablets	3		Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	29/01/2009
Bonviva	Polyfarma Ltd	13/08/2007	150 mg	Film-coated tablet	1	EU/1/03/265/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Poland, Portugal, Spain, The Netherlands	Ireland, United Kingdom	03/12/2009
Bonviva	Singad Pharma ApS	05/05/2008	150 mg	Film-coated tablet	1	EU/1/03/265/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	19/05/2009
Busilvex	CC Pharma	04/08/2009	6 mg/ml	Concentrate for solution for infusion	8 vials	EU/1/03/254/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	28/08/2009
BYETTA	Kohlpharma GmbH	15/04/2009	10 micrograms	Solution for injection	1 pre-filled pen	EU/1/06/362/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/06/2009
BYETTA	Kohlpharma GmbH	15/04/2009	10 micrograms	Solution for injection	3 pre-filled pens	EU/1/06/362/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/06/2009
BYETTA	Kohlpharma GmbH	15/04/2009	5 micrograms	Solution for injection	1 pre-filled pen	EU/1/06/362/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/06/2009
Caelyx	Aaston Healthcare GmbH	09/04/2009	2.0 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	26/05/2009

Caelyx	Delfarma SpZoo	18/06/2009	2.0 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	28/08/2009
Caelyx	Inopha GmbH	29/12/2008	2.0 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Iceland, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	06/03/2009
Caelyx	Inopha GmbH	28/07/2009	2.0 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	30/07/2009
Caelyx	MedicoPharm AG	24/06/2009	2.0 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/96/011/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	20/07/2009
Cancidas	CC Pharma	20/01/2009	50mg	Powder for concentrate for solution for infusion	1	EU/1/01/196/001	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Romania	Germany	25/02/2009
Cancidas	CC Pharma	20/01/2009	70mg	Powder for concentrate for solution for infusion	1	EU/1/01/196/003	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Romania	Germany	25/02/2009
Cancidas	Orifarm AS	05/05/2009	50 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania, Cyprus, Lithuania	Denmark	17/06/2009
Cancidas	Orifarm AS	05/05/2009	70 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania, Cyprus	Denmark	17/06/2009

Cancidas	Pharma Gerke GmbH	02/09/2009	50 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/196/001	Austria, Belgium, France, Finland, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	26/11/2009
CellCept	Beragena Arzneimittel GmbH	14/04/2009	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	22/06/2009
CellCept	Beragena Arzneimittel GmbH	14/04/2009	500 mg	Film-coated tablet	150	EU/1/96/005/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	22/06/2009
CellCept	BR Pharma International Ltd	09/02/2009	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/02/2009
CellCept	BR Pharma International Ltd	09/02/2009	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/02/2009
CellCept	BR Pharma International Ltd	31/03/2009	500 mg	Film-coated tablet	150	EU/1/96/005/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/04/2009
CellCept	BR Pharma International Ltd	01/04/2009	250 mg	Capsule, hard	300	EU/1/96/005/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/04/2009
CellCept	Combiphar Europe BV	22/01/2009	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/05/2009
CellCept	Haemato Pharm AG	13/08/2009	250 mg	Capsule, hard	100	EU/1/96/005/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany	18/08/2009

CellCept	Haemato Pharm AG	07/09/2009	500 mg	Film-coated tablet	50	EU/1/96/005/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany	03/11/2009
CellCept	Haemato Pharm AG	07/09/2009	500 mg	Film-coated tablet	150	EU/1/96/005/004	Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Hungary, Germany, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	03/11/2009
CellCept	Haemato Pharm AG	11/09/2009	250 mg	Capsule, hard	300	EU/1/96/005/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany	22/09/2009
CellCept	Imed Healthcare Ltd	17/04/2009	250 mg	Capsule, hard	100	EU/1/96/005/001	United Kingdom, Belgium, Cyprus, Denmark, Finland, France, Germany, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Sweden, The Netherlands	Ireland, Malta, United Kingdom	30/04/2009
CellCept	Kohlpharma GmbH	27/10/2009	1 g/5 ml	Powder for oral suspension	1 bottle	EU/1/96/005/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/11/2009
CellCept	Milinda Arzneimittel GmbH	22/10/2009	500 mg	Film-coated tablet	50	EU/1/96/005/002	Ireland, United Kingdom	Germany	30/11/2009
CellCept	Milinda Arzneimittel GmbH	22/10/2009	500 mg	Film-coated tablet	150	EU/1/96/005/004	Ireland, United Kingdom	Germany	30/11/2009
CellCept	Orifarm AS	02/07/2009	250 mg	Capsule, hard	300	EU/1/96/005/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Denmark	28/08/2009
CellCept	Paranova Danmark AS	21/09/2009	500 mg	Film-coated tablet	150	EU/1/96/005/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/10/2009

CellCept	Pharma Gerke GmbH	23/12/2008	500 mg	Film-coated tablet	150	EU/1/96/005/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland, Romania	Germany	02/02/2009
CellCept	Pharma Gerke GmbH	09/02/2009	250 mg	Capsule, hard	300	EU/1/96/005/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/02/2009
Celsentri	CC Pharma	19/01/2009	150 mg	Film-coated tablet	60	EU/1/07/418/003	Norway, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	27/01/2009
Celsentri	CC Pharma	19/01/2009	300 mg	Film-coated tablet	60	EU/1/07/418/008	Norway, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, Czech Republic	Germany	27/01/2009
Celsentri	Haemato Pharm AG	01/10/2009	150 mg	Film-coated tablet	60	EU/1/07/418/003	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	Germany, Austria	23/10/2009
Celsentri	Haemato Pharm AG	01/10/2009	300 mg	Film-coated tablet	60	EU/1/07/418/008	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Germany	Germany, Austria	23/10/2009
Celsentri	Kohlpharma GmbH	15/06/2009	150 mg	Film-coated tablet	60	EU/1/07/418/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/09/2009
Celsentri	Kohlpharma GmbH	20/07/2009	300 mg	Film-coated tablet	60	EU/1/07/418/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/09/2009
Celsentri	Medcor Pharmaceuticals BV	30/03/2009	150 mg	Film-coated tablet	60	EU/1/07/418/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	08/04/2009
Celsentri	Pharma Gerke GmbH	26/10/2009	150 mg	Film-coated tablet	60	EU/1/07/418/003	Austria, Belgium, Finland, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	03/11/2009

Cervarix	CC Pharma	09/03/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/07/419/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	20/04/2009
Cervarix	Dr Fisher Farma BV	12/05/2009	- -	Suspension for injection	1 pre-filled syringe + 1 needle	EU/1/07/419/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	18/09/2009
Cervarix	Kohlpharma GmbH	24/03/2009	- -	Suspension for injection	1 pre-filled syringe + 1 needle	EU/1/07/419/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Norway, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/04/2009
Cervarix	Pharma Westen GmbH	27/01/2009	- -	Suspension for injection	10 pre-filled syringes + 10 needles	EU/1/07/419/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Slovenia, Slovak Republic, Czech Republic	Germany	11/02/2009
Champix	Extrapharm BV	26/03/2009	0.5 mg	Film-coated tablet	56 (bottle)	EU/1/06/360/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/05/2009
Champix	Extrapharm BV	26/03/2009	1.0 mg	Film-coated tablet	56 (blister)	EU/1/06/360/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/05/2009
Champix	G-Pharma Ltd	10/06/2009	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	05/08/2009
Champix	Kohlpharma GmbH	30/11/2009	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Germany	11/12/2009
Champix	Medcor Pharmaceuticals BV	26/05/2009	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, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	The Netherlands	06/08/2009

Champix	Medcor Pharmaceuticals BV	26/05/2009	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, Sweden, United Kingdom, Poland	The Netherlands	06/08/2009
Champix	MPT Pharma	25/02/2009	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, Hungary, Greece, Germany, France, Finland, Estonia, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Romania, Portugal, Poland, Norway, Malta, Luxembourg, United Kingdom, The Netherlands, Sweden, Spain	Ireland, United Kingdom	01/04/2009
Champix	O.P.D. Laboratories Ltd	03/12/2007	0.5mg	Film-coated tablets	56	EU/1/06/360/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	03/12/2009
Champix	O.P.D. Laboratories Ltd	03/12/2007	0.5mg + 1mg	Film-coated tablets	11+14	EU/1/06/360/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	03/12/2009
Champix	O.P.D. Laboratories Ltd	03/12/2007	1mg	Film-coated tablets	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	03/12/2009
Champix	O.P.D. Laboratories Ltd	03/12/2007	1mg	Film-coated tablets	56	EU/1/06/360/005	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	03/12/2009
Champix	Orifarm AB	23/11/2009	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, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom	Sweden	08/12/2009
Champix	Orifarm AB	23/11/2009	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, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom	Sweden	08/12/2009
Champix	Orifarm AB	23/11/2009	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom	Sweden	08/12/2009
Champix	Orifarm AB	23/11/2009	1.0 mg	Film-coated tablet	28	EU/1/06/360/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom	Sweden	08/12/2009

Champix	Orifarm AB	23/11/2009	1.0 mg	Film-coated tablet	112	EU/1/06/360/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom, Cyprus	Sweden	08/12/2009
Champix	PCO Manufacturing	30/09/2008	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, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	United Kingdom, Ireland	08/04/2009
Champix	PCO Manufacturing	30/09/2008	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	United Kingdom, Ireland	08/04/2009
Champix	PCO Manufacturing	06/07/2009	1.0 mg	Film-coated tablet	56	EU/1/06/360/005	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	20/07/2009
Champix	Pharma Westen GmbH	27/03/2009	1.0 mg	Film-coated tablet	56 (blister in a card)	EU/1/06/360/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	07/05/2009
Champix	Polyfarma Ltd	03/10/2007	1.0 mg	Film-coated tablet	28	EU/1/06/360/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain	United Kingdom, Ireland	11/03/2009
Champix	Waymade PLC	19/03/2009	1.0 mg	Film-coated tablet	28 in a carton	EU/1/06/360/009	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Italy, The Netherlands, Norway, Poland, Portugal, Spain, Sweden	United Kingdom	08/05/2009
Cholestagel	Docpharm GmbH&CoKGaA	31/07/2009	625 mg	Film-coated tablet	180	EU/1/03/268/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	18/09/2009
Cholestagel	Kohlpharma GmbH	17/10/2008	625 mg	Film-coated tablet	180	EU/1/03/268/003	Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	10/02/2009
Cholestagel	Kohlpharma GmbH	02/06/2009	625 mg	Film-coated tablet	180 (no outer carton)	EU/1/03/268/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/07/2009
Cialis	2 Care 4	14/09/2009	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/12/2009

Cialis	2 Care 4	14/09/2009	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/12/2009
Cialis	2 Care 4	02/10/2009	20 mg	Film-coated tablet	8	EU/1/02/237/004	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/12/2009
Cialis	Abis-Pharma	14/08/2009	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	Germany	12/10/2009
Cialis	Abis-Pharma	14/08/2009	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	Germany	12/10/2009
Cialis	Abis-Pharma	14/08/2009	20 mg	Film-coated tablet	12	EU/1/02/237/005	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	12/10/2009
Cialis	AN Pharmacy Sp. z o.o.	07/04/2009	20 mg	Film-coated tablet	2	EU/1/02/237/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	06/05/2009
Cialis	AN Pharmacy Sp. z o.o.	07/04/2009	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, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	06/05/2009

Cialis	AN Pharmacy Sp. z o.o.	07/04/2009	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, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	06/05/2009
Cialis	AN Pharmacy Sp. z o.o.	07/04/2009	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, Spain, Sweden, The Netherlands, United Kingdom	Slovenia	06/05/2009
Cialis	AN Pharmacy Sp. z o.o.	24/08/2009	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, Sweden, The Netherlands, United Kingdom	Austria, Germany	26/11/2009
Cialis	AN Pharmacy Sp. z o.o.	24/08/2009	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, Sweden, The Netherlands, United Kingdom	Austria, Germany	26/11/2009
Cialis	BR Lewis Ltd	04/12/2008	20 mg	Film-coated tablet	2	EU/1/02/237/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Romania, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	27/04/2009
Cialis	Docpharm GmbH&CoKGaA	05/03/2009	10 mg	Film-coated tablet	4	EU/1/02/237/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	31/03/2009
Cialis	Dr Fisher Farma BV	27/04/2009	20 mg	Film-coated tablet	8	EU/1/02/237/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/05/2009

Cialis	Emra-Med	08/05/2009	20 mg	Film-coated tablet	12	EU/1/02/237/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Romania, Poland, Slovak Republic, Slovenia	Germany	18/09/2009
Cialis	Emra-Med	18/08/2009	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/09/2009
Cialis	Extrapharm BV	20/04/2009	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/05/2009
Cialis	Gervasi Farmacia SL	24/11/2008	20mg	Film-coated tablets	2	EU/1/02/237/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	26/02/2009
Cialis	Gervasi Farmacia SL	24/11/2008	20mg	Film-coated tablets	4	EU/1/02/237/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	26/02/2009
Cialis	Gervasi Farmacia SL	24/11/2008	20mg	Film-coated tablets	8	EU/1/02/237/004	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	26/02/2009
Cialis	Ichem SpZoo	18/03/2009	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, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	16/06/2009

Cialis	InPharm Spzoo	01/04/2009	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, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	11/05/2009
Cialis	Kohlpharma GmbH	22/12/2008	5 mg	Film-coated tablet	28	EU/1/02/237/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/01/2009
Cialis	MedicoPharm AG	08/07/2009	20 mg	Film-coated tablet	4	EU/1/02/237/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2009
Cialis	Pharma Westen GmbH	10/02/2009	10mg	Film-coated tablets	4	EU/1/02/237/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	27/02/2009
Circadin	B&S Healthcare Ltd	21/08/2009	2 mg	Prolonged-release tablet	21	EU/1/07/392/001	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	09/09/2009
Circadin	BR Lewis Ltd	16/12/2008	2 mg	Prolonged-release tablet	21	EU/1/07/392/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	06/01/2009
Circadin	CC Pharma	01/06/2009	2 mg	Prolonged release tablet	21	EU/1/07/392/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	09/07/2009
Circadin	Eureco-Pharma B.V.	15/06/2009	2 mg	Prolonged release tablet	21	EU/1/07/392/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/06/2009
Circadin	Kohlpharma GmbH	31/03/2009	2 mg	Prolonged release tablet	21	EU/1/07/392/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Norway, Portugal, Spain, Sweden, United Kingdom, The Netherlands, Malta	Germany	12/05/2009

Circadin	O.P.D. Laboratories Ltd	12/02/2009	2mg	Prolonged released tablets	21	EU/1/07/392/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	27/02/2009
Circadin	Swingward Ltd TA Medihealth	03/06/2009	2 mg	Prolonged release tablet	21	EU/1/07/392/001	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	03/07/2009
Circadin	Waymade PLC	13/08/2009	2 mg	Prolonged-release tablet	21	EU/1/07/392/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, Ireland, Malta, United Kingdom	United Kingdom, Ireland, Malta	28/08/2009
CoAprovel	2 Care 4	02/06/2009	300 mg/12.50 mg	Film-coated tablet	98	EU/1/98/086/020	Austria, Belgium, France, Germany, Greece, Hungary, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/08/2009
CoAprovel	2 Care 4	02/06/2009	300 mg/25 mg	Film-coated tablet	98	EU/1/98/086/027	Austria, Belgium, France, Germany, Greece, Hungary, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	07/08/2009
CoAprovel	2 Care 4	23/06/2009	150 mg/12.50 mg	Film-coated tablet	98	EU/1/98/086/015	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/08/2009
CoAprovel	Docpharm GmbH&CoKGaA	12/06/2009	150 mg/12.50 mg	Film-coated tablet	28	EU/1/98/086/012	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Ireland, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	28/08/2009
CoAprovel	Docpharm GmbH&CoKGaA	12/06/2009	150 mg/12.50 mg	Film-coated tablet	56	EU/1/98/086/013	Austria, Belgium, Denmark, Finland, France, Greece, Italy, Ireland, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	28/08/2009
CoAprovel	Docpharm GmbH&CoKGaA	12/06/2009	150 mg/12.50 mg	Film-coated tablet	98	EU/1/98/086/015	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	28/08/2009
CoAprovel	Emra-Med	14/05/2009	150 mg/12.50 mg	Film-coated tablet	56	EU/1/98/086/013	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/06/2009

CoAprovel	Emra-Med	14/05/2009	150 mg/12.50 mg	Film-coated tablet	28	EU/1/98/086/012	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/06/2009
CoAprovel	Emra-Med	14/05/2009	150 mg/12.50 mg	Film-coated tablet	98	EU/1/98/086/015	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/06/2009
CoAprovel	Emra-Med	06/07/2009	300 mg/25 mg	Film-coated tablet	98	EU/1/98/086/027	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	17/07/2009
CoAprovel	Emra-Med	06/07/2009	300 mg/25 mg	Film-coated tablet	56	EU/1/98/086/025	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	17/07/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	150/12.5mg	Film-coated tablets	30		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	150/12.5mg	Film-coated tablets	90		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	300/12.5mg	Film-coated tablets	30		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	300/12.5mg	Film-coated tablets	90		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	300/25mg	Film-coated tablets	30		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Euro Registratie Collectief BV	14/10/2008	300/25mg	Film-coated tablets	90		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	15/01/2009
CoAprovel	Pharma Westen GmbH	19/01/2009	300 mg/12.50 mg	Film-coated tablet	98	EU/1/98/086/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/05/2009

CoAprovel	Pharma Westen GmbH	28/05/2009	150 mg/12.50 mg	Tablet	98	EU/1/98/086/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	23/07/2009
CoAprovel	Swingward Ltd TA Medihealth	01/02/2008	300 mg/12.50 mg	Tablet	28	EU/1/98/086/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	26/05/2009
Combivir	Aaston Healthcare GmbH	01/04/2009	150 mg/300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Cyprus, Denmark, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, France, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Germany	13/05/2009
Combivir	InPharm Spzoo	09/07/2009	150 mg/300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	31/07/2009
Combivir	SIA 'Elpis'	12/01/2009	150 mg/300 mg	Film-coated tablet	60	EU/1/98/058/001	Austria, Belgium, Bulgaria, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	24/03/2009
Competact	B&S Healthcare Ltd	01/09/2009	15 mg/850 mg	Film-coated tablet	60	EU/1/06/354/006	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	25/11/2009
Competact	CC Pharma	08/06/2009	15 mg/850 mg	Film-coated tablet	98	EU/1/06/354/008	Austria, Belgium, France, Greece, Italy, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	11/06/2009
Competact	Chemilines Ltd	03/08/2009	15 mg/850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	15/09/2009

Competact	Doncaster Pharmaceuticals Group Ltd	22/07/2009	15 mg/850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	03/09/2009
Competact	Emra-Med	11/11/2009	15 mg/850 mg	Film-coated tablet	98	EU/1/06/354/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Competact	Interport Ltd	08/09/2009	15 mg/850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	14/10/2009
Competact	Kohlpharma GmbH	28/09/2009	15 mg/850 mg	Film-coated tablet	98	EU/1/06/354/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/10/2009
Competact	Lexon (UK) Ltd	16/04/2009	15 mg/850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	24/04/2009
Competact	Pharma Westen GmbH	27/04/2009	15 mg/850 mg	Film-coated tablet	98	EU/1/06/354/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	02/06/2009
Competact	Swingward Ltd TA Medihealth	12/02/2009	15 mg/850 mg	Film-coated tablet	56	EU/1/06/354/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	24/03/2009
Competact	Waymade PLC	17/12/2008	15/850mg	Film-coated tablets	56		Austria, Belgium, Denmark, Finland, France, Germany, Greece, Italy, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	13/01/2009
Comtess	Axicorp Pharma GmbH	13/08/2009	200 mg	Film-coated tablet	100	EU/1/98/082/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/10/2009
Comtess	CC Pharma	19/01/2009	200 mg	Film-coated tablet	100	EU/1/98/082/003	United Kingdom, Italy	Germany	27/01/2009
Comtess	Euro Registratie Collectief BV	18/03/2009	200 mg	Film-coated tablet	30	EU/1/98/082/001	Austria, Belgium, France, Germany, Ireland, Italy, Greece, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/03/2009

Comtess	PCO Manufacturing	12/05/2009	200 mg	Film-coated tablet	100	EU/1/98/082/003	Austria, Belgium, Cyprus, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Denmark, Italy	United Kingdom, Ireland	15/10/2009
Comtess	Pharma Gerke GmbH	15/01/2009	200mg	Film-coated tablets	100		Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/01/2009
Cymbalta	Aaston Healthcare GmbH	06/04/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	19/05/2009
Cymbalta	Axicorp Pharma GmbH	03/04/2009	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Lithuania, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/05/2009
Cymbalta	B&S Healthcare Ltd	02/07/2009	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, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Ireland, United Kingdom	28/08/2009
Cymbalta	Beragena Arzneimittel GmbH	22/05/2009	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	11/06/2009
Cymbalta	Beragena Arzneimittel GmbH	22/05/2009	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	11/06/2009
Cymbalta	Emra-Med	06/04/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/07/2009
Cymbalta	Haemato Pharm AG	12/11/2008	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/02/2009
Cymbalta	Haemato Pharm AG	12/11/2008	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/02/2009

Cymbalta	Haemato Pharm AG	12/11/2008	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/02/2009
Cymbalta	Interport Ltd	08/12/2008	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/296/006	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	03/02/2009
Cymbalta	Mevita HandelsGmbH	06/04/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	29/04/2009
Cymbalta	Mevita HandelsGmbH	06/04/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	29/04/2009
Cymbalta	Milinda Arzneimittel GmbH	04/09/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	France, Ireland, Italy, United Kingdom	Germany	27/11/2009
Cymbalta	Milinda Arzneimittel GmbH	04/09/2009	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	France, Ireland, Italy, United Kingdom	Germany	27/11/2009
Cymbalta	Milinda Arzneimittel GmbH	04/09/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	27/11/2009
Cymbalta	Milinda Arzneimittel GmbH	04/09/2009	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	27/11/2009
Cymbalta	O.P.D. Laboratories Ltd	27/10/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/296/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	03/11/2009
Cymbalta	PCO Manufacturing	07/05/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Italy	Ireland, United Kingdom	29/05/2009
Cymbalta	PCO Manufacturing	08/06/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	02/07/2009
Cymbalta	Pharma Gerke GmbH	27/05/2008	30 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Finland, Ireland, Sweden	Germany	06/07/2009
Cymbalta	Veron Pharma	13/02/2009	30mg	Gastro-resistant capsules	28	EU/1/04/296/001	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom, Sweden, Finland	Germany	27/02/2009

Cymbalta	Veron Pharma	07/04/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/296/002	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom, Finland, Sweden	Germany	15/05/2009
Cymbalta	Veron Pharma	08/04/2009	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/296/004	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom, Finland, Sweden	Germany	15/05/2009
Dukoral	Eurim-Pharm	05/06/2008	-	Suspension of vaccine and effervescent granules for oral solution	2x1		Italy	Germany	29/01/2009
Dukoral	Kohlpharma GmbH	19/02/2009	- -	Suspension of vaccine and effervescent granules for oral solution	2 x 1 dose	EU/1/03/263/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Sweden	Germany	23/03/2009
DuoTrav	B&S Healthcare Ltd	23/01/2009	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	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	United Kingdom, Ireland	21/07/2009
DuoTrav	Kohlpharma GmbH	08/05/2009	40 µg /ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/338/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Poland	Germany	26/06/2009
DuoTrav	Kohlpharma GmbH	15/06/2009	40 µg /ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/338/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Poland	Germany	06/07/2009
DuoTrav	Medartuum AB	24/02/2009	40 µg /ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/338/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	26/03/2009
Dynastat	CC Pharma	23/06/2009	40 mg	Powder for solution for injection	10 vials	EU/1/02/209/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	21/09/2009
Dynepo	Kohlpharma GmbH	04/11/2008	3.000IU	Solution for injection in a pre-filled syringe	6		Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/01/2009

Dynepo	Kohlpharma GmbH	04/11/2008	6.000IU	Solution for injection in a pre-filled syringe	6		Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/01/2009
Ebixa	BR Pharma International Ltd	29/10/2008	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Poland	06/02/2009
ECALTA	Haemato Pharm AG	01/10/2009	100 mg	Powder and solvent for concentrate for solution for infusion	1 vial + 1 vial	EU/1/07/416/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/11/2009
Efient	Docpharm GmbH&CoKGaA	10/08/2009	10 mg	Film-coated tablet	28	EU/1/08/503/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/09/2009
Efient	Docpharm GmbH&CoKGaA	10/08/2009	10 mg	Film-coated tablet	98	EU/1/08/503/014	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/09/2009
Efient	Docpharm GmbH&CoKGaA	10/08/2009	5 mg	Film-coated tablet	28	EU/1/08/503/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/09/2009
Efient	Docpharm GmbH&CoKGaA	10/08/2009	5 mg	Film-coated tablet	98	EU/1/08/503/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/09/2009
Efient	Pharma Westen GmbH	02/07/2009	10 mg	Film-coated tablet	98	EU/1/08/503/014	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2009
Elaprase	CC Pharma	08/06/2009	2 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/06/365/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	11/06/2009
Emend	Abacus Medicine A/S	22/05/2009	125 mg + 80 mg	Capsule, hard	1 (125 mg) + 2 (80 mg)	EU/1/03/262/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic	Denmark	15/07/2009

Emend	Dr Fisher Farma BV	11/11/2009	80 mg	Capsule, hard	2	EU/1/03/262/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Luxembourg	The Netherlands	24/11/2009
Emend	Medcor Pharmaceuticals BV	15/10/2008	80 mg	Capsule, hard	2	EU/1/03/262/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/02/2009
Emend	PCO Manufacturing	01/12/2009	125 mg + 80 mg	Capsule, hard	1 (125 mg) + 2 (80 mg)	EU/1/03/262/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	18/12/2009
Emselex	CC Pharma	01/10/2009	15 mg	Prolonged-release tablet	28	EU/1/04/294/023	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/10/2009
Emselex	CC Pharma	01/10/2009	15 mg	Prolonged-release tablet	98	EU/1/04/294/026	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/10/2009
Emselex	CC Pharma	01/10/2009	7.5 mg	Prolonged-release tablet	28	EU/1/04/294/017	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/10/2009
Emselex	CC Pharma	01/10/2009	7.5 mg	Prolonged-release tablet	98	EU/1/04/294/020	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/10/2009
Emselex	Emra-Med	08/05/2009	15 mg	Prolonged release tablet	49	EU/1/04/294/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2009
Emselex	Emra-Med	08/05/2009	15 mg	Prolonged release tablet	28	EU/1/04/294/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2009
Emselex	Emra-Med	08/05/2009	7.5 mg	Prolonged release tablet	28	EU/1/04/294/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2009
Emselex	Emra-Med	08/05/2009	7.5 mg	Prolonged release tablet	49	EU/1/04/294/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/05/2009

Emselex	Kohlpharma GmbH	31/03/2009	7.5 mg	Prolonged release tablet	28	EU/1/04/294/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	17/04/2009
Emselex	Kohlpharma GmbH	31/03/2009	7.5 mg	Prolonged release tablet	49	EU/1/04/294/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/04/2009
Emselex	Pharma Westen GmbH	18/02/2009	15 mg	Prolonged release tablet	98	EU/1/04/294/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/03/2009
Emselex	Veron Pharma	12/06/2009	15 mg	Prolonged release tablet	28	EU/1/04/294/023	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	21/09/2009
Emselex	Veron Pharma	12/06/2009	15 mg	Prolonged release tablet	49	EU/1/04/294/024	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	21/09/2009
Emselex	Veron Pharma	12/06/2009	15 mg	Prolonged release tablet	98	EU/1/04/294/026	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	21/09/2009
Emselex	Veron Pharma	12/06/2009	7.5 mg	Prolonged release tablet	28	EU/1/04/294/017	Austria, Belgium, France, Greece, Italy, Portugal, The Netherlands, Spain, United Kingdom	Germany	21/09/2009
Emselex	Veron Pharma	12/06/2009	7.5 mg	Prolonged release tablet	49	EU/1/04/294/018	Austria, Belgium, France, Italy, Portugal, Spain, The Netherlands, United Kingdom, Greece	Germany	21/09/2009
Emselex	Veron Pharma	12/06/2009	7.5 mg	Prolonged release tablet	98	EU/1/04/294/020	Austria, Belgium, France, Greece, Italy, Portugal, The Netherlands, Spain, United Kingdom	Germany	21/09/2009
Emtriva	SIA 'Elpis'	10/02/2009	200 mg	Capsule, hard	30	EU/1/03/261/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	06/04/2009

Enbrel	Aaston Healthcare GmbH	20/04/2009	25 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Italy, Ireland, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	06/07/2009
Enbrel	Abacus Medicine A/S	13/07/2009	25 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/013	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	23/07/2009
Enbrel	Abacus Medicine A/S	13/07/2009	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	23/07/2009
Enbrel	Dr Fisher Farma BV	10/02/2009	50 mg	Solution for injection in a 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, Sweden, United Kingdom	The Netherlands	02/07/2009
Enbrel	Emra-Med	01/04/2008	25mg	Solution for injection in a pre-filled syringe	8+16		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/01/2009
Enbrel	Emra-Med	01/04/2008	25mg	Solution for injection in a pre-filled syringe	24+48		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/01/2009
Enbrel	EuroPharma DK	01/06/2009	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/09/2009
Enbrel	Haemato Pharm AG	30/09/2008	50 mg	Solution for injection in a pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	Germany, Austria	16/02/2009

Enbrel	Haemato Pharm AG	07/11/2008	25 mg	Solution for injection in a pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	Germany, Austria	02/03/2009
Enbrel	Haemato Pharm AG	02/12/2009	25 mg	Solution for injection in pre-filled syringe	24 pre-filled syringes + 48 alcohol swabs	EU/1/99/126/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany, Austria	16/12/2009
Enbrel	Haemato Pharm AG	02/12/2009	25 mg	Solution for injection in pre-filled syringe	8 pre-filled syringes + 16 alcohol swabs	EU/1/99/126/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany, Austria	16/12/2009
Enbrel	Haemato Pharm AG	02/12/2009	50 mg	Solution for injection in pre-filled syringe	12 pre-filled syringes + 24 alcohol swabs	EU/1/99/126/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany, Austria	16/12/2009
Enbrel	Kohlpharma GmbH	11/11/2009	25 mg	Powder and solvent for solution for injection for paediatric use	4 vials + 4 pre-filled syringes + 8 syringes + 20 needles + 24 alcohol swabs	EU/1/99/126/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Enbrel	r.p. pharma GmbH	25/06/2008	50 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/017	Austria, Belgium, France, Greece, Italy, Portugal, Spain	Germany	12/01/2009
Enbrel	Veron Pharma	20/04/2009	25 mg	Solution for injection in pre-filled syringe	4 pre-filled syringes + 8 alcohol swabs	EU/1/99/126/013	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Germany	13/10/2009
Epivir	Haemato Pharm AG	09/10/2009	300 mg	Film-coated tablet	30	EU/1/96/015/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Latvia, Lithuania, Slovenia, Bulgaria, Czech Republic, Estonia, Hungary, Poland, Romania, Slovak Republic	Germany, Austria	20/10/2009

Epivir	Haemato Pharm AG	09/11/2009	150 mg	Film-coated tablet	60	EU/1/96/015/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovenia, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Poland, Romania, Slovak Republic	Germany, Austria	25/11/2009
Erbitux	Aaston Healthcare GmbH	09/04/2009	5 mg/ml	Solution for infusion	1 vial	EU/1/04/281/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	27/05/2009
Erbitux	Aaston Healthcare GmbH	08/06/2009	5 mg/ml	Solution for infusion	1 vial	EU/1/04/281/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	15/06/2009
Erbitux	Inopha GmbH	28/07/2009	5 mg/ml	Solution for infusion	1 vial	EU/1/04/281/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/08/2009
Erbitux	Paranova Danmark AS	02/10/2009	5 mg/ml	Solution for infusion	1 vial	EU/1/04/281/005	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	11/11/2009
Eucreas	Dr Fisher Farma BV	30/03/2009	50 mg / 1000 mg	Film-coated tablet	60	EU/1/07/425/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/05/2009
Eucreas	Dr Fisher Farma BV	30/03/2009	50 mg / 850 mg	Film-coated tablet	60	EU/1/07/425/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/04/2009
Eucreas	Dr Fisher Farma BV	22/06/2009	50 mg / 1000 mg	Film-coated tablet	30	EU/1/07/425/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/07/2009

Eucreas	Dr Fisher Farma BV	22/06/2009	50 mg / 850 mg	Film-coated tablet	30	EU/1/07/425/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/07/2009
Eucreas	Pharmachim AB	31/12/2008	50 mg / 1000 mg	Film-coated tablet	60	EU/1/07/425/009	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/02/2009
Eucreas	Pharmachim AB	31/12/2008	50 mg / 850 mg	Film-coated tablet	60	EU/1/07/425/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/02/2009
Evista	BR Pharma International Ltd	14/09/2009	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/09/2009
Evista	CC Pharma	24/07/2009	60 mg	Film-coated tablet	28	EU/1/98/073/002	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Romania	Germany	02/09/2009
Evista	CC Pharma	24/07/2009	60 mg	Film-coated tablet	84	EU/1/98/073/003	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Romania	Germany	02/09/2009
EVRA	Emra-Med	07/09/2009	- -	Transdermal patch	9 transdermal patches	EU/1/02/223/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/09/2009
Exelon	2 Care 4	06/11/2008	1.5 mg	Capsule, hard	56	EU/1/98/066/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom, Norway, Sweden, Finland, Iceland	Denmark	16/01/2009
Exelon	2 Care 4	13/11/2008	1.5 mg	Capsule, hard	112	EU/1/98/066/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom, Norway, Sweden, Finland, Iceland	Denmark	16/01/2009
Exelon	B&S Healthcare Ltd	15/09/2009	4.6 mg / 24 hours	Transdermal patch	20 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	05/10/2009

Exelon	B&S Healthcare Ltd	11/11/2009	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, United Kingdom	26/11/2009
Exelon	BR Lewis Ltd	27/02/2009	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	12/05/2009
Exelon	CC Pharma	19/01/2009	1.5mg	Hard capsules	56	EU/1/98/066/002	Spain, Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, United Kingdom	Germany	27/01/2009
Exelon	Delfarma SpZoo	16/11/2009	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, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	25/11/2009
Exelon	Dr Fisher Farma BV	01/09/2009	4.5 mg	Capsule, hard	28	EU/1/98/066/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	The Netherlands	09/09/2009
Exelon	Euro Registratie Collectief BV	17/12/2008	9.5 mg/24 h	Transdermal patches	30		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	12/01/2009
Exelon	EuroPharma DK	04/06/2009	1.5 mg	Capsule, hard	28	EU/1/98/066/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	01/07/2009
Exelon	EuroPharma DK	04/06/2009	1.5 mg	Capsule, hard	56	EU/1/98/066/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	01/07/2009
Exelon	EuroPharma DK	04/06/2009	1.5 mg	Capsule, hard	112	EU/1/98/066/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	01/07/2009

Exelon	Kohlpharma GmbH	04/02/2009	6 mg	Capsule, hard	112	EU/1/98/066/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Iceland	Germany	07/04/2009
Exelon	Kohlpharma GmbH	04/02/2009	9.5 mg / 24 hours	Transdermal patch	60 sachets (30 x 2 sachets)	EU/1/98/066/025	Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Cyprus, Denmark, Finland, Iceland, Liechtenstein, Luxembourg, Malta	Germany	15/04/2009
Exelon	Kohlpharma GmbH	04/02/2009	9.5 mg / 24 hours	Transdermal patch	90 sachets (30 x 3 sachets)	EU/1/98/066/026	Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Cyprus, Denmark, Finland, Iceland, Liechtenstein, Luxembourg, Malta	Germany	15/04/2009
Exelon	Kohlpharma GmbH	06/05/2009	4.6 mg / 24 hours	Transdermal patch	90 (30 x 3) sachets	EU/1/98/066/022	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/06/2009
Exelon	Lexon (UK) Ltd	31/03/2009	4.6 mg / 24 hours	Transdermal patch	20 sachets	EU/1/98/066/020	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	21/05/2009
Exelon	Lexon (UK) Ltd	31/03/2009	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland	21/05/2009
Exelon	Paranova Danmark AS	17/08/2009	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/11/2009
Exelon	Paranova Danmark AS	28/08/2009	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/11/2009
Exelon	Paranova Danmark AS	02/09/2009	6 mg	Capsule, hard	112	EU/1/98/066/012	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/10/2009
Exelon	Swingward Ltd TA Medihealth	13/02/2009	4.6 mg/24 h	Transdermal patches	30	EU/1/98/066/020	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	10/03/2009

Exelon	Swingward Ltd TA Medihealth	13/02/2009	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Bulgaria, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	10/03/2009
Exelon	Swingward Ltd TA Medihealth	28/04/2009	6 mg	Capsule, hard	56	EU/1/98/066/011	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain	United Kingdom	06/07/2009
Exelon	Waymade PLC	01/12/2009	4.6 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/020	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Malta, Ireland, United Kingdom	16/12/2009
Exelon	Waymade PLC	01/12/2009	9.5 mg / 24 hours	Transdermal patch	30 sachets	EU/1/98/066/024	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Malta, Ireland, United Kingdom	16/12/2009
Exforge	B&S Healthcare Ltd	04/03/2009	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Cyprus, Bulgaria, Czech 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	24/03/2009
Exforge	B&S Healthcare Ltd	07/09/2009	10 mg/160 mg	Film-coated tablet	30	EU/1/06/370/020	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	United Kingdom, Ireland	17/09/2009
Exforge	Dr Fisher Farma BV	16/03/2009	5 mg/80 mg	Film-coated tablet	14	EU/1/06/370/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/04/2009
Exforge	Eureco-Pharma B.V.	20/04/2009	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	03/06/2009
Exforge	Eureco-Pharma B.V.	20/04/2009	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	03/06/2009

Exforge	Eureco-Pharma B.V.	20/04/2009	5 mg/80 mg	Film-coated tablet	28	EU/1/06/370/003	Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	03/06/2009
Exforge	EuroPharma DK	01/06/2009	10 mg/160 mg	Film-coated tablet	30	EU/1/06/370/020	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/07/2009
Exforge	EuroPharma DK	01/06/2009	5 mg/160 mg	Film-coated tablet	30	EU/1/06/370/012	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/07/2009
Exforge	EuroPharma DK	01/06/2009	5 mg/160 mg	Film-coated tablet	90	EU/1/06/370/014	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/07/2009
Exforge	EuroPharma DK	02/06/2009	10 mg/160 mg	Film-coated tablet	90	EU/1/06/370/022	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	21/07/2009
Exforge	Kohlpharma GmbH	18/05/2009	10 mg/160 mg	Film-coated tablet	28	EU/1/06/370/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	10 mg/160 mg	Film-coated tablet	56	EU/1/06/370/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	10 mg/160 mg	Film-coated tablet	98	EU/1/06/370/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	5 mg/160 mg	Film-coated tablet	28	EU/1/06/370/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009

Exforge	Kohlpharma GmbH	18/05/2009	5 mg/160 mg	Film-coated tablet	56	EU/1/06/370/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	5 mg/160 mg	Film-coated tablet	98	EU/1/06/370/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	5 mg/80 mg	Film-coated tablet	56	EU/1/06/370/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Kohlpharma GmbH	18/05/2009	5 mg/80 mg	Film-coated tablet	98	EU/1/06/370/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/06/2009
Exforge	Orifarm AS	18/04/2008	10/160mg	Film-coated tablets	28	EU/1/06/370/019	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/03/2009
Exforge	Orifarm AS	18/04/2008	10/160mg	Film-coated tablets	98	EU/1/06/370/023	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/03/2009
Exforge	Orifarm AS	18/04/2008	5/160mg	Film-coated tablets	28	EU/1/06/370/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/03/2009
Exforge	Orifarm AS	18/04/2008	5/160mg	Film-coated tablets	98	EU/1/06/370/015	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/03/2009
Exforge	Pharma Westen GmbH	25/06/2009	10 mg/160 mg	Film-coated tablet	98	EU/1/06/370/023	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	23/07/2009

Exforge	Pharma Westen GmbH	25/06/2009	5 mg/160 mg	Film-coated tablet	98	EU/1/06/370/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	23/07/2009
Exjade	CC Pharma	16/01/2009	125 mg	Dispersible tablet	84	EU/1/06/356/002	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Denmark	Germany	15/05/2009
Exjade	CC Pharma	16/01/2009	250 mg	Dispersible tablet	84	EU/1/06/356/004	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Denmark, Slovak Republic, Sweden, Finland	Germany	15/05/2009
Exjade	CC Pharma	16/01/2009	500 mg	Dispersible tablet	84	EU/1/06/356/006	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, United Kingdom, Romania, Denmark, Finland, Spain, Sweden, Slovak Republic	Germany	15/05/2009
Exjade	Combiphar Europe BV	20/07/2009	500 mg	Dispersible tablet	28	EU/1/06/356/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/11/2009
Exjade	Emra-Med	24/10/2008	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/04/2009
Exjade	Haemato Pharm AG	01/06/2009	250 mg	Dispersible tablet	84	EU/1/06/356/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Romania, Slovak Republic, Slovenia, Poland	Germany	15/06/2009
Exjade	Haemato Pharm AG	01/06/2009	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Germany, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany, Austria	15/06/2009
Exjade	Kohlpharma GmbH	20/07/2009	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/07/2009

Exjade	Pharma Westen GmbH	29/04/2009	500 mg	Dispersible tablet	84	EU/1/06/356/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Germany	13/05/2009
Extavia	CC Pharma	22/05/2009	250 µg / ml	Powder and solvent for solution for injection	15 vials + 15 pre-filled syringes	EU/1/08/454/002	Norway, Poland, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Latvia, Czech Republic, Bulgaria, Slovak Republic, Denmark, Sweden, Finland	Germany	23/06/2009
Extavia	Emra-Med	27/04/2009	250 µg / ml	Powder and solvent for solution for injection	15 vials + 15 pre-filled syringes	EU/1/08/454/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany	16/07/2009
Extavia	Haemato Pharm AG	16/10/2009	250 µg / ml	Powder and solvent for solution for injection	15 vials + 15 pre-filled syringes	EU/1/08/454/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Germany, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany, Austria	10/11/2009
Extavia	Haemato Pharm AG	26/11/2009	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, 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, Austria	17/12/2009
Extavia	Kohlpharma GmbH	06/07/2009	250 µg / ml	Powder and solvent for solution for injection	15 vials + 15 pre-filled syringes	EU/1/08/454/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic	Germany	22/07/2009
Extavia	Pharma Gerke GmbH	18/05/2009	250 µg / ml	Powder and solvent for solution for injection	15 vials + 15 pre-filled syringes	EU/1/08/454/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland, Lithuania, Slovak Republic, Romania, Czech Republic	Germany	01/10/2009
Extavia	Pharma Gerke GmbH	02/09/2009	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, Finland, France, Greece, Italy, The Netherlands, Norway, Poland, Portugal, Spain, United Kingdom, Slovak Republic, Romania, Czech Republic	Germany	01/10/2009

Fabrazyme	Abacus Medicine A/S	05/05/2009	35 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/188/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/05/2009
Fabrazyme	Abacus Medicine A/S	05/05/2009	5 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/01/188/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	13/05/2009
Fabrazyme	CC Pharma	05/02/2009	35mg	Powder for concentrate for solution for infusion	5	EU/1/01/188/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/06/2009
Faslodex	Haemato Pharm AG	01/06/2009	250 mg/5 ml	Solution for injection	1 pre-filled syringe + 1 safety needle	EU/1/03/269/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	09/06/2009
Faslodex	Kohlpharma GmbH	16/02/2009	250 mg/5 ml	Solution for injection	1 pre-filled syringe + 1 safety needle	EU/1/03/269/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Hungary	Germany	06/05/2009
Faslodex	Pharma Westen GmbH	04/02/2009	250 mg/5 ml	Solution for injection	1 pre-filled syringe + 1 safety needle	EU/1/03/269/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Hungary, Lithuania, Bulgaria	Germany	06/04/2009
Firazyr	CC Pharma	29/06/2009	30 mg	Solution for injection	1 pre-filled syringe + 1 needle	EU/1/08/461/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	06/08/2009
Fosavance	CC Pharma	29/06/2009	70 mg / 5600 IU	Tablet	12	EU/1/05/310/008	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Denmark, Hungary, Finland, Sweden	Germany	28/08/2009
Fosavance	CC Pharma	29/06/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Denmark, Finland, Hungary, Sweden	Germany	28/08/2009

FOSAVANCE	Emra-Med	09/06/2008	70 mg/2800 IU	Tablet	1 x 4	EU/1/05/310/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	14/01/2009
FOSAVANCE	Emra-Med	09/06/2008	70 mg/2800 IU	Tablet	3 x 4	EU/1/05/310/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	14/01/2009
Fosavance	Emra-Med	24/06/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	09/07/2009
Fosavance	Emra-Med	24/06/2009	70 mg / 5600 IU	Tablet	12	EU/1/05/310/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	09/07/2009
Fosavance	Eureco-Pharma B.V.	15/06/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	18/06/2009
Fosavance	Eureco-Pharma B.V.	15/06/2009	70 mg / 5600 IU	Tablet	12	EU/1/05/310/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	18/06/2009
Fosavance	Euro Registratie Collectief BV	14/08/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/09/2009
Fosavance	Europharma DK	22/09/2008	70 mg/2800 IU	Tablets	12		Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	12/01/2009

Fosavance	Europharma DK	22/09/2008	70 mg/2800 IU	Tablets	4		Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	12/01/2009
Fosavance	Europharma Sverige AB	26/01/2009	70 mg/2800 IU	Tablet	3 x 4	EU/1/05/310/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	24/03/2009
Fosavance	Kohlpharma GmbH	02/07/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/07/2009
Fosavance	Kohlpharma GmbH	02/07/2009	70 mg / 5600 IU	Tablet	12	EU/1/05/310/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/07/2009
Fosavance	Lex ano UAB	23/01/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Italy, Latvia, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Lithuania	25/03/2009
Fosavance	Mediwin Ltd	11/02/2009	70 mg/2800 IU	Tablet	3 x 4	EU/1/05/310/004	Austria, Belgium, Denmark, Finland, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	France	06/05/2009
Fosavance	Mediwin Ltd	11/02/2009	70 mg/5600 IU	Tablets	12	EU/1/05/310/008	Austria, Belgium, Denmark, Finland, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	France	06/05/2009
Fosavance	MPT Pharma	30/03/2009	70 mg/2800 IU	Tablet	1 x 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, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic, Slovenia	United Kingdom, Ireland	11/05/2009
FOSAVANCE	Paranova Läkemedel AB	16/12/2008	70 mg/2800 IU	Tablet	3 x 4	EU/1/05/310/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Iceland, Liechtenstein	Sweden	06/01/2009
Fosavance	Pharma Gerke GmbH	05/02/2009	70 mg/2800 IU	Tablet	3 x 4	EU/1/05/310/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria	Germany	14/04/2009

Fosavance	Pharma Lab	09/02/2009	70 mg/2800 IU	Tablets	12	EU/1/05/310/004	Austria, Belgium, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	France	13/08/2009
Fosavance	Pharma Lab	30/06/2009	70 mg / 5600 IU	Tablet	4	EU/1/05/310/007	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	France	05/08/2009
FOSAVANCE	Pharma Westen GmbH	12/12/2008	70 mg / 5600 IU	Tablet	12	EU/1/05/310/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/01/2009
Fosavance	SIA 'Elpis'	27/10/2008	70 mg/2800 IU	Tablet	1 x 4	EU/1/05/310/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	06/04/2009
Fuzeon	CC Pharma	19/01/2009	90 mg/ml	Powder and solvent for solution for injection	60 vials + 60 vials + 60 3 ml syringes + 60 1 ml syringes	EU/1/03/252/001	Norway, Austria, Belgium, France, Greece, Italy, Romania, The Netherlands, Portugal, Spain, United Kingdom	Germany	27/01/2009
Fuzeon	Haemato Pharm AG	01/10/2009	90 mg/ml	Powder and solvent for solution for injection	60 vials + 60 vials + 60 3 ml syringes + 60 1 ml syringes	EU/1/03/252/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Germany	Germany, Austria	25/11/2009
Galvus	CC Pharma	01/06/2009	50 mg	Tablet	90	EU/1/07/414/007	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	01/07/2009
Galvus	Extrapharm BV	18/03/2009	50 mg	Tablet	56	EU/1/07/414/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009

Ganfort	2 Care 4	26/10/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden, United Kingdom, Poland	Denmark	10/12/2009
Ganfort	2 Care 4	26/10/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden, United Kingdom, Poland	Denmark	10/12/2009
Ganfort	Amimed Direct Ltd	09/07/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/09/2009
Ganfort	B&S Healthcare Ltd	23/01/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Bulgaria, Cyprus, Denmark, 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	United Kingdom, Ireland	04/06/2009
Ganfort	Cross Pharma AB	29/10/2008	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	12/02/2009
Ganfort	Parallell Pharma AB	01/04/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Ireland, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	29/07/2009
Ganfort	Paranova Danmark AS	07/10/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/11/2009
Ganfort	Pharmachim AB	06/05/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	1 bottle	EU/1/06/340/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom, Poland	Sweden	30/06/2009
Ganfort	Pharmachim AB	06/05/2009	300 µg/ml + 5 mg/ml	Eye drops, solution	3 bottles	EU/1/06/340/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom, Poland	Sweden	26/06/2009

Gardasil	Aaston Healthcare GmbH	23/04/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	29/06/2009
Gardasil	CC Pharma	22/12/2008	- -	Suspension for injection	10 pre-filled syringes with needle guard + 20 needles	EU/1/06/357/016	Norway	Germany	17/02/2009
Gardasil	CC Pharma	22/12/2008	- -	Suspension for injection	1 pre-filled syringe with needle guard + 2 needles	EU/1/06/357/015	Norway	Germany	17/02/2009
Gardasil	CC Pharma	08/06/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	11/06/2009
Gardasil	Emra-Med	01/12/2008	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/01/2009
Gardasil	Haemato Pharm AG	24/10/2008	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Norway, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	08/01/2009
Gardasil	Haemato Pharm AG	11/05/2009	- -	Suspension for injection	10 pre-filled syringes + 20 needles	EU/1/06/357/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Norway	Germany	03/06/2009

Gardasil	Medartuum AB	30/01/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	26/05/2009
Gardasil	MedicoPharm AG	12/10/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/10/2009
Gardasil	Orifarm AS	22/12/2008	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/01/2009
Gardasil	Paranova Danmark AS	22/05/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	02/07/2009
Gardasil	Paranova Läkemedel AB	09/03/2009	- -	Suspension for injection	1 pre-filled syringe + 2 needles	EU/1/06/357/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	10/06/2009
Gardasil	Pharma Westen GmbH	21/01/2009	- -	Suspension for injection	10 pre-filled syringes + 20 needles	EU/1/06/357/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/02/2009
Glivec	Abacus Medicine A/S	05/05/2009	100 mg	Film-coated tablet	60 film-coated	EU/1/01/198/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Romania, Slovak Republic	Denmark	13/05/2009
Glivec	Abacus Medicine A/S	05/05/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Romania, Slovak Republic	Denmark	13/05/2009
Glivec	Beragena Arzneimittel GmbH	06/04/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	15/04/2009

Glivec	Beragena Arzneimittel GmbH	06/04/2009	400 mg	Film-coated tablet	90 film-coated	EU/1/01/198/013	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	15/04/2009
Glivec	Combiphar Europe BV	27/07/2009	100 mg	Film-coated tablet	60 film-coated	EU/1/01/198/008	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/09/2009
Glivec	Delphi Pharmaceuticals BV	22/07/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	11/09/2009
Glivec	Docpharm GmbH&CoKGaA	10/08/2009	100 mg	Film-coated tablet	20 film-coated	EU/1/01/198/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/09/2009
Glivec	Docpharm GmbH&CoKGaA	10/08/2009	100 mg	Film-coated tablet	60 film-coated	EU/1/01/198/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/09/2009
Glivec	Dr Fisher Farma BV	12/01/2009	100 mg	Film-coated tablet	60 film-coated	EU/1/01/198/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/01/2009
Glivec	Dr Fisher Farma BV	12/01/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/01/2009
Glivec	Emra-Med	24/10/2008	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Glivec	Emra-Med	24/10/2008	400 mg	Film-coated tablet	90 film-coated	EU/1/01/198/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Glivec	Haemato Pharm AG	25/08/2009	400 mg	Film-coated tablet	90 film-coated	EU/1/01/198/013	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, United Kingdom	Germany	26/11/2009
Glivec	Paranova Danmark AS	06/05/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/06/2009

Glivec	Pharma Gerke GmbH	12/12/2008	400 mg	Film-coated tablet	90 film-coated	EU/1/01/198/013	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland	Germany	13/01/2009
Glivec	Pharma Gerke GmbH	12/12/2008	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland	Germany	13/01/2009
Glivec	Pharma Westen GmbH	19/01/2009	400 mg	Film-coated tablet	30 film-coated	EU/1/01/198/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic	Germany	03/02/2009
Glivec	Pharmachim AB	09/12/2008	400mg	Film-coated tablets	30		Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	05/01/2009
Glivec	Pharmachim AB	22/12/2008	100mg	Film-coated tablets	60		Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	19/01/2009
Glivec	Pharmachim AB	22/01/2009	100 mg	Film-coated tablet	120 film-coated	EU/1/01/198/011	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The NetherlandsPortugal, The Netherlands, Portugal, Spain, United Kingdom	Sweden	03/02/2009
Gonal-f	CC Pharma	06/11/2008	300 IU/0.5 ml	Solution for injection in a pre-filled pen	1+5	EU/1/95/001/033	The Netherlands, Italy, Norway, Austria, Belgium, France, Greece, Portugal, Spain, United Kingdom, Denmark, Finland, Poland, Sweden	Germany	04/09/2009
Gonal-f	CC Pharma	12/01/2009	450 IU/0.5 ml	powder and solvent for solution for injection	1+1+6	EU/1/95/001/031	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	29/01/2009
GONAL-f	CC Pharma	25/08/2009	75 IU (5.5 micrograms)	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe	EU/1/95/001/025	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	12/10/2009
Gonal-f	Emra-Med	28/01/2008	900 IU/1.5 ml	Solution for injection in a pre-filled pen	1+14		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/01/2009
GONAL-f	Emra-Med	16/10/2008	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, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/01/2009
GONAL-f	Emra-Med	01/05/2009	300 IU/0.5 ml (22 micrograms/0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 5 needles	EU/1/95/001/033	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/05/2009

GONAL-f	Emra-Med	28/05/2009	1050 IU/1.75 ml (77 micrograms/1.75 ml)	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 15 disposable syringes	EU/1/95/001/021	Austria, Belgium, Cyprus, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/06/2009
GONAL-f	Kohlpharma GmbH	19/12/2008	1050 IU/1.75 ml (77 micrograms/1.75 ml)	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 15 disposable syringes	EU/1/95/001/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/02/2009
GONAL-f	Kohlpharma GmbH	04/02/2009	450 IU/0.75 ml (33 micrograms/0.75 ml)	Powder and solvent for solution for injection	1 vial + 1 pre-filled syringe + 6 disposable syringes	EU/1/95/001/031	Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Cyprus, Denmark, Finland, Iceland, Liechtenstein, Luxembourg, Malta	Germany	07/04/2009
GONAL-f	Orifarm AS	26/01/2009	900 IU/1.5 ml (66 micrograms/1.5 ml)	Solution for injection in a pre-filled pen	1 pre-filled pen + 14 needles	EU/1/95/001/035	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, Cyprus, Iceland, Liechtenstein, Luxembourg, Malta, United Kingdom, Czech Republic, Hungary, Lithuania, Poland, Romania	Denmark	13/03/2009
GONAL-f	Orifarm AS	17/06/2009	300 IU/0.5 ml (22 micrograms/0.5 ml)	Solution for injection in pre-filled pen	1 pre-filled pen + 5 needles	EU/1/95/001/033	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	23/09/2009
GONAL-f	Orifarm AS	17/06/2009	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, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	23/09/2009
HBVAXPRO	Dr Fisher Farma BV	03/03/2009	40 µg/ml	Suspension for injection	1 vial	EU/1/01/183/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	31/03/2009
HBVAXPRO	Emra-Med	21/08/2008	5 µg/0.5 ml	Suspension for injection	1 vial	EU/1/01/183/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/01/2009

HBVAXPRO	Emra-Med	21/08/2008	5µg/0.5ml	Suspension for injection in a vial	10		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/01/2009
Hepsera	Delphi Pharmaceuticals BV	22/07/2009	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	30/07/2009
Hepsera	Dr Fisher Farma BV	04/12/2009	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	15/12/2009
Hepsera	Haemato Pharm AG	29/06/2009	10 mg	Tablet	3 x 30	EU/1/03/251/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/09/2009
Hepsera	Inopha GmbH	24/03/2009	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	30/04/2009
Hepsera	Kohlpharma GmbH	15/06/2009	10 mg	Tablet	3 x 30	EU/1/03/251/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/06/2009
Hepsera	Orifarm AS	10/08/2009	10 mg	Tablet	30	EU/1/03/251/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	14/08/2009
Hepsera	Pharma Westen GmbH	04/02/2009	10 mg	Tablet	3 x 30	EU/1/03/251/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/03/2009
Herceptin	Aaston Healthcare GmbH	17/03/2009	150 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/00/145/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Romania, Slovak Republic, Slovenia	Germany	06/04/2009

Humalog	CC Pharma	29/09/2008	100 U/ml	Solution for injection	1 vial	EU/1/96/007/002	Spain, Italy, Austria, Belgium, France, Greece, The Netherlands, Norway, Portugal, United Kingdom	Germany	05/02/2009
Humalog	CC Pharma	04/02/2009	100 U/ml	Solution for injection	5 x 1 vial	EU/1/96/007/021	Italy, Spain, Austria, Belgium, France, Greece, The Netherlands, Norway, Portugal, United Kingdom	Germany	16/02/2009
Humalog	Pharma Westen GmbH	03/03/2009	100 U/ml	Solution for injection	5 cartridges	EU/1/96/007/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria	Germany	18/05/2009
Humalog	Pharma Westen GmbH	03/03/2009	100 U/ml	Solution for injection	2 x 5 cartridges	EU/1/96/007/023	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria	Germany	18/05/2009
Humalog	Swingward Ltd TA Medihealth	26/03/2009	100 U/ml	Solution for injection	1 vial	EU/1/96/007/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	17/04/2009
Humalog KwikPen	CC Pharma	04/11/2009	100 U/ml	Solution for injection	5 pens	EU/1/96/007/031	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	26/11/2009
Humalog KwikPen	CC Pharma	04/11/2009	100 U/ml	Solution for injection	2 x 5 pens	EU/1/96/007/032	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009
Humalog KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Solution for injection	5 pens	EU/1/96/007/031	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humalog KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Solution for injection	2 x 5 pens	EU/1/96/007/032	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humalog Mix 25 KwikPen	CC Pharma	04/11/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/033	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009
Humalog Mix 25 KwikPen	CC Pharma	04/11/2009	100 U/ml	Suspension for injection	2 x 5 pens	EU/1/96/007/034	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009

Humalog Mix 25 Pen	Emra-Med	21/09/2009	100 U/ml	Suspension for injection	2 x 5 pens	EU/1/96/007/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	13/11/2009
Humalog Mix 25 Pen	Pharma Westen GmbH	02/04/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/016	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/05/2009
Humalog Mix 50 KwikPen	CC Pharma	04/11/2009	100 U/ml	Suspension for injection	2 x 5 pens	EU/1/96/007/036	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009
Humalog Mix 50 KwikPen	CC Pharma	04/11/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/035	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/11/2009
Humalog Mix 50 Pen	Cross Healthcare Ltd	26/08/2008	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/017	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/02/2009
Humalog Mix25	CC Pharma	03/08/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/016	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	21/09/2009
Humalog Mix25	CC Pharma	03/08/2009	100 U/ml	Suspension for injection	5 cartridges	EU/1/96/007/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	21/09/2009
Humalog Mix25 KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/033	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humalog Mix25 KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Suspension for injection	2 x 5 pens	EU/1/96/007/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humalog Mix50	CC Pharma	03/08/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/017	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	21/09/2009
Humalog Mix50	CC Pharma	03/08/2009	100 U/ml	Suspension for injection	5 cartridges	EU/1/96/007/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	21/09/2009

Humalog Mix50 KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Suspension for injection	5 pens	EU/1/96/007/035	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humalog Mix50 KwikPen	Kohlpharma GmbH	21/09/2009	100 U/ml	Suspension for injection	2 x 5 pens	EU/1/96/007/036	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	27/11/2009
Humira	Aaston Healthcare GmbH	26/03/2009	40 mg	Solution for injection	2 pre-filled syringes + 2 alcohol pads	EU/1/03/256/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/05/2009
Humira	Haemato Pharm AG	15/01/2009	40 mg	Solution for injection	2 pre-filled syringes + 2 alcohol pads	EU/1/03/256/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany, Austria	27/04/2009
Humira	Haemato Pharm AG	14/08/2009	40 mg	Solution for injection	6 pre-filled syringes + 6 alcohol pads	EU/1/03/256/005	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	01/10/2009
Humira	Haemato Pharm AG	04/09/2009	40 mg	Solution for injection	6 pre-filled pens + 6 alcohol pads in a blister	EU/1/03/256/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	01/10/2009
Hycamtin	Aaston Healthcare GmbH	11/06/2009	4 mg	Powder for concentrate for solution for infusion	5 vials (17 ml)	EU/1/96/027/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	20/07/2009
Hycamtin	Pharma Gerke GmbH	03/11/2009	4 mg	Powder for concentrate for solution for infusion	5 vials (17 ml)	EU/1/96/027/001	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	10/12/2009

Infanrix hexa	Kohlpharma GmbH	19/01/2009	- -	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/02/2009
Inovelon	CC Pharma	14/07/2009	400 mg	Film-coated tablet	100	EU/1/06/378/015	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	10/09/2009
Inovelon	CC Pharma	15/07/2009	200 mg	Film-coated tablet	50	EU/1/06/378/008	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	10/09/2009
Insulatard	Pharma Westen GmbH	23/03/2009	100 IU/ml	Suspension for injection	10 cartridges	EU/1/02/233/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Hungary	Germany	12/05/2009
Insulatard	Pharmachim AB	31/12/2008	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/02/233/011	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	28/01/2009
Insulatard FlexPen	Kohlpharma GmbH	02/03/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/03/2009
Insulatard FlexPen	Kohlpharma GmbH	02/06/2009	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	09/07/2009
Insulatard FlexPen	Pharma Westen GmbH	10/06/2009	100 IU/ml	Suspension for injection	5 pre-filled pens	EU/1/02/233/014	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Sweden, Spain, United Kingdom	Germany	13/08/2009
Insulatard FlexPen	Pharma Westen GmbH	10/06/2009	100 IU/ml	Suspension for injection	10 pre-filled pens	EU/1/02/233/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	13/08/2009
Insulatard Penfill	Kohlpharma GmbH	31/03/2009	100 IU/ml	Suspension for injection	5 cartridges	EU/1/02/233/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary, Poland	Germany	17/04/2009

Insulatard Penfill	Kohlpharma GmbH	31/03/2009	100 IU/ml	Suspension for injection	10 cartridges	EU/1/02/233/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary, Poland	Germany	17/04/2009
Insulatard Penfill	Medcor Pharmaceuticals BV	09/03/2009	100 IU/ml	Suspension for injection	5 cartridges	EU/1/02/233/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/05/2009
Insulatard Penfill	Pharma Westen GmbH	23/03/2009	100 IU/ml	Suspension for injection	5 cartridges	EU/1/02/233/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Hungary	Germany	12/05/2009
Insuman Basal	Kohlpharma GmbH	31/03/2009	100 IU/ml	Suspension for injection	5 cartridges	EU/1/97/030/035	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Germany	17/04/2009
Insuman Basal	Kohlpharma GmbH	31/03/2009	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/058	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom, Czech Republic	Germany	17/04/2009
Insuman Basal	Kohlpharma GmbH	15/06/2009	100 IU/ml	OptiSet, suspension for injection	5 pens	EU/1/97/030/071	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/07/2009
Insuman Basal	Kohlpharma GmbH	15/06/2009	100 IU/ml	OptiSet, suspension for injection	10 pens	EU/1/97/030/072	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/07/2009
Insuman Basal	Pharma Westen GmbH	08/12/2008	100 IU/ml	Suspension for injection	5 cartridges	EU/1/97/030/035	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary, Romania, Poland, Czech Republic, Slovak Republic	Germany	04/03/2009
Insuman Basal	Pharma Westen GmbH	12/11/2009	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/058	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania, Poland, Czech Republic, Slovak Republic	Germany	27/11/2009

Insuman Comb 25	Axicorp Pharma GmbH	26/01/2009	100IU/ml	Suspension for injection in a cartridge	10x3ml	EU/1/97/030/062	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary	Germany	25/02/2009
Insuman Comb 25	Axicorp Pharma GmbH	26/01/2009	100IU/ml	Suspension for injection in a cartridge	5x3ml	EU/1/97/030/045	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary	Germany	25/02/2009
Insuman Comb 25	Kohlpharma GmbH	18/05/2009	100 IU/ml	OptiSet, suspension for injection	5 pens	EU/1/97/030/079	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/06/2009
Insuman Comb 25	Pharma Westen GmbH	12/11/2009	100 IU/ml	Suspension for injection	10 cartridges	EU/1/97/030/062	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania, Czech Republic, Slovak Republic, Poland, Bulgaria	Germany	27/11/2009
Insuman Combi 25	Kohlpharma GmbH	02/07/2009	100 IU/ml	OptiSet, suspension for injection	10 pens	EU/1/97/030/080	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/07/2009
Insuman Rapid	Axicorp Pharma GmbH	26/01/2009	100 IU/ml	Solution for injection	5 cartridges	EU/1/97/030/030	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Hungary	Germany	23/02/2009
Insuman Rapid	Axicorp Pharma GmbH	26/01/2009	100 IU/ml	Solution for injection	10 cartridges	EU/1/97/030/056	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Hungary	Germany	23/02/2009
Insuman Rapid	Kohlpharma GmbH	06/07/2009	100 IU/ml	OptiSet, solution for injection	5 pens	EU/1/97/030/067	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania	Germany	22/10/2009
Insuman Rapid	Kohlpharma GmbH	06/07/2009	100 IU/ml	OptiSet, solution for injection	10 pens	EU/1/97/030/068	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania	Germany	22/10/2009

Insuman Rapid	Pharma Westen GmbH	08/12/2008	100 IU/ml	Solution for injection	5 cartridges	EU/1/97/030/030	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary, Romania, Czech Republic, Slovak Republic, Poland	Germany	04/03/2009
Insuman Rapid	Pharma Westen GmbH	20/01/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	03/02/2009
Insuman Rapid	Pharma Westen GmbH	12/11/2009	100 IU/ml	Solution for injection	10 cartridges	EU/1/97/030/056	Austria, Belgium, Denmark, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	27/11/2009
Integrilin	Lex ano UAB	03/03/2009	0.75 mg/ml	Solution for infusion	1 vial	EU/1/99/109/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, 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	Lithuania	30/04/2009
INTELENCE	CC Pharma	06/03/2009	100 mg	Tablet	120	EU/1/08/468/001	Norway	Germany	04/05/2009
Intelence	Haemato Pharm AG	13/02/2009	100mg	Tablets	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden, United Kingdom, Bulgaria, Czech Republic, Denmark, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Spain, Germany	Germany, Austria	24/02/2009
INTELENCE	Kohlpharma GmbH	04/12/2009	100 mg	Tablet	120	EU/1/08/468/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/12/2009
Intrinsa	Cross Healthcare Ltd	19/01/2009	300 µ/24 hours	Transdermal patch	8 patches	EU/1/06/352/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	16/02/2009

Invanz	Delfarma SpZoo	08/06/2009	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, Malta, Luxembourg, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, The Netherlands, Sweden, United Kingdom	Poland	16/07/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	3 mg	Prolonged release tablet	28	EU/1/07/395/021	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, United Kingdom, Spain	Germany	04/06/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	3 mg	Prolonged release tablet	49	EU/1/07/395/023	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	3 mg	Prolonged release tablet	98	EU/1/07/395/025	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	6 mg	Prolonged release tablet	28	EU/1/07/395/026	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	6 mg	Prolonged release tablet	49	EU/1/07/395/028	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009
Invega	Beragena Arzneimittel GmbH	19/05/2009	6 mg	Prolonged release tablet	98	EU/1/07/395/030	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	04/06/2009
Invega	BR Pharma International Ltd	11/12/2008	3 mg	Prolonged release tablet	28	EU/1/07/395/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Invega	BR Pharma International Ltd	11/12/2008	6 mg	Prolonged release tablet	28	EU/1/07/395/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Invega	BR Pharma International Ltd	04/03/2009	3 mg	Prolonged release tablet	49	EU/1/07/395/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Invega	BR Pharma International Ltd	04/03/2009	3 mg	Prolonged release tablet	98	EU/1/07/395/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009

Invega	BR Pharma International Ltd	04/03/2009	6 mg	Prolonged release tablet	49	EU/1/07/395/028	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Invega	BR Pharma International Ltd	04/03/2009	6 mg	Prolonged release tablet	98	EU/1/07/395/030	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Invega	CC Pharma	22/06/2009	3 mg	Prolonged release tablet	98	EU/1/07/395/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	3 mg	Prolonged release tablet	49	EU/1/07/395/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	3 mg	Prolonged release tablet	28	EU/1/07/395/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	6 mg	Prolonged release tablet	98	EU/1/07/395/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	6 mg	Prolonged release tablet	49	EU/1/07/395/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	6 mg	Prolonged release tablet	28	EU/1/07/395/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	9 mg	Prolonged release tablet	98	EU/1/07/395/015	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	9 mg	Prolonged release tablet	49	EU/1/07/395/013	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	CC Pharma	22/06/2009	9 mg	Prolonged release tablet	28	EU/1/07/395/011	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/07/2009
Invega	Emra-Med	29/06/2009	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009

Invega	Emra-Med	29/06/2009	9 mg	Prolonged-release tablet	49	EU/1/07/395/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	29/06/2009	9 mg	Prolonged-release tablet	98	EU/1/07/395/015	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	01/07/2009	3 mg	Prolonged-release tablet	28	EU/1/07/395/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	01/07/2009	3 mg	Prolonged-release tablet	49	EU/1/07/395/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	01/07/2009	3 mg	Prolonged-release tablet	98	EU/1/07/395/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	02/07/2009	6 mg	Prolonged-release tablet	98	EU/1/07/395/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	02/07/2009	6 mg	Prolonged-release tablet	49	EU/1/07/395/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Emra-Med	02/07/2009	6 mg	Prolonged-release tablet	28	EU/1/07/395/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/07/2009
Invega	Euro Registratie Collectief BV	14/08/2009	9 mg	Prolonged-release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Denmark	The Netherlands	15/09/2009

Invega	Ingala Healthcare Ltd	19/01/2009	9 mg	Prolonged release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	06/02/2009
Invega	Kohlpharma GmbH	19/12/2008	6 mg	Prolonged-release tablet	28	EU/1/07/395/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/01/2009
Invega	Kohlpharma GmbH	19/12/2008	6 mg	Prolonged-release tablet	49	EU/1/07/395/028	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/01/2009
Invega	Kohlpharma GmbH	19/12/2008	6 mg	Prolonged-release tablet	98	EU/1/07/395/030	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/01/2009
Invega	Kohlpharma GmbH	04/02/2009	3 mg	Prolonged release tablet	28 (white blister)	EU/1/07/395/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/03/2009
Invega	Kohlpharma GmbH	04/02/2009	3 mg	Prolonged release tablet	49 (white blister)	EU/1/07/395/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/03/2009
Invega	Kohlpharma GmbH	10/02/2009	6mg	Prolonged-release tablets	28 (transparent blister)	EU/1/07/395/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	31/03/2009
Invega	Kohlpharma GmbH	10/02/2009	6mg	Prolonged-release tablets	49 (transparent blister)	EU/1/07/395/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	31/03/2009
Invega	Kohlpharma GmbH	10/02/2009	6mg	Prolonged-release tablets	98 (transparent blister)	EU/1/07/395/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	31/03/2009

Invega	Kohlpharma GmbH	15/06/2009	9 mg	Prolonged release tablet	28	EU/1/07/395/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	18/06/2009
Invega	Medcor Pharmaceuticals BV	15/10/2008	6 mg	Prolonged-release tablet	28	EU/1/07/395/026	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Sweden	The Netherlands	27/01/2009
Invega	Pharma Westen GmbH	24/12/2008	3 mg	Prolonged release tablet	98	EU/1/07/395/025	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/03/2009
Invega	Pharma Westen GmbH	19/01/2009	6 mg	Prolonged release tablet	98	EU/1/07/395/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/03/2009
Invega	Pharma Westen GmbH	10/06/2009	3 mg	Prolonged release tablet	98	EU/1/07/395/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/06/2009
Invega	Pharma Westen GmbH	10/06/2009	6 mg	Prolonged release tablet	98	EU/1/07/395/030	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/06/2009
Invirase	Haemato Pharm AG	04/09/2009	500 mg	Film-coated tablet	1 bottle	EU/1/96/026/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Bulgaria, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	05/10/2009
Invirase	SIA 'Elpis'	21/04/2009	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, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Latvia	29/05/2009
Iscover	Axicorp Pharma GmbH	22/12/2008	75 mg	Film-coated tablet	100	EU/1/98/070/004a	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	02/02/2009

Iscover	Docpharm GmbH&CoKGaA	05/08/2008	75 mg	Film-coated tablet	28	EU/1/98/070/001a	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/02/2009
Iscover	Docpharm GmbH&CoKGaA	05/08/2008	75 mg	Film-coated tablet	30	EU/1/98/070/005a	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/02/2009
Iscover	Docpharm GmbH&CoKGaA	05/08/2008	75 mg	Film-coated tablet	90	EU/1/98/070/006a	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/02/2009
Iscover	Docpharm GmbH&CoKGaA	05/08/2008	75 mg	Film-coated tablet	100	EU/1/98/070/004a	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/02/2009
Iscover	Kohlpharma GmbH	18/05/2009	75 mg	Film-coated tablet	28	EU/1/98/070/001a	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/06/2009
Isentress	Abacus Medicine A/S	22/05/2009	400 mg	Film-coated tablet	60	EU/1/07/436/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Denmark	Denmark	16/06/2009
Isentress	Euro Registratie Collectief BV	18/03/2009	400 mg	Film-coated tablet	60	EU/1/07/436/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/03/2009
Isentress	Haemato Pharm AG	21/10/2009	400 mg	Film-coated tablet	60	EU/1/07/436/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	03/11/2009
Janumet	Eureco-Pharma B.V.	08/06/2009	50 mg/1000 mg	Film-coated tablet	28	EU/1/08/455/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	30/06/2009

Janumet	Eureco-Pharma B.V.	08/06/2009	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, United Kingdom	The Netherlands	30/06/2009
Janumet	Eureco-Pharma B.V.	08/06/2009	50 mg/850 mg	Film-coated tablet	28	EU/1/08/455/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	30/06/2009
Janumet	Eureco-Pharma B.V.	08/06/2009	50 mg/850 mg	Film-coated tablet	56	EU/1/08/455/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Norway, Spain, Sweden, The Netherlands	The Netherlands	30/06/2009
Janumet	Medcor Pharmaceuticals BV	12/02/2009	50/1000mg	Film-coated tablets	56	EU/1/08/455/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	The Netherlands	24/02/2009
Janumet	Medcor Pharmaceuticals BV	12/02/2009	50/850mg	Film-coated tablets	56	EU/1/08/455/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	The Netherlands	24/02/2009
Januvia	Doncaster Pharmaceuticals Group Ltd	06/03/2009	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	04/05/2009
Januvia	Dr Fisher Farma BV	04/12/2009	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, United Kingdom	The Netherlands	17/12/2009
Januvia	Eureco-Pharma B.V.	22/06/2009	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, Luxembourg, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	08/10/2009
Januvia	Kohlpharma GmbH	31/03/2009	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/04/2009

Januvia	Kohlpharma GmbH	31/03/2009	100 mg	Film-coated tablet	98	EU/1/07/383/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/04/2009
Januvia	PCO Manufacturing	29/05/2009	100 mg	Film-coated tablet	28	EU/1/07/383/014	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland	02/07/2009
Januvia	Singad Pharma ApS	26/03/2008	100mg	Film-coated tablets	98	EU/1/07/383/017	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom	Denmark	24/02/2009
Kaletra	Aaston Healthcare GmbH	15/05/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Germany	30/06/2009
Kaletra	CC Pharma	04/02/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/005	France, Austria, Belgium, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/02/2009
Kaletra	CC Pharma	03/11/2009	200 mg / 50 mg	Film-coated tablet	360 (3 x 120)	EU/1/01/172/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Sweden, Denmark, Czech Republic, Finland, Slovak Republic	Germany	10/12/2009
Kaletra	Delphi Pharmaceuticals BV	23/07/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	31/07/2009
Kaletra	Emra-Med	10/07/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/005	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	22/07/2009
Kaletra	Haemato Pharm AG	08/06/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic	Germany, Austria	26/06/2009

Kaletra	Kohlpharma GmbH	16/02/2009	200 mg / 50 mg	Film-coated tablet	120 (blister)	EU/1/01/172/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/04/2009
Kaletra	Mevita HandelsGmbH	13/11/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	10/12/2009
Kaletra	Mevita HandelsGmbH	13/11/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	10/12/2009
Kaletra	Paranova Danmark AS	04/11/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	13/11/2009
Kaletra	Pharma Gerke GmbH	17/09/2008	100 mg / 25mg	Film-coated tablet	60	EU/1/01/172/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/04/2009
Kaletra	Pharma Gerke GmbH	13/02/2009	200/50mg	Film coated tablets	120 (blister)	EU/1/01/172/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Netherlands	Germany	23/02/2009
Kaletra	Pharma Westen GmbH	27/01/2009	200 mg / 50 mg	Film-coated tablet	120	EU/1/01/172/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/02/2009
Karvezide	Emra-Med	27/10/2006	150 mg/12.50 mg	Tablet	28	EU/1/98/085/001	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Karvezide	Emra-Med	27/10/2006	150 mg/12.50 mg	Tablet	56	EU/1/98/085/002	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Karvezide	Emra-Med	27/10/2006	150 mg/12.50 mg	Tablet	98	EU/1/98/085/003	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Karvezide	Emra-Med	27/11/2006	300 mg/12.50 mg	Tablet	28	EU/1/98/085/004	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, The Netherlands	Germany	09/02/2009

Karvezide	Emra-Med	27/11/2006	300 mg/12.50 mg	Tablet	56	EU/1/98/085/005	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/02/2009
Karvezide	Emra-Med	27/11/2006	300 mg/12.50 mg	Tablet	98	EU/1/98/085/006	Italy, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/02/2009
Karvezide	Emra-Med	01/05/2009	300 mg/25 mg	Film-coated tablet	28	EU/1/98/085/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/05/2009
Karvezide	Emra-Med	01/05/2009	300 mg/25 mg	Film-coated tablet	98	EU/1/98/085/027	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/05/2009
Karvezide	Pharma Westen GmbH	17/09/2007	300 mg/25 mg	Film-coated tablet	28	EU/1/98/085/024	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Karvezide	Pharma Westen GmbH	17/09/2007	300 mg/25 mg	Film-coated tablet	56	EU/1/98/085/025	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Karvezide	Pharma Westen GmbH	17/09/2007	300 mg/25 mg	Film-coated tablet	98	EU/1/98/085/027	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/03/2009
Keppra	Axicorp Pharma GmbH	04/12/2008	500 mg	Film-coated tablet	200	EU/1/00/146/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009
Keppra	Axicorp Pharma GmbH	06/07/2009	1000 mg	Film-coated tablet	200	EU/1/00/146/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/08/2009
Keppra	BR Lewis Ltd	10/02/2009	1000mg	Film-coated tablets	60	EU/1/00/146/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/06/2009

Keppra	BR Pharma International Ltd	19/12/2008	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/03/2009
Keppra	BR Pharma International Ltd	19/12/2008	750 mg	Film-coated tablet	200	EU/1/00/146/028	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/03/2009
Keppra	CC Pharma	20/01/2009	100 mg/ml	Oral solution	1 bottle + 1 oral syringe	EU/1/00/146/027	Austria, Italy, Spain, Norway, Belgium, France, Greece, The Netherlands, Portugal, United Kingdom	Germany	09/02/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	1000 mg	Film-coated tablet	50	EU/1/00/146/023	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	02/07/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	1000 mg	Film-coated tablet	100	EU/1/00/146/025	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	02/07/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	1000 mg	Film-coated tablet	200	EU/1/00/146/026	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	07/08/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/07/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	500 mg	Film-coated tablet	100	EU/1/00/146/011	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	23/07/2009
Keppra	Docpharm GmbH&CoKGaA	12/06/2009	500 mg	Film-coated tablet	200	EU/1/00/146/013	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	07/08/2009
Keppra	Docpharm GmbH&CoKGaA	03/08/2009	250 mg	Film-coated tablet	200	EU/1/00/146/029	Austria, Belgium, Denmark, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, United Kingdom, Finland, France, Spain, The Netherlands	Germany	12/10/2009
Keppra	Docpharm GmbH&CoKGaA	03/08/2009	250 mg	Film-coated tablet	100	EU/1/00/146/005	Austria, Belgium, Denmark, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, United Kingdom, Finland, France, Spain, The Netherlands	Germany	18/09/2009

Keppra	Docpharm GmbH&CoKGaA	03/08/2009	250 mg	Film-coated tablet	50	EU/1/00/146/003	Austria, Belgium, Denmark, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, United Kingdom, Finland, France, Malta, Spain, The Netherlands	Germany	18/09/2009
Keppra	Doncaster Pharmaceuticals Group Ltd	21/10/2009	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, Ireland, Italy, Liechtenstein, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Luxembourg	United Kingdom, Ireland, Malta	24/11/2009
Keppra	Dr Fisher Farma BV	30/04/2009	1000 mg	Film-coated tablet	60	EU/1/00/146/024	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	05/05/2009
Keppra	Dr Fisher Farma BV	30/04/2009	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Cyprus	The Netherlands	05/05/2009
Keppra	Emra-Med	11/05/2009	250 mg	Film-coated tablet	50	EU/1/00/146/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/06/2009
Keppra	Emra-Med	11/05/2009	250 mg	Film-coated tablet	100	EU/1/00/146/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/06/2009
Keppra	Emra-Med	11/05/2009	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/06/2009
Keppra	Emra-Med	14/05/2009	750 mg	Film-coated tablet	200	EU/1/00/146/028	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/06/2009
Keppra	Emra-Med	07/07/2009	500 mg	Film-coated tablet	200	EU/1/00/146/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Hungary, Estonia, Latvia, Lithuania, Romania, Poland, Slovenia, Slovak Republic	Germany	28/08/2009

Keppra	Euro Registratie Collectief BV	18/03/2009	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/03/2009
Keppra	EuroPharma DK	01/05/2009	100 mg/ml	Oral solution	1 bottle + 1 oral syringe	EU/1/00/146/027	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	13/05/2009
Keppra	Haemato Pharm AG	17/11/2009	1000 mg	Film-coated tablet	200	EU/1/00/146/026	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
Keppra	Haemato Pharm AG	17/11/2009	1000 mg	Film-coated tablet	100	EU/1/00/146/025	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
Keppra	Haemato Pharm AG	17/11/2009	1000 mg	Film-coated tablet	50	EU/1/00/146/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
Keppra	Haemato Pharm AG	17/11/2009	500 mg	Film-coated tablet	50	EU/1/00/146/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	24/11/2009
Keppra	Haemato Pharm AG	17/11/2009	500 mg	Film-coated tablet	100	EU/1/00/146/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
Keppra	Haemato Pharm AG	17/11/2009	500 mg	Film-coated tablet	200	EU/1/00/146/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/11/2009
Keppra	Interport Ltd	17/08/2009	100 mg/ml	Oral solution	1 bottle + 1 oral syringe 10 ml	EU/1/00/146/027	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain, The Netherlands	United Kingdom	23/10/2009

Keppra	Kohlpharma GmbH	02/12/2008	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009
Keppra	Kohlpharma GmbH	02/12/2008	750 mg	Film-coated tablet	200	EU/1/00/146/028	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	26/05/2009
Keppra	Kohlpharma GmbH	19/05/2009	250 mg	Film-coated tablet	200	EU/1/00/146/029	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Luxembourg	Germany	11/06/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	1000 mg	Film-coated tablet	50	EU/1/00/146/023	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	25/11/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	1000 mg	Film-coated tablet	200	EU/1/00/146/026	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	10/12/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	1000 mg	Film-coated tablet	100	EU/1/00/146/025	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	25/11/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	500 mg	Film-coated tablet	50	EU/1/00/146/009	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	25/11/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	500 mg	Film-coated tablet	100	EU/1/00/146/011	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	25/11/2009
Keppra	Milinda Arzneimittel GmbH	21/08/2009	500 mg	Film-coated tablet	200	EU/1/00/146/013	France, Ireland, Italy, Portugal, Spain, United Kingdom	Germany	10/12/2009
Keppra	PCO Manufacturing	19/01/2009	250 mg	Film-coated tablet	60	EU/1/00/146/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	15/05/2009
Keppra	Pharma First Limited	26/06/2009	500 mg	Film-coated tablet	60	EU/1/00/146/010	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland	07/09/2009
Keppra	Pharma Westen GmbH	24/07/2009	750 mg	Film-coated tablet	100	EU/1/00/146/019	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	31/07/2009

Keppra	Pharmachim AB	17/08/2009	100 mg/ml	Oral solution	1 bottle + 1 oral syringe 10 ml	EU/1/00/146/027	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	18/09/2009
Kineret	CC Pharma	10/02/2009	100mg	Solution for injection in a pre-filled syringe	28	EU/1/02/203/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	21/07/2009
Kinzalkomb	Medcor Pharmaceuticals BV	30/11/2009	40 mg/12.5 mg	Tablet	28	EU/1/02/214/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	11/12/2009
Kinzalkomb	Medcor Pharmaceuticals BV	30/11/2009	80 mg/12.5 mg	Tablet	28	EU/1/02/214/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	11/12/2009
Kinzalmono	Medcor Pharmaceuticals BV	16/03/2009	40 mg	Tablet	14	EU/1/98/091/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	20/04/2009
Kinzalmono	Medcor Pharmaceuticals BV	16/03/2009	80 mg	Tablet	14	EU/1/98/091/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	20/04/2009
Kivexa	Aaston Healthcare GmbH	29/05/2009	600mg/300mg	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Germany	30/06/2009
Kivexa	Abacus Medicine A/S	05/05/2009	600mg/300mg	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Hungary, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	14/05/2009
Kivexa	Combiphar Europe BV	29/01/2009	600/300mg	Film-coated tablets	30	EU/1/04/298/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/07/2009

Kivexa	Haemato Pharm AG	04/09/2009	600mg/300mg	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/10/2009
Kivexa	Pharmacodane Aps	17/02/2009	- -	Film-coated tablet	30	EU/1/04/298/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	07/04/2009
Lantus	Axicorp Pharma GmbH	22/12/2008	100IU/ml	Solution for injection in a cartridge	6x3ml		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/01/2009
Lantus	Axicorp Pharma GmbH	22/12/2008	100IU/ml	Solution for injection in a cartridge	9x3ml		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/01/2009
Lantus	Cross Healthcare Ltd	29/09/2008	100 Units/ml	Solution for injection in a vial	1 vial	EU/1/00/134/012	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/01/2009
Lantus	Eureco-Pharma B.V.	19/01/2009	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	11/03/2009
Lantus	Pharmachim AB	23/12/2008	100IU/ml	Solution for injection in a cartridge	5x3ml		Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	19/01/2009
Lantus Opticlick	Cross Healthcare Ltd	29/09/2008	100 Units/ml	Solution for injection in cartridge	5 cartridges	EU/1/00/134/025	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/01/2009
Lantus OptiClick	Euro Registratie Collectief BV	17/12/2008	100IU/ml	Solution for injection in a cartridge	5x3ml		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	12/01/2009
Lantus Optiset	Cross Healthcare Ltd	02/09/2008	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/010	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/01/2009
Lantus Optiset	Pharma Westen GmbH	12/02/2009	100 Units/ml	Solution for injection	9 pre-filled pens	EU/1/00/134/021	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/03/2009

Lantus OptiSet	Pharmachim AB	23/12/2008	100IU/ml	Solution for injection in a pre-filled pen	5 pens		Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	19/01/2009
Lantus Solostar	Euro Registratie Collectief BV	17/12/2008	100IU/ml	Solution for injection in a pre-filled pen	5 pens		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	12/01/2009
Lantus Solostar	Orifarm AS	16/10/2008	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	14/04/2009
Lantus Solostar	Pharma Westen GmbH	12/02/2009	100 Units/ml	Solution for injection	9 pre-filled pens	EU/1/00/134/036	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/03/2009
Lantus Solostar	Pharmachim AB	29/01/2009	100 Units/ml	Solution for injection	5 pre-filled pens	EU/1/00/134/033	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	14/04/2009
Levemir	Kohlpharma GmbH	01/12/2008	100 U/ml	Solution for injection	5 cartridges	EU/1/04/278/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009
Levemir FlexPen	Kohlpharma GmbH	04/02/2009	100IU/ml	Solution for injection in a pre-filled pen	5x3ml	EU/1/04/278/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/03/2009
Levemir FlexPen	Kohlpharma GmbH	04/02/2009	100IU/ml	Solution for injection in a pre-filled pen	10x3ml	EU/1/04/278/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/03/2009
Levemir FlexPen	Pharma Westen GmbH	17/02/2009	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/04/278/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/06/2009
Levemir FlexPen	Pharma Westen GmbH	17/02/2009	100 U/ml	Solution for injection	10 pre-filled pens	EU/1/04/278/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/06/2009
Levitra	2 Care 4	28/09/2009	20 mg	Film-coated tablet	4	EU/1/03/248/010	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	08/10/2009

Levitra	2 Care 4	19/10/2009	10 mg	Film-coated tablet	12	EU/1/03/248/008	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	28/10/2009
Levitra	2 Care 4	19/10/2009	20 mg	Film-coated tablet	12	EU/1/03/248/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	28/10/2009
Levitra	Abis-Pharma	14/08/2009	20 mg	Film-coated tablet	4	EU/1/03/248/010	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	12/10/2009
Levitra	Abis-Pharma	14/08/2009	20 mg	Film-coated tablet	8	EU/1/03/248/011	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	12/10/2009
Levitra	Abis-Pharma	14/08/2009	20 mg	Film-coated tablet	12	EU/1/03/248/012	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	12/10/2009
Levitra	AN Pharmacy Sp. z o.o.	08/09/2009	20 mg	Film-coated tablet	2	EU/1/03/248/009	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	14/12/2009
Levitra	AN Pharmacy Sp. z o.o.	08/09/2009	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, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	14/12/2009

Levitra	AN Pharmacy Sp. z o.o.	08/09/2009	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, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Slovenia	14/12/2009
Levitra	Euro Registratie Collectief BV	27/01/2009	20mg	Film-coated tablets	4	EU/1/03/248/010	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	17/02/2009
Levitra	Gervasi Farmacia SL	25/02/2009	10 mg	Film-coated tablet	2	EU/1/03/248/005	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009
Levitra	Gervasi Farmacia SL	25/02/2009	10 mg	Film-coated tablet	4	EU/1/03/248/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009
Levitra	Gervasi Farmacia SL	25/02/2009	10 mg	Film-coated tablet	8	EU/1/03/248/007	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009
Levitra	Gervasi Farmacia SL	25/02/2009	20 mg	Film-coated tablet	2	EU/1/03/248/009	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009
Levitra	Gervasi Farmacia SL	25/02/2009	20 mg	Film-coated tablet	4	EU/1/03/248/010	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009

Levitra	Gervasi Farmacia SL	25/02/2009	20 mg	Film-coated tablet	8	EU/1/03/248/011	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, The Netherlands, United Kingdom	Spain	28/04/2009
Levitra	Paranova Danmark AS	16/02/2009	20 mg	Film-coated tablet	12	EU/1/03/248/012	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/03/2009
Lucentis	Abacus Medicine A/S	05/05/2009	10 mg/ml	Solution for injection	1 vial	EU/1/06/374/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	14/05/2009
Lumigan	2 Care 4	02/10/2009	0.3 mg/ml	Eye drops, solution	3 bottles	EU/1/02/205/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Spain, Sweden, The Netherlands, United Kingdom	Denmark	22/10/2009
Lumigan	2 Care 4	16/10/2009	0.3 mg/ml	Eye drops, solution	1 bottle	EU/1/02/205/001	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Spain, Sweden, The Netherlands, United Kingdom	Denmark	03/12/2009
Lumigan	Orifarm AS	11/05/2009	0.3 mg/ml	Eye drops, solution	3 bottles	EU/1/02/205/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/06/2009
Lumigan	PCO Manufacturing	12/05/2009	0.3 mg/ml	Eye drops, solution	1 bottle	EU/1/02/205/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	United Kingdom, Ireland	10/09/2009
Lyrica	A1 Pharmaceuticals	22/12/2008	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	03/02/2009

Lyrica	A1 Pharmaceuticals	22/12/2008	200 mg	Capsule, hard	84	EU/1/04/279/021	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	03/02/2009
Lyrica	A1 Pharmaceuticals	22/12/2008	25 mg	Capsule, hard	56	EU/1/04/279/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	03/02/2009
Lyrica	A1 Pharmaceuticals	05/05/2009	25 mg	Capsule, hard	14	EU/1/04/279/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	07/07/2009
Lyrica	A1 Pharmaceuticals	05/05/2009	75 mg	Capsule, hard	14	EU/1/04/279/011	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	07/07/2009
Lyrica	A1 Pharmaceuticals	06/05/2009	50 mg	Capsule, hard	21	EU/1/04/279/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands	United Kingdom	07/07/2009
Lyrica	BeachCourse Limited	19/10/2009	75 mg	Capsule, hard	56	EU/1/04/279/012	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	03/11/2009
Lyrica	BR Lewis Ltd	16/10/2009	25 mg	Capsule, hard	100	EU/1/04/279/036	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom, Malta	04/12/2009
Lyrica	CC Pharma	14/11/2008	75mg	Hard capsules	100x1	EU/1/04/279/013	Norway	Germany	25/03/2009

Lyrica	Crystal Pharma Ltd	16/10/2009	25 mg	Capsule, hard	56	EU/1/04/279/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	26/11/2009
Lyrica	Doncaster Pharmaceuticals Group Ltd	15/04/2009	100 mg	Capsule, hard	21	EU/1/04/279/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	05/06/2009
Lyrica	Emra-Med	21/09/2007	300 mg	Capsule, hard	56	EU/1/04/279/024	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	26/01/2009
Lyrica	Interport Ltd	22/09/2009	25 mg	Capsule, hard	100	EU/1/04/279/036	Austria, Belgium, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Romania, Slovak Republic, Spain	United Kingdom	13/11/2009
Lyrica	Kohlpharma GmbH	06/05/2009	150 mg	Capsule, hard	100 x 1	EU/1/04/279/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/07/2009
Lyrica	Manichem Limited	04/02/2009	25 mg	Capsule, hard	56	EU/1/04/279/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Cyprus, Iceland, Liechtenstein, Norway	United Kingdom, Ireland	16/03/2009
Lyrica	Manichem Limited	28/05/2009	25 mg	Capsule, hard	14	EU/1/04/279/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Cyprus, Iceland, Liechtenstein, Norway	United Kingdom, Ireland	30/06/2009
Lyrica	MedicoPharm AG	08/07/2009	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/11/2009
Lyrica	MedicoPharm AG	08/07/2009	75 mg	Capsule, hard	56	EU/1/04/279/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/11/2009

Lyrica	Mediwin Ltd	24/06/2009	25 mg	Capsule, hard	84	EU/1/04/279/004	Austria, Belgium, France, Germany, Greece, Italy, Malta, The Netherlands, Portugal, Spain, Czech Republic, Latvia, Poland, Romania, Slovak Republic, Slovenia, Norway, Denmark	United Kingdom	30/07/2009
Lyrica	Mediwin Ltd	24/06/2009	50 mg	Capsule, hard	84	EU/1/04/279/009	Austria, Belgium, France, Germany, Greece, Italy, Malta, The Netherlands, Portugal, Spain, Cyprus, Denmark, Finland, Sweden	United Kingdom	30/07/2009
Lyrica	Milinda Arzneimittel GmbH	22/10/2009	75 mg	Capsule, hard	56	EU/1/04/279/012	France, Italy, Portugal	Germany	09/12/2009
Lyrica	MPT Pharma	29/01/2009	75mg	Hard capsules	14	EU/1/04/279/011	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	United Kingdom, Ireland	19/03/2009
Lyrica	MPT Pharma	29/01/2009	75mg	Hard capsules	56	EU/1/04/279/012	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	United Kingdom, Ireland	19/03/2009
Lyrica	MPT Pharma	23/02/2009	100 mg	Capsule, hard	84	EU/1/04/279/015	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	19/03/2009
Lyrica	MPT Pharma	23/02/2009	150 mg	Capsule, hard	14	EU/1/04/279/017	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	08/04/2009
Lyrica	MPT Pharma	23/02/2009	150 mg	Capsule, hard	56	EU/1/04/279/018	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	08/04/2009

Lyrica	MPT Pharma	16/03/2009	50 mg	Capsule, hard	21	EU/1/04/279/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	19/03/2009
Lyrica	MPT Pharma	16/03/2009	50 mg	Capsule, hard	56	EU/1/04/279/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom, Slovak Republic	United Kingdom, Ireland	19/03/2009
Lyrica	MPT Pharma	16/03/2009	50 mg	Capsule, hard	84	EU/1/04/279/009	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom, Slovak Republic	United Kingdom, Ireland	19/03/2009
Lyrica	MPT Pharma	14/04/2009	25 mg	Capsule, hard	14	EU/1/04/279/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	18/05/2009
Lyrica	MPT Pharma	14/04/2009	25 mg	Capsule, hard	56	EU/1/04/279/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	18/05/2009
Lyrica	MPT Pharma	14/04/2009	25 mg	Capsule, hard	84	EU/1/04/279/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom, Slovak Republic	Ireland, United Kingdom	18/05/2009
Lyrica	MPT Pharma	23/04/2009	100 mg	Capsule, hard	21	EU/1/04/279/014	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Romania, Spain, Sweden, The Netherlands, United Kingdom, Slovak Republic	Ireland, United Kingdom	04/05/2009

Lyrica	MPT Pharma	08/10/2009	25 mg	Capsule, hard	21	EU/1/04/279/002	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, Slovak Republic	Ireland, United Kingdom	03/11/2009
Lyrica	N B Enterprises Ltd	04/03/2009	75 mg	Capsule, hard	56	EU/1/04/279/012	Austria, Belgium, Bulgaria, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	16/07/2009
Lyrica	O.P.D. Laboratories Ltd	28/07/2009	25 mg	Capsule, hard	100	EU/1/04/279/036	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom, Ireland	06/08/2009
Lyrica	Pharma Westen GmbH	17/02/2009	75 mg	Capsule, hard	100 x 1	EU/1/04/279/013	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/07/2009
Lyrica	Pharma Westen GmbH	21/04/2009	300 mg	Capsule, hard	100	EU/1/04/279/043	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Romania	Germany	06/07/2009
Lyrica	Veron Pharma	08/05/2009	100 mg	Capsule, hard	21	EU/1/04/279/014	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	30/06/2009
Lyrica	Veron Pharma	08/05/2009	200 mg	Capsule, hard	21	EU/1/04/279/020	Austria, Belgium, France, Greece, Italy, Portugal, Spain, The Netherlands, United Kingdom	Germany	30/06/2009
Lyrica	Waymade PLC	13/05/2009	25 mg	Capsule, hard	100	EU/1/04/279/036	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Italy, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	28/05/2009
Lyrica	Waymade PLC	23/10/2009	150 mg	Capsule, hard	14	EU/1/04/279/017	Austria, Belgium, Bulgaria, Czech Republic, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Latvia, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	06/11/2009
Lysodren	CC Pharma	25/08/2009	500 mg	Tablet	100	EU/1/04/273/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	17/09/2009
Lysodren	Kohlpharma GmbH	02/03/2009	500 mg	Tablet	100	EU/1/04/273/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/03/2009

MabCampath	CC Pharma	04/08/2009	30 mg/ml	Concentrate for solution for infusion	3 vials	EU/1/01/193/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	28/08/2009
MabCampath	Emra-Med	31/12/2008	30mg/ml	Concentrate for solution for infusion	3	EU/1/01/193/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	11/03/2009
MabCampath	Haemato Pharm AG	07/05/2009	30 mg/ml	Concentrate for solution for infusion	3 vials	EU/1/01/193/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany, Austria	17/06/2009
Mabthera	Aaston Healthcare GmbH	20/03/2009	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	12/06/2009
Mabthera	Kohlpharma GmbH	15/09/2008	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Italy, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/02/2009
Mabthera	Kohlpharma GmbH	15/09/2008	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Italy, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	10/02/2009
Mabthera	Orifarm OY	25/03/2009	100 mg	Concentrate for solution for infusion	2 vials	EU/1/98/067/001	Austria, Belgium, Cyprus, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Denmark, Czech Republic, Poland, Romania, Bulgaria, Hungary, Latvia, Lithuania	Finland	18/05/2009
Mabthera	Orifarm OY	25/03/2009	500 mg	Concentrate for solution for infusion	1 vial	EU/1/98/067/002	Austria, Belgium, Cyprus, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Denmark, Czech Republic, Poland, Romania, Bulgaria, Hungary, Latvia, Lithuania, Slovak Republic	Finland	18/05/2009
Micardis	2 Care 4	11/02/2008	80mg	Tablets	98		Czech Republic, Bulgaria	Denmark	21/01/2009

Micardis	Abis-Pharma	14/08/2009	40 mg	Tablet	28	EU/1/98/090/002	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	Abis-Pharma	14/08/2009	40 mg	Tablet	56	EU/1/98/090/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	Abis-Pharma	14/08/2009	40 mg	Tablet	98	EU/1/98/090/004	Slovak Republic, Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	Abis-Pharma	14/08/2009	80 mg	Tablet	28	EU/1/98/090/006	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	Abis-Pharma	14/08/2009	80 mg	Tablet	56	EU/1/98/090/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	Abis-Pharma	14/08/2009	80 mg	Tablet	98	EU/1/98/090/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	24/09/2009
Micardis	EuroPharma DK	20/04/2009	20 mg	Tablet	28	EU/1/98/090/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/06/2009
Micardis	G-Pharma Ltd	12/01/2009	80mg	Tablets	14	EU/1/98/090/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	28/01/2009

Micardis	G-Pharma Ltd	12/01/2009	40mg	Tablets	14	EU/1/98/090/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	28/01/2009
Micardis Plus	Pharma Gerke GmbH	22/12/2008	80/25mg	Tablets	98		Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	19/01/2009
Micardis Plus	Pharma Gerke GmbH	22/12/2008	80/25mg	Tablets	56		Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	19/01/2009
MicardisPlus	Beragena Arzneimittel GmbH	07/10/2009	40 mg/12.5 mg	Tablet	98	EU/1/02/213/005	Austria, Belgium, France, Greece, Italy, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	03/11/2009
MicardisPlus	Crystal Pharma Ltd	27/10/2006	80 mg/12.5 mg	Tablet	28	EU/1/02/213/007	Austria, Belgium, Cyprus, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, Denmark, Finland, Iceland, Liechtenstein, Luxembourg, Romania, Slovak Republic	United Kingdom	15/05/2009
MicardisPlus	Doncaster Pharmaceuticals Group Ltd	02/04/2009	80 mg/25 mg	Tablet	28	EU/1/02/213/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	18/05/2009
MicardisPlus	Doncaster Pharmaceuticals Group Ltd	26/10/2009	40 mg/12.5 mg	Tablet	14	EU/1/02/213/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/11/2009
MicardisPlus	Emra-Med	26/01/2009	80 mg/25 mg	Tablet	98	EU/1/02/213/023	Austria, Belgium, Cyprus, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovenia, Spain, Sweden, United Kingdom, Czech Republic	Germany	03/02/2009
MicardisPlus	G-Pharma Ltd	31/07/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	28/08/2009
MicardisPlus	Kohlpharma GmbH	10/02/2009	80 mg/25 mg	Tablet	28	EU/1/02/213/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/04/2009

MicardisPlus	Kohlpharma GmbH	10/02/2009	80 mg/25 mg	Tablet	56	EU/1/02/213/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/04/2009
MicardisPlus	Kohlpharma GmbH	10/02/2009	80 mg/25 mg	Tablet	98	EU/1/02/213/023	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/04/2009
MicardisPlus	Orifarm AS	05/12/2008	80 mg/25 mg	Tablet	28	EU/1/02/213/018	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/01/2009
MicardisPlus	Orifarm AS	05/12/2008	80 mg/25 mg	Tablet	98	EU/1/02/213/023	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/01/2009
MicardisPlus	Pharma Gerke GmbH	15/05/2008	80 mg/12.5 mg	Tablet	28	EU/1/02/213/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	14/09/2009
MicardisPlus	Pharma Gerke GmbH	15/05/2008	80 mg/12.5 mg	Tablet	56	EU/1/02/213/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	14/09/2009
MicardisPlus	Pharma Gerke GmbH	15/05/2008	80 mg/12.5 mg	Tablet	98	EU/1/02/213/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	14/09/2009
MicardisPlus	Polyfarma Ltd	04/02/2009	40 mg/12.5 mg	Tablet	28	EU/1/02/213/002	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom, Ireland	12/02/2009
MicardisPlus	Samforlab SA	09/02/2009	80 mg/12.5 mg	Tablet	28	EU/1/02/213/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Sweden, United Kingdom, Greece	Spain	02/06/2009
MicardisPlus	Swingward Ltd TA Medihealth	30/06/2009	40 mg/12.5 mg	Tablet	14	EU/1/02/213/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden	United Kingdom	08/12/2009

Mimpara	Abacus Medicine A/S	22/05/2009	30 mg	Film-coated tablet	28	EU/1/04/292/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Slovak Republic, Slovenia, Bulgaria, Romania	Denmark	17/06/2009
Mimpara	Abacus Medicine A/S	22/05/2009	60 mg	Film-coated tablet	28	EU/1/04/292/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Slovak Republic, Slovenia, Bulgaria, Romania	Denmark	17/06/2009
Mimpara	Abacus Medicine A/S	22/05/2009	90 mg	Film-coated tablet	28	EU/1/04/292/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Slovak Republic, Slovenia, Bulgaria, Romania	Denmark	17/06/2009
Mimpara	Lex ano UAB	03/03/2009	60 mg	Film-coated tablet	28	EU/1/04/292/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, 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	Lithuania	30/04/2009
Mimpara	Lex ano UAB	06/04/2009	30 mg	Film-coated tablet	28	EU/1/04/292/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Liechtenstein, Latvia, Italy, Ireland, Iceland, Luxembourg, Malta, Norway, Poland, Portugal, Romania, United Kingdom, The Netherlands, Sweden, Spain, Slovenia, Slovak Republic	Lithuania	29/04/2009
Mirapexin	BR Pharma International Ltd	10/09/2009	0.7 mg	Tablet	100	EU/1/97/051/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/10/2009
Mirapexin	Euro Registratie Collectief BV	18/03/2009	0.088 mg	Tablet	30	EU/1/97/051/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/03/2009
Mirapexin	Europharma DK	17/12/2007	0.7 mg	Tablet	100	EU/1/97/051/006	Austria, Belgium, France, Germany, Greece, Italy, NT, Portugal, Spain, United Kingdom, The Netherlands	Denmark	14/05/2009

Mirapexin	MedicoPharm AG	02/09/2009	0.18 mg	Tablet	30	EU/1/97/051/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/11/2009
Mirapexin	Milinda Arzneimittel GmbH	21/08/2009	0.7 mg	Tablet	100	EU/1/97/051/006	Italy, Spain	Germany	26/11/2009
Mirapexin	Orifarm AB	26/10/2009	0.18 mg	Tablet	100	EU/1/97/051/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	04/11/2009
Mirapexin	Orifarm AB	26/10/2009	0.7 mg	Tablet	100	EU/1/97/051/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	04/11/2009
Mirapexin	PCO Manufacturing	07/05/2009	0.18 mg	Tablet	30	EU/1/97/051/003	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Italy	Ireland, United Kingdom	15/10/2009
Mirapexin	PCO Manufacturing	07/05/2009	0.7 mg	Tablet	30	EU/1/97/051/005	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Italy	Ireland, United Kingdom	15/10/2009
Mirapexin	Pharma Gerke GmbH	24/11/2006	0.18 mg	Tablet	100	EU/1/97/051/004	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	15/10/2009
Mircera	CC Pharma	02/07/2009	100 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Latvia	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	120 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/020	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Latvia	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	150 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/011	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Romania, Denmark, Finland, Sweden	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	200 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/012	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Latvia	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	250 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/013	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	05/08/2009

Mircera	CC Pharma	02/07/2009	30 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/017	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Latvia	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	50 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Spain, United Kingdom, Portugal, Bulgaria, Latvia	Germany	05/08/2009
Mircera	CC Pharma	02/07/2009	75 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Latvia	Germany	05/08/2009
Mircera	Kohlpharma GmbH	06/05/2009	100 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/06/2009
Mircera	Kohlpharma GmbH	06/05/2009	50 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/06/2009
Mircera	Kohlpharma GmbH	06/05/2009	75 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	18/06/2009
Mircera	Kohlpharma GmbH	18/05/2009	150 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	18/06/2009
Mircera	Kohlpharma GmbH	03/08/2009	200 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/08/2009
Mircera	Kohlpharma GmbH	04/08/2009	250 µg/0.3 ml	Solution for injection	1 pre-filled syringe	EU/1/07/400/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/08/2009
Mixtard 30 Penfill	Pharma Westen GmbH	03/12/2008	100IU/ml	Suspension for injection in a cartridge	5x3ml		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009

Mixtard 30 Penfill	Pharma Westen GmbH	03/12/2008	100IU/ml	Suspension for injection in a cartridge	10x3ml		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009
M-M-RVAXPRO	Emra-med	28/01/2009	- -	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, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/04/2009
M-M-RVAXPRO	Emra-med	28/01/2009	- -	Powder and solvent for suspension for injection in pre-filled syringe	10 vials + 10 pre-filled syringes + 20 needles	EU/1/06/337/012	Belgium, Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/04/2009
Myocet	Haemato Pharm AG	11/09/2009	50 mg	Powder and pre-admixtures for concentrate for liposomal dispersion for infusion	Two sets of one vial for each of the three components	EU/1/00/141/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	12/11/2009
Neoclarityn	B&S Healthcare Ltd	18/08/2009	5 mg	Film-coated tablet	30	EU/1/00/161/011	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, United Kingdom	08/09/2009
Neoclarityn	BR Lewis Ltd	04/12/2008	0.5mg/ml	Syrup	100ml		Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	05/01/2009
Neoclarityn	Doncaster Pharmaceuticals Group Ltd	02/07/2009	0.5 mg/ml	Oral solution	1 bottle	EU/1/00/161/062	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	04/08/2009
Neoclarityn	Euro Registratie Collectief BV	14/04/2009	5 mg	Film-coated tablet	30	EU/1/00/161/011	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/05/2009
Neoclarityn	PCO Manufacturing	19/01/2009	0.5 mg/ml	Syrup	100 ml - 1 bottle with 1 spoon	EU/1/00/161/017	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland	06/07/2009

NeoRecormon	Haemato Pharm AG	09/04/2009	10000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/036	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2009
NeoRecormon	Haemato Pharm AG	09/04/2009	30000 IU	Solution for injection	4 pre-filled syringes	EU/1/97/031/046	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2009
NeoRecormon	Haemato Pharm AG	09/04/2009	5000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Germany	Germany, Austria	04/06/2009
NeoRecormon	Haemato Pharm AG	09/04/2009	6000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/044	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	04/06/2009
NeoRecormon	Haemato Pharm AG	23/06/2009	2000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/030	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic	Germany	10/08/2009
NeoRecormon	Paranova Läkemedel AB	02/03/2009	5000 IU	Solution for injection	6 pre-filled syringes	EU/1/97/031/034	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	06/04/2009
Neulasta	Aaston Healthcare GmbH	23/03/2009	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	22/04/2009
Neulasta	Cancernova GmbH onkologische Arzneimittel	29/10/2009	6 mg	Solution for injection	1 pre-filled syringe (non-blister)	EU/1/02/227/002	Hungary, Poland	Germany	23/11/2009
Neulasta	CC Pharma	11/08/2009	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/002	Austria, Belgium, France, Greece, Hungary, Italy, Norway, Portugal, Spain, United Kingdom, Poland, Czech Republic	Germany	16/09/2009

Neulasta	eurorx Arzneimittel GmbH	18/05/2009	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, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	25/06/2009
Neulasta	Haemato Pharm AG	29/06/2009	6 mg	Solution for injection	1 pre-filled syringe (non blister)	EU/1/02/227/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Germany, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany, Austria	15/07/2009
Neulasta	Inopha GmbH	18/02/2009	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	13/05/2009
Neulasta	Kohlpharma GmbH	01/06/2009	6 mg	Solution for injection	1 pre-filled syringe (non-blister)	EU/1/02/227/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/06/2009
Neulasta	Medartuum AB	24/08/2009	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/002	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	12/10/2009
Neulasta	MedicoPharm AG	13/05/2009	6 mg	Solution for injection	1 pre-filled syringe (non blister)	EU/1/02/227/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	28/05/2009
Neulasta	NewNeopharm BV	04/09/2009	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	The Netherlands	16/09/2009

Neulasta	Pharma Westen GmbH	23/12/2008	6 mg	Solution for injection	1 pre-filled syringe	EU/1/02/227/002	Poland, Bulgaria	Germany	10/02/2009
Neupro	BR Pharma International Ltd	17/08/2009	2 mg/24h	Transdermal patch	28 patches	EU/1/05/331/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, Malta, United Kingdom	12/11/2009
Neupro	BR Pharma International Ltd	17/08/2009	4 mg/24h	Transdermal patch	28 patches	EU/1/05/331/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, Malta, United Kingdom	12/11/2009
Neupro	BR Pharma International Ltd	17/08/2009	6 mg/24h	Transdermal patch	28 patches	EU/1/05/331/008	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, Malta, United Kingdom	12/11/2009
Neupro	BR Pharma International Ltd	17/08/2009	8 mg/24h	Transdermal patch	28 patches	EU/1/05/331/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, Liechtenstein, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, Malta, United Kingdom	12/11/2009
Neupro	O.P.D. Laboratories Ltd	16/10/2009	2 mg/24h	Transdermal patch	28 patches	EU/1/05/331/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	28/10/2009
Neupro	O.P.D. Laboratories Ltd	16/10/2009	4 mg/24h	Transdermal patch	28 patches	EU/1/05/331/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	28/10/2009
Neupro	O.P.D. Laboratories Ltd	16/10/2009	6 mg/24h	Transdermal patch	28 patches	EU/1/05/331/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	28/10/2009
Neupro	O.P.D. Laboratories Ltd	16/10/2009	8 mg/24h	Transdermal patch	28 patches	EU/1/05/331/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	Ireland, United Kingdom	28/10/2009
Neupro	Waymade PLC	19/08/2009	4 mg/24h	Transdermal patch	28 patches	EU/1/05/331/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, Malta, United Kingdom	United Kingdom, Malta, Ireland	10/09/2009
Neupro	Waymade PLC	19/08/2009	8 mg/24h	Transdermal patch	28 patches	EU/1/05/331/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, Malta, United Kingdom	United Kingdom, Malta, Ireland	10/09/2009
Nevanac	Interport Ltd	05/11/2009	1 mg/ml	Eye drops, suspension	1 x 5 ml	EU/1/07/433/001	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	03/12/2009

Nexavar	Haemato Pharm AG	11/06/2009	200 mg	Film-coated tablet	112	EU/1/06/342/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany, Hungary, Lithuania, Latvia, Slovenia, Slovak Republic, Poland, Romania	Germany, Austria, Bulgaria, Czech Republic, Estonia	04/08/2009
Norvir	Aaston Healthcare GmbH	08/06/2009	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	11/06/2009
Norvir	Emra-Med	24/06/2009	100 mg	Capsule, soft	4 bottles x 84	EU/1/96/016/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/10/2009
Norvir	Emra-Med	24/06/2009	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/10/2009
Norvir	Haemato Pharm AG	28/05/2009	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/08/2009
Norvir	Haemato Pharm AG	07/09/2009	100 mg	Capsule, soft	4 bottles x 84	EU/1/96/016/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Slovak Republic, Spain, Sweden, United Kingdom	Germany	22/10/2009
Norvir	Medcor Pharmaceuticals BV	17/02/2009	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Sweden, Spain, United Kingdom	The Netherlands	30/04/2009
Norvir	Paranova Läkemedel AB	02/03/2009	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	14/04/2009
Norvir	Pharma Gerke GmbH	04/08/2008	100 mg	Capsule, soft	84 soft	EU/1/96/016/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	26/01/2009

NovoMix 30 FlexPen	Ginova Ltd	15/03/2007	100IU/ml	Suspension for injection in a pre-filled pen	5x3ml		Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	14/01/2009
NovoMix 30 Penfill	Ginova Ltd	15/03/2007	100IU/ml	Suspension for injection in a cartridge	5x3ml		Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	14/01/2009
NovoNorm	abis-Pharma	13/11/2009	0.5 mg	Tablet	30	EU/1/98/076/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
NovoNorm	Abis-Pharma	13/11/2009	1 mg	Tablet	30	EU/1/98/076/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
NovoNorm	Abis-Pharma	13/11/2009	2 mg	Tablet	30	EU/1/98/076/018	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	04/12/2009
NovoNorm	Europharma Sverige AB	02/04/2009	1 mg	Tablet	90	EU/1/98/076/012	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/05/2009
NovoNorm	Europharma Sverige AB	02/04/2009	2 mg	Tablet	90	EU/1/98/076/019	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	06/05/2009
NovoNorm	Orifarm AB	09/10/2009	2 mg	Tablet	120	EU/1/98/076/020	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom, Czech Republic, Romania, Slovak Republic	Sweden	22/10/2009
NovoNorm	Pharmachim AB	20/07/2009	1 mg	Tablet	120	EU/1/98/076/013	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/07/2009
NovoNorm	Pharmachim AB	20/07/2009	2 mg	Tablet	120	EU/1/98/076/020	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Portugal, Spain, The Netherlands, United Kingdom	Sweden	27/07/2009
NovoRapid	CC Pharma	16/10/2009	100 U/ml	Solution for injection	5 vials	EU/1/99/119/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/11/2009

NovoRapid FlexPen	Dr Fisher Farma BV	29/01/2009	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/99/119/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/02/2009
NovoRapid Flexpen	Medcor Pharmaceuticals BV	09/03/2009	100 U/ml	Solution for injection	5 pre-filled pens	EU/1/99/119/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	04/05/2009
NovoRapid Penfill	CC Pharma	16/10/2009	100 U/ml	Solution for injection	5 cartridges	EU/1/99/119/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	25/11/2009
Noxafil	CC Pharma	10/02/2009	40mg/ml	Oral suspension	1 bottle	EU/1/05/320/001	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	25/02/2009
Nplate	CC Pharma	01/10/2009	250 µg	Powder for solution for injection	1 vial	EU/1/08/497/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/11/2009
Nplate	CC Pharma	01/10/2009	500 µg	Powder for solution for injection	1 vial	EU/1/08/497/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/11/2009
Nplate	Haemato Pharm AG	14/09/2009	250 µg	Powder for solution for injection	1 vial	EU/1/08/497/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	30/11/2009
Nplate	Haemato Pharm AG	14/09/2009	250 µg	Powder for solution for injection	4 vials	EU/1/08/497/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	30/11/2009
Nplate	Haemato Pharm AG	14/09/2009	500 µg	Powder for solution for injection	1 vial	EU/1/08/497/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	30/11/2009

Nplate	Haemato Pharm AG	14/09/2009	500 µg	Powder for solution for injection	4 vials	EU/1/08/497/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	30/11/2009
NutropinAq	Paranova Läkemedel AB	09/03/2009	10 mg/2 ml	Solution for injection	3 cartridges	EU/1/00/164/004	Belgium, Austria, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	07/04/2009
Opatanol	AN Pharmacy Sp. z o.o.	07/04/2009	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	09/06/2009
Opatanol	AN Pharmacy Sp. z o.o.	07/04/2009	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	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, Spain, Sweden, The Netherlands, United Kingdom, Poland	Slovenia	09/06/2009
Opatanol	Gervasi Farmacia SL	24/11/2008	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	15/01/2009
Opatanol	Kohlpharma GmbH	09/10/2009	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/11/2009
Opatanol	PCO Manufacturing	07/05/2009	1 mg/ml	Eye drops, solution	1 bottle	EU/1/02/217/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Hungary, Italy	Ireland, United Kingdom	14/05/2009

Orencia	Abacus Medicine A/S	22/05/2009	250 mg	Powder for concentrate for solution for infusion	1 vial + 1 syringe	EU/1/07/389/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic	Denmark	24/06/2009
Orencia	CC Pharma	04/02/2009	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/07/389/002	Norway, United Kingdom, Greece, Austria, Belgium, France, Italy, The Netherlands, Portugal, Spain, Romania	Germany	23/02/2009
Orencia	CC Pharma	04/02/2009	250 mg	Powder for concentrate for solution for infusion	3 vials + 3 syringes	EU/1/07/389/003	Norway, United Kingdom, Greece, Austria, Belgium, France, Italy, The Netherlands, Portugal, Spain, Romania	Germany	23/02/2009
Orencia	Haemato Pharm AG	01/06/2009	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/07/389/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Lithuania, Romania, Slovak Republic, Slovenia	Germany	14/07/2009
Orencia	Haemato Pharm AG	01/06/2009	250 mg	Powder for concentrate for solution for infusion	3 vials + 3 syringes	EU/1/07/389/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Lithuania, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	14/07/2009
Orencia	Kohlpharma GmbH	23/07/2009	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/07/389/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/10/2009
Orencia	Kohlpharma GmbH	23/07/2009	250 mg	Powder for concentrate for solution for infusion	3 vials + 3 syringes	EU/1/07/389/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/10/2009
Orencia	Orifarm AS	10/08/2009	250 mg	Powder for concentrate for solution for infusion	1 vial + 1 syringe	EU/1/07/389/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania, Czech Republic, Slovak Republic, Slovenia	Denmark	27/11/2009

Orencia	Pharma Gerke GmbH	17/04/2009	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/07/389/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/04/2009
Orencia	Pharma Gerke GmbH	17/04/2009	250 mg	Powder for concentrate for solution for infusion	3 vials + 3 syringes	EU/1/07/389/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/04/2009
Orencia	Pharma Westen GmbH	08/04/2009	250 mg	Powder for concentrate for solution for infusion	2 vials + 2 syringes	EU/1/07/389/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/04/2009
Orencia	Pharma Westen GmbH	08/04/2009	250 mg	Powder for concentrate for solution for infusion	3 vials + 3 syringes	EU/1/07/389/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/04/2009
Orgalutran	Dr Fisher Farma BV	26/01/2009	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	06/02/2009
Ovitrelle	Emra-Med	08/12/2008	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/01/2009
Ovitrelle	Pharmachim AB	10/03/2009	250 micrograms/0.5 ml	Solution for injection	1 pre-filled syringe	EU/1/00/165/007	Austria, Belgium, Bulgaria, Czech Republic, France, Germany, Greece, Hungary, Ireland, Italy, Latvia, Lithuania, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Spain, United Kingdom	Sweden	22/04/2009
Pegasys	Axicorp Pharma GmbH	23/10/2009	180 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	02/12/2009
Pegasys	Combiphar Europe BV	01/09/2009	180 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/008	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	22/09/2009
Pegasys	Dr Fisher Farma BV	18/02/2009	180 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	25/06/2009

Pegasys	Emra-Med	01/04/2008	180 µg	Solution for injection	1 pre-filled syringe + 1 injection needle	EU/1/02/221/007	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	12/02/2009
Pegasys	Emra-Med	01/04/2008	180 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/008	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	12/02/2009
Pegasys	Emra-Med	01/04/2008	180 µg	Solution for injection	12 pre-filled syringes + 12 injection needles	EU/1/02/221/010	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	12/02/2009
Pegasys	Kohlpharma GmbH	15/06/2009	135 µg	Solution for injection	12 pre-filled syringes + 12 injection needles	EU/1/02/221/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	18/06/2009
Pegasys	Medartuum AB	02/03/2009	180 µg	Solution for injection	4 pre-filled syringes + 4 injection needles	EU/1/02/221/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	20/03/2009
Pegasys	Pharma Gerke GmbH	18/02/2008	180 µg	Solution for injection	12 pre-filled syringes + 12 injection needles	EU/1/02/221/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania, Latvia	Germany	12/10/2009
Pegasys	Pharma Westen GmbH	18/09/2008	180µg	Solution for injection in a pre-filled syringe	4+4	EU/1/02/221/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Iceland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania, Latvia, Lithuania, Estonia	Germany	16/02/2009
PegIntron	Axicorp Pharma GmbH	23/10/2009	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, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/12/2009

PegIntron	Axicorp Pharma GmbH	23/10/2009	120 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/044	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/12/2009
PegIntron	Axicorp Pharma GmbH	23/10/2009	150 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/048	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	03/12/2009
PegIntron	Kohlpharma GmbH	01/12/2008	100IU/ml	Solution for injection in a cartridge	5x3ml		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/01/2009
PegIntron	Kohlpharma GmbH	14/10/2009	150 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/048	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Iceland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/12/2009
PegIntron	Orifarm AS	13/07/2009	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, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	15/07/2009
PegIntron	Orifarm AS	13/07/2009	80 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/036	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	15/07/2009
PegIntron	Paranova Läkemedel AB	28/02/2007	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	Belgium, Finland, Greece, The Netherlands, Norway, Portugal, Austria, Cyprus, Denmark, France, Germany, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Spain, United Kingdom	Sweden	16/03/2009
PegIntron	Paranova Läkemedel AB	19/01/2009	80 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/131/036	Belgium, Finland, Greece, The Netherlands, Norway, Portugal, Spain, Austria, Cyprus, Denmark, France, Germany, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, United Kingdom	Sweden	16/03/2009

Plavix	B&S Healthcare Ltd	20/07/2009	75 mg	Film-coated tablet	30	EU/1/98/069/005a	Austria, Belgium, Bulgaria, Cyprus, Czech 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	28/08/2009
Plavix	CC Pharma	26/08/2008	75 mg	Film-coated tablet	100	EU/1/98/069/004a	Italy, Norway, Portugal, Spain, Austria, Belgium, France, Greece, The Netherlands, United Kingdom	Germany	26/01/2009
Plavix	CC Pharma	26/08/2008	75 mg	Film-coated tablet	28	EU/1/98/069/001a	Italy, Norway, Portugal, Spain, Austria, Belgium, France, Greece, The Netherlands, United Kingdom	Germany	26/01/2009
Plavix	Doncaster Pharmaceuticals Group Ltd	07/05/2009	75 mg	Film-coated tablet	30	EU/1/98/069/005a	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Ireland, Malta, United Kingdom	18/06/2009
Plavix	Dr Fisher Farma BV	10/03/2009	300 mg	Film-coated tablet	30 x 1	EU/1/98/069/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/04/2009
Plavix	Eureco-Pharma B.V.	24/02/2009	75 mg	Film-coated tablet	30	EU/1/98/069/005a	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	06/07/2009
Plavix	Euro Registratie Collectief BV	27/01/2009	300mg	Film-coated tablets	30	EU/1/98/069/009	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	17/02/2009
Plavix	Imed Healthcare Ltd	13/10/2009	75 mg	Film-coated tablet	30	EU/1/98/069/005a	United Kingdom, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden	Ireland, United Kingdom, Malta	03/11/2009
Plavix	Paranova Danmark AS	01/06/2009	75 mg	Film-coated tablet	28	EU/1/98/069/001a	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	26/06/2009
Plavix	Pharma First Limited	30/06/2009	75 mg	Film-coated tablet	28	EU/1/98/069/001a	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Ireland	21/10/2009

Plavix	Profind Wholesale Ltd.	09/02/2009	75 mg	Film-coated tablet	30	EU/1/98/069/005a	United Kingdom	Ireland	26/03/2009
Pradaxa	Interport Ltd	16/02/2009	110 mg	Capsule, hard	10 x 1	EU/1/08/442/005	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Poland, Spain	United Kingdom	09/03/2009
Pradaxa	Interport Ltd	16/02/2009	75 mg	Capsule, hard	10 x 1	EU/1/08/442/001	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Poland, Portugal, Spain	United Kingdom	09/03/2009
Prandin	Emra-Med	29/09/2008	1 mg	Tablet	120	EU/1/00/162/011	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/01/2009
Prandin	Emra-Med	20/10/2008	2 mg	Tablet	120	EU/1/00/162/017	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/01/2009
Preotact	CC Pharma	01/10/2009	100 µg	Powder and solvent for solution for injection	2 cartridges	EU/1/06/339/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/11/2009
Preotact	CC Pharma	01/10/2009	100 µg	Powder and solvent for solution for injection	6 cartridges	EU/1/06/339/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/11/2009
Prevenar	Eurim-Pharm	24/02/2009	- -	Suspension for injection	1 pre-filled syringe with separate needle	EU/1/00/167/006	France	Germany	11/05/2009
Prevenar	InPharm Spzoo	21/04/2009	- -	Suspension for injection	1 pre-filled syringe with separate needle	EU/1/00/167/006	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Sweden, The Netherlands, United Kingdom, Portugal, Spain	Poland	24/06/2009
Prevenar	Kohlpharma GmbH	16/07/2009	- -	Suspension for injection	1 pre-filled syringe with separate needle	EU/1/00/167/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/08/2009
Prevenar	Pharmachim AB	03/03/2009	- -	Suspension for injection	1 pre-filled syringe with separate needle	EU/1/00/167/006	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	06/05/2009
Prezista	CC Pharma	23/12/2008	300mg	Filmcoated tablets	120	EU/1/06/380/001	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Romania	Germany	03/03/2009

Prezista	CC Pharma	06/10/2009	400 mg	Film-coated tablet	60	EU/1/06/380/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Denmark, Romania, Finland, Sweden, Slovak Republic, Czech Republic	Germany	15/10/2009
Prezista	CC Pharma	06/10/2009	600 mg	Film-coated tablet	60	EU/1/06/380/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	15/10/2009
Prezista	Haemato Pharm AG	05/10/2009	400 mg	Film-coated tablet	60	EU/1/06/380/003	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	15/10/2009
Prezista	Haemato Pharm AG	05/10/2009	600 mg	Film-coated tablet	60	EU/1/06/380/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	15/10/2009
Prezista	Kohlpharma GmbH	08/10/2009	400 mg	Film-coated tablet	60	EU/1/06/380/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/10/2009
Prezista	Kohlpharma GmbH	08/10/2009	600 mg	Film-coated tablet	60	EU/1/06/380/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/10/2009
Prezista	Pharma Westen GmbH	26/03/2009	300 mg	Film-coated tablet	120	EU/1/06/380/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Romania	Germany	15/04/2009
Prezista	Pharma Westen GmbH	09/10/2009	400 mg	Film-coated tablet	60	EU/1/06/380/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	28/10/2009
Pritor	BR Lewis Ltd	17/09/2008	40 mg	Tablet	28	EU/1/98/089/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Romania, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/02/2009

Pritor	BR Lewis Ltd	25/09/2008	80 mg	Tablet	28	EU/1/98/089/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Romania, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	09/02/2009
PritorPlus	BR Lewis Ltd	01/10/2008	80 mg/12.5 mg	Tablet	28	EU/1/02/215/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Romania	United Kingdom, Ireland, Malta	25/11/2009
PritorPlus	Interport Ltd	10/08/2009	80 mg/12.5 mg	Tablet	28	EU/1/02/215/007	Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	28/08/2009
PritorPlus	Paranova Danmark AS	25/02/2009	80 mg/12.5 mg	Tablet	98	EU/1/02/215/010	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	29/04/2009
PritorPlus	Pharma Westen GmbH	07/09/2009	40 mg/12.5 mg	Tablet	98	EU/1/02/215/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/09/2009
PritorPlus	Pharma Westen GmbH	07/09/2009	80 mg/12.5 mg	Tablet	98	EU/1/02/215/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/09/2009
Procoralan	Emra-Med	17/06/2009	5 mg	Film-coated tablet	56	EU/1/05/316/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/06/2009
Procoralan	Emra-Med	17/06/2009	5 mg	Film-coated tablet	98	EU/1/05/316/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/06/2009
Procoralan	Kohlpharma GmbH	03/08/2009	5 mg	Film-coated tablet	98	EU/1/05/316/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, Sweden, United Kingdom	Germany	28/08/2009
Procoralan	MTK Pharma GmbH	27/10/2009	7.5 mg	Film-coated tablet	28	EU/1/05/316/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/12/2009

Procoralan	MTK Pharma GmbH	27/10/2009	7.5 mg	Film-coated tablet	56	EU/1/05/316/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/12/2009
Procoralan	MTK Pharma GmbH	19/11/2009	7.5 mg	Film-coated tablet	98	EU/1/05/316/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/12/2009
Prometax	Kohlpharma GmbH	10/02/2009	1.5 mg	Capsule, hard	56	EU/1/98/092/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/03/2009
Prometax	Kohlpharma GmbH	10/02/2009	1.5 mg	Capsule, hard	112	EU/1/98/092/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/03/2009
Prometax	Kohlpharma GmbH	10/02/2009	3 mg	Capsule, hard	56	EU/1/98/092/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/03/2009
Prometax	Kohlpharma GmbH	10/02/2009	3 mg	Capsule, hard	112	EU/1/98/092/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	25/03/2009
Prometax	Kohlpharma GmbH	02/03/2009	4.5 mg	Capsule, hard	56	EU/1/98/092/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/03/2009
Prometax	Kohlpharma GmbH	02/03/2009	4.5 mg	Capsule, hard	112	EU/1/98/092/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/03/2009
Prometax	Kohlpharma GmbH	16/03/2009	6 mg	Capsule, hard	112	EU/1/98/092/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Sweden, Spain, The Netherlands, United Kingdom	Germany	25/03/2009

Prometax	Paranova Danmark AS	10/09/2009	1.5 mg	Capsule, hard	56	EU/1/98/092/002	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/09/2009
Prometax	Paranova Danmark AS	21/09/2009	4.5 mg	Capsule, hard	112	EU/1/98/092/009	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	27/11/2009
Protelos	Pharmachim AB	14/01/2009	2 g	Granules for oral suspension	28 sachets	EU/1/04/288/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	04/02/2009
Protelos	Pharmachim AB	14/01/2009	2 g	Granules for oral suspension	84 sachets	EU/1/04/288/005	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	04/02/2009
Protopic	Kohlpharma GmbH	19/05/2009	0.1%	Ointment	1 tube	EU/1/02/201/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	07/07/2009
Protopic	Paranova Danmark AS	01/07/2009	0.1%	Ointment	1 tube	EU/1/02/201/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/09/2009
Protopic	Pharma Westen GmbH	17/04/2009	0.1%	Ointment	1 tube 30 g	EU/1/02/201/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/07/2009
Puregon	Beragena Arzneimittel GmbH	22/12/2008	300 IU/0.36 ml	Solution for injection	1 cartridge + 2 packs with 3 pen needles	EU/1/96/008/038	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	18/05/2009
Puregon	Beragena Arzneimittel GmbH	22/12/2008	600 IU/0.72 ml	Solution for injection	1 cartridge + 2 packs with 3 pen needles	EU/1/96/008/039	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	18/05/2009
Puregon	Beragena Arzneimittel GmbH	22/12/2008	900 IU/1.08 ml	Solution for injection	1 cartridge + 3 packs with 3 pen needles	EU/1/96/008/041	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	18/05/2009
Puregon	Cross Pharma AB	20/11/2008	900 IU/1.08 ml	Solution for injection in a cartridge	1+3+3		Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	09/01/2009

Puregon	EuroPharma DK	04/06/2009	900 IU/1.08 ml	Solution for injection	1 cartridge + 3 packs with 3 pen needles	EU/1/96/008/041	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	10/07/2009
Puregon	Pharma Gerke GmbH	25/03/2009	300 IU/0.36 ml	Solution for injection	1 cartridge + 2 packs with 3 pen needles	EU/1/96/008/038	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	02/04/2009
Puregon	Pharma Westen GmbH	21/11/2008	300 IU/0.36 ml	Solution for injection in a cartridge	1+2+3		Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/01/2009
Rapamune	Abacus Medicine A/S	22/05/2009	1 mg	Coated tablet	30	EU/1/01/171/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/07/2009
Rapamune	Abacus Medicine A/S	22/05/2009	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/07/2009
Rapamune	Abacus Medicine A/S	22/05/2009	2 mg	Coated tablet	30	EU/1/01/171/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	22/07/2009
Rapamune	Abacus Medicine A/S	22/05/2009	2 mg	Coated tablet	100	EU/1/01/171/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	22/07/2009
Rapamune	Beragena Arzneimittel GmbH	16/10/2009	2 mg	Coated tablet	100	EU/1/01/171/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	05/11/2009
Rapamune	Mevita HandelsGmbH	27/04/2009	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	12/05/2009
Rapamune	Mevita HandelsGmbH	27/04/2009	2 mg	Coated tablet	100	EU/1/01/171/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	12/05/2009
Rapamune	r.p. pharma GmbH	30/04/2008	1mg	Coated tablets	100		Austria, France, Greece, Italy, Norway, Portugal, Spain	Germany	12/01/2009

Rapamune	Veron Pharma	26/01/2009	1 mg	Coated tablet	30	EU/1/01/171/007	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Germany	19/05/2009
Rapamune	Veron Pharma	26/01/2009	1 mg	Coated tablet	100	EU/1/01/171/008	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Germany	19/05/2009
Rasilez	2 Care 4	03/11/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/12/2009
Rasilez	2 Care 4	03/11/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/12/2009
Rasilez	B&S Healthcare Ltd	12/05/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	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	United Kingdom, Ireland	16/09/2009
Rasilez	Doncaster Pharmaceuticals Group Ltd	04/09/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	16/09/2009
Rasilez	Doncaster Pharmaceuticals Group Ltd	10/09/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	16/09/2009
Rasilez	Emra-Med	07/09/2009	300 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/09/2009
Rasilez	Kohlpharma GmbH	16/03/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Norway, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/03/2009
Rasilez	Kohlpharma GmbH	31/03/2009	150 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/04/2009

Rasilez	Kohlpharma GmbH	20/07/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2009
Rasilez	Kohlpharma GmbH	20/07/2009	300 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/019	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	28/08/2009
Rasilez	O.P.D. Laboratories Ltd	14/08/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom, Ireland	15/09/2009
Rasilez	Orifarm AS	10/08/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/09/2009
Rasilez	Orifarm AS	10/08/2009	150 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/09/2009
Rasilez	Orifarm AS	10/08/2009	300 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/019	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	09/09/2009
Rasilez	Paranova Danmark AS	02/09/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	28/09/2009
Rasilez	Pharma Gerke GmbH	09/12/2009	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez	Pharma Gerke GmbH	09/12/2009	150 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/009	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez	Pharma Gerke GmbH	10/12/2009	150 mg	Film-coated tablet	56	EU/1/07/405/006	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez	Pharma Gerke GmbH	14/12/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009

Rasilez	Pharma Gerke GmbH	14/12/2009	300 mg	Film-coated tablet	56	EU/1/07/405/016	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez	Pharma Gerke GmbH	14/12/2009	300 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/019	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez	Pharma Westen GmbH	04/11/2009	150 mg	Film-coated tablet	98 (2 x 49)	EU/1/07/405/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/11/2009
Rasilez	Samforlab SA	10/11/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Sweden, United Kingdom	Spain	08/12/2009
Rasilez	Swingward Ltd TA Medihealth	14/03/2008	150 mg	Film-coated tablet	28	EU/1/07/405/003	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Slovak Republic, Slovenia, Spain, Sweden	United Kingdom	06/04/2009
Rasilez	Swingward Ltd TA Medihealth	06/04/2009	300 mg	Film-coated tablet	28	EU/1/07/405/013	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	04/06/2009
Rasilez HCT	Dr Fisher Farma BV	07/09/2009	300 mg / 12.5 mg	Film-coated tablet	28	EU/1/08/491/052	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/09/2009
Rasilez HCT	Dr Fisher Farma BV	07/09/2009	300 mg / 25 mg	Film-coated tablet	28	EU/1/08/491/072	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/09/2009
Rasilez HCT	Pharma Gerke GmbH	09/12/2009	300 mg / 12.5 mg	Film-coated tablet	56	EU/1/08/491/055	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	09/12/2009	300 mg / 12.5 mg	Film-coated tablet	98	EU/1/08/491/058	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	09/12/2009	300 mg / 12.5 mg	Film-coated tablet	28	EU/1/08/491/052	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009

Rasilez HCT	Pharma Gerke GmbH	10/12/2009	150 mg / 12.5 mg	Film-coated tablet	28	EU/1/08/491/012	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	10/12/2009	150 mg / 12.5 mg	Film-coated tablet	56	EU/1/08/491/015	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	10/12/2009	150 mg / 12.5 mg	Film-coated tablet	98	EU/1/08/491/018	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	14/12/2009	300 mg / 25 mg	Film-coated tablet	28	EU/1/08/491/072	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	14/12/2009	300 mg / 25 mg	Film-coated tablet	56	EU/1/08/491/075	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rasilez HCT	Pharma Gerke GmbH	14/12/2009	300 mg / 25 mg	Film-coated tablet	98	EU/1/08/491/078	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	16/12/2009
Rebetol	Axicorp Pharma GmbH	13/08/2009	200 mg	Capsule, hard	168 hard	EU/1/99/107/003	Austria, Belgium, Denmark, Estonia, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Germany	17/09/2009
Rebetol	Axicorp Pharma GmbH	13/08/2009	200 mg	Capsule, hard	84 hard	EU/1/99/107/001	Austria, Belgium, Denmark, Estonia, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Germany	17/09/2009
Rebetol	Delfarma SpZoo	16/11/2009	200 mg	Capsule, hard	140 hard	EU/1/99/107/002	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	25/11/2009
Rebetol	Dr Fisher Farma BV	16/12/2008	200 mg	Capsule, hard	84 hard	EU/1/99/107/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	29/01/2009
Rebetol	Medcor Pharmaceuticals BV	16/03/2009	200 mg	Capsule, hard	84 hard	EU/1/99/107/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	14/04/2009

Rebetol	Mevita HandelsGmbH	31/10/2008	200 mg	Capsule, hard	168 hard	EU/1/99/107/003	Belgium, France, Greece, Italy, The Netherlands, Portugal, United Kingdom, Norway, Czech Republic	Germany	26/01/2009
Rebetol	Orifarm AS	14/07/2009	200 mg	Capsule, hard	168 hard	EU/1/99/107/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Lithuania	Denmark	11/08/2009
Rebif	Aaston Healthcare GmbH	23/03/2009	22 µg (6 million IU)	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	21/05/2009
Rebif	Aaston Healthcare GmbH	23/03/2009	44 µg (12 million IU)	Solution for injection	12 pre-filled syringes	EU/1/98/063/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	21/05/2009
Rebif	Haemato Pharm AG	15/01/2009	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Latvia, Romania, Slovak Republic, Germany	Germany, Austria	14/04/2009
Rebif	Haemato Pharm AG	25/08/2009	44 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/006	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Latvia, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, United Kingdom, Germany	Germany, Austria	26/11/2009
Rebif	Inopha GmbH	20/10/2009	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	11/12/2009
Rebif	Mevita HandelsGmbH	04/02/2009	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	27/05/2009
Rebif	Mevita HandelsGmbH	04/02/2009	44 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/006	Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	27/05/2009

Rebif	Pharma Gerke GmbH	29/05/2008	44 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland, Slovak Republic, Romania, Ireland	Germany	24/06/2009
Rebif	Pharma Gerke GmbH	16/06/2008	22 µg	Solution for injection	12 pre-filled syringes	EU/1/98/063/003	Austria, Belgium, Czech Republic, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Slovak Republic, Spain, United Kingdom	Germany	14/10/2009
Remicade	Aaston Healthcare GmbH	26/03/2009	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, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	26/06/2009
Remicade	Abacus Medicine A/S	06/07/2009	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/99/116/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	17/07/2009
Remicade	CC Pharma	11/08/2009	100 mg	Powder for concentrate for solution for infusion	4 vials	EU/1/99/116/004	Austria, Belgium, Czech Republic, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Slovak Republic	Germany	16/10/2009
Remicade	CC Pharma	11/08/2009	100 mg	Powder for concentrate for solution for infusion	5 vials	EU/1/99/116/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic, Slovak Republic	Germany	16/10/2009
Remicade	Haemato Pharm AG	01/06/2009	100 mg	Powder for concentrate for solution for infusion	2 vials	EU/1/99/116/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Germany	Germany, Austria	09/07/2009
Remicade	Haemato Pharm AG	01/06/2009	100 mg	Powder for concentrate for solution for infusion	3 vials	EU/1/99/116/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Germany	Germany, Austria	09/07/2009
Remicade	Haemato Pharm AG	19/10/2009	100 mg	Powder for concentrate for solution for infusion	5 vials	EU/1/99/116/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic, Romania	Germany	11/12/2009

Remicade	Haemato Pharm AG	19/10/2009	100 mg	Powder for concentrate for solution for infusion	4 vials	EU/1/99/116/004	Austria, Belgium, Cyprus, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Spain, Sweden, United Kingdom	Germany	11/12/2009
Remicade	Orifarm AS	02/12/2009	100 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/99/116/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Spain, Sweden, United Kingdom	Denmark	17/12/2009
Remicade	Pharma Gerke GmbH	03/08/2009	100 mg	Powder for concentrate for solution for infusion	4 vials	EU/1/99/116/004	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/10/2009
Remicade	Pharma Gerke GmbH	03/08/2009	100 mg	Powder for concentrate for solution for infusion	5 vials	EU/1/99/116/005	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/10/2009
Remicade	Pharma Westen GmbH	28/09/2009	100 mg	Powder for concentrate for solution for infusion	4 vials	EU/1/99/116/004	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	05/10/2009
Remicade	Pharma Westen GmbH	28/09/2009	100 mg	Powder for concentrate for solution for infusion	5 vials	EU/1/99/116/005	Austria, Belgium, Czech Republic, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/10/2009
Renagel	Combiphar Europe BV	31/10/2008	800 mg	Film-coated tablet	1 bottle of 180 with outer carton	EU/1/99/123/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, United Kingdom	The Netherlands	23/06/2009
Renagel	Cross Pharma AB	28/11/2008	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	04/02/2009
Renagel	Delphi Pharmaceuticals BV	24/07/2009	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	31/07/2009
Renagel	Europharma Sverige AB	14/05/2009	800 mg	Film-coated tablet	1 bottle of 180 with outer carton	EU/1/99/123/008	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	24/06/2009

Renagel	Medartuum AB	24/12/2008	800 mg	Film-coated tablet	1 bottle of 180 with outer carton	EU/1/99/123/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	05/05/2009
Renagel	Milinda Arzneimittel GmbH	28/08/2009	800 mg	Film-coated tablet	1 bottle of 180 with outer carton	EU/1/99/123/008	Ireland, Italy, United Kingdom	Germany	01/12/2009
Renagel	NewNeopharm BV	30/06/2009	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	11/08/2009
Renagel	Orifarm AB	13/05/2009	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	12/06/2009
Renagel	Orifarm AB	01/06/2009	800 mg	Film-coated tablet	1 bottle of 180 with outer carton	EU/1/99/123/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Sweden	12/06/2009
Renagel	Oy Cross Pharma AB	18/02/2009	800 mg	Film-coated tablet	1 bottle of 180 without outer carton	EU/1/99/123/012	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Finland	06/07/2009
Revatio	Beragena Arzneimittel GmbH	02/06/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	12/06/2009
Revatio	BR Pharma International Ltd	27/01/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/02/2009

Revatio	Delfarma SpZoo	08/06/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Latvia, Lithuania, Luxembourg, Malta, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	29/06/2009
Revatio	Docpharm GmbH&CoKGaA	15/12/2008	20mg	Film-coated tablets	90		Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	26/01/2009
Revatio	Haemato Pharm AG	04/09/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	05/10/2009
Revatio	Kohlpharma GmbH	20/07/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	21/07/2009
Revatio	MedicoPharm AG	09/11/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Austria, Belgium, Cyprus, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Denmark	Germany	24/11/2009
Revatio	Milinda Arzneimittel GmbH	04/09/2009	20 mg	Film-coated tablet	90	EU/1/05/318/001	Ireland, United Kingdom	Germany	27/11/2009
Reyataz	Aaston Healthcare GmbH	06/04/2009	150 mg	Capsule, hard	60	EU/1/03/267/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	02/07/2009
Reyataz	Aaston Healthcare GmbH	29/05/2009	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	02/07/2009

Reyataz	Abacus Medicine A/S	22/05/2009	150 mg	Capsule, hard	60 (bottle)	EU/1/03/267/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	29/05/2009
Reyataz	Abacus Medicine A/S	22/05/2009	150 mg	Capsule, hard	60 (blister)	EU/1/03/267/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	29/05/2009
Reyataz	Abacus Medicine A/S	22/05/2009	300 mg	Capsule, hard	30 (bottle)	EU/1/03/267/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	29/05/2009
Reyataz	Abacus Medicine A/S	21/10/2009	200 mg	Capsule, hard, blister	60	EU/1/03/267/006	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Denmark	03/11/2009
Reyataz	Axicorp Pharma GmbH	04/12/2008	300 mg	Capsule, hard	30 (bottle)	EU/1/03/267/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/01/2009
Reyataz	CC Pharma	20/04/2009	300 mg	Capsule, hard	30	EU/1/03/267/010	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Romania, Sweden, Denmark, Latvia	Germany	08/09/2009
Reyataz	Delphi Pharmaceuticals BV	22/07/2009	150 mg	Capsule, hard	60	EU/1/03/267/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	30/07/2009
Reyataz	Emra-Med	24/08/2009	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic, Slovenia, Lithuania, Romania	Germany	14/12/2009

Reyataz	Haemato Pharm AG	25/03/2009	150 mg	Capsule, hard	60	EU/1/03/267/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Norway	Germany	26/05/2009
Reyataz	Haemato Pharm AG	25/03/2009	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, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	26/05/2009
Reyataz	Haemato Pharm AG	24/04/2009	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	06/07/2009
Reyataz	Haemato Pharm AG	02/12/2009	300 mg	Capsule, hard	30 x 3	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	16/12/2009
Reyataz	Kohlpharma GmbH	19/12/2008	300 mg	Capsule, hard	30	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/01/2009
Reyataz	Medcor Pharmaceuticals BV	30/01/2009	300 mg	Capsule, hard	30 (bottle)	EU/1/03/267/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	The Netherlands	17/03/2009
Reyataz	Mevita HandelsGmbH	06/07/2009	150 mg	Capsule, hard	60	EU/1/03/267/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/07/2009
Reyataz	Mevita HandelsGmbH	06/07/2009	200 mg	Capsule, hard	60	EU/1/03/267/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/07/2009
Reyataz	Orifarm AS	18/05/2009	150 mg	Capsule, hard	60 (bottle)	EU/1/03/267/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	04/06/2009

Reyataz	Orifarm AS	18/05/2009	200 mg	Capsule, hard	60 (bottle)	EU/1/03/267/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	17/06/2009
Reyataz	Orifarm AS	19/05/2009	300 mg	Capsule, hard	30 (bottle)	EU/1/03/267/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	17/06/2009
Reyataz	Paranova Danmark AS	17/08/2009	150 mg	Capsule, hard	60	EU/1/03/267/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	20/10/2009
Reyataz	Pharma Westen GmbH	12/01/2009	200 mg	Capsule, hard	60	EU/1/03/267/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/01/2009
Rilutek	Milinda Arzneimittel GmbH	26/08/2009	50 mg	Film-coated tablet	56	EU/1/96/010/001	France, Ireland, United Kingdom	Germany	06/11/2009
Rilutek	PCO Manufacturing	12/05/2009	50 mg	Film-coated tablet	56	EU/1/96/010/001	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	United Kingdom, Ireland	13/10/2009
RoActemra 200 mg/10 ml	CC Pharma	11/08/2009	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Denmark, Sweden	Germany	28/08/2009
RoActemra 400 mg/20 ml	CC Pharma	11/08/2009	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Denmark, Sweden, Hungary	Germany	28/08/2009
RoActemra 80 mg/4 ml	CC Pharma	11/08/2009	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/08/492/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Denmark	Germany	28/08/2009
Sifrol	CC Pharma	06/07/2009	0.35 mg	Tablet	100	EU/1/97/050/012	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Sweden, Denmark	Germany	07/10/2009
Sifrol	Chemilines Ltd	29/07/2009	0.18 mg	Tablet	30	EU/1/97/050/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	03/08/2009
Sifrol	Chemilines Ltd	29/07/2009	0.18 mg	Tablet	100	EU/1/97/050/004	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	United Kingdom, Ireland	03/08/2009
Sifrol	Chemilines Ltd	29/07/2009	0.7 mg	Tablet	30	EU/1/97/050/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	03/08/2009

Sifrol	Chemilines Ltd	29/07/2009	0.7 mg	Tablet	100	EU/1/97/050/006	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	03/08/2009
Sifrol	Euro Registratie Collectief BV	27/01/2009	0.7 mg	Tablet	100	EU/1/97/050/006	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	13/03/2009
Sifrol	Euro Registratie Collectief BV	10/08/2009	0.7 mg	Tablet	30	EU/1/97/050/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	20/10/2009
Sifrol	Lexon (UK) Ltd	13/03/2009	0.18 mg	Tablet	30	EU/1/97/050/003	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	02/04/2009
Sifrol	Lexon (UK) Ltd	13/03/2009	0.7 mg	Tablet	30	EU/1/97/050/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Spain, Sweden, United Kingdom, Liechtenstein, Portugal	United Kingdom, Ireland	02/04/2009
Sifrol	Paranova Danmark AS	02/09/2009	0.18 mg	Tablet	100	EU/1/97/050/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/10/2009
Sifrol	Swingward Ltd TA Medihealth	08/09/2009	0.18 mg	Tablet	100	EU/1/97/050/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	14/10/2009
Sifrol	Swingward Ltd TA Medihealth	08/09/2009	0.7 mg	Tablet	100	EU/1/97/050/006	Austria, Belgium, Denmark, Finland, France, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, Greece, The Netherlands, Cyprus, Germany, Liechtenstein, Malta	United Kingdom	14/10/2009
Sifrol	Waymade PLC	18/06/2009	0.18 mg	Tablet	30	EU/1/97/050/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	06/08/2009
Sifrol	Waymade PLC	18/06/2009	0.35 mg	Tablet	30	EU/1/97/050/011	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	06/08/2009
Sifrol	Waymade PLC	18/06/2009	0.7 mg	Tablet	30	EU/1/97/050/005	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	06/08/2009

Simulect	Orifarm AS	27/10/2009	20 mg	Powder and solvent for solution for injection or infusion	1 vial + 1 ampoule	EU/1/98/084/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	08/12/2009
Somavert	Abacus Medicine A/S	12/06/2009	10 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	06/07/2009
Somavert	Abacus Medicine A/S	12/06/2009	15 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	06/07/2009
Somavert	Abacus Medicine A/S	12/06/2009	20 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	06/07/2009
Somavert	CC Pharma	06/03/2009	10 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Spain, United Kingdom, Portugal	Germany	06/07/2009
Somavert	CC Pharma	06/03/2009	20 mg	Powder and solvent for solution for injection	30 vials + 30 vials	EU/1/02/240/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	06/07/2009
Sprycel	Abacus Medicine A/S	22/05/2009	100 mg	Film-coated tablet	30	EU/1/06/363/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	03/07/2009
Sprycel	Abacus Medicine A/S	22/05/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	03/07/2009

Sprycel	Abacus Medicine A/S	22/05/2009	70 mg	Film-coated tablet	60	EU/1/06/363/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	03/07/2009
Sprycel	CC Pharma	12/02/2009	20 mg	Film-coated tablet	60	EU/1/06/363/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Greece	Germany	22/04/2009
Sprycel	CC Pharma	12/02/2009	70 mg	Film-coated tablet	60	EU/1/06/363/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Greece	Germany	22/04/2009
Sprycel	CC Pharma	06/05/2009	20 mg	Film-coated tablet	56	EU/1/06/363/004	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	14/12/2009
Sprycel	CC Pharma	06/05/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Romania, Lithuania, Bulgaria, Denmark, Finland, Sweden	Germany	14/12/2009
Sprycel	CC Pharma	06/05/2009	70 mg	Film-coated tablet	56	EU/1/06/363/006	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	14/12/2009
Sprycel	Haemato Pharm AG	16/03/2009	50 mg	Film-coated tablet	56	EU/1/06/363/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	Germany, Austria	06/07/2009
Sprycel	Kohlpharma GmbH	14/08/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/09/2009
Sprycel	Kohlpharma GmbH	14/08/2009	70 mg	Film-coated tablet	60	EU/1/06/363/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden	Germany	15/09/2009
Sprycel	Kohlpharma GmbH	27/10/2009	20 mg	Film-coated tablet	60	EU/1/06/363/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/11/2009

Sprycel	Orifarm AB	10/08/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Sweden	05/10/2009
Sprycel	Orifarm AS	10/08/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Lithuania, Latvia, Slovenia, Slovak Republic, Poland	Denmark	15/09/2009
Sprycel	Pharma Gerke GmbH	27/04/2009	50 mg	Film-coated tablet	56	EU/1/06/363/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/05/2009
Sprycel	Pharma Westen GmbH	02/03/2009	50 mg	Film-coated tablet	56	EU/1/06/363/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	31/03/2009
Sprycel	Pharma Westen GmbH	07/09/2009	50 mg	Film-coated tablet	60	EU/1/06/363/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Hungary, Romania, Poland, Bulgaria, Czech Republic, Estonia, Lithuania, Latvia, Slovak Republic, Slovenia	Germany	07/10/2009
Sprycel	Pharma Westen GmbH	28/09/2009	70 mg	Film-coated tablet	60	EU/1/06/363/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Germany	08/12/2009
Stalevo	Docpharm GmbH&CoKGaA	06/05/2009	50 mg / 12.5 mg / 200 mg	Film-coated tablet	30	EU/1/03/260/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, The Netherlands, United Kingdom, Spain	Germany	25/06/2009
Stalevo	Docpharm GmbH&CoKGaA	06/05/2009	50 mg / 12.5 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, The Netherlands, United Kingdom, Spain	Germany	25/06/2009
Stalevo	Docpharm GmbH&CoKGaA	01/06/2009	100 mg / 25 mg / 200 mg	Film-coated tablet	30	EU/1/03/260/006	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	25/06/2009
Stalevo	Docpharm GmbH&CoKGaA	01/06/2009	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, The Netherlands, United Kingdom	Germany	25/06/2009

Stalevo	Dr Fisher Farma BV	07/09/2009	200 mg / 50 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/021	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	16/09/2009
Stalevo	Generfarma, S.L.	13/08/2009	100 mg / 25 mg / 200 mg	Film-coated tablet	100	EU/1/03/260/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Norway, Sweden, United Kingdom	Spain	01/12/2009
Stalevo	Haemato Pharm AG	18/05/2009	150 mg / 37.5 mg / 200 mg	Film-coated tablet	30	EU/1/03/260/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany, Austria	04/06/2009
Stalevo	Kohlpharma GmbH	19/01/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/02/2009
Stalevo	Kohlpharma GmbH	08/12/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/12/2009
Stalevo	Kohlpharma GmbH	08/12/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/12/2009
Starlix	Emra-Med	12/01/2009	120mg	Film-coated tablets	120		Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	21/01/2009
Starlix	Kohlpharma GmbH	21/07/2008	120mg	Film-coated tablets	120		Austria, Belgium, France, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	23/01/2009
Starlix	Kohlpharma GmbH	04/02/2009	60mg	Film-coated tablets	120	EU/1/01/174/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/03/2009

Stocrin	Abacus Medicine A/S	06/05/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	28/05/2009
Stocrin	Beragena Arzneimittel GmbH	08/10/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Latvia	Germany	03/11/2009
Stocrin	Emra-Med	07/09/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	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	16/09/2009
Stocrin	Medartuum AB	18/03/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	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	29/05/2009
Stocrin	Pharma Gerke GmbH	11/05/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, France, Greece, Latvia, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	24/06/2009
Stocrin	Pharma Westen GmbH	20/01/2009	600 mg	Film-coated tablet	30	EU/1/99/111/008	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Romania, Latvia	Germany	03/02/2009
Suboxone	Orifarm AB	17/02/2009	8 mg/2 mg	Sublingual tablet	28	EU/1/06/359/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Italy, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	14/04/2009
Suboxone	Swingward Ltd TA Medihealth	28/09/2009	2 mg/0.5 mg	Sublingual tablet	28	EU/1/06/359/002	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	05/10/2009
Suboxone	Waymade PLC	18/08/2009	2 mg/0.5 mg	Sublingual tablet	28	EU/1/06/359/002	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	09/09/2009

Suboxone	Waymade PLC	18/08/2009	8 mg/2 mg	Sublingual tablet	28	EU/1/06/359/004	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Norway, Portugal, Spain, Sweden, The Netherlands	United Kingdom	09/09/2009
Sustiva	CC Pharma	22/05/2009	600 mg	Film-coated tablet	30	EU/1/99/110/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	05/08/2009
Sustiva	CC Pharma	31/07/2009	600 mg	Film-coated tablet	90 (3 x 30 x 1)	EU/1/99/110/010	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	06/08/2009
Sustiva	Haemato Pharm AG	06/08/2009	600 mg	Film-coated tablet	30	EU/1/99/110/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/08/2009
Sustiva	Haemato Pharm AG	06/08/2009	600 mg	Film-coated tablet	90 (3 x 30 x 1)	EU/1/99/110/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	28/08/2009
Sutent	Abacus Medicine A/S	05/05/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/06/2009
Sutent	Abacus Medicine A/S	05/05/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/06/2009
Sutent	Abacus Medicine A/S	05/05/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	15/06/2009
Sutent	Beragena Arzneimittel GmbH	14/09/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	03/11/2009
Sutent	Beragena Arzneimittel GmbH	14/09/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	03/11/2009
Sutent	Beragena Arzneimittel GmbH	14/09/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	03/11/2009

Sutent	CC Pharma	23/01/2009	25 mg	Capsule, hard	30	EU/1/06/347/002	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Romania, Spain, United Kingdom	Germany	24/03/2009
Sutent	CC Pharma	23/01/2009	50 mg	Capsule, hard	30	EU/1/06/347/003	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom, Lithuania, Romania, Hungary, Bulgaria	Germany	24/03/2009
Sutent	CC Pharma	19/03/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	14/04/2009
Sutent	CC Pharma	19/03/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	14/04/2009
Sutent	CC Pharma	19/03/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	14/04/2009
Sutent	CC Pharma	21/04/2009	12.5 mg	Capsule, hard	30	EU/1/06/347/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic, Romania, Bulgaria	Germany	05/05/2009
Sutent	Combiphar Europe BV	04/06/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	13/11/2009
Sutent	Delfarma SpZoo	28/09/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	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	14/10/2009
Sutent	Delfarma SpZoo	28/09/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	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	14/10/2009
Sutent	Delfarma SpZoo	28/09/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	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	14/10/2009

Sutent	Dr Fisher Farma BV	01/10/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/10/2009
Sutent	Dr Fisher Farma BV	01/10/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/10/2009
Sutent	Dr Fisher Farma BV	01/10/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	12/10/2009
Sutent	Haemato Pharm AG	01/06/2009	12.5 mg	Capsule, hard	30	EU/1/06/347/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Estonia, Latvia, Lithuania, Hungary, Poland, Romania, Bulgaria, Slovenia, Slovak Republic, Germany	Germany, Austria	06/08/2009
Sutent	Haemato Pharm AG	01/06/2009	25 mg	Capsule, hard	30	EU/1/06/347/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Romania, Slovenia, Slovak Republic, Poland, Latvia, Lithuania, Hungary, Estonia, Germany	Germany, Austria	06/08/2009
Sutent	Haemato Pharm AG	01/06/2009	50 mg	Capsule, hard	30	EU/1/06/347/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Estonia, Bulgaria, Romania, Latvia, Lithuania, Poland, Slovenia, Slovak Republic, Hungary, Germany	Germany, Austria	06/08/2009
Sutent	Haemato Pharm AG	12/06/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Norway, The Netherlands, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Latvia, Lithuania, Hungary, Poland, Slovenia, Slovak Republic, Romania, Bulgaria	Germany	06/08/2009

Sutent	Haemato Pharm AG	12/06/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Estonia, Latvia, Lithuania, Hungary, Poland, Slovenia, Slovak Republic, Romania, Bulgaria	Germany	06/08/2009
Sutent	Haemato Pharm AG	12/06/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Czech Republic, Estonia, Latvia, Lithuania, Hungary, Poland, Slovenia, Slovak Republic, Romania, Bulgaria	Germany	06/08/2009
Sutent	Pharma Westen GmbH	06/07/2009	12.5 mg	Capsule, hard	28	EU/1/06/347/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	08/07/2009
Sutent	Pharma Westen GmbH	06/07/2009	25 mg	Capsule, hard	28	EU/1/06/347/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	08/07/2009
Sutent	Pharma Westen GmbH	06/07/2009	50 mg	Capsule, hard	28	EU/1/06/347/006	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	08/07/2009
Sutent	Pharma Westen GmbH	28/09/2009	12.5 mg	Capsule, hard	30	EU/1/06/347/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	19/10/2009
Sutent	Pharma Westen GmbH	28/09/2009	25 mg	Capsule, hard	30	EU/1/06/347/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	19/10/2009
Sutent	Pharma Westen GmbH	28/09/2009	50 mg	Capsule, hard	30	EU/1/06/347/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	19/10/2009
Synagis	CC Pharma	13/10/2008	100 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/99/117/002	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	12/01/2009
Synagis	CC Pharma	13/10/2008	50 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/99/117/001	Norway	Germany	12/01/2009

Synagis	Pharma Westen GmbH	12/01/2009	100 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/99/117/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	29/04/2009
Synflorix	Pharma Westen GmbH	23/07/2009	- -	Suspension for injection	1 pre-filled syringe + 1 needle	EU/1/09/508/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/10/2009
Synflorix	Pharma Westen GmbH	23/07/2009	- -	Suspension for injection	10 pre-filled syringes + 10 needles	EU/1/09/508/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	05/10/2009
Synflorix	Veron Pharma	28/07/2009	- -	Suspension for injection	1 pre-filled syringe	EU/1/09/508/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	13/08/2009
TachoSil	Abacus Medicine A/S	22/05/2009	- -	Medicated sponge	1 sponge of 9.5 x 4.8 cm	EU/1/04/277/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	01/07/2009
TachoSil	Abacus Medicine A/S	22/05/2009	- -	Medicated sponge	2 sponges of 4.8 x 4.8 cm	EU/1/04/277/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	01/07/2009
Tamiflu	A1 Pharmaceuticals	22/06/2009	75 mg	Capsule, hard	10	EU/1/02/222/001	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	20/07/2009
Tamiflu	Abacus Medicine A/S	10/08/2009	75 mg	Capsule, hard	10	EU/1/02/222/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Denmark	15/09/2009

Tamiflu	CC Pharma	09/06/2009	75 mg	Capsule, hard	10	EU/1/02/222/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria	Germany	29/06/2009
Tamiflu	Delphi Pharmaceuticals BV	22/06/2009	75 mg	Capsule, hard	10	EU/1/02/222/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	22/09/2009
Tamiflu	Emra-Med	14/08/2009	75 mg	Capsule, hard	10	EU/1/02/222/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	18/09/2009
Tarceva	Aaston Healthcare GmbH	12/05/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	12/08/2009
Tarceva	Abacus Medicine A/S	05/05/2009	100 mg	Film-coated tablet	30	EU/1/05/311/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/05/2009
Tarceva	Abacus Medicine A/S	05/05/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/05/2009
Tarceva	Abacus Medicine A/S	05/05/2009	25 mg	Film-coated tablet	30	EU/1/05/311/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	27/05/2009
Tarceva	Emra-Med	27/04/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	16/07/2009
Tarceva	Haemato Pharm AG	01/06/2009	100 mg	Film-coated tablet	30	EU/1/05/311/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Germany	Germany, Austria	04/06/2009

Tarceva	Haemato Pharm AG	01/06/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	04/06/2009
Tarceva	Orifarm AS	26/11/2009	100 mg	Film-coated tablet	30	EU/1/05/311/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Spain, Sweden, United Kingdom	Denmark	15/12/2009
Tarceva	Pharma Gerke GmbH	03/11/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	10/12/2009
Tarceva	Pharma Westen GmbH	09/07/2009	150 mg	Film-coated tablet	30	EU/1/05/311/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	20/07/2009
Tasigna	Abacus Medicine A/S	22/05/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Slovak Republic, Romania	Denmark	17/06/2009
Tasigna	Beragena Arzneimittel GmbH	22/10/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	03/12/2009
Tasigna	CC Pharma	18/02/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	United Kingdom, Norway	Germany	15/04/2009
Tasigna	Emra-Med	11/11/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/11/2009
Tasigna	Haemato Pharm AG	10/08/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	15/09/2009

Tasigna	Kohlpharma GmbH	28/10/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/11/2009
Tasigna	Pharma Westen GmbH	15/10/2009	200 mg	Capsule, hard	112	EU/1/07/422/003	Belgium, Austria, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	26/11/2009
Tasmar	Axicorp Pharma GmbH	13/08/2009	100 mg	Film-coated tablet	100	EU/1/97/044/003	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/09/2009
Taxotere	Aaston Healthcare GmbH	01/04/2009	80 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Germany, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	13/05/2009
Taxotere	Cross Healthcare Ltd	23/01/2009	20 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Malta, Ireland	24/03/2009
Taxotere	InPharm Spzoo	03/06/2009	20 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/001	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	28/08/2009
Taxotere	InPharm Spzoo	03/06/2009	80 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/002	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	28/08/2009
Taxotere	Lex ano UAB	14/08/2009	20 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/001	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, 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	Lithuania	22/09/2009

Taxotere	Lex ano UAB	14/08/2009	80 mg	Concentrate and solvent for solution for infusion	1 vial + 1 vial	EU/1/95/002/002	Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, 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	Lithuania	22/09/2009
Temodal	Abacus Medicine A/S	22/05/2009	100 mg	Capsule, hard	5	EU/1/98/096/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	140 mg	Capsule, hard	5	EU/1/98/096/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	180 mg	Capsule, hard	5	EU/1/98/096/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	20 mg	Capsule, hard	20	EU/1/98/096/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	250 mg	Capsule, hard	5	EU/1/98/096/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	5 mg	Capsule, hard	5	EU/1/98/096/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009
Temodal	Abacus Medicine A/S	22/05/2009	5 mg	Capsule, hard	20	EU/1/98/096/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	03/07/2009

Temodal	Axicorp Pharma GmbH	10/09/2009	100 mg	Capsule, hard	5	EU/1/98/096/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/11/2009
Temodal	Axicorp Pharma GmbH	10/09/2009	140 mg	Capsule, hard	5	EU/1/98/096/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/11/2009
Temodal	Axicorp Pharma GmbH	10/09/2009	250 mg	Capsule, hard	5	EU/1/98/096/007	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/11/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	100 mg	Capsule, hard	5	EU/1/98/096/005	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	100 mg	Capsule, hard	20	EU/1/98/096/006	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	140 mg	Capsule, hard	5	EU/1/98/096/009	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	180 mg	Capsule, hard	5	EU/1/98/096/011	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	20 mg	Capsule, hard	20	EU/1/98/096/004	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	250 mg	Capsule, hard	5	EU/1/98/096/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	Beragena Arzneimittel GmbH	14/11/2008	5 mg	Capsule, hard	20	EU/1/98/096/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Spain, United Kingdom, Portugal	Germany	18/02/2009
Temodal	CC Pharma	22/12/2006	100 mg	Capsule, hard	20	EU/1/98/096/006	Greece, Norway	Germany	29/07/2009
Temodal	Combiphar Europe BV	11/06/2009	140 mg	Capsule, hard	5	EU/1/98/096/009	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	10/09/2009
Temodal	Emra-Med	08/12/2008	250 mg	Capsule, hard	100	EU/1/04/279/038	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/01/2009

Temodal	Emra-Med	01/06/2009	140 mg	Capsule, hard	5	EU/1/98/096/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	25/06/2009
Temodal	Euro Registratie Collectief BV	27/01/2009	250mg	Hard capsules	5	EU/1/98/096/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Norway, Portugal, Spain, United Kingdom	The Netherlands	20/02/2009
Temodal	Haemato Pharm AG	12/11/2008	250 mg	Capsule, hard	5	EU/1/98/096/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	09/02/2009
Temodal	Kohlpharma GmbH	23/11/2009	20 mg	Capsule, hard	20	EU/1/98/096/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	08/12/2009
Temodal	Pharma Westen GmbH	11/09/2008	100 mg	Capsule, hard	20	EU/1/98/096/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	03/11/2009
Temodal	Singad Pharma ApS	27/04/2009	100 mg	Capsule, hard	5	EU/1/98/096/005	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	23/06/2009
Torisel	CC Pharma	20/01/2009	25 mg/ml	Concentrate and diluent for solution for infusion	1 vial + 1 vial	EU/1/07/424/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom, Denmark, Finland, Romania, Sweden	Germany	16/10/2009
Torisel	Haemato Pharm AG	30/03/2009	25 mg/ml	Concentrate and diluent for solution for infusion	1 vial + 1 vial	EU/1/07/424/001	Austria, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Belgium	Germany, Austria	06/07/2009
Toviaz	Euro Registratie Collectief BV	17/06/2009	4 mg	Prolonged release tablet	28	EU/1/07/386/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/06/2009
Toviaz	Euro Registratie Collectief BV	17/06/2009	8 mg	Prolonged release tablet	28	EU/1/07/386/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/06/2009

Toviaz	Orifarm AS	14/05/2009	4 mg	Prolonged release tablet	84	EU/1/07/386/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Denmark	20/05/2009
Toviaz	Orifarm AS	14/05/2009	4 mg	Prolonged release tablet	28	EU/1/07/386/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	20/05/2009
Toviaz	Orifarm AS	14/05/2009	8 mg	Prolonged release tablet	84	EU/1/07/386/012	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	20/05/2009
Toviaz	Orifarm AS	14/05/2009	8 mg	Prolonged release tablet	28	EU/1/07/386/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Denmark	20/05/2009
Toviaz	Pharmachim AB	03/08/2009	8 mg	Prolonged-release tablet	28	EU/1/07/386/008	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	16/09/2009
Toviaz	Pharmachim AB	03/08/2009	8 mg	Prolonged-release tablet	84	EU/1/07/386/012	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	16/09/2009
Tractocile	Abacus Medicine A/S	22/05/2009	7.5 mg/ml	Solution for injection	1 vial	EU/1/99/124/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	03/07/2009

Tractocile	Abacus Medicine A/S	22/05/2009	7.5 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/99/124/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	03/07/2009
Travatan	2 Care 4	22/06/2009	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Denmark	17/07/2009
Travatan	2 Care 4	06/07/2009	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Spain, Sweden, United Kingdom	Denmark	17/07/2009
Travatan	Emra-Med	01/12/2008	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	19/01/2009
Travatan	Emra-Med	08/12/2008	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	19/01/2009
Travatan	Lexon (UK) Ltd	09/01/2009	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland	22/01/2009
Travatan	Orifarm AB	15/05/2009	40 µg/ml	Eye drops, solution	1 bottle	EU/1/01/199/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Poland, Hungary, Czech Republic	Sweden	09/06/2009
Travatan	Pharma Westen GmbH	18/02/2009	40 µg/ml	Eye drops, solution	3 bottles	EU/1/01/199/002	Austria, Belgium, Denmark, France, Finland, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Poland	Germany	14/04/2009

Trizivir	Aaston Healthcare GmbH	27/03/2009	300 mg/150 mg/300 mg	Film-coated tablet	60 (blister)	EU/1/00/156/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom, Sweden, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Germany	30/04/2009
Trizivir	Haemato Pharm AG	25/03/2009	- -	Film-coated tablet	60	EU/1/00/156/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Norway, Spain, Sweden, The Netherlands, United Kingdom	Germany, Austria	14/05/2009
Trizivir	Medicopharm AG	31/10/2008	- -	Film-coated tablet	60 (bottle)	EU/1/00/156/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Italy, Ireland, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Luxembourg	Germany	25/11/2009
Trizivir	SIA 'Elpis'	31/07/2009	300mg/150mg /300mg	Film-coated tablet	60	EU/1/00/156/003	Austria, Belgium, Bulgaria, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	14/09/2009
Truvada	Aaston Healthcare GmbH	27/03/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/06/2009
Truvada	CC Pharma	05/02/2009	200 mg/ 245 mg	Film-coated tablet	3 x 30	EU/1/04/305/002	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, United Kingdom, Norway, The Netherlands, Czech Republic, Denmark, Finland, Sweden	Germany	02/03/2009
Truvada	Delphi Pharmaceuticals BV	22/07/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Portugal, Spain, United Kingdom	The Netherlands	30/07/2009
Truvada	Haemato Pharm AG	02/03/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Germany	Germany, Austria	11/03/2009
Truvada	Haemato Pharm AG	29/06/2009	200 mg/ 245 mg	Film-coated tablet	3 x 30	EU/1/04/305/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	15/07/2009

Truvada	NewNeopharm BV	30/06/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	11/08/2009
Truvada	Orifarm AS	14/04/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Bulgaria, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Denmark	07/05/2009
Truvada	Parallell Pharma AB	06/04/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, The Netherlands, United Kingdom	Sweden	14/05/2009
Truvada	Pharma Westen GmbH	21/04/2009	200 mg/ 245 mg	Film-coated tablet	3 x 30	EU/1/04/305/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/05/2009
Truvada	SIA 'Elpis'	10/02/2009	200 mg/ 245 mg	Film-coated tablet	30	EU/1/04/305/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	31/03/2009
Twinrix Adult	ACA Müller ADAG Pharma AG	17/10/2006	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Norway, Estonia, Lithuania, Latvia	Germany	05/05/2009
Twinrix Adult	Kohlpharma GmbH	09/06/2009	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Poland	Germany	22/07/2009
Twinrix Adult	Kohlpharma GmbH	09/06/2009	- -	Suspension for injection	10 syringes with separate needle	EU/1/96/020/008	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	22/07/2009

Twinrix Adult	Kohlpharma GmbH	09/06/2009	- -	Suspension for injection	10 pre-filled syringes	EU/1/96/020/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	22/07/2009
Twinrix Adult	Medcor Pharmaceuticals BV	28/05/2009	- -	Suspension for injection	1 syringe with separate needle	EU/1/96/020/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic	The Netherlands	13/07/2009
Tysabri	Aaston Healthcare GmbH	20/03/2009	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	12/06/2009
Tysabri	Haemato Pharm AG	16/03/2009	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	23/04/2009
Tysabri	Kohlpharma GmbH	21/07/2009	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/09/2009
Tysabri	MedicoPharm AG	30/09/2009	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	12/11/2009
Tysabri	Pharma Westen GmbH	27/03/2009	300 mg	Concentrate for solution for infusion	1 vial	EU/1/06/346/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	09/06/2009
Tyverb	Aaston Healthcare GmbH	23/04/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	19/05/2009
Tyverb	Beragena Arzneimittel GmbH	08/10/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/11/2009
Tyverb	CC Pharma	16/01/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Norway, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, Bulgaria, Czech Republic	Germany	28/01/2009

Tyverb	Parallell Pharma AB	08/05/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, The Netherlands, United Kingdom	Sweden	30/06/2009
Tyverb	Pharma Gerke GmbH	02/04/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Romania	Germany	28/05/2009
Tyverb	Pharma Westen GmbH	12/12/2008	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/01/2009
Tyverb	Singad Pharma ApS	08/05/2009	250 mg	Film-coated tablet	70	EU/1/07/440/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	11/12/2009
Vaniqa	PCO Manufacturing	08/06/2009	11.5%	Cream	1 tube	EU/1/01/173/002	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Italy	Ireland, United Kingdom	06/08/2009
Vectibix	CC Pharma	13/01/2009	20 mg/ml	Concentrate for solution for infusion	1 vial - 5 ml	EU/1/07/423/001	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	21/01/2009
Vectibix	CC Pharma	22/05/2009	20 mg/ml	Concentrate for solution for infusion	1 vial - 20 ml	EU/1/07/423/003	Norway, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	25/06/2009
Vectibix	Haemato Pharm AG	15/01/2009	20 mg/ml	Concentrate for solution for infusion	1 vial	EU/1/07/423/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	19/02/2009
VELCADE	Aaston Healthcare GmbH	01/04/2009	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	14/05/2009
VELCADE	Abacus Medicine A/S	22/05/2009	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/06/2009

VELCADE	CC Pharma	23/01/2009	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	Norway, Poland, United Kingdom, Romania, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, Slovak Republic, Denmark, Finland, Sweden	Germany	24/03/2009
VELCADE	Emra-Med	11/06/2009	3.5 mg	Powder for solution for injection	1 vial	EU/1/04/274/001	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	Germany	28/09/2009
VELCADE	Haemato Pharm AG	09/07/2009	1.0 mg	Powder for solution for injection	1 vial	EU/1/04/274/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	20/07/2009
VELCADE	Inopha GmbH	29/12/2008	3.5 mg	Powder for solution for injection	1 vial (10 ml)	EU/1/04/274/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia, Germany	Germany, Austria	06/03/2009
Vfend	2 Care 4	03/09/2008	200 mg	Tablet	30	EU/1/02/212/018	The Netherlands, Austria, Belgium, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden	Denmark	27/01/2009
Vfend	2 Care 4	07/09/2009	50 mg	Tablet	30	EU/1/02/212/006	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	06/10/2009
Vfend	Axicorp Pharma GmbH	06/11/2009	200 mg	Tablet	100	EU/1/02/212/021	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Germany	24/11/2009
Vfend	Beragena Arzneimittel GmbH	07/07/2009	200 mg	Tablet	100	EU/1/02/212/021	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	03/09/2009
Vfend	CC Pharma	23/06/2009	200 mg	Powder for solution for infusion	1 vial	EU/1/02/212/025	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	02/09/2009

Vfend	CC Pharma	22/07/2009	40 mg/ml	Powder for oral suspension	1 bottle	EU/1/02/212/026	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	22/09/2009
Vfend	EuroPharma DK	10/06/2009	200 mg	Tablet	30	EU/1/02/212/018	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	30/06/2009
Vfend	Mevita HandelsGmbH	03/03/2009	200 mg	Tablet	30	EU/1/02/212/018	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	11/05/2009
Vfend	Orifarm AS	26/06/2009	200 mg	Powder for solution for infusion	1 vial	EU/1/02/212/025	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Romania	Denmark	04/08/2009
Vfend	Orifarm AS	13/07/2009	200 mg	Tablet	30	EU/1/02/212/018	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/08/2009
Vfend	Orifarm AS	13/07/2009	50 mg	Tablet	30	EU/1/02/212/006	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	05/08/2009
Vfend	Paranova Danmark AS	29/04/2009	200 mg	Tablet	28	EU/1/02/212/017	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	13/05/2009
Vfend	SIA 'Elpis'	14/01/2009	200 mg	Tablet	14	EU/1/02/212/015	Austria, Belgium, Bulgaria, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	24/03/2009
Vfend	SIA 'Elpis'	10/02/2009	200 mg	Powder for solution for infusion	1 vial	EU/1/02/212/025	Austria, Belgium, Czech Republic, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Lithuania, Luxembourg, The Netherlands, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom, Bulgaria, Estonia, Liechtenstein, Malta	Latvia	16/04/2009

Viagra	2 Care 4	14/10/2008	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Germany, Greece, Italy, The Netherlands, Spain, United Kingdom	Denmark	15/09/2009
Viagra	2 Care 4	14/10/2008	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Germany, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Denmark	15/09/2009
Viagra	2 Care 4	12/11/2008	25 mg	Film-coated tablet	12	EU/1/98/077/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	12/06/2009
Viagra	2 Care 4	12/11/2008	50 mg	Film-coated tablet	12	EU/1/98/077/008	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/11/2009
Viagra	2 Care 4	24/11/2008	25 mg	Film-coated tablet	4	EU/1/98/077/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/11/2009
Viagra	2 Care 4	11/12/2008	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/11/2009
Viagra	Abacus Medicine A/S	15/08/2008	25 mg	Film-coated tablet	8	EU/1/98/077/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/01/2009
Viagra	Abacus Medicine A/S	15/08/2008	25 mg	Film-coated tablet	12	EU/1/98/077/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	09/01/2009
Viagra	Abis-Pharma	14/08/2009	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/10/2009
Viagra	Abis-Pharma	14/08/2009	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Poland, Portugal, Slovak Republic, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/10/2009

Viagra	AN Pharmacy Sp. z o.o.	21/07/2009	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, Spain, Sweden, The Netherlands, United Kingdom	Slovenia	08/09/2009
Viagra	AN Pharmacy Sp. z o.o.	02/09/2009	100 mg	Film-coated tablet	8	EU/1/98/077/011	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	15/09/2009
Viagra	AN Pharmacy Sp. z o.o.	02/09/2009	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, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Poland	15/09/2009
Viagra	Beragena Arzneimittel GmbH	07/04/2008	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, France, Greece, Italy, Portugal, Spain, United Kingdom, The Netherlands	Germany	14/04/2009
Viagra	Beragena Arzneimittel GmbH	07/04/2008	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	14/04/2009
Viagra	Ichem SpZoo	20/10/2008	100mg	Film-coated tablets	4		Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	20/01/2009
Viagra	Ichem SpZoo	20/10/2008	100mg	Film-coated tablets	8		Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Poland	20/01/2009

Viagra	Kohlpharma GmbH	16/03/2009	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	27/03/2009
Viagra	PCO Manufacturing	07/05/2009	50 mg	Film-coated tablet	4	EU/1/98/077/006	Austria, Belgium, Cyprus, Denmark, France, Germany, Greece, Luxembourg, Malta, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Italy	Ireland, United Kingdom	06/07/2009
Viagra	Singad Pharma ApS	11/05/2009	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	16/09/2009
Viagra	Singad Pharma ApS	11/05/2009	100 mg	Film-coated tablet	8	EU/1/98/077/011	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	16/09/2009
Viagra	Veron Pharma	05/03/2009	100 mg	Film-coated tablet	4	EU/1/98/077/010	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Germany	16/06/2009
Viagra	Veron Pharma	05/03/2009	100 mg	Film-coated tablet	12	EU/1/98/077/012	Austria, Belgium, France, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Germany	16/06/2009
Vidaza	CC Pharma	28/10/2009	25 mg/ml	Powder for suspension for injection	1 vial	EU/1/08/488/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	04/11/2009
Vidaza	Haemato Pharm AG	25/11/2009	25 mg/ml	Powder for suspension for injection	1 vial	EU/1/08/488/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovenia, Slovak Republic, Germany	Germany, Austria	16/12/2009
Vimpat	CC Pharma	25/08/2009	10 mg/ml	Solution for infusion	1 vial	EU/1/08/470/016	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	100 mg	Film-coated tablet	14	EU/1/08/470/004	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009

Vimpat	CC Pharma	25/08/2009	15 mg/ml	Syrup	1 bottle	EU/1/08/470/014	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	150 mg	Film-coated tablet	56	EU/1/08/470/008	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	200 mg	Film-coated tablet	56	EU/1/08/470/011	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	50 mg	Film-coated tablet	56	EU/1/08/470/002	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	CC Pharma	25/08/2009	50 mg	Film-coated tablet	14	EU/1/08/470/001	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	15/09/2009
Vimpat	Pharma Westen GmbH	08/09/2009	100 mg	Film-coated tablet	56	EU/1/08/470/005	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/12/2009
Vimpat	Pharma Westen GmbH	08/09/2009	100 mg	Film-coated tablet	168	EU/1/08/470/006	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/12/2009
ViraferonPeg	Orifarm AS	09/07/2009	100 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/132/040	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	02/09/2009
ViraferonPeg	Orifarm AS	09/07/2009	150 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/132/048	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	13/08/2009
ViraferonPeg	Orifarm AS	09/07/2009	80 micrograms	Powder and solvent for solution for injection in pre-filled pen	4 pens + 4 injection needles + 8 cleansing swabs	EU/1/00/132/036	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Ireland, Iceland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	13/08/2009
Viramune	Milinda Arzneimittel GmbH	21/08/2009	200 mg	Tablet	60	EU/1/97/055/001	France	Germany	25/11/2009
Viramune	Milinda Arzneimittel GmbH	21/08/2009	200 mg	Tablet	120	EU/1/97/055/003	France	Germany	25/11/2009

Viramune	Pharma Gerke GmbH	18/01/2008	50mg/5ml	Oral suspension	240ml		Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	29/01/2009
Viramune	Pharma Gerke GmbH	09/10/2009	200 mg	Tablet	120	EU/1/97/055/003	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Romania, Spain, United Kingdom	Germany	03/11/2009
Viread	Aaston Healthcare GmbH	29/05/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	30/06/2009
Viread	Beragena Arzneimittel GmbH	15/06/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/07/2009
Viread	Beragena Arzneimittel GmbH	15/06/2009	245 mg	Film-coated tablet	3 x 30	EU/1/01/200/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	23/07/2009
Viread	CC Pharma	10/02/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, France, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Greece, Romania, Czech Republic, Denmark, Sweden, Finland, Bulgaria	Germany	25/02/2009
Viread	CC Pharma	05/05/2009	245 mg	Film-coated tablet	3 x 30	EU/1/01/200/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic, Sweden, Finland, Bulgaria	Germany	06/07/2009
Viread	Haemato Pharm AG	29/06/2009	245 mg	Film-coated tablet	3 x 30	EU/1/01/200/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Latvia, Poland, Romania, Slovak Republic, Slovenia	Germany	09/07/2009
Viread	Kohlpharma GmbH	12/05/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	16/06/2009
Viread	NewNeopharm BV	30/06/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/08/2009

Viread	Pharma Gerke GmbH	22/05/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom, Czech Republic	Germany	30/11/2009
Viread	Pharma Westen GmbH	19/05/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	29/06/2009
Viread	Pharma Westen GmbH	12/11/2009	245 mg	Film-coated tablet	3 x 30	EU/1/01/200/002	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Romania	Germany	26/11/2009
Viread	SIA 'Elpis'	10/02/2009	245 mg	Film-coated tablet	30	EU/1/01/200/001	Austria, Belgium, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Latvia	06/04/2009
Xagrid	Haemato Pharm AG	06/07/2009	0.5 mg	Capsule, hard	100	EU/1/04/295/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Lithuania, Latvia, Poland, Romania, Slovak Republic, Slovenia	Germany	06/08/2009
Xagrid	Kohlpharma GmbH	08/06/2009	0.5 mg	Capsule, hard	100	EU/1/04/295/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Germany	09/07/2009
Xagrid	Orifarm AS	25/06/2009	0.5 mg	Capsule, hard	100	EU/1/04/295/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	23/07/2009
Xarelto	Euro Registratie Collectief BV	17/06/2009	10 mg	Film-coated tablet	10	EU/1/08/472/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/06/2009
Xarelto	Euro Registratie Collectief BV	17/06/2009	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	26/06/2009

Xarelto	Orifarm AS	22/05/2009	10 mg	Film-coated tablet	30	EU/1/08/472/007	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Poland, Romania	Denmark	30/06/2009
Xarelto	Orifarm AS	22/05/2009	10 mg	Film-coated tablet	100	EU/1/08/472/008	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Poland, Romania	Denmark	30/06/2009
Xeloda	Aaston Healthcare GmbH	01/04/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	05/06/2009
Xeloda	Beragena Arzneimittel GmbH	04/02/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	23/03/2009
Xeloda	Emra-Med	26/01/2009	150mg	Film-coated tablets	60		Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Liechtenstein, Lithuania, Luxembourg, Malta, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	Germany	30/01/2009
Xeloda	Inopha GmbH	29/04/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Germany	13/05/2009
Xeloda	Kohlpharma GmbH	22/05/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Romania	Germany	24/06/2009
Xeloda	NewNeopharm BV	30/06/2009	150 mg	Film-coated tablet	60	EU/1/00/163/001	Austria, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	11/08/2009

Xeloda	NewNeopharm BV	30/06/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	11/08/2009
Xeloda	Pharma Gerke GmbH	24/07/2009	500 mg	Film-coated tablet	120	EU/1/00/163/002	Austria, Belgium, Finland, France, Greece, Italy, Norway, Portugal, Romania, Spain, The Netherlands, United Kingdom	Germany	08/09/2009
Xenical	B&S Healthcare Ltd	15/12/2008	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, The Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, United Kingdom	United Kingdom, Ireland	18/02/2009
Xenical	BCN Farma SL	22/12/2008	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, The Netherlands, Norway, Portugal, Romania, Slovak Republic, Slovenia, Sweden, United Kingdom	Spain	29/01/2009
Xenical	BR Pharma International Ltd	23/07/2008	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	06/04/2009
Xenical	Generfarma, S.L.	31/07/2009	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Denmark, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Portugal, Sweden, United Kingdom	Spain	06/11/2009
Xenical	Imed Healthcare Ltd	17/11/2008	120 mg	Capsule, hard	84	EU/1/98/071/003	United Kingdom, Belgium, Cyprus, Denmark, Finland, France, Germany, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden	Ireland, United Kingdom, Malta	12/01/2009
Xenical	Mediwin Ltd	06/08/2009	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Denmark, Finland, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Estonia, Latvia, Lithuania, Hungary, Poland, Slovenia, Slovak Republic, Bulgaria, Romania	France	28/08/2009

Xenical	MPT Pharma	27/02/2009	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Liechtenstein, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Ireland, United Kingdom	02/04/2009
Xenical	Pharma Lab	06/07/2009	120 mg	Capsule, hard	84	EU/1/98/071/003	Austria, Belgium, Cyprus, Denmark, Finland, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Sweden, United Kingdom	Spain	14/09/2009
Xeristar	B&S Healthcare Ltd	17/06/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	25/06/2009
Xeristar	Chemilines Ltd	06/01/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	11/12/2009
Xeristar	Doncaster Pharmaceuticals Group Ltd	01/07/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	13/08/2009
Xeristar	Emra-Med	05/12/2008	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	27/01/2009
Xeristar	G-Pharma Ltd	14/08/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	21/09/2009
Xeristar	Interport Ltd	02/12/2008	30mg	Hard gastro-resistant capsules	7		Austria, Belgium, France, Germany, Greece, Italy, The Netherlands, Portugal, Spain	United Kingdom	05/01/2009
Xeristar	Kohlpharma GmbH	19/01/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/01/2009

Xeristar	Kohlpharma GmbH	19/01/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	26/01/2009
Xeristar	Kohlpharma GmbH	10/02/2009	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, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/02/2009
Xeristar	O.P.D. Laboratories Ltd	09/10/2009	30 mg	Gastro-resistant capsule, hard	7	EU/1/04/297/006	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Italy, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom, Ireland	19/10/2009
Xeristar	Orifarm AS	28/09/2009	60 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	22/10/2009
Xeristar	Orifarm AS	28/09/2009	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	22/10/2009
Xeristar	Paranova Danmark AS	01/07/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	26/11/2009
Xeristar	Paranova Danmark AS	01/07/2009	60 mg	Gastro-resistant capsule, hard	98	EU/1/04/297/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	26/11/2009
Xeristar	Pharmacodane Aps	07/09/2009	30 mg	Gastro-resistant capsule, hard	28	EU/1/04/297/001	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	22/10/2009
Xolair	Abacus Medicine A/S	22/05/2009	150 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/05/319/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Czech Republic, Latvia, Lithuania, Poland, Slovak Republic, Slovenia, Romania, Hungary, Bulgaria	Denmark	24/06/2009

Xolair	CC Pharma	06/02/2009	150 mg	Powder and solvent for solution for injection	4 intermediate packs (1 vial + 1 ampoule)	EU/1/05/319/003	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/05/2009
Xolair	CC Pharma	06/02/2009	150 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/05/319/002	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	27/05/2009
Xolair	CC Pharma	15/10/2009	150 mg	Powder and solvent for solution for injection	10 intermediate packs (1 vial + 1 ampoule)	EU/1/05/319/004	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	09/12/2009
Xolair	Pharmachim AB	26/11/2008	150 mg	Powder and solvent for solution for injection	1 vial + 1 ampoule	EU/1/05/319/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom	Sweden	09/01/2009
Yentreve	B&S Healthcare Ltd	05/10/2009	20 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/007	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	28/10/2009
Yentreve	B&S Healthcare Ltd	05/10/2009	40 mg	Gastro-resistant capsule, hard	56	EU/1/04/280/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Ireland, United Kingdom	28/10/2009
Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	20 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/007	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009
Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	20 mg	Gastro-resistant capsule, hard	56	EU/1/04/280/001	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009
Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	20 mg	Gastro-resistant capsule, hard	98	EU/1/04/280/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009
Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	40 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/002	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009

Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	40 mg	Gastro-resistant capsule, hard	56	EU/1/04/280/003	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009
Yentreve	Docpharm GmbH&CoKGaA	19/12/2008	40 mg	Gastro-resistant capsule, hard	98	EU/1/04/280/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/04/2009
Yentreve	Emra-Med	08/05/2009	20 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/11/2009
Yentreve	Emra-Med	08/05/2009	40 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	12/11/2009
Yentreve	Pharma Gerke GmbH	02/06/2008	20 mg	Gastro-resistant capsule, hard	28	EU/1/04/280/007	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/11/2009
Yentreve	Pharma Gerke GmbH	02/06/2008	20 mg	Gastro-resistant capsule, hard	98	EU/1/04/280/008	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/11/2009
Yentreve	Pharmachim AB	08/12/2008	20mg	Hard gastro-resistant capsules	28		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	26/01/2009
Yentreve	Pharmachim AB	08/12/2008	40mg	Hard gastro-resistant capsules	56		Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	26/01/2009
Yondelis	Haemato Pharm AG	12/10/2009	0.25 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/07/417/001	Austria, Belgium, Bulgaria, Cyprus, 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, Czech Republic, Germany	Germany, Austria	20/10/2009
Yondelis	Haemato Pharm AG	12/10/2009	1 mg	Powder for concentrate for solution for infusion	1 vial	EU/1/07/417/002	Austria, Belgium, Bulgaria, Cyprus, 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, Czech Republic, Germany	Germany, Austria	20/10/2009

Zeffix	Axicorp Pharma GmbH	01/06/2009	100 mg	Film-coated tablet	28	EU/1/99/114/001	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	23/07/2009
Zerit	CC Pharma	12/10/2009	40 mg	Capsule, hard	56	EU/1/96/009/008	Austria, Belgium, France, Greece, Italy, Norway, Portugal, Spain, The Netherlands, United Kingdom	Germany	20/10/2009
Ziagen	SIA 'Elpis'	14/01/2009	300mg	Film-coated tablets	60	EU/1/99/112/001	Austria, Belgium, Bulgaria, Czech Republic, Estonia, Finland, France, Germany, Greece, Hungary, Ireland, Italy, Liechtenstein, Lithuania, Luxembourg, Malta, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, The Netherlands, United Kingdom	Latvia	24/03/2009
Zometa	CC Pharma	19/01/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Norway, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	27/01/2009
Zometa	CC Pharma	04/02/2009	4 mg/5 ml	Concentrate for solution for infusion	4	EU/1/01/176/005	Norway, United Kingdom, Austria, Belgium, France, Greece, Italy, The Netherlands, Portugal, Spain	Germany	24/02/2009
Zometa	Dr Fisher Farma BV	05/01/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	27/01/2009
Zometa	Haemato Pharm AG	25/08/2009	4 mg	Concentrate for solution for infusion	4 vials	EU/1/01/176/005	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	13/11/2009
Zometa	Kohlpharma GmbH	20/03/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	04/05/2009
Zometa	Orifarm AB	14/05/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	30/06/2009
Zometa	Orifarm AS	14/05/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Spain, Sweden, United Kingdom, Portugal	Denmark	30/06/2009

Zometa	Pharma Gerke GmbH	22/06/2009	4 mg	Concentrate for solution for infusion	1 vial	EU/1/01/176/004	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	13/07/2009
Zonegran	B&S Healthcare Ltd	14/09/2009	50 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	03/12/2009
Zonegran	B&S Healthcare Ltd	14/09/2009	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	03/12/2009
Zonegran	BR Lewis Ltd	24/02/2009	25 mg	Capsule, hard	14	EU/1/04/307/001	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	United Kingdom, Ireland, Malta	04/05/2009
Zonegran	BR Lewis Ltd	27/02/2009	100 mg	Capsule, hard	56	EU/1/04/307/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	04/05/2009
Zonegran	Chemilines Ltd	12/06/2009	100 mg	Capsule, hard	56	EU/1/04/307/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	27/07/2009
Zonegran	Chemilines Ltd	12/06/2009	25 mg	Capsule, hard	14	EU/1/04/307/001	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	27/07/2009
Zonegran	Chemilines Ltd	12/06/2009	50 mg	Capsule, hard	28	EU/1/04/307/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden	Ireland, United Kingdom	27/07/2009
Zonegran	Europharma Sverige AB	17/02/2009	100 mg	Capsule, hard	98	EU/1/04/307/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	23/03/2009
Zonegran	Europharma Sverige AB	17/02/2009	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	23/03/2009
Zonegran	Europharma Sverige AB	17/02/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	23/03/2009

Zonegran	Interport Ltd	08/10/2009	100 mg	Capsule, hard	56	EU/1/04/307/004	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	30/11/2009
Zonegran	Interport Ltd	08/10/2009	25 mg	Capsule, hard	14	EU/1/04/307/001	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	30/11/2009
Zonegran	Interport Ltd	08/10/2009	50 mg	Capsule, hard	28	EU/1/04/307/009	Austria, Belgium, France, Germany, Greece, Italy, Portugal, Spain	United Kingdom	30/11/2009
Zonegran	Medartuum AB	06/07/2009	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovak Republic, Slovenia	Sweden	05/08/2009
Zonegran	Medartuum AB	06/07/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Bulgaria, Czech Republic, Estonia, Hungary, Latvia, Lithuania, Poland, Romania, Slovenia, Slovak Republic	Sweden	05/08/2009
Zonegran	Orifarm AS	24/06/2009	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	17/07/2009
Zonegran	Orifarm AS	24/06/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	17/07/2009
Zonegran	Paranova Danmark AS	26/08/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	27/11/2009
Zonegran	Paranova Läkemedel AB	09/03/2009	100 mg	Capsule, hard	98	EU/1/04/307/007	Austria, Belgium, Denmark, Finland, France, Germany, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom, Cyprus, Greece, Liechtenstein	Sweden	10/06/2009

Zonegran	Paranova Läkemedel AB	18/06/2009	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	16/07/2009
Zonegran	Pharmachim AB	29/07/2009	100 mg	Capsule, hard	98	EU/1/04/307/007	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom, Czech Republic	Sweden	05/08/2009
Zonegran	Pharmachim AB	29/07/2009	25 mg	Capsule, hard	56	EU/1/04/307/002	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom, Czech Republic	Sweden	05/08/2009
Zonegran	Pharmachim AB	29/07/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, United Kingdom, Czech Republic	Sweden	05/08/2009
Zonegran	Swingward Ltd TA Medihealth	06/04/2009	50 mg	Capsule, hard	56	EU/1/04/307/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	06/07/2009
Zyprexa	2 Care 4	24/11/2008	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, Iceland	Denmark	12/03/2009
Zyprexa	2 Care 4	24/11/2008	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, Iceland	Denmark	12/03/2009
Zyprexa	2 Care 4	24/11/2008	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, Iceland	Denmark	12/03/2009
Zyprexa	2 Care 4	24/11/2008	7.5 mg	Coated tablet	28	EU/1/96/022/011	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, Iceland	Denmark	12/03/2009
Zyprexa	2 Care 4	24/11/2008	7.5 mg	Coated tablet	56	EU/1/96/022/006	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom, Finland, Iceland	Denmark	12/03/2009
Zyprexa	2 Care 4	06/05/2009	2.5 mg	Coated tablet	28	EU/1/96/022/002	Austria, Belgium, France, Germany, Greece, Hungary, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom, Iceland, Finland	Denmark	12/05/2009

Zyprexa	2 Care 4	07/09/2009	15 mg	Coated tablet	28	EU/1/96/022/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	12/11/2009
Zyprexa	2 Care 4	07/09/2009	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	12/11/2009
Zyprexa	A-Pharma ApS	09/06/2009	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	08/09/2009
Zyprexa	A-Pharma ApS	09/06/2009	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	08/09/2009
Zyprexa	A-Pharma ApS	09/06/2009	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	08/09/2009
Zyprexa	A-Pharma ApS	09/06/2009	7.5 mg	Coated tablet	56	EU/1/96/022/006	Austria, Belgium, Finland, France, Germany, Greece, Ireland, Italy, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	08/09/2009
Zyprexa	Axicorp Pharma GmbH	30/06/2009	20 mg	Coated tablet	35	EU/1/96/022/028	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	06/08/2009
Zyprexa	BR Lewis Ltd	02/03/2009	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Ireland, Malta, United Kingdom	11/03/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	10 mg	Coated tablet	35	EU/1/96/022/026	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	10 mg	Coated tablet	70	EU/1/96/022/032	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	15 mg	Coated tablet	35	EU/1/96/022/027	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	15 mg	Coated tablet	70	EU/1/96/022/033	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009

Zyprexa	BR Pharma International Ltd	06/04/2009	2.5 mg	Coated tablet	35	EU/1/96/022/023	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	20 mg	Coated tablet	35	EU/1/96/022/028	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	5 mg	Coated tablet	35	EU/1/96/022/024	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	5 mg	Coated tablet	70	EU/1/96/022/030	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	06/04/2009	7.5 mg	Coated tablet	70	EU/1/96/022/031	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	08/04/2009	2.5 mg	Coated tablet	70	EU/1/96/022/029	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	20/04/2009
Zyprexa	BR Pharma International Ltd	03/12/2009	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/12/2009
Zyprexa	BR Pharma International Ltd	03/12/2009	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/12/2009
Zyprexa	BR Pharma International Ltd	03/12/2009	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/12/2009
Zyprexa	BR Pharma International Ltd	03/12/2009	5 mg	Coated tablet	56	EU/1/96/022/020	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	16/12/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	15 mg	Coated tablet	28	EU/1/96/022/012	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009

Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	15 mg	Coated tablet	35	EU/1/96/022/027	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	15 mg	Coated tablet	56	EU/1/96/022/021	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	15 mg	Coated tablet	70	EU/1/96/022/033	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	20 mg	Coated tablet	35	EU/1/96/022/028	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	20 mg	Coated tablet	56	EU/1/96/022/022	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Docpharm GmbH&CoKGaA	03/08/2009	20 mg	Coated tablet	70	EU/1/96/022/034	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	14/09/2009
Zyprexa	Dr Fisher Farma BV	31/07/2009	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	28/08/2009
Zyprexa	Emra-Med	09/07/2007	10 mg	Coated tablet	35	EU/1/96/022/026	Austria, Belgium, France, Italy, The Netherlands, Portugal, Spain, United Kingdom	Germany	06/07/2009
Zyprexa	Emra-Med	09/07/2007	15 mg	Coated tablet	35	EU/1/96/022/027	The Netherlands, United Kingdom, Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden	Germany	06/07/2009
Zyprexa	Emra-Med	09/07/2007	20 mg	Coated tablet	35	EU/1/96/022/028	The Netherlands, Ireland, Malta, United Kingdom	Germany	06/07/2009

Zyprexa	EuroPharma DK	28/11/2008	15 mg	Coated tablet	28	EU/1/96/022/012	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Denmark	30/04/2009
Zyprexa	EuroPharma DK	28/11/2008	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, The Netherlands, Portugal, Spain, Sweden, United Kingdom	Denmark	30/04/2009
Zyprexa	EuroPharma DK	05/03/2009	2.5 mg	Coated tablet	28	EU/1/96/022/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	31/03/2009
Zyprexa	EuroPharma DK	15/05/2009	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	15/06/2009
Zyprexa	EuroPharma DK	15/05/2009	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	15/06/2009
Zyprexa	EuroPharma DK	15/05/2009	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	16/06/2009
Zyprexa	G-Pharma Ltd	26/05/2009	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden	United Kingdom	30/06/2009
Zyprexa	Imed Healthcare Ltd	17/11/2008	10 mg	Coated tablet	28	EU/1/96/022/009	United Kingdom, Greece, Italy, Belgium, Cyprus, Denmark, Finland, France, Germany, Iceland, Ireland, Luxembourg, Malta, Norway, Portugal, Sweden, The Netherlands	Ireland, Malta, United Kingdom	26/01/2009
Zyprexa	Medcor Pharmaceuticals BV	12/03/2009	10 mg	Coated tablet	28	EU/1/96/022/009	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009
Zyprexa	Medcor Pharmaceuticals BV	12/03/2009	15 mg	Coated tablet	28	EU/1/96/022/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009

Zyprexa	Medcor Pharmaceuticals BV	12/03/2009	2.5 mg	Coated tablet	28	EU/1/96/022/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009
Zyprexa	Medcor Pharmaceuticals BV	12/03/2009	20 mg	Coated tablet	28	EU/1/96/022/014	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009
Zyprexa	Medcor Pharmaceuticals BV	12/03/2009	5 mg	Coated tablet	28	EU/1/96/022/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	07/04/2009
Zyprexa	Orifarm AB	28/09/2009	10 mg	Powder for solution for injection	1 vial	EU/1/96/022/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Sweden	22/10/2009
Zyprexa	Parallell Pharma AB	22/05/2009	10 mg	Coated tablet	56	EU/1/96/022/010	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	30/06/2009
Zyprexa	Pharma Gerke GmbH	21/04/2009	10 mg	Coated tablet	35	EU/1/96/022/026	Austria, Belgium, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	07/05/2009
Zyprexa	Pharma Gerke GmbH	22/04/2009	10 mg	Coated tablet	70	EU/1/96/022/032	Austria, Belgium, France, Greece, Italy, Norway, The Netherlands, Portugal, Spain, United Kingdom	Germany	07/05/2009
Zyprexa	Pharma Gerke GmbH	31/07/2009	7.5 mg	Coated tablet	70	EU/1/96/022/031	Austria, Belgium, Finland, France, Greece, Italy, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	28/08/2009
Zyprexa	Pharma Westen GmbH	02/04/2009	2.5 mg	Coated tablet	70	EU/1/96/022/029	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	17/04/2009
Zyprexa	Pharmacodane Aps	09/03/2009	2.5 mg	Coated tablet	28	EU/1/96/022/002	Austria, Belgium, Cyprus, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	21/04/2009

Zyprexa Velotab	2 Care 4	03/11/2009	10 mg	Orodispersible tablet	28	EU/1/99/125/002	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/12/2009
Zyprexa Velotab	2 Care 4	03/11/2009	15 mg	Orodispersible tablet	28	EU/1/99/125/003	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/12/2009
Zyprexa Velotab	2 Care 4	03/11/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Denmark	10/12/2009
Zyprexa Velotab	Axicorp Pharma GmbH	05/10/2009	15 mg	Orodispersible tablet	35	EU/1/99/125/011	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/10/2009
Zyprexa Velotab	Axicorp Pharma GmbH	05/10/2009	20 mg	Orodispersible tablet	35	EU/1/99/125/012	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/10/2009
Zyprexa Velotab	Axicorp Pharma GmbH	05/10/2009	20 mg	Orodispersible tablet	70	EU/1/99/125/016	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/10/2009
Zyprexa Velotab	Axicorp Pharma GmbH	06/10/2009	15 mg	Orodispersible tablet	70	EU/1/99/125/015	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Luxembourg, Norway, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Germany	20/10/2009
Zyprexa Velotab	BR Lewis Ltd	27/02/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Liechtenstein, Luxembourg, Portugal, Spain, Sweden, The Netherlands, United Kingdom	Malta, United Kingdom, Ireland	11/03/2009
Zyprexa Velotab	BR Pharma International Ltd	18/09/2009	10 mg	Orodispersible tablet	70	EU/1/99/125/014	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/09/2009
Zyprexa Velotab	BR Pharma International Ltd	18/09/2009	20 mg	Orodispersible tablet	70	EU/1/99/125/016	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/09/2009
Zyprexa Velotab	BR Pharma International Ltd	18/09/2009	5 mg	Orodispersible tablet	70	EU/1/99/125/013	Austria, Belgium, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	30/09/2009

Zyprexa Velotab	Docpharm GmbH&CoKGaA	29/05/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/06/2009
Zyprexa Velotab	Docpharm GmbH&CoKGaA	29/05/2009	20 mg	Orodispersible tablet	35	EU/1/99/125/012	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/06/2009
Zyprexa Velotab	Docpharm GmbH&CoKGaA	29/05/2009	20 mg	Orodispersible tablet	56	EU/1/99/125/008	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/06/2009
Zyprexa Velotab	Docpharm GmbH&CoKGaA	29/05/2009	20 mg	Orodispersible tablet	70	EU/1/99/125/016	Austria, Belgium, Denmark, Finland, France, Greece, Ireland, Italy, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, United Kingdom	Germany	09/06/2009
Zyprexa Velotab	Dr Fisher Farma BV	31/07/2009	15 mg	Orodispersible tablet	28	EU/1/99/125/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	28/08/2009
Zyprexa Velotab	Dr Fisher Farma BV	31/07/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	28/08/2009
Zyprexa Velotab	Emra-Med	23/11/2009	20 mg	Orodispersible tablet	35	EU/1/99/125/012	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/12/2009
Zyprexa Velotab	Emra-Med	23/11/2009	20 mg	Orodispersible tablet	70	EU/1/99/125/016	Austria, Belgium, Cyprus, Denmark, Finland, France, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, The Netherlands, Norway, Portugal, Spain, Sweden, United Kingdom	Germany	15/12/2009
Zyprexa Velotab	EuroPharma DK	20/11/2008	15 mg	Orodispersible tablet	28	EU/1/99/125/003	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Sweden	06/05/2009
Zyprexa Velotab	EuroPharma DK	20/11/2008	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Denmark, Finland, France, Germany, Greece, Ireland, Italy, Luxembourg, The Netherlands, Portugal, Spain, United Kingdom	Sweden	06/05/2009

Zyprexa Velotab	Medcor Pharmaceuticals BV	18/03/2009	10 mg	Orodispersible tablet	28	EU/1/99/125/002	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/04/2009
Zyprexa Velotab	Medcor Pharmaceuticals BV	18/03/2009	15 mg	Orodispersible tablet	28	EU/1/99/125/003	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/04/2009
Zyprexa Velotab	Medcor Pharmaceuticals BV	18/03/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Ireland, Italy, Liechtenstein, Luxembourg, Malta, Norway, Portugal, Spain, Sweden, United Kingdom	The Netherlands	17/04/2009
Zyprexa Velotab	Pharmachim AB	14/01/2009	20 mg	Orodispersible tablet	28	EU/1/99/125/004	Austria, Belgium, France, Germany, Greece, Ireland, Italy, Malta, The Netherlands, Portugal, Spain, United Kingdom	Sweden	29/01/2009