



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

9 June 2016
EMA/415297/2016
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: methylphenidate

Procedure no.: PSUSA/00002024/201510



Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Methylphenidate "Sandoz", depottabletter 27 mg	DK/H/2138/001	49666	SANDOZ A/S	DK
Difumenil, 27 mg, tabletki o przedłużonym uwalnianiu	DK/H/2138/001	20921	SANDOZ GMBH	PL
Ritalin LA 20 mg - Kapseln	not available	1-24270	NOVARTIS PHARMA GMBH	AT
Ritalin LA 30 mg - Kapseln	not available	1-24271	NOVARTIS PHARMA GMBH	AT
Ritalin LA 40 mg - Kapseln	not available	1-24272	NOVARTIS PHARMA GMBH	AT
Ritalin 10 mg - Tabletten	not available	1-22028	NOVARTIS PHARMA GMBH (ART57)	AT

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin LA 10 mg - Kapseln	not available	1-31212	NOVARTIS PHARMA GMBH (ART57)	AT
RITALIN TABLETY 10 MG	not available	06/1179/97-C	NOVARTIS, S.R.O. (ART57)	CZ
Ritalin LA 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	77239.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin LA 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67249.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin LA 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67250.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin LA 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67251.00.00	NOVARTIS PHARMA GMBH (ART57)	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin® 10 mg Tabletten	not available	6094573.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin Uno	not available	53335	NOVARTIS HEALTHCARE A/S (ART57)	DK
Ritalin	not available	01870	NOVARTIS HEALTHCARE A/S (ART57)	DK
Ritalin Uno	not available	34355	NOVARTIS HEALTHCARE A/S (ART57)	DK
Ritalin Uno	not available	34356	NOVARTIS HEALTHCARE A/S (ART57)	DK
Ritalin Uno	not available	34357	NOVARTIS HEALTHCARE A/S (ART57)	DK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin Uno	not available	44662	NOVARTIS HEALTHCARE A/S (ART57)	DK
RITALIN 60 mg säädellysti vapauttava kapseli, kova	not available	31914	NOVARTIS FINLAND OY (ART57)	FI
Ritalin 40 mg depotkapsel, hård	not available	23150	NOVARTIS FINLAND OY (ART57)	FI
Ritalin 10 mg säädellysti vapauttava kapseli, kovakova	not available	29462	NOVARTIS FINLAND OY (ART57)	FI
Ritalin 20 mg depotkapsel, hård	not available	23148	NOVARTIS FINLAND OY (ART57)	FI
Ritalin 30 mg säädellysti vapauttava kapseli, kova	not available	23149	NOVARTIS FINLAND OY (ART57)	FI

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RITALINE 10 mg comprimé	not available	3400933929404	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE 10 mg comprimé	not available	3400933942410	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE 10 mg comprimé	not available	3400933942649	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE L.P. 30mg, gélule à libération prolongée	not available	3400936231504	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE L.P. 30mg, gélule à libération prolongée	not available	3400936535015	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE L.P. 30mg, gélule à libération prolongée	not available	3400956497164	NOVARTIS PHARMA S.A.S. (ART57)	FR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritaline L.P. 40 mg, gélule à libération prolongée	not available	3400936231733	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritaline L.P. 40 mg, gélule à libération prolongée	not available	3400936535183	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritaline L.P. 40 mg, gélule à libération prolongée	not available	3400956497225	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritaline LP 10 mg, gélule à libération prolongée	not available	3400941686535	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritaline LP 10 mg, gélule à libération prolongée	not available	3400941686764	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritaline LP 10 mg, gélule à libération prolongée	not available	3400957974572	NOVARTIS PHARMA S.A.S. (ART57)	FR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400936231382	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400936534933	NOVARTIS PHARMA S.A.S. (ART57)	FR
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400956496914	NOVARTIS PHARMA S.A.S. (ART57)	FR
Ritalin LA 60 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/05	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Ritalin 10 mg tabletta	not available	OGYI-T-6954/01	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Ritalin LA 20 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/02	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin LA 30 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/03	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
Ritalin LA 40 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/04	NOVARTIS HUNGÁRIA KFT. PHARMA (ART57)	HU
®RITALIN 10MG TABLETS	not available	PA 13/66/1	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
RITALIN®LA 20MG, PROLONGED-RELEASE CAPSULES	not available	PA 13/66/2	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
RITALIN®LA 30MG, PROLONGED-RELEASE CAPSULES	not available	PA 13/66/3	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
RITALIN®LA 40MG, PROLONGED-RELEASE CAPSULES	not available	PA 13/66/4	NOVARTIS PHARMACEUTICALS UK LIMITED	IE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin Uno 60 mg hylki með breyttan losunarhraða, hart	not available	IS/1/15/020/01	NOVARTIS HEALTHCARE A/S (ART57)	IS
Ritalin 10 mg töflur	not available	640044	NOVARTIS HEALTHCARE A/S (ART57)	IS
Ritalin Uno 10 mg hylki með breyttan losunarhraða, hart	not available	IS/1/10/133/01	NOVARTIS HEALTHCARE A/S (ART57)	IS
Ritalin Uno 20 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/01	NOVARTIS HEALTHCARE A/S (ART57)	IS
Ritalin Uno 30 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/02	NOVARTIS HEALTHCARE A/S (ART57)	IS
Ritalin Uno 40 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/03	NOVARTIS HEALTHCARE A/S (ART57)	IS

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RITALIN 10 mg compresse	not available	035040017	NOVARTIS FARMA S.P.A. (ART57)	IT
RITALIN® 10 MG TABLETES	not available	00-0118	NOVARTIS FINLAND OY (ART57)	LV
RITALIN® 10 MG TABLETS	not available	088/03101	NOVARTIS PHARMACEUTICALS UK LIMITED	MT
Ritalin, tabletten 10 mg	not available	RVG 03957	NOVARTIS PHARMA B.V. (ART57)	NL
Ritalin 60 mg kapsel med modifiert frisetting, hard	not available	13-9867	NOVARTIS NORGE AS (ART57)	NO
Ritalin 10 mg kapsler med modifisert frisetting, harde	not available	09-6501	NOVARTIS NORGE AS (ART57)	NO

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin 20 mg kapsler med modifisert frisetting, harde	not available	04-2876	NOVARTIS NORGE AS (ART57)	NO
Ritalin 30 mg kapsler med modifisert frisetting, harde	not available	05-3197	NOVARTIS NORGE AS (ART57)	NO
Ritalin 40 mg kapsler med modifisert frisetting, harde	not available	05-3198	NOVARTIS NORGE AS (ART57)	NO
RITALIN Novartis	not available	3449	NOVARTIS NORGE AS (ART57)	NO
Ritalina LA 60 mg cápsulas de libertação modificada	not available	5635115	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 60 mg cápsulas de libertação modificada	not available	5635123	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalina LA 20 mg cápsulas de libertação modificada	not available	3761780	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 20 mg cápsulas de libertação modificada	not available	3761889	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 30 mg cápsulas de libertação modificada	not available	3761988	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 30 mg cápsulas de libertação modificada	not available	3762085	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 40 mg cápsulas de libertação modificada	not available	3762184	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT
Ritalina LA 40 mg cápsulas de libertação modificada	not available	3762283	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A. ART57	PT

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RITALIN 60 MG HÅRDA KAPSLAR MED MODIFIERAD FRISÄTTNING	not available	50408	NOVARTIS SVERIGE AB (ART57)	SE
RITALIN 10 MG HÅRDA KAPSLAR MED MODIFIERAD FRISÄTTNING	not available	41556	NOVARTIS SVERIGE AB (ART57)	SE
Ritalin 10 mg tabletter	not available	20606	NOVARTIS SVERIGE AB (ART57)	SE
RITALIN 20 MG HÅRDA KAPSLAR MED MODIFIERAD FRISÄTTNING	not available	20607	NOVARTIS SVERIGE AB (ART57)	SE
RITALIN 30 MG HÅRDA KAPSLAR MED MODIFIERAD FRISÄTTNING	not available	20608	NOVARTIS SVERIGE AB (ART57)	SE
RITALIN 40 MG HÅRDA KAPSLAR MED MODIFIERAD FRISÄTTNING	not available	20609	NOVARTIS SVERIGE AB (ART57)	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin®	not available	PL 00101/0539	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Equasym 30 mg cápsulas duras de liberación modificada	UK/H/0819/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	MA 19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym XL 50 mg modified-release capsules, hard	UK/H/0819/005	PA 1575/1/8	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym Depot	UK/H/0819/003	38536	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym 5mg, Tablets	UK/H/0416/001	PA 1575/1/4	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym Depot, hårde kapsler med modificeret udløsning 10 mg	UK/H/0819/001	38534	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 20 mg	UK/H/0819/002	38535	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym® 10mg, Tablets	UK/H/0416/002	PA 1575/1/5	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym Depot 10 mg hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym® 20mg, Tablets	UK/H/0416/003	PA 1575/1/6	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym Depot 20 mg hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/001	64003.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/002	64004.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 10 mg Tabletten	UK/H/0416/002	RVG 26494	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym Depot 30 mg hylki með breyttan losunarhraða	UK/H/0819/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/003	64005.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	MA 19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Retard 20 mg depotkapseli, kova	UK/H/0819/002	MA 19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 5 mg tablets	UK/H/0416/001	PL 27303/0001	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 10 mg tablets	UK/H/0416/002	PL 27303/0002	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 20 mg tablets	UK/H/0416/003	PL 27303/0003	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 5 mg Tabletten	UK/H/0416/001	51447.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 5 mg töflur	UK/H/0416/001	IS/1/03/036/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/1/1	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym 10 mg Tabletten	UK/H/0416/002	51447.01.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 10 mg töflur	UK/H/0416/002	IS/1/03/036/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 20 mg töflur	UK/H/0416/003	IS/1/03/036/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 20 mg Tabletten	UK/H/0416/003	51447.02.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 5 mg tabletter	UK/H/0416/001	35416	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym 10 mg tabletter	UK/H/0416/002	35417	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PA 1575/1/2	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym 20 mg tabletter	UK/H/0416/003	35418	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 20 mg capsule met gereguleerde afgifte, hard	UK/H/0819/002	RVG 33228	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XL 30 mg capsule met gereguleerde afgifte, hard	UK/H/0819/003	RVG 33229	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PL 27303/0005	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PL 27303/0006	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 10 mg cápsulas duras de liberación modificada	UK/H/0819/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym 20 mg cápsulas duras de liberación modificada	UK/H/0819/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Equasym Depot, hårde kapsler med modificeret udløsning 50 mg	UK/H/0819/005	50038	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Depot, hårde kapsler med modificeret udløsning 60 mg	UK/H/0819/006	50039	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 40 mg	UK/H/0819/004	50037	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot 40 mg kapsler med modificeret frisætning, hårde	UK/H/0819/004	11-8780	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 50 mg kapsler med modificeret frisætning, hårde	UK/H/0819/005	11-8781	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 60 mg kapsler med modificeret frisætning, hårde	UK/H/0819/006	11-8782	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XL 40 mg, capsule met gereguleerde afgifte, hard.	UK/H/0819/004	RVG 111228	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 50 mg, capsule met gereguleerde afgifte, hard.	UK/H/0819/005	RVG 111232	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 60 mg, capsule met gereguleerde afgifte, hard	UK/H/0819/006	RVG 111233	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 40 mg modified-release capsules, hard	UK/H/0819/004	PA 1575/1/7	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 60 mg modified-release capsules, hard	UK/H/0819/006	PA 1575/1/9	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XL 40 mg modified-release capsules, hard	UK/H/0819/04	PL 27303/0007	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 50 mg modified-release capsules, hard	UK/H/0819/005	PL 27303/0008	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 60 mg modified-release capsules, hard	UK/H/0819/006	PL 27303/0009	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Quasym 40 mg cápsulas de libertação modificada	UK/H/0819/004	5550462	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 50 mg cápsulas de libertação modificada	UK/H/0819/005	5550454	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Quasym 60 mg cápsulas de libertação modificada	UK/H/0819/006	5550470	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Equasym® Retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/004	86841.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym® Retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/005	86842.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym® Retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/006	86843.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 40 mg depotkapseli, kova	UK/H/0819/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Retard 50 mg depotkapseli, kova	UK/H/0819/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg depotkapseli, kova	UK/H/0819/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym 40 mg cápsulas duras de liberación modificada	UK/H/0819/004	77163	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 50 mg cápsulas duras de liberación modificada	UK/H/0819/005	77164	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 60 mg cápsulas duras de liberación modificada	UK/H/0819/006	77165	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XR 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889041/M	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889104/M	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg capsule rigide a rilascio modificato	UK/H/0819/003	041889155/M	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
QUASYM 40 mg, gélule à libération modifiée	UK/H/0819/004	585 147-0	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
QUASYM 50 mg, gélule à libération modifiée	UK/H/0819/005	585 149-3	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM 60 mg, gélule à libération modifiée	UK/H/0819/006	585 151-8	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
Methylphenidat "Alternova", tabletter	not available	54924	ALTERNOVA A/S	DK
Methylphenidat "Alternova", tabletter	not available	54926	ALTERNOVA A/S	DK
MEFINAD 5 mg καψάκια ελεγχόμενης αποδέσμευσης, σκληρά	not available	48340/19-6-2013	PROTON PHARMA S.A.	GR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MEFINAD 10 mg καψάκια ελεγχόμενης αποδέσμευσης, σκληρά	not available	48341/19-6-2013	PROTON PHARMA S.A.	GR
MEFINAD 20 mg καψάκια ελεγχόμενης αποδέσμευσης, σκληρά	not available	48342/19-6-2013	PROTON PHARMA S.A.	GR
MEFINAD 30 mg καψάκια ελεγχόμενης αποδέσμευσης, σκληρά	not available	48343/19-6-2013	PROTON PHARMA S.A.	GR
MEFINAD 40 mg καψάκια ελεγχόμενης αποδέσμευσης, σκληρά	not available	48344/19-6-2013	PROTON PHARMA S.A.	GR
Equasym Depot 30 mg hård kapsel med modifierad frisättning	not available	20409	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Depot 10 mg hård kapsel med modifierad frisättning	not available	20407	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 20 mg hård kapsel med modifierad frisättning	not available	20408	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 40 mg kapslar med modifierad frisättning, hårda	not available	47785	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 50 mg kapslar med modifierad frisättning, hårda	not available	47786	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 60 mg kapslar med modifierad frisättning, hårda	not available	47787	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 36 mg Retardtabletten	UK/H/0544/002	BE242697	JANSSEN-CILAG NV	BE
Concerta 54 mg Retardtabletten	UK/H/0544/003	BE242706	JANSSEN-CILAG NV	BE
Concerta® 54 mg depottabletter	UK/H/0544/003	17410	JANSSEN-CILAG OY	FI
Concerta® 36 mg depottabletter	UK/H/0544/002	17409	JANSSEN-CILAG OY	FI
Concerta 27 mg, Retardtabletten	UK/H/0544/004	0162/08120051	JANSSEN-CILAG NV	LU
Concerta 54 mg, Retardtabletten	UK/H/0544/003	2008100012	JANSSEN-CILAG NV	LU

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 36 mg, Retardtabletten	UK/H/0544/002	2008100011	JANSSEN-CILAG NV	LU
Concerta	UK/H/0544/001	42318	JANSSEN-CILAG A/S	DK
Concerta	UK/H/0544/002	42319	JANSSEN-CILAG A/S	DK
Concerta 36 mg depottabletter	UK/H/0544/002	02-1222	JANSSEN-CILAG A/S	NO
Concerta, 18 mg, tabletki o przedłużonym uwalnianiu	UK/H/544/001/R/002	14751	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta 18 mg depottabletter	UK/H/0544/001	02-1221	JANSSEN-CILAG A/S	NO

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 18 mg depottabletter	UK/H/0544/001	18541	JANSSEN-CILAG AB	SE
Concerta 36 mg depottabletter	UK/H/0544/002	18542	JANSSEN-CILAG AB	SE
Concerta 18 mg forðatöflur	UK/H/0544/001	IS/1/02/032/01	JANSSEN-CILAG AB	IS
Concerta, 36 mg, tabletki o przedłużonym uwalnianiu	UK/H/0544/002/R/002	14752	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta 36 mg forðatöflur	UK/H/0544/002	IS/1/02/032/02	JANSSEN-CILAG AB	IS
Concerta 36 mg tablety s predĺženým uvoľňovaním	UK/H/0544/002	78/0127/08-S	JOHNSON & JOHNSON, S.R.O	SK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 18 mg tablety s predĺženým uvoľňovaním	UK/H/0544/001	78/0126/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 54 mg tablety s predĺženým uvoľňovaním	UK/H/0544/003	78/0128/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 54 mg tabletten met verlengde afgifte.	UK/H/0544/003	RVG 28075	JANSSEN-CILAG BV	NL
Concerta 27 mg tabletten met verlengde afgifte.	UK/H/0544/004	RVG 101739	JANSSEN-CILAG BV	NL
Concerta 36 mg tabletten met verlengde afgifte.	UK/H/0544/002	RVG 28074	JANSSEN-CILAG BV	NL
Concerta 18 mg tabletten met verlengde afgifte.	UK/H/0544/001	RVG 28073	JANSSEN-CILAG BV	NL

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 54 mg prolonged-release tablets	UK/H/0544/003	MA 018/02603	JANSSEN-CILAG INTERNATIONAL NV	MT
CONCERTA 18 mg prolonged-release tablets	UK/H/0544/001	MA 018/02601	JANSSEN-CILAG INTERNATIONAL NV	MT
CONCERTA 36 mg prolonged-release tablets	UK/H/0544/002	MA 018/02602	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 27 mg depottabletter	UK/H/0544/004	08-5646	JANSSEN-CILAG A/S	NO
Concerta 27 mg forðatöflur	UK/H/0544/004	IS/1/08/032/01	JANSSEN-CILAG AB	IS
Concerta 27 mg depottabletter	UK/H/0544/004	26554	JANSSEN-CILAG AB	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 54 mg forðatöflur	UK/H/0544/003	IS/1/02/032/03	JANSSEN-CILAG AB	IS
Concerta 27 mg depottabletit	UK/H/0544/004	24631	JANSSEN-CILAG OY	FI
Concerta 54 mg depottabletter	UK/H/0544/003	18543	JANSSEN-CILAG AB	SE
Concerta	UK/H/0544/003	42320	JANSSEN-CILAG A/S	DK
Concerta 54 mg tabletten met verlengde afgifte	UK/H/0544/003	BE242706	JANSSEN-CILAG NV	BE
Concerta 27 mg tabletten met verlengde afgifte	UK/H/0544/004	BE324913	JANSSEN-CILAG NV	BE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta, 18 mg toimeainet prolongeeritult vabastavad tabletid	UK/H/0544/001	583608	UAB JOHNSON & JOHNSON	EE
Concerta 54 mg comprimate cu eliberare prelungită	UK/H/0544/003	5341/2013/01-02	JANSSEN PHARMACEUTICA NV	RO
Concerta 18 mg prolonged release tablets	UK/H/0544/001	20060458	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	BG
CONCERTA 54 mg pailginto atpalaidavimo tabletės	UK/H/544/003	LT/1/08/1152/005-006	UAB JOHNSON & JOHNSON	LT
CONCERTA 54 mg ilgstošās darbības tabletes	UK/H/0544/003	08-0132	UAB JOHNSON & JOHNSON	LV
Concerta, 54 mg toimeainet prolongeeritult vabastavad tabletid	UK/H/0544/003	583808	UAB JOHNSON & JOHNSON	EE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 36 mg pailginto atpalaidavimo tabletės	UK/H/544/002	LT/1/08/1152/003-004	UAB JOHNSON & JOHNSON	LT
CONCERTA 36 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/002	20340	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta, 36 mg toimeainet prolongeeritult vabastavad tabletid	UK/H/0544/002	583708	UAB JOHNSON & JOHNSON	EE
Concerta 36 mg comprimate cu eliberare prelungită	UK/H/0544/002	5340/2013/01-02	JANSSEN PHARMACEUTICA NV	RO
Concerta 36 mg tablety s prodlouženým uvolňováním	UK/H/0544/002	06/408/08-C	JANSSEN-CILAG S.R.O	CZ
CONCERTA 18 mg ilgstošās darbības tabletes	UK/H/0544/001	08-0130	UAB JOHNSON & JOHNSON	LV

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 54 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/003	20336	JANSSEN-CILAG INTERNATIONAL NV	CY
CONCERTA 36 mg ilgstošās darbības tabletes	UK/H/0544/002	08-0131	UAB JOHNSON & JOHNSON	LV
Concerta 18 mg comprimate cu eliberare prelungită	UK/H/0544/001	5339/2013/01-02	JANSSEN PHARMACEUTICA NV	RO
CONCERTA 18 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/001	20339	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 18 mg Retardtabletten	UK/H/0544/001	54861.00.00	JANSSEN-CILAG GMBH	DE
Concerta XL 36 mg prolonged-release tablets.	UK/H/0544/002	PL 00242/0373	JANSSEN-CILAG LIMITED	UK

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 36 mg Retardtabletten	UK/H/0544/002	54861.01.00	JANSSEN-CILAG GMBH	DE
Concerta 54 mg Retardtabletten	UK/H/0544/003	54861.02.00	JANSSEN-CILAG GMBH	DE
Concerta XL 18 mg prolonged-release tablets.	UK/H/0544/001	PL 00242/0372	JANSSEN-CILAG LIMITED	UK
Concerta XL 27 mg prolonged-release tablets.	UK/H/0544/004	PL 00242/0400	JANSSEN-CILAG LIMITED	UK
Concerta 27 mg Retardtabletten	UK/H/544/004	72263.00.00	JANSSEN-CILAG GMBH	DE
CONCERTA LP 36 mg, comprimé à libération prolongée	UK/H/0544/002	34009 361 555 8 9	JANSSEN-CILAG	FR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA LP 54 mg, comprimé à libération prolongée	UK/H/0544/003	34009 361 556 4 0	JANSSEN-CILAG	FR
CONCERTA LP 18 mg, comprimé à libération prolongée	UK/H/0544/001	34009 361 554 1 1	JANSSEN-CILAG	FR
Concerta 18 mg, comprimés à libération prolongée	UK/H/0544/001	2008100010	JANSSEN-CILAG NV	LU
Concerta 36 mg tablete s podaljšanim sproščanjem	UK/H/0544/002	5363-I-2140/12	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
Concerta 18 mg tablete s podaljšanim sproščanjem	UK/H/0544/001	5363-2139/12	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 18 mg Retardtabletten	UK/H/0544/001	1-24812	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 27 mg Retardtabletten	UK/H/0544/004	1-27727	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 36 mg Retardtabletten	UK/H/0544/002	1-24813	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 36 mg comprimidos de liberación prolongada.	UK/H/0544/002	65.170	JANSSEN-CILAG S.A.	ES
Concerta XL 36 mg prolonged-release tablets	UK/H/0544/02	PA 748/49/2	JANSSEN-CILAG LIMITED	IE
Concerta 18 mg tablety s prodlouženým uvolňováním	UK/H/0544/001/E/001	06/407/08-C	JANSSEN-CILAG S.R.O	CZ

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta XL 27 mg prolonged-release tablets	UK/H/0544/004	PA 748/49/4	JANSSEN-CILAG LIMITED	IE
Concerta XL 18 mg prolonged-release tablets	UK/H/0544/01	PA 748/49/1	JANSSEN-CILAG LIMITED	IE
Concerta 18 mg comprimidos de liberación prolongada.	UK/H/0544/001	65.148	JANSSEN-CILAG S.A.	ES
CONCERTA 36 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/002	88895	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
CONCERTA 18 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/001	88894/09/20.1.2010	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
ΙΟΝΟΕΪΤΑ 54 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/003	784	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta XL 54 mg prolonged-release tablets.	UK/H/0544/003	PL 00242/0374	JANSSEN-CILAG LIMITED	UK
CONCERTA 27 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/004	785	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 54 mg comprimidos de liberación prolongada.	UK/H/0544/003	65.171	JANSSEN-CILAG S.A.	ES
Concerta 54 mg Retardtabletten	UK/H/0544/003	1-24814	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 54 mg tablety s prodlouženým uvolňováním	UK/H/0544/003/E/001	06/409/08-C	JANSSEN-CILAG S.R.O	CZ
Concerta XL 54 mg prolonged-release tablets	UK/H/0544/03	PA 748/49/3	JANSSEN-CILAG LIMITED	IE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 54 mg comprimidos de libertação prolongada	UK/H/0544/003	4261186	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
CONCERTA 36 mg comprimidos de libertação prolongada	UK/H/0544/002	4260980	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 54 mg tablete s podaljšanim sproščanjem	UK/H/0544/003	5363-I-2141/12	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
CONCERTA 54 mg comprimidos de libertação prolongada	UK/H/0544/003	4261285	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
CONCERTA 18 mg comprimidos de libertação prolongada	UK/H/0544/001	4260881	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
CONCERTA 27 mg comprimidos de libertação prolongada	UK/H/0544/004	5205307	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 27 mg comprimidos de libertação prolongada	UK/H/0544/004	5205273	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
CONCERTA 36 mg comprimidos de libertação prolongada	UK/H/0544/002	4261087	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Methylphenidate "Hexal", depottabletter 27 mg	DK/H/2139/001	49667	HEXAL A/S	DK
Methylphenidate Alternova 5 mg tabletter	not available	51909	ALTERNOVA A/S	SE
Methylphenidate Alternova 10 mg tabletter	not available	51910	ALTERNOVA A/S	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Methylphenidate Alternova 20 mg tabletter	not available	51911	ALTERNOVA A/S	SE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 40 mg gélules à libération modifiée	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XR 50 mg gélules à libération modifiée	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Methylphenidat "Alternova", tablette	not available	54925	ALTERNOVA A/S	DK
Concerta 18 mg tablete	not available	UP/I-530-09/11-02/141	Johnson & Johnson S.E. d.o.o. - hR	HR
Concerta 36 mg tablete	not available	UP/I-530-09/11-02/142	Johnson & Johnson S.E. d.o.o. - hR	HR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rilatine Modified Release 20mg	not available	BE241534	Novartis Pharma N.V.	BE
Rilatine Modified Release 30mg	not available	BE241543	Novartis Pharma N.V.	BE
Rilatine Modified Release 40mg	not available	BE241552	Novartis Pharma N.V.	BE
Rilatine 10mg	not available	BE051597	Novartis Pharma N.V.	BE
Rilatine Modified Release 10mg	not available	BE426413	Novartis Pharma N.V.	BE
Ritalin Adult 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	77240.00.00	NOVARTIS PHARMA GMBH (ART57)	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin Adult 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67252.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin Adult 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67253.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Ritalin Adult 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67254.00.00	NOVARTIS PHARMA GMBH (ART57)	DE
Rilatine Modified release 10 mg	not available	2014060160	Novartis Pharma N.V.	LU
Rilatine Modified release 20 mg	not available	0010/10/03/0097	Novartis Pharma N.V.	LU
Rilatine Modified release 30 mg	not available	0010/10/03/0098	Novartis Pharma N.V.	LU

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rilatine Modified release 40 mg	not available	0010/10/08/0034	Novartis Pharma N.V.	LU
Rilatine 10 mg	not available	2002096657	Novartis Pharma N.V.	LU
Metilfenidato Sandoz 18 mg comprimidos de liberación prolongada EFG	DK/H/2028	75.518	Sandoz Farmacéutica S.	ES
Metilfenidato Sandoz 27 mg comprimidos de liberación prolongada EFG	DK/H/2138	77.14	Sandoz Farmacéutica S.	ES
Metilfenidato Sandoz 36mg comprimidos de liberación prolongada EFG	DK/H/2028	75.519	Sandoz Farmacéutica S.	ES
Metilfenidato Sandoz 36mg comprimidos de liberación prolongada EFG	DK/H/2028	75.520	Sandoz Farmacéutica S.	ES
QUASYM LP 10 mg gélule à libération modifiée		NL32247	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
QUASYM LP 20 mg gélule à libération modifiée		NL32248	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM LP 30 mg gélule à libération modifiée		NL32249	SHIRE PHARMACEUTICALS IRELAND LIMITED	fr
Medikinet XL	DE/H/2223/001- 005	10-0448,10-0461, 10-0462,10-0463, 10-0464	Medice Arzneimittel Putter GmbH & Co.KG	LV
Medikinet	DE/H/2222/001, 003	10-0447, 10-0460	Medice Arzneimittel Putter GmbH & Co.KG	LV
Medikinet retard 5 mg	DE/H/0690/008	66625.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 20 mg	DE/H/0690/005	54569.01.01	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet retard 10 mg	DE/H/0690/004	54569.00.01	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 30 mg	DE/H/0690/006	62567.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 40 mg	DE/H/0690/007	62568.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 5 mg	not available	66747.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 5 mg		86181.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 20 mg	not available	63891.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 10 mg	not available	63890.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 10 mg		86182.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet adult 20 mg		86183.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 30 mg		65964.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 30 mg		86184.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 40 mg	not available	65965.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 40 mg		86185.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 50 mg		88862.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 60 mg		88863.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medicebran 20 mg comprimido	DE/H/2222/003/DC	5397757	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medicebran 5 mg comprimido	DE/H/2222/001/DC	5397732	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 5 mg	DE/H/2222/001/DC	75791.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet 10 mg	DE/H/2222/002/DC	75792.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet 20 mg	DE/H/2222/003/DC	75793.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 5 mg	DE/H/2223/001/DC	75794.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 10 mg	DE/H/2223/002/DC	75795.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 20 mg	DE/H/2223/003/DC	75796.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet retard 30 mg	DE/H/2223/004/DC	75797.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 40 mg	DE/H/2223/005/DC	75798.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/009/DC	88767.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/010/DC	88768.00.00	Medice Arzneimittel Pütter GmbH & Co.KG	DE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MEDIKINET 10 MG		23836	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 20 MG		23837	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 5 MG		23835	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 10 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		23840	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 20 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		23841	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MEDIKINET 30 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		23842	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 40 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		23843	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 5 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		44810	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 50 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		48667	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
MEDIKINET 60 MG KAPSEL MED MODIFIERAD FRISÄTTNING, HÅRD		48668	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/007	1-26728	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet retard 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/008	1-30056	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/010	135389	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/009	135358	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/005	1-26726	Medice Arzneimittel Pütter GmbH & Co KG	

Product Name (in authorisation country)	MRP / DCP or CP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Medikinet retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/004	1-26725	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/006	1-26727	Medice Arzneimittel Pütter GmbH & Co KG	
Medikinet XL Modified-Release Capsules, hard 5 mg	DE/H/0690/001-010	PL 11243/0010	MEDICE ARZNEIMITTEL PUTTER GMBH & CO KG	UK
Medikinet XL Modified-Release Capsules, hard 50 mg	DE/H/0690/001-010	PL 11243/0011	MEDICE ARZNEIMITTEL PUTTER GMBH & CO KG	UK
Medikinet XL Modified-Release Capsules, hard 60 mg	DE/H/0690/001-010	PL 11243/0012	MEDICE ARZNEIMITTEL PUTTER GMBH & CO KG	UK