14 June 2018
EMA/415385/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): methylphenidate

Procedure No.: PSUSA/00002024/201710

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 18 mg comprimate cu eliberare prelungită	UK/H/0544/001	5339/2013/01-02	JANSSEN PHARMACEUTICA NV	RO
Concerta 18 mg comprimidos de liberación prolongada.	UK/H/0544/001	65.148	JANSSEN-CILAG S.A.	ES
Concerta 18 mg comprimidos de libertação prolongada	UK/H/0544/ 001	4260881	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 18 mg comprimidos de libertação prolongada.	UK/H/0544/ 001	4260782	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 18 mg depottabletit	UK/H/0544/001	17408	JANSSEN-CILAG OY	FI
Concerta 18 mg depottabletter	UK/H/0544/001	02-1221	JANSSEN-CILAG A/S	NO
Concerta 18 mg depottabletter.	UK/H/0544/ 001	18541	JANSSEN-CILAG AB	SE
Concerta 18 mg forðatöflur.	UK/H/0544/001	IS/1/02/032/01	JANSSEN-CILAG AB	IS
Concerta 18 mg ilgstošās darbības tabletes	UK/H/0544/001	08-0130	UAB JOHNSON & JOHNSON	LV
CONCERTA 18 mg pailginto atpalaidavimo tabletės	UK/H/0544/001	LT/1/08/1152/002	UAB JOHNSON & JOHNSON	LT
CONCERTA 18 mg pailginto atpalaidavimo tabletės	UK/H/0544/001	LT/1/08/1152/001	UAB JOHNSON & JOHNSON	LT
Concerta 18 mg prolonged release tablets	UK/H/0544/ 001	20060458	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	BG
Concerta 18 mg prolonged-release tablets	UK/H/0544/001	MA018/02601	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 18 mg Retardtabletten	UK/H/0544/001	1-24812	JANSSEN-CILAG PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP 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	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg Retardtabletten	UK/H/0544/001	54861.00.00	JANSSEN-CILAG GMBH	DE
Concerta 18 mg tablete s podaljšanim sproščanjem	UK/H/0544/ 001	H/08/00411/001	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
Concerta 18 mg tablete s produljenim oslobađanjem	not available	HR-H-078951991	JOHNSON & JOHNSON S.E. D.O.O.	HR
Concerta 18 mg tabletten met verlengde afgifte	UK/H/0544/ 001	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg tabletten met verlengde afgifte	UK/H/0544/ 001	RVG 28073	JANSSEN-CILAG BV	NL
Concerta 18 mg tablety s predĺženým uvoľňovaním	UK/H/0544/001	78/0126/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 18 mg tablety s prodlouženým uvolňováním	UK/H/0544/001	06/407/08-C	JANSSEN-CILAG S.R.O	CZ
CONCERTA 18 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/001	12582	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 18 mg δισκία παρατεταμένης αποδέσμευσης.	UK/H/0544/001	20339	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 18 mg, comprimés à libération prolongée	UK/H/0544/ 001	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg, comprimés à libération prolongée	UK/H/0544/ 001	2008100010	JANSSEN-CILAG NV	LU
Concerta 18 mg,	UK/H/0544/ 001	2008100010	JANSSEN-CILAG NV	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Retardtabletten				
Concerta 27 mg comprimidos de liberación prolongada.	UK/H/0544/004	69.988	JANSSEN-CILAG S.A.	ES
Concerta 27 mg comprimidos de libertação prolongada.	UK/H/0544/ 004	5205307	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 27 mg comprimidos de libertação prolongada.	UK/H/0544/ 004	5205273	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 27 mg depottabletit	UK/H/0544/004	24631	JANSSEN-CILAG OY	FI
Concerta 27 mg depottabletter	UK/H/0544/004	08-5646	JANSSEN-CILAG A/S	NO
Concerta 27 mg depottabletter.	UK/H/0544/ 004	26554	JANSSEN-CILAG AB	SE
Concerta 27 mg forðatöflur.	UK/H/0544/004	IS/1/08/032/01	JANSSEN-CILAG AB	IS
Concerta 27 mg Retardtabletten	UK/H/0544/004	1-27727	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 27 mg Retardtabletten	UK/H/0544/ 004	BE324913	JANSSEN-CILAG NV	BE
Concerta 27 mg Retardtabletten	UK/H/0544/004	72263.00.00	JANSSEN-CILAG GMBH	DE
Concerta 27 mg tabletten met verlengde afgifte	UK/H/0544/ 004	BE324913	JANSSEN-CILAG NV	BE
Concerta 27 mg tabletten met verlengde afgifte.	UK/H/0544/ 004	RVG 101739	JANSSEN-CILAG BV	NL
Concerta 27 mg δισκία παρατεταμένης αποδέσμευσης.	UK/H/0544/004	12585/19.02.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 27 mg, comprimés à libération	UK/H/0544/ 004	BE324913	JANSSEN-CILAG NV	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
Concerta 27 mg, comprimés à libération prolongée	UK/H/0544/ 004	2008120051	JANSSEN-CILAG NV	LU
Concerta 27 mg, Retardtabletten	UK/H/0544/ 004	2008120051	JANSSEN-CILAG NV	LU
Concerta 36 mg comprimate cu eliberare prelungită	UK/H/0544/002	5340/2013/02	JANSSEN PHARMACEUTICA NV	RO
Concerta 36 mg comprimate cu eliberare prelungită	UK/H/0544/002	5340/2013/01	JANSSEN PHARMACEUTICA NV	RO
Concerta 36 mg comprimidos de liberación prolongada.	UK/H/0544/002	65.170	JANSSEN-CILAG S.A.	ES
Concerta 36 mg comprimidos de libertação prolongada	UK/H/0544/ 002	4260980	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 36 mg comprimidos de libertação prolongada	UK/H/0544/ 002	4261087	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 36 mg depottabletit	UK/H/0544/002	17409	JANSSEN-CILAG OY	FI
Concerta 36 mg depottabletter	UK/H/0544/002	02-1222	JANSSEN-CILAG A/S	NO
Concerta 36 mg depottabletter.	UK/H/0544/ 002	18542	JANSSEN-CILAG AB	SE
Concerta 36 mg forðatöflur.	UK/H/0544/002	IS/1/02/032/02	JANSSEN-CILAG AB	IS
Concerta 36 mg ilgstošās darbības tabletes	UK/H/0544/002	08-0131	UAB JOHNSON & JOHNSON	LV
CONCERTA 36 mg pailginto atpalaidavimo tabletės	UK/H/0544/002	LT/1/08/1152/004	UAB JOHNSON & JOHNSON	LT
CONCERTA 36 mg	UK/H/0544/002	LT/1/08/1152/003	UAB JOHNSON & JOHNSON	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pailginto atpalaidavimo tabletės				
Concerta 36 mg prolonged-release tablets	UK/H/0544/002	MA018/02602	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 36 mg Retardtabletten	UK/H/0544/002	1-24813	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 36 mg Retardtabletten	UK/H/0544/ 002	BE242697	JANSSEN-CILAG NV	BE
Concerta 36 mg Retardtabletten	UK/H/0544/002	54861.01.00	JANSSEN-CILAG GMBH	DE
Concerta 36 mg tablete s podaljšanim sproščanjem	UK/H/0544/ 002	H/08/00411/005	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
Concerta 36 mg tablete s produljenim oslobađanjem	not available	HR-H-413338677	JOHNSON & JOHNSON S.E. D.O.O.	HR
Concerta 36 mg tabletten met verlengde afgifte	UK/H/0544/ 002	BE242697	JANSSEN-CILAG NV	BE
Concerta 36 mg tabletten met verlengde afgifte	UK/H/0544/002	RVG 28074	JANSSEN-CILAG BV	NL
Concerta 36 mg tablety s predĺženým uvoľňovaním	UK/H/0544/002	78/0127/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 36 mg tablety s prodlouženým uvolňováním	UK/H/0544/002	06/408/08-C	JANSSEN-CILAG S.R.O	CZ
CONCERTA 36 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/002	12583	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 36 mg δισκία παρατεταμένης αποδέσμευσης.	UK/H/0544/002	20340	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 36 mg,	UK/H/0544/ 002	BE242697	JANSSEN-CILAG NV	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimés à libération prolongée				
Concerta 36 mg, comprimés à libération prolongée	UK/H/0544/ 002	2008100011	JANSSEN-CILAG NV	LU
Concerta 36 mg, Retardtabletten	UK/H/0544/ 002	2008100011	JANSSEN-CILAG NV	LU
Concerta 54 mg comprimate cu eliberare prelungită	UK/H/0544/003	5341/2013/02	JANSSEN PHARMACEUTICA NV	RO
Concerta 54 mg comprimate cu eliberare prelungită	UK/H/0544/003	5341/2013/01	JANSSEN PHARMACEUTICA NV	RO
Concerta 54 mg comprimidos de liberación prolongada.	UK/H/0544/003	65.171	JANSSEN-CILAG S.A.	ES
Concerta 54 mg comprimidos de libertação prolongada	UK/H/0544/003	4261186	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 54 mg comprimidos de libertação prolongada	UK/H/0544/003	4261285	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 54 mg depottabletit	UK/H/0544/003	17410	JANSSEN-CILAG OY	FI
Concerta 54 mg depottabletter	UK/H/0544/003	02-1223	JANSSEN-CILAG A/S	NO
Concerta 54 mg depottabletter	UK/H/0544/003	18543	JANSSEN-CILAG AB	SE
Concerta 54 mg forðatöflur.	UK/H/0544/003	IS/1/02/032/03	JANSSEN-CILAG AB	IS
Concerta 54 mg ilgstošās darbības tabletes	UK/H/0544/003	08-0132	UAB JOHNSON & JOHNSON	LV
CONCERTA 54 mg pailginto atpalaidavimo tabletės	UK/H/0544/003	LT/1/08/1152/006	UAB JOHNSON & JOHNSON	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CONCERTA 54 mg pailginto atpalaidavimo tabletės	UK/H/0544/003	LT/1/08/1152/005	UAB JOHNSON & JOHNSON	LT
Concerta 54 mg prolonged-release tablets	UK/H/0544/003	MA018/02603	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 54 mg Retardtabletten	UK/H/0544/003	1-24814	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 54 mg Retardtabletten	UK/H/0544/ 003	BE242706	JANSSEN-CILAG NV	BE
Concerta 54 mg Retardtabletten	UK/H/0544/003	54861.02.00	JANSSEN-CILAG GMBH	DE
Concerta 54 mg tablete s podaljšanim sproščanjem	UK/H/0544/ 003	H/08/00411/009	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
Concerta 54 mg tabletten met verlengde afgifte	UK/H/0544/ 003	BE242706	JANSSEN-CILAG NV	BE
Concerta 54 mg tabletten met verlengde afgifte	UK/H/0544/003	RVG 28075	JANSSEN-CILAG BV	NL
Concerta 54 mg tablety s predĺženým uvoľňovaním	UK/H/0544/003	78/0128/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 54 mg tablety s prodlouženým uvolňováním	UK/H/0544/003	06/409/08-C	JANSSEN-CILAG S.R.O	CZ
CONCERTA 54 mg δισκία παρατεταμένης αποδέσμευσης	UK/H/0544/003	12584	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 54 mg δισκία παρατεταμένης αποδέσμευσης.	UK/H/0544/003	20336	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 54 mg, comprimés à libération prolongée	UK/H/0544/ 003	BE242706	JANSSEN-CILAG NV	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Concerta 54 mg, comprimés à libération prolongée	UK/H/0544/ 003	2008100012	JANSSEN-CILAG NV	LU
Concerta 54 mg, Retardtabletten	UK/H/0544/ 003	2008100012	JANSSEN-CILAG NV	LU
CONCERTA LP 18 mg, comprimé à libération prolongée	UK/H/0544/001	34009 361 554 1 1	JANSSEN-CILAG	FR
CONCERTA LP 36 mg, comprimé à libération prolongée	UK/H/0544/002	34009 361 555 8 9	JANSSEN-CILAG	FR
CONCERTA LP 54 mg, comprimé à libération prolongée	UK/H/0544/003	34009 361 556 4 0	JANSSEN-CILAG	FR
Concerta XL 18 mg prolonged-release tablets	UK/H/0544/001	PA 0748/049/001	JANSSEN-CILAG LIMITED	IE
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	PA 0748/049/004	JANSSEN-CILAG LIMITED	IE
Concerta XL 27 mg prolonged-release tablets.	UK/H/0544/004	PL 00242/0400	JANSSEN-CILAG LIMITED	UK
Concerta XL 36 mg prolonged-release tablets	UK/H/0544/002	PA 0748/049/002	JANSSEN-CILAG LIMITED	IE
Concerta XL 36 mg prolonged-release tablets.	UK/H/0544/002	PL 00242/0373	JANSSEN-CILAG LIMITED	UK
Concerta XL 54 mg prolonged-release tablets	UK/H/0544/003	PA 0748/049/003	JANSSEN-CILAG LIMITED	IE
Concerta XL 54 mg prolonged-release tablets.	UK/H/0544/003	PL 00242/0374	JANSSEN-CILAG LIMITED	UK
Concerta, 18 mg	UK/H/0544/001	583608	UAB JOHNSON & JOHNSON	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
toimeainet prolongeeritult vabastavad tabletid				
Concerta, 18 mg, tabletki o przedłużonym uwalnianiu	UK/H/0544/001	14751	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta, 36 mg toimeainet prolongeeritult vabastavad tabletid	UK/H/0544/002	583708	UAB JOHNSON & JOHNSON	EE
Concerta, 36 mg, tabletki o przedłużonym uwalnianiu	UK/H/0544/002	14752	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta, 54 mg toimeainet prolongeeritult vabastavad tabletid	UK/H/0544/003	583808	UAB JOHNSON & JOHNSON	EE
Concerta, depottabletter	UK/H/0544/001	42318	JANSSEN-CILAG A/S	DK
Concerta, depottabletter	UK/H/0544/002	42319	JANSSEN-CILAG A/S	DK
Concerta, depottabletter	UK/H/0544/003	42320	JANSSEN-CILAG A/S	DK
Concerta® 18 mg depottabletter	UK/H/0544/001	17408	JANSSEN-CILAG OY	FI
Concerta® 27 mg depottabletter	UK/H/0544/004	24631	JANSSEN-CILAG OY	FI
Concerta® 36 mg depottabletter	UK/H/0544/002	17409	JANSSEN-CILAG OY	FI
Concerta® 54 mg depottabletter	UK/H/0544/003	17410	JANSSEN-CILAG OY	FI
DIFUMENIL, 27 MG, TABLETKI O PRZEDŁUŻONYM UWALNIANIU	DK/H/2138/001	20921	SANDOZ GMBH	PL
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 Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym 10 mg cápsulas duras de liberación modificada	UK/H/0819/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas duras de liberación modificada	UK/H/0819/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas duras de liberación modificada	UK/H/0819/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas duras de liberación modificada	UK/H/0819/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889015	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889039	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889041	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889054	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	UK/H/0819/001	041889066	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg tablets	UK/H/0416/002	PL 27303/0002	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 10 mg tablets	UK/H/0416/002	PL 27303/0002	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 10 mg tablets	UK/H/0416/002	PL 27303/0002	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 10 mg Tabletten	UK/H/0416/002	51447.01.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE

Product Name (in authorisation country)	MRP/DCP 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 Tabletten	UK/H/0416/002	51447.01.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 10 mg Tabletten	UK/H/0416/002	RVG 26494	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym 10 mg Tabletten	UK/H/0416/002	RVG 26494	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym 10 mg Tabletten	UK/H/0416/002	RVG 26494	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym 10 mg tabletter	UK/H/0416/002	35417	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 10 mg tabletter	UK/H/0416/002	35417	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 10 mg tabletter	UK/H/0416/002	35417	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
EQUASYM 10 mg tabletter	UK/H/0416/002	03-1987	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 10 mg tabletter	UK/H/0416/002	03-1987	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 10 mg tabletter	UK/H/0416/002	03-1987	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 10 mg töflur	UK/H/0416/002	IS/1/03/036/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 10 mg töflur	UK/H/0416/002	IS/1/03/036/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 10 mg töflur	UK/H/0416/002	IS/1/03/036/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 20 mg cápsulas duras de liberación modificada	UK/H/0819/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg cápsulas duras de liberación modificada	UK/H/0819/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES

Product Name (in authorisation country)	MRP/DCP 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
Equasym 20 mg cápsulas duras de liberación modificada	UK/H/0819/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg cápsulas duras de liberación modificada	UK/H/0819/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889080	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889092	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889104	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889116	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	UK/H/0819/002	041889128	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg tablets	UK/H/0416/003	PL 27303/0003	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 20 mg tablets	UK/H/0416/003	PL 27303/0003	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 20 mg tablets	UK/H/0416/003	PL 27303/0003	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 20 mg Tabletten	UK/H/0416/003	51447.02.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 20 mg Tabletten	UK/H/0416/003	51447.02.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 20 mg	UK/H/0416/003	51447.02.00	SHIRE PHARMACEUTICALS	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tabletten			IRELAND LIMITED	
Equasym 20 mg tabletter	UK/H/0416/003	35418	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 20 mg tabletter	UK/H/0416/003	35418	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 20 mg tabletter	UK/H/0416/003	35418	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
EQUASYM 20 mg tabletter	UK/H/0416/003	03-1988	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 20 mg tabletter	UK/H/0416/003	03-1988	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 20 mg tabletter	UK/H/0416/003	03-1988	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 20 mg töflur	UK/H/0416/003	IS/1/03/036/03	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 töflur	UK/H/0416/003	IS/1/03/036/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 30 mg cápsulas duras de liberación modificada	UK/H/0819/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	UK/H/0819/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	UK/H/0819/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	UK/H/0819/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg capsule rigide a rilascio modificato	UK/H/0819/003	041889130	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg capsule	UK/H/0819/003	041889142	SHIRE PHARMACEUTICALS	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rigide a rilascio modificato			IRELAND LIMITED	
Equasym 30 mg capsule rigide a rilascio modificato	UK/H/0819/003	041889155	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg capsule rigide a rilascio modificato	UK/H/0819/003	041889167	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 40 mg cápsulas duras de liberación modificada	UK/H/0819/004	77163	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 40 mg cápsulas duras de liberación modificada	UK/H/0819/004	77163	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 40 mg capsule rigide a rilascio modificato	UK/H/0819/004	041889179	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 40 mg capsule rigide a rilascio modificato	UK/H/0819/004	041889181	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 5 mg tablets	UK/H/0416/001	PL 27303/0001	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 5 mg tablets	UK/H/0416/001	PL 27303/0001	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 5 mg tablets	UK/H/0416/001	PL 27303/0001	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym 5 mg Tabletten	UK/H/0416/001	51447.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 5 mg Tabletten	UK/H/0416/001	51447.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 5 mg Tabletten	UK/H/0416/001	RVG 26493	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym 5 mg Tabletten	UK/H/0416/001	RVG 26493	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym 5 mg Tabletten	UK/H/0416/001	RVG 26493	SHIRE PHARMACEUTICALS	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			IRELAND LIMITED	
Equasym 5 mg tabletter	UK/H/0416/001	35416	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 5 mg tabletter	UK/H/0416/001	35416	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym 5 mg tabletter	UK/H/0416/001	35416	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
EQUASYM 5 mg tabletter	UK/H/0416/001	03-1986	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 5 mg tabletter	UK/H/0416/001	03-1986	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
EQUASYM 5 mg tabletter	UK/H/0416/001	03-1986	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym 5 mg töflur	UK/H/0416/001	IS/1/03/036/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 5 mg töflur	UK/H/0416/001	IS/1/03/036/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 5 mg töflur	UK/H/0416/001	IS/1/03/036/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym 5 mg Tabletten	UK/H/0416/001	51447.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym 50 mg cápsulas duras de liberación modificada	UK/H/0819/005	77164	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 50 mg cápsulas duras de liberación modificada	UK/H/0819/005	77164	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 50 mg capsule rigide a rilascio modificato	UK/H/0819/005	041889193	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 50 mg capsule rigide a rilascio modificato	UK/H/0819/005	041889205/M	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 60 mg cápsulas duras de liberación	UK/H/0819/006	77165	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modificada				
Equasym 60 mg cápsulas duras de liberación modificada	UK/H/0819/006	77165	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 60 mg capsule rigide a rilascio modificato	UK/H/0819/006	041889217	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 60 mg capsule rigide a rilascio modificato	UK/H/0819/006	041889229	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða	UK/H/0819/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 10 mg	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS	NO

Product Name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
kapsler med modifisert frisetting, harde			IRELAND LIMITED	
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 10 mg hård kapsel med modifierad frisättning	not available	20407	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 10 mg hård kapsel med modifierad frisättning	not available	20407	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða	UK/H/0819/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 20 mg hård kapsel med modifierad frisättning	not available	20408	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 20 mg hård kapsel med modifierad frisättning	not available	20408	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða	UK/H/0819/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða	UK/H/0819/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða	UK/H/0819/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða	UK/H/0819/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 30 mg kapsler med modifisert	UK/H/0819/003	05-3713	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
frisetting, harde				
Equasym Depot 30 mg hård kapsel med modifierad frisättning	not available	20409	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 30 mg hård kapsel med modifierad frisättning	not available	20409	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 40 mg kapslar med modifierad frisättning, hårda	not available	47785	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 40 mg kapslar med modifierad frisättning, hårda	not available	47785	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 40 mg kapsler med modifisert frisetting, harde	UK/H/0819/004	11-8780	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 40 mg kapsler med modifisert frisetting, harde	UK/H/0819/004	11-8780	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 50 mg kapslar med modifierad frisättning, hårda	not available	47786	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 50 mg kapslar med modifierad frisättning, hårda	not available	47786	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 50 mg kapsler med modifisert frisetting, harde	UK/H/0819/005	11-8781	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 50 mg kapsler med modifisert frisetting, harde	UK/H/0819/005	11-8781	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 60 mg kapslar med modifierad frisättning, hårda	not available	47787	SHIRE PHARMACEUTICALS IRELAND LIMITED	SE
Equasym Depot 60 mg	not available	47787	SHIRE PHARMACEUTICALS	SE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kapslar med modifierad frisättning, hårda			IRELAND LIMITED	
Equasym Depot 60 mg kapsler med modifisert frisetting, harde	UK/H/0819/006	11-8782	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
Equasym Depot 60 mg kapsler med modifisert frisetting, harde	UK/H/0819/006	11-8782	SHIRE PHARMACEUTICALS IRELAND LIMITED	NO
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 10 mg	UK/H/0819/001	38534	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
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 10 mg	UK/H/0819/001	38534	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
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
Equasym Depot, hårde kapsler med modificeret udløsning 20 mg	UK/H/0819/002	38535	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 20 mg	UK/H/0819/002	38535	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 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 20 mg	UK/H/0819/002	38535	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 30 mg	UK/H/0819/003	38536	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 30 mg	UK/H/0819/003	38536	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 30 mg	UK/H/0819/003	38536	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 30 mg	UK/H/0819/003	38536	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, hårde kapsler med modificeret udløsning 40 mg	UK/H/0819/004	50037	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 50 mg	UK/H/0819/005	50038	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Depot, hårde kapsler med modificeret udløsning 50 mg	UK/H/0819/005	50038	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
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 60 mg	UK/H/0819/006	50039	SHIRE PHARMACEUTICALS IRELAND LIMITED	DK
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/001	64003.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/001	64003.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/001	64003.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	UK/H/0819/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP / DCP 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	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/002	64004.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/002	64004.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/002	64004.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg	UK/H/0819/002	19258	SHIRE PHARMACEUTICALS	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kapslar med modifierad frisättning, hårda			IRELAND LIMITED	
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	MA 19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/003	64005.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	UK/H/0819/003	64005.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	UK/H/0819/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg depotkapseli, kova	UK/H/0819/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg	UK/H/0819/004	30420	SHIRE PHARMACEUTICALS	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
depotkapseli, kova			IRELAND LIMITED	
Equasym Retard 40 mg kapslar med modifierad frisättning, hårda	UK/H/0819/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg kapslar med modifierad frisättning, hårda	UK/H/0819/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg depotkapseli, kova	UK/H/0819/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg depotkapseli, kova	UK/H/0819/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg kapslar med modifierad frisättning, hårda	UK/H/0819/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg kapslar med modifierad frisättning, hårda	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 Retard 60 mg depotkapseli, kova	UK/H/0819/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg kapslar med modifierad frisättning, hårda	UK/H/0819/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg kapslar med modifierad frisättning, hårda	UK/H/0819/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard	UK/H/0819/001	RVG 33227	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/001/001	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/001/001	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/001/001	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/001/001	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PA 1575/001/001	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 10 mg modified-release	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
capsules, hard				
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 10 mg modified-release capsules, hard	UK/H/0819/001	PL 27303/0004	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 20 mg capsule met gereguleerde afgifte, hard	UK/H/0819/002	RVG 33228	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
Equasym XL 20 mg capsule met gereguleerde afgifte, hard	UK/H/0819/002	RVG 33228	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
Equasym XL 20 mg capsule met gereguleerde afgifte, hard	UK/H/0819/002	RVG 33228	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PA 1575/001/002	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PA 1575/001/002	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 20 mg modified-release	UK/H/0819/002	PA 1575/001/002	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
capsules, hard				
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PA 1575/001/002	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PA 1575/001/002	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 20 mg modified-release capsules, hard	UK/H/0819/002	PL 27303/0005	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 20 mg modified-release capsules, hard	UK/H/0819/002	PL 27303/0005	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 20 mg modified-release capsules, hard	UK/H/0819/002	PL 27303/0005	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 30 mg capsule met gereguleerde afgifte, hard	UK/H/0819/003	RVG 33229	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard	UK/H/0819/003	RVG 33229	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard	UK/H/0819/003	RVG 33229	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 30 mg	UK/H/0819/003	RVG 33229	SHIRE PHARMACEUTICALS	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
capsule met gereguleerde afgifte, hard			IRELAND LIMITED	
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PA 1575/001/003	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PA 1575/001/003	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PA 1575/001/003	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PA 1575/001/003	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 30 mg modified-release capsules, hard	UK/H/0819/003	PL 27303/0006	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 XL 30 mg modified-release capsules, hard	UK/H/0819/003	PL 27303/0006	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 XL 40 mg modified-release capsules, hard	UK/H/0819/004	PA 1575/001/007	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 40 mg modified-release capsules, hard	UK/H/0819/004	PA 1575/001/007	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 40 mg modified-release	UK/H/0819/04	PL 27303/0007	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
capsules, hard				
Equasym XL 40 mg modified-release capsules, hard	UK/H/0819/04	PL 27303/0007	SHIRE PHARMACEUTICALS IRELAND LIMITED	UK
Equasym XL 40 mg, capsule met gereguleerde afgifte, hard	UK/H/0819/004	RVG 111228	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 40 mg, capsule met gereguleerde afgifte, hard	UK/H/0819/004	RVG 111228	SHIRE PHARMACEUTICALS IRELAND LIMITED	NL
Equasym XL 50 mg modified-release capsules, hard	UK/H/0819/005	PA 1575/001/008	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 50 mg modified-release capsules, hard	UK/H/0819/005	PA 1575/001/008	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 50 mg modified-release capsules, hard	UK/H/0819/005	PL 27303/0008	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 50 mg, capsule met gereguleerde afgifte, hard	UK/H/0819/005	RVG 111232	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 modified-release capsules, hard	UK/H/0819/006	PA 1575/001/009	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE

Product Name (in authorisation country)	**MRP / DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
Equasym XL 60 mg modified-release capsules, hard	UK/H/0819/006	PA 1575/001/009	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym XL 60 mg modified-release capsules, hard	UK/H/0819/006	PL 27303/0009	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
Equasym XL 60 mg, capsule met gereguleerde afgifte, hard	UK/H/0819/006	RVG 111233	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 XR 10 mg capsules met gereguleerde afgifte, hard	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg capsules met	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
gereguleerde afgifte, hard				
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 10 mg gélules à libération modifiée	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 20 mg capsules met gereguleerde afgifte,	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
hard				
Equasym XR 20 mg capsules met gereguleerde afgifte, hard	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 20 mg capsules met gereguleerde afgifte, hard	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 20 mg capsules met gereguleerde afgifte, hard	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 20 mg capsules met gereguleerde afgifte, hard	UK/H/0819/002	BE423595	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 20 mg gélules à libération modifiée	UK/H/0819/002	BE423595	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 20 mg gélules à libération modifiée	UK/H/0819/002	BE423595	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 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 20 mg gélules à libération modifiée	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 30 mg capsules met gereguleerde afgifte, hard	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg capsules met gereguleerde afgifte, hard	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg capsules met gereguleerde afgifte, hard	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg capsules met gereguleerde afgifte, hard	UK/H/0819/003	BE423604	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 30 mg gélules à libération modifiée	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
gélules à libération modifiée			IRELAND LIMITED	
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 30 mg gélules à libération modifiée	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 40 mg capsules met gereguleerde afgifte, hard	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 40 mg capsules met gereguleerde afgifte, hard	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 40 mg gélules à libération modifiée	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 40 mg gélules à libération modifiée	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 40 mg gélules à libération modifiée	UK/H/0819/004	2013120618	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 40 mg	UK/H/0819/004	2013120618	SHIRE PHARMACEUTICALS	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
gélules à libération modifiée			IRELAND LIMITED	
Equasym XR 50 mg capsules met gereguleerde afgifte, hard	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 50 mg capsules met gereguleerde afgifte, hard	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 50 mg gélules à libération modifiée	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 50 mg gélules à libération modifiée	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 50 mg gélules à libération modifiée	UK/H/0819/005	2013120619	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 50 mg gélules à libération modifiée	UK/H/0819/005	2013120619	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 60 mg capsules met gereguleerde afgifte, hard	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 60 mg capsules met gereguleerde afgifte, hard	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 60 mg gélules à libération modifiée	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym XR 60 mg gélules à libération	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modifiée				
Equasym XR 60 mg gélules à libération modifiée	UK/H/0819/006	2013120620	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 60 mg gélules à libération modifiée	UK/H/0819/006	2013120620	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® 10mg, Tablets	UK/H/0416/002	PA 1575/1/5	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 10mg, Tablets	UK/H/0416/002	PA 1575/1/5	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 10mg, Tablets	UK/H/0416/002	PA 1575/1/5	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 20mg Tablets	UK/H/0416/003	PA 1575/1/6	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 20mg Tablets	UK/H/0416/003	PA 1575/1/6	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 20mg Tablets	UK/H/0416/003	PA 1575/1/6	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 5mg, Tablets	UK/H/0416/001	PA 1575/1/4	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 5mg, Tablets	UK/H/0416/001	PA 1575/1/4	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
Equasym® 5mg, Tablets	UK/H/0416/001	PA 1575/1/4	SHIRE PHARMACEUTICALS IRELAND LIMITED	IE
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	UK/H/0819/006	86843.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
veränderter Wirkstofffreisetzung				
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	BE423586	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 10 mg Hartkapseln mit veränderter	UK/H/0819/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	BE423595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	UK/H/0819/002	2013120616	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 30 mg Hartkapseln mit	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
veränderter				
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	BE423604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 30 mg Hartkapseln mit veränderter	UK/H/0819/003	2013120617	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 40 mg Hartkapseln mit veränderter	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 40 mg Hartkapseln mit veränderter	UK/H/0819/004	BE437595	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 40 mg Hartkapseln mit veränderter	UK/H/0819/004	2013120618	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 50 mg Hartkapseln mit veränderter	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 50 mg Hartkapseln mit veränderter	UK/H/0819/005	BE437604	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 50 mg Hartkapseln mit veränderter	UK/H/0819/005	2013120619	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 50 mg	UK/H/0819/005	2013120619	SHIRE PHARMACEUTICALS	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Hartkapseln mit veränderter			IRELAND LIMITED	
Equasym® XR 60 mg Hartkapseln mit veränderter	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 60 mg Hartkapseln mit veränderter	UK/H/0819/006	BE437613	SHIRE PHARMACEUTICALS IRELAND LIMITED	BE
Equasym® XR 60 mg Hartkapseln mit veränderter	UK/H/0819/006	2013120620	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym® XR 60 mg Hartkapseln mit veränderter	UK/H/0819/006	2013120620	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Medanef 20 mg tabletter	UK/H/5833/003	14-10264	MYLAN AB	NO
Medanef 20 mg tabletter	UK/H/5833/003	51607	MYLAN AB	SE
Medanef 5 mg tabletter	UK/H/5833/001	14-10262	MYLAN AB	NO
Medanef 5 mg tabletter	UK/H/5833/001	51605	MYLAN AB	SE
Medanef, tabletter	UK/H/5833/001	54633	MYLAN AB	DK
Medanef, tabletter	UK/H/5833/003	54635	MYLAN AB	DK
Medikinet 10 mg cápsulas de libertação modificada	DE/H/2223/002	5361555	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 10 mg cápsulas duras de liberación modificada	DE/H/0690/004	68.542	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 10 mg capsule rigide a rilascio modificato	DE/H/2223/002	041438033	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 10 mg capsule rigide a rilascio modificato	DE/H/2223/002	041438045	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 10 mg kapsler med modifisert frisetting, harde	DE/H/0690/004	06-4119	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MEDIKINET 10 mg, gélules à libération modifiée	DE/H/2223/002	CIS: 6 300 053 3	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Medikinet 10 mg, kapsel med modifierad frisättning, hård	DE/H/0690/004	23840	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 20 mg cápsulas de libertação modificada	DE/H/2223/003	5361563	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 20 mg cápsulas duras de liberación modificada	DE/H/0690/005	68.543	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 20 mg capsule rigide a rilascio modificato	DE/H/2223/003	041438058	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 20 mg capsule rigide a rilascio modificato	DE/H/2223/003	041438060	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 20 mg kapsler med modifisert frisetting, harde	DE/H/0690/005	06-4120	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
MEDIKINET 20 mg, gélules à libération modifiée	DE/H/2223/003	CIS: 6 365 782 8	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Medikinet 20 mg, kapsel med modifierad frisättning, hård	DE/H/0690/005	23841	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 30 mg cápsulas de libertação modificada	DE/H/2223/004	5361571	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 30 mg cápsulas duras de liberación modificada	DE/H/0690/006	68.544	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 30 mg capsule rigide a rilascio	DE/H/2223/004	041438072	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modificato				
Medikinet 30 mg capsule rigide a rilascio modificato	DE/H/2223/004	041438084	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 30 mg kapsler med modifisert frisetting, harde	DE/H/0690/006	06-4121	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
MEDIKINET 30 mg, gélules à libération modifiée	DE/H/2223/004	CIS: 6 659 298 3	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Medikinet 30 mg, kapsel med modifierad frisättning, hård	DE/H/0690/006	23842	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 40 mg cápsulas de libertação modificada	DE/H/2223/005	5361605	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 40 mg cápsulas duras de liberación modificada	DE/H/0690/007	68.545	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 40 mg capsule rigide a rilascio modificato	DE/H/2223/005	041438096	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 40 mg capsule rigide a rilascio modificato	DE/H/2223/005	041438108	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 40 mg kapsler med modifisert frisetting, harde	DE/H/0690/007	06-4122	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
MEDIKINET 40 mg, gélules à libération modifiée	DE/H/2223/005	CIS: 6 421 119 1	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Medikinet 40 mg, kapsel med modifierad frisättning, hård	DE/H/0690/007	23843	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 5 mg cápsulas	DE/H/2223/001	5361605	MEDICE ARZNEIMITTEL	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
de libertação modificada			PÜTTER GMBH & CO. KG	
Medikinet 5 mg cápsulas duras de liberación modificada	DE/H/0690/008	73.308	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 5 mg capsule rigide a rilascio modificato	DE/H/2223/001/DC	041438019/M; 041438021/M	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 5 mg capsule rigide a rilascio modificato	DE/H/2223/001	041438019	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 5 mg capsule rigide a rilascio modificato	DE/H/2223/001	041438021	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 5 mg kapsler med modifisert frisetting, harde	DE/H/0690/008	10/7785	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
MEDIKINET 5 mg, gélules à libération modifiée	DE/H/2223/001	CIS: 6 114 564 0	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Medikinet 5 mg, kapsel med modifierad frisättning, hård	DE/H/0690/008	44810	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 50 mg cápsulas duras de liberación modificada	DE/H/0690/009	78.453	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 50 mg kapsler med modifisert frisetting, harde	DE/H/0690/009	12-9267	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
Medikinet 50 mg, kapsel med modifierad frisättning, hård	DE/H/0690/009	48667	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 60 mg cápsulas duras de liberación modificada	DE/H/0690/010	78.454	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 60 mg kapsler	DE/H/0690/010	12-9268	MEDICE ARZNEIMITTEL	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
med modifisert frisetting, harde			PÜTTER GMBH & CO. KG	
Medikinet 60 mg, kapsel med modifierad frisättning, hård	DE/H/0690/010	48668	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet adult 10 mg Hartkapseln, retardiert	not available	86182.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 10 mg Hartkapseln, retardiert	not available	63890.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 20 mg Hartkapseln, retardiert	not available	86183.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 20 mg Hartkapseln, retardiert	not available	63891.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 30 mg Hartkapseln, retardiert	not available	65964.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 30 mg Hartkapseln, retardiert	not available	86184.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 40 mg Hartkapseln, retardiert	not available	86185.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 40 mg Hartkapseln, retardiert	not available	65965.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 5 mg Hartkapseln, retardiert	not available	66747.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 5 mg Hartkapseln, retardiert	not available	86181.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 50 mg Hartkapseln, retardiert	not available	88862.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet adult 60 mg Hartkapseln, retardiert	not available	88863.00.00	MEDICE PHARMA GMBH & CO. KG	DE
Medikinet CR 10 mg hörð hylki með breyttan losunarhraða	DE/H/0690/004/E/001	IS/1/14/087/005	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 10 mg kapsułki o zmodyfikowanym	DE/H/0690/004	12844	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
uwalnianiu, twarde				
Medikinet CR 10 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/004	RVG 34027	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 20 mg hörð hylki með breyttan losunarhraða	DE/H/0690/005/E/001	IS/1/14/087/006	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 20 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/005	12845	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 20 mg säädellysti vapauttava kapseli, kova	DE/H/0690/005	22213	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 20 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/005	RVG 34028	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 30 mg hörð hylki með breyttan losunarhraða	DE/H/0690/006/E/001	IS/1/14/087/007	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 30 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/006	12846	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 30 mg säädellysti vapauttava kapseli, kova	DE/H/0690/006	22214	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 30 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/006	RVG 34029	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 40 mg hörð hylki með breyttan	DE/H/0690/007/E/001	IS/1/14/087/008	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
losunarhraða				
Medikinet CR 40 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/007	12847	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 40 mg säädellysti vapauttava kapseli, kova	DE/H/0690/007	22215	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 40 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/007	RVG 34030	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 5 mg hörð hylki með breyttan losunarhraða	DE/H/0690/008/E/001	IS/1/14/087/004	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 5 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/008	18355	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 5 mg säädellysti vapauttava kapseli, kova	DE/H/0690/008	28970	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 5 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/008	RVG 108026	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 50 mg hörð hylki með breyttan losunarhraða	DE/H/0690/009/E/001	IS/1/14/087/009	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 50 mg säädellysti vapauttava kapseli, kova	DE/H/0690/009	31130	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 50 mg, 50 mg, kapsułki o zmodyfikowanym	DE/H/0690/009	21934	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
uwalnianiu, twarde				
Medikinet CR 50 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/009	RVG 112771	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR 60 mg hörð hylki með breyttan losunarhraða	DE/H/0690/010/E/001	IS/1/14/087/010	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 60 mg säädellysti vapauttava kapseli, kova	DE/H/0690/010	31131	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet CR 60 mg, 60 mg, kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/010	21935	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 60 mg, capsules met gereguleerde afgifte, hard	DE/H/0690/010	RVG 112772	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
Medikinet CR, hårde kapsler med modificeret udløsning	DE/H/0690/007	39461	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR, hårde kapsler med modificeret udløsning	DE/H/0690/006	39460	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR, hårde kapsler med modificeret udløsning	DE/H/0690/004	39458	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR, hårde kapsler med modificeret udløsning	DE/H/0690/005	39459	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR, hårde kapsler med modificeret udløsning 5 mg	DE/H/0690/008	47393	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR, hårde	DE/H/0690/009	51662	MEDICE ARZNEIMITTEL	DK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kapsler med modificeret udløsning 50 mg			PÜTTER GMBH & CO. KG	
Medikinet CR, hårde kapsler med modificeret udløsning 60 mg	DE/H/0690/010	51662	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Medikinet CR10 mg säädellysti vapauttava kapseli, kova	DE/H/0690/004	22212	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Medikinet EM 10 mg capsule cu eliberare modificată	DE/H/2223/002/DC	6241/2014/01, 6241/2014/02	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	RO
Medikinet EM 20 mg capsule cu eliberare modificată	DE/H/2223/003/DC	6242/2014/01, 6242/2014/02	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	RO
Medikinet EM 30 mg capsule cu eliberare modificată	DE/H/2223/004/DC	6243/2014/01, 6243/2014/02	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	RO
Medikinet EM 40 mg capsule cu eliberare modificată	DE/H/2223/005/DC	6244/2014/01, 6244/2014/02	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	RO
Medikinet MR 10 mg modified-release capsules, hard	DE/H/2223/002	PA 1555/1/2	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet MR 20 mg modified-release capsules, hard	DE/H/2223/003	PA1555/001/003	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet MR 30 mg modified-release capsules, hard	DE/H/2223/004	PA1555/001/004	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet MR 40 mg modified-release capsules, hard	DE/H/2223/005	PA1555/001/005	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet MR 5 mg modified-release capsules, hard	DE/H/2223/001	PA1555/001/001	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE

Product Name (in authorisation country)	MRP / DCP 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	AT
Medikinet retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/2223/002	75795.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/004	54569.00.01	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/004	1363/07020042	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 10 mg, Capsules met gereguleerde afgifte, hard	DE/H/2223/002	BE 381577	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/005	1-26726	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/2223/003	75796.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/005	54569.01.01	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 20 mg Hartkapseln mit veränderter	DE/H/0690/005	1363/07020043	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Wirkstofffreisetzung				
Medikinet retard 20 mg, gélules à libération modifiée	DE/H/2223/003	BE 381586	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/006	1-26727	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/2223/004	75797.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/006	62567.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/006	1363/07020044	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 30 mg, Capsules met gereguleerde afgifte, hard	DE/H/2223/004	BE 381595	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/007	1-26728	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/2223/005	75798.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 40 mg Hartkapseln mit veränderter	DE/H/0690/007	62568.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Wirkstofffreisetzung				
Medikinet retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/007	1363/07020045	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 40 mg, Capsules met gereguleerde afgifte, hard	DE/H/2223/005	BE381604	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/008	1-30056	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/008	66625.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/2223/001	75794.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/008	1754/11030018	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 5 mg, Capsules met gereguleerde afgifte, hard	DE/H/2223/001	BE 381561	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/009	135358	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 50 mg Hartkapseln mit	DE/H/0690/009	88767.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
veränderter Wirkstofffreisetzung				
Medikinet retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	97387.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/009	2014060146	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/010	135389	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	97388.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/010	88768.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet retard 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/0690/010	2014060147	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet XL 10 mg ilgstošās darbības cietās kapsulas	DE/H/2223/002	10-0461	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet XL 10 mg modified-release capsules, hard	DE/H/0690/004	PL 11243/0005	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL 10 mg modifikuoto atpalaidavimo kietosios	DE/H/2223/002	N28 – LT/1/10/2274/003, N30 – LT/1/10/2274/004	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kapsulės				
Medikinet XL 20 mg ilgstošās darbības cietās kapsulas	DE/H/2223/003	10-0462	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet XL 20 mg modified-release capsules, hard	DE/H/0690/005	PL 11243/0006	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL 20 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/003	N28 – LT/1/10/2274/005, N30 – LT/1/10/2274/006	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 30 mg ilgstošās darbības cietās kapsulas	DE/H/2223/004	10-0463	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet XL 30 mg modified-release capsules, hard	DE/H/0690/006	PL 11243/0007	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL 30 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/004	N28 – LT/1/10/2274/007, N30 – LT/1/10/2274/008	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 40 mg ilgstošās darbības cietās kapsulas	DE/H/2223/005	10-0464	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet XL 40 mg modified-release capsules, hard	DE/H/0690/007	PL 11243/0008	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL 40 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/005	N28 – LT/1/10/2274/009, N30 – LT/1/10/2274/010	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 5 mg ilgstošās darbības cietās kapsulas	DE/H/2223/001	10-0448	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet XL 5 mg	DE/H/0690/008	PL 11243/0010	MEDICE ARZNEIMITTEL	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modified-release capsules, hard			PÜTTER GMBH & CO. KG	
Medikinet XL 5 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/001	N28 – LT/1/10/2274/001, N30 – LT/1/10/2274/002	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 5 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/001	705610	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet XL 50 mg modified-release capsules, hard	DE/H/0690/009	PL 11243/0011	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL 60 mg modified-release capsules, hard	DE/H/0690/010	PL 11243/0012	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	UK
Medikinet XL10 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/002	705110	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet XL20 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/003	705010	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet XL30 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/004	705310	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet XL40 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/005	705710	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Methylfenidaat HCl Mylan 5 mg, tabletten	UK/H/5833/001	RVG 116232	MYLAN B.V.	NL
Methylphenidat	not available	54924	ALTERNOVA A/S	DK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
"Alternova", tabletter				
Methylphenidat "Alternova", tabletter	not available	54926	ALTERNOVA A/S	DK
Methylphenidate ”Sandoz”, depottabletter 27 mg	DK/H/2138/001	49666	SANDOZ A/S	DK
Methylphenidate Hydrochloride 20 mg Tablets	UK/H/5833/003	PL 04569/1498	GENERICS [UK] LIMITED	UK
Methylphenidate Hydrochloride 5 mg Tablets	UK/H/5833/001	PL 04569/1496	GENERICS [UK] LIMITED	UK
Methylphenidate Mylan 20 mg tabletten	UK/H/5833/003	BE489413	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 5 mg tabletten	UK/H/5833/001	BE489395	MYLAN BVBA/SPRL	BE
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
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 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 40 mg cápsulas de libertação modificada	UK/H/0819/004	5550462	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 40 mg cápsulas de libertação modificada	UK/H/0819/004	5550462	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
QUASYM 40 mg, gélule à libération modifiée	UK/H/0819/004	585 147-0	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
Quasym 50 mg cápsulas de libertação modificada	UK/H/0819/005	5550454	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 50 mg cápsulas de libertação modificada	UK/H/0819/005	5550454	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
QUASYM 50 mg, gélule à libération modifiée	UK/H/0819/005	585 149-3	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
Quasym 60 mg cápsulas de libertação modificada	UK/H/0819/006	5550470	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
Quasym 60 mg cápsulas de libertação modificada	UK/H/0819/006	5550470	SHIRE PHARMACEUTICALS IRELAND LIMITED	PT
QUASYM 60 mg, gélule à libération modifiée	UK/H/0819/006	585 151-8	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	UK/H/0819/001	377 617-8	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	UK/H/0819/001	497 261-6	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
QUASYM L.P. 10 mg, gélule à libération modifiée	UK/H/0819/001	377 618-4	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	UK/H/0819/001	570 268-1	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	UK/H/0819/001	570 269-8	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	UK/H/0819/002	377 620-9	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	UK/H/0819/002	377 619-0	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	UK/H/0819/002	497 262-2	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	UK/H/0819/002	378 313-2	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	UK/H/0819/002	570 271-2	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	UK/H/0819/003	377 621-5	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	UK/H/0819/003	497 267-4	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	UK/H/0819/003	377 622-1	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR
QUASYM L.P. 30 mg, gélule à libération	UK/H/0819/003	378 314-9	SHIRE PHARMACEUTICALS IRELAND LIMITED	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modifiée				
Rilatine 10 mg comprimés	not available	2002096657	NOVARTIS PHARMA N.V.	LU
RILATINE 10 mg tabletten	not available	BE051597	NOVARTIS PHARMA N.V.	BE
Rilatine 10 mg Tabletten	not available	BE051597	NOVARTIS PHARMA N.V.	BE
Rilatine 10 mg Tabletten	not available	2002096657	NOVARTIS PHARMA N.V.	LU
Rilatine 10 mg, comprimés	not available	BE051597	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE426413	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2014060160	NOVARTIS PHARMA N.V.	LU
RILATINE Modified Release 10 mg, capsules met gereguleerde afgifte, hard	not available	BE426413	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 10 mg, gélules à libération modifiée	not available	BE426413	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 10 mg, gélules à libération modifiée	not available	2014060160	NOVARTIS PHARMA N.V.	LU
Rilatine Modified Release 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE241534	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	0010/10/03/0097	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RILATINE Modified Release 20 mg, capsules met gereguleerde afgifte, hard	not available	BE241534	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 20 mg, gélules à libération modifiée	not available	BE241534	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 20 mg, gélules à libération modifiée	not available	0010/10/03/0097	NOVARTIS PHARMA N.V.	LU
Rilatine Modified Release 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE241543	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	0010/10/03/0098	NOVARTIS PHARMA N.V.	LU
RILATINE Modified Release 30 mg, capsules met gereguleerde afgifte, hard	not available	BE241543	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 30 mg, gélules à libération modifiée	not available	BE241543	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 30 mg, gélules à libération modifiée	not available	0010/10/03/0098	NOVARTIS PHARMA N.V.	LU
Rilatine Modified Release 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE241552	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 40 mg Hartkapseln mit veränderter	not available	0010/10/08/0034	NOVARTIS PHARMA N.V.	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
WirkstofffreisetzungMethylphenidat				
RILATINE Modified Release 40 mg, capsules met gereguleerde afgifte, hard	not available	BE241552	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 40 mg, gélules à libération modifiée	not available	BE241552	NOVARTIS PHARMA N.V.	BE
Rilatine Modified Release 40 mg, gélules à libération modifiée	not available	0010/10/08/0034	NOVARTIS PHARMA N.V.	LU
Ritalin	not available	01870	NOVARTIS HEALTHCARE A/S	DK
Ritalin 10 mg - Tabletten	not available	1-22028	NOVARTIS PHARMA GMBH	AT
RITALIN 10 mg compresse	Not Applicable	035040017	Novartis Farma S.p.A.	IT
Ritalin 10 mg hårda kapslar med modifierad frisättning	not available	41556	NOVARTIS SVERIGE AB	SE
Ritalin 10 mg kapsler med modifisert frisetting, harde	not available	09-6501	NOVARTIS NORGE AS	NO
Ritalin 10 mg tabletta	not available	OGYI-T-6954/01	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Ritalin 10 mg tabletter	not available	3449	NOVARTIS NORGE AS	NO
Ritalin 10 mg tabletter	not available	20606	NOVARTIS SVERIGE AB	SE
Ritalin 10 mg tablety	not available	06/1179/97-C	NOVARTIS, S.R.O.	CZ
Ritalin 10 mg töflur	not available	640044	NOVARTIS HEALTHCARE A/S	IS
RITALIN 10MG TABLETS	not available	PA 13/66/1	NOVARTIS PHARMACEUTICALS UK LIMITED	IE
Ritalin 20 mg hårda kapslar med modifierad frisättning	not available	20607	NOVARTIS SVERIGE AB	SE

Product Name (in authorisation country)	MRP/DCP 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	NO
Ritalin 30 mg hårda kapslar med modifierad frisättning	not available	20608	NOVARTIS SVERIGE AB	SE
Ritalin 30 mg kapsler med modifisert frisetting, harde	not available	05-3197	NOVARTIS NORGE AS	NO
Ritalin 40 mg hårda kapslar med modifierad frisättning	not available	20609	NOVARTIS SVERIGE AB	SE
Ritalin 40 mg kapsler med modifisert frisetting, harde	not available	05-3198	NOVARTIS NORGE AS	NO
Ritalin 60 mg hårda kapslar med modifierad frisättning	not available	50408	NOVARTIS SVERIGE AB	SE
Ritalin 60 mg kapsel med modifiert frisetting, hard	not available	13-9867	NOVARTIS NORGE AS	NO
Ritalin LA 10 mg - Kapseln	not available	1-31212	NOVARTIS PHARMA GMBH	AT
Ritalin LA 10 mg, harde capsules met gereguleerde afgifte	not available	RVG 116377	NOVARTIS PHARMA B.V.	NL
Ritalin LA 20 mg - Kapseln	not available	1-24270	NOVARTIS PHARMA GMBH	AT
Ritalin LA 20 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/02	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Ritalin LA 20 mg, harde capsules met gereguleerde afgifte	not available	RVG 116379	NOVARTIS PHARMA B.V.	NL
Ritalin LA 30 mg -	not available	1-24271	NOVARTIS PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kapseln				
Ritalin LA 30 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/03	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Ritalin LA 30 mg, harde capsules met gereguleerde afgifte	not available	RVG 116380	NOVARTIS PHARMA B.V.	NL
Ritalin LA 40 mg - Kapseln	not available	1-24272	NOVARTIS PHARMA GMBH	AT
Ritalin LA 40 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/04	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Ritalin LA 40 mg, harde capsules met gereguleerde afgifte	not available	RVG 116381	NOVARTIS PHARMA B.V.	NL
Ritalin LA 60 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/05	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Ritalin LA 60 mg, harde capsules met gereguleerde afgifte	not available	RVG 116382	NOVARTIS PHARMA B.V.	NL
Ritalin Uno	not available	34357	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno	not available	34355	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno	not available	53335	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno	not available	34356	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno	not available	44662	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno 10 mg hylki með breyttan losunarhraða, hart	not available	IS/1/10/133/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 20 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/01	NOVARTIS HEALTHCARE A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin Uno 30 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/02	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 40 mg hylki með breyttan losunarhraða, hart	not available	IS/1/02/127/03	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 60 mg hylki með breyttan losunarhraða, hart	not available	IS/1/15/020/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin, tabletten 10 mg	not available	RVG 03957	NOVARTIS PHARMA B.V.	NL
Ritalin®	not available	PL 00101/0539	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
RITALIN® 10 MG TABLETES	not available	00-0118	NOVARTIS FINLAND OY	LV
RITALIN® 10 mg Tablets	not available	088/03101	NOVARTIS PHARMACEUTICALS UK LIMITED	MT
Ritalin® 10 mg Tabletten	not available	6094573.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® Adult 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	77240.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® Adult 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67252.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® Adult 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67253.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® Adult 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67254.00.00	NOVARTIS PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ritalin® Adult 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	91388.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	77239.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67249.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67250.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67251.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	91387.00.00	NOVARTIS PHARMA GMBH	DE
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
Ritalina LA 20 mg cápsulas de libertação	Not Applicable	3761780	Novartis Farma - Produtos Farmaceuticos S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
modificada				
Ritalina LA 20 mg cápsulas de libertação modificada	Not Applicable	3761889	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 30 mg cápsulas de libertação modificada	Not Applicable	3761988	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 30 mg cápsulas de libertação modificada	Not Applicable	3762085	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 30 mg cápsulas de libertação modificada	Not Applicable	3762084	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 40 mg cápsulas de libertação modificada	Not Applicable	3762283	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 60 mg cápsulas de libertação modificada	Not Applicable	5635115	Novartis Farma - Produtos Farmaceuticos S.A.	PT
Ritalina LA 60 mg cápsulas de libertação modificada	Not Applicable	5635123	Novartis Farma - Produtos Farmaceuticos S.A.	PT
RITALINE 10 mg comprimé	not available	3400933929404	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg comprimé	not available	3400933942410	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg comprimé	not available	3400933942649	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	3400941686764	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	3400957974572	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg,	not available	3400941686535	NOVARTIS PHARMA S.A.S.	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
gélule à libération prolongée				
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400936231382	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400956496914	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 20 mg gélule à libération prolongée	not available	3400936534933	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 30 mg, gélule à libération prolongée	not available	3400936535015	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 30 mg, gélule à libération prolongée	not available	3400936231504	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 30 mg, gélule à libération prolongée	not available	3400956497164	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 40 mg, gélule à libération prolongée	not available	3400956497225	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 40 mg, gélule à libération prolongée	not available	3400936535183	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 40 mg, gélule à libération prolongée	not available	3400936231733	NOVARTIS PHARMA S.A.S.	FR