EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

23 June 2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): methylphenidate

Procedure No. PSUSA/00002024/202110

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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
®Ritalin 10mg Tablets	not available	PA0896/026/001	NOVARTIS IRELAND LIMITED	IE
®Ritalin 10mg Tablets	not available	PA0896/026/001	NOVARTIS IRELAND LIMITED	IE
Concerta 18 mg comprimate cu eliberare prelungită	DE/H/6007 001	5339/2013/01	JANSSEN PHARMACEUTICA NV	RO
Concerta 18 mg comprimate cu eliberare prelungită	DE/H/6007 001	5339/2013/02	JANSSEN PHARMACEUTICA NV	RO
Concerta 18 mg comprimidos de liberación prolongada.	DE/H/6007/001	65.148	JANSSEN-CILAG S.A.	ES
Concerta 18 mg comprimidos de libertação prolongada	DE/H/6007 001	4260881	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 18 mg comprimidos de libertação prolongada.	DE/H/6007 001	4260782	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 18 mg depottabletit	DE/H/6007001	17408	JANSSEN-CILAG OY	FI
Concerta 18 mg depottabletter	DE/H/6007001	17408	JANSSEN-CILAG OY	FI
Concerta 18 mg depottabletter	DE/H/6007/001	02-1221	JANSSEN-CILAG A/S	NO
Concerta 18 mg depottabletter.	DE/H/6007/001	18541	JANSSEN-CILAG AB	SE
Concerta 18 mg forðatöflur.	DE/H/6007/001	IS/1/02/032/01	JANSSEN-CILAG AB	IS
Concerta 18 mg ilgstošās darbības tabletes	DE/H/6007/001	08-0130	UAB "JOHNSON & JOHNSON"	LV
CONCERTA 18 mg pailginto atpalaidavimo tabletės	DE/H/6007/001	LT/1/08/1152/002	UAB "JOHNSON & JOHNSON"	LT
CONCERTA 18 mg pailginto atpalaidavimo	DE/H/6007/001	LT/1/08/1152/001	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
tabletės				
Concerta 18 mg prolonged-release tablets.	DE/H/6007/001	MA018/02601	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 18 mg Retardtabletten	DE/H/6007 001	1-24812	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 18 mg Retardtabletten	DE/H/6007 001	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg Retardtabletten	DE/H/6007/001	54861.00.00	JANSSEN-CILAG GMBH	DE
Concerta 18 mg Retardtabletten	DE/H/6007/001	2008100010	JANSSEN-CILAG NV	LU
Concerta 18 mg tablete s podaljšanim sproščanjem	DE/H/6007/001	H/08/00411/001	JOHNSON & JOHNSON D.O.O.	SI
Concerta 18 mg tablete s podaljšanim sproščanjem	DE/H/6007/001	H/08/00411/002	JOHNSON & JOHNSON D.O.O.	SI
Concerta 18 mg tablete s podaljšanim sproščanjem	DE/H/6007/001	H/08/00411/003	JOHNSON & JOHNSON D.O.O.	SI
Concerta 18 mg tablete s podaljšanim sproščanjem	DE/H/6007/001	H/08/00411/004	JOHNSON & JOHNSON 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	DE/H/6007/001	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg tabletten met verlengde afgifte.	DE/H/6007/001	RVG 28073	JANSSEN-CILAG BV	NL
Concerta 18 mg tablety s predĺženým uvoľňovaním	DE/H/6007 001	78/0126/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 18 mg tablety s prodlouženým uvolňováním	DE/H/6007 001	06/407/08-C	JANSSEN-CILAG S.R.O	CZ
Concerta 18 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007/001	20339	JANSSEN-CILAG INTERNATIONAL NV	CY

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 δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007 001	12582/19.02.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 18 mg, comprimés à libération prolongée	DE/H/6007/001	BE242681	JANSSEN-CILAG NV	BE
Concerta 18 mg, comprimés à libération prolongée	DE/H/6007/001	2008100010	JANSSEN-CILAG NV	LU
Concerta 27 mg comprimidos de liberación prolongada.	DE/H/6007 004	69.988	JANSSEN-CILAG S.A.	ES
Concerta 27 mg comprimidos de libertação prolongada.	DE/H/6007/004	5205307	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 27 mg comprimidos de libertação prolongada.	DE/H/6007/004	5205273	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 27 mg depottabletit	DE/H/6007004	24631	JANSSEN-CILAG OY	FI
Concerta 27 mg depottabletter	DE/H/6007004	24631	JANSSEN-CILAG OY	FI
Concerta 27 mg depottabletter	DE/H/6007/004	08-5646	JANSSEN-CILAG A/S	NO
Concerta 27 mg depottabletter.	DE/H/6007/004	26554	JANSSEN-CILAG AB	SE
Concerta 27 mg forðatöflur.	DE/H/6007/004	IS/1/08/032/01	JANSSEN-CILAG AB	IS
Concerta 27 mg Retardtabletten	DE/H/6007 004	1-27727	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 27 mg Retardtabletten	DE/H/6007 004	BE324913	JANSSEN-CILAG NV	BE
Concerta 27 mg Retardtabletten	DE/H/6007 004	72263.00.00	JANSSEN-CILAG GMBH	DE
Concerta 27 mg Retardtabletten	DE/H/6007/004	2008120051	JANSSEN-CILAG NV	LU
Concerta 27 mg	DE/H/6007/004	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
tabletten met verlengde afgifte				
Concerta 27 mg tabletten met verlengde afgifte.	DE/H/6007/004	RVG 101739	JANSSEN-CILAG BV	NL
Concerta 27 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007 004	12585/19.02.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 27 mg, comprimés à libération prolongée	DE/H/6007/004	BE324913	JANSSEN-CILAG NV	BE
Concerta 27 mg, comprimés à libération prolongée	DE/H/6007/004	2008120051	JANSSEN-CILAG NV	LU
Concerta 36 mg comprimate cu eliberare prelungită	DE/H/6007 002	5340/2013/02	JANSSEN PHARMACEUTICA NV	RO
Concerta 36 mg comprimate cu eliberare prelungită	DE/H/6007 002	5340/2013/01	JANSSEN PHARMACEUTICA NV	RO
Concerta 36 mg comprimidos de liberación prolongada.	DE/H/6007 002	65.170	JANSSEN-CILAG S.A.	ES
Concerta 36 mg comprimidos de libertação prolongada.	DE/H/6007/002	4260980	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 36 mg comprimidos de libertação prolongada.	DE/H/6007/002	4261087	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 36 mg depottabletit	DE/H/6007002	17409	JANSSEN-CILAG OY	FI
Concerta 36 mg depottabletter	DE/H/6007002	17409	JANSSEN-CILAG OY	FI
Concerta 36 mg depottabletter	DE/H/6007/002	02-1222	JANSSEN-CILAG A/S	NO
Concerta 36 mg depottabletter.	DE/H/6007/002	18542	JANSSEN-CILAG 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
Concerta 36 mg forðatöflur.	DE/H/6007/002	IS/1/02/032/02	JANSSEN-CILAG AB	IS
Concerta 36 mg ilgstošās darbības tabletes	DE/H/6007/002	08-0131	UAB "JOHNSON & JOHNSON"	LV
CONCERTA 36 mg pailginto atpalaidavimo tabletės	DE/H/6007/002	LT/1/08/1152/004	UAB "JOHNSON & JOHNSON"	LT
CONCERTA 36 mg pailginto atpalaidavimo tabletės	DE/H/6007/002	LT/1/08/1152/003	UAB "JOHNSON & JOHNSON"	LT
Concerta 36 mg prolonged-release tablets.	DE/H/6007/002	MA018/02602	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 36 mg Retardtabletten	DE/H/6007 002	1-24813	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 36 mg Retardtabletten	DE/H/6007/002	BE242697	JANSSEN-CILAG NV	BE
Concerta 36 mg Retardtabletten	DE/H/6007/002	54861.01.00	JANSSEN-CILAG GMBH	DE
Concerta 36 mg Retardtabletten	DE/H/6007/002	2008100011	JANSSEN-CILAG NV	LU
Concerta 36 mg tablete s podaljšanim sproščanjem	DE/H/6007/002	H/08/00411/005	JOHNSON & JOHNSON D.O.O.	SI
Concerta 36 mg tablete s podaljšanim sproščanjem	DE/H/6007/002	H/08/00411/007	JOHNSON & JOHNSON D.O.O.	SI
Concerta 36 mg tablete s podaljšanim sproščanjem	DE/H/6007/002	H/08/00411/008	JOHNSON & JOHNSON D.O.O.	SI
Concerta 36 mg tablete s podaljšanim sproščanjem	DE/H/6007/002	H/08/00411/006	JOHNSON & JOHNSON 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	DE/H/6007/002	BE242697	JANSSEN-CILAG NV	BE
Concerta 36 mg tabletten met verlengde	DE/H/6007/002	RVG 28074	JANSSEN-CILAG BV	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
afgifte.				
Concerta 36 mg tablety s predĺženým uvoľňovaním	DE/H/6007 002	78/0127/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 36 mg tablety s prodlouženým uvolňováním	DE/H/6007/002	06/408/08-C	JANSSEN-CILAG S.R.O	CZ
Concerta 36 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007/002	20340	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 36 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007 002	12583/19.02.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 36 mg, comprimés à libération prolongée	DE/H/6007/002	BE242697	JANSSEN-CILAG NV	BE
Concerta 36 mg, comprimés à libération prolongée	DE/H/6007 002	2008100011	JANSSEN-CILAG NV	LU
Concerta 54 mg comprimate cu eliberare prelungită	DE/H/6007 003	5341/2013/02	JANSSEN PHARMACEUTICA NV	RO
Concerta 54 mg comprimate cu eliberare prelungită	DE/H/6007 003	5341/2013/01	JANSSEN PHARMACEUTICA NV	RO
Concerta 54 mg comprimidos de liberación prolongada.	DE/H/6007 003	65.171	JANSSEN-CILAG S.A.	ES
Concerta 54 mg comprimidos de libertação prolongada.	DE/H/6007/003	4261186	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 54 mg comprimidos de libertação prolongada.	DE/H/6007/003	4261285	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Concerta 54 mg depottabletit	DE/H/6007003	17410	JANSSEN-CILAG OY	FI
Concerta 54 mg depottabletter	DE/H/6007003	17410	JANSSEN-CILAG OY	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
Concerta 54 mg depottabletter	DE/H/6007/003	02-1223	JANSSEN-CILAG A/S	NO
Concerta 54 mg depottabletter.	DE/H/6007/003	18543	JANSSEN-CILAG AB	SE
Concerta 54 mg forðatöflur.	DE/H/6007/003	IS/1/02/032/03	JANSSEN-CILAG AB	IS
Concerta 54 mg ilgstošās darbības tabletes	DE/H/6007/003	08-0132	UAB "JOHNSON & JOHNSON"	LV
CONCERTA 54 mg pailginto atpalaidavimo tabletės	DE/H/6007/003	LT/1/08/1152/006	UAB "JOHNSON & JOHNSON"	LT
CONCERTA 54 mg pailginto atpalaidavimo tabletės	DE/H/6007/003	LT/1/08/1152/005	UAB "JOHNSON & JOHNSON"	LT
Concerta 54 mg prolonged-release tablets.	DE/H/6007/003	MA018/02603	JANSSEN-CILAG INTERNATIONAL NV	MT
Concerta 54 mg Retardtabletten	DE/H/6007 003	1-24814	JANSSEN-CILAG PHARMA GMBH	AT
Concerta 54 mg Retardtabletten	DE/H/6007 003	BE242706	JANSSEN-CILAG NV	BE
Concerta 54 mg Retardtabletten	DE/H/6007/003	54861.02.00	JANSSEN-CILAG GMBH	DE
Concerta 54 mg Retardtabletten	DE/H/6007/003	2008100012	JANSSEN-CILAG NV	LU
Concerta 54 mg tablete s podaljšanim sproščanjem	DE/H/6007/003	H/08/00411/009	JOHNSON & JOHNSON D.O.O.	SI
Concerta 54 mg tablete s podaljšanim sproščanjem	DE/H/6007/003	H/08/00411/010	JOHNSON & JOHNSON D.O.O.	SI
Concerta 54 mg tablete s podaljšanim sproščanjem	DE/H/6007/003	H/08/00411/011	JOHNSON & JOHNSON D.O.O.	SI
Concerta 54 mg tablete s podaljšanim sproščanjem	DE/H/6007/003	H/08/00411/012	JOHNSON & JOHNSON D.O.O.	SI
Concerta 54 mg tabletten met verlengde afgifte	DE/H/6007/003	BE242706	JANSSEN-CILAG NV	BE
Concerta 54 mg	DE/H/6007/003	RVG 28075	JANSSEN-CILAG BV	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
tabletten met verlengde afgifte.				
Concerta 54 mg tablety s predĺženým uvoľňovaním	DE/H/6007 003	78/0128/08-S	JOHNSON & JOHNSON, S.R.O	SK
Concerta 54 mg tablety s prodlouženým uvolňováním	DE/H/6007 003	06/409/08-C	JANSSEN-CILAG S.R.O	CZ
Concerta 54 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007/003	20336	JANSSEN-CILAG INTERNATIONAL NV	CY
Concerta 54 mg δισκία παρατεταμένης αποδέσμευσης.	DE/H/6007 003	12584/19.02.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Concerta 54 mg, comprimés à libération prolongée	DE/H/6007/003	BE242706	JANSSEN-CILAG NV	BE
Concerta 54 mg, comprimés à libération prolongée	DE/H/6007 003	2008100012	JANSSEN-CILAG NV	LU
CONCERTA LP 18 mg, comprimé à libération prolongée	DE/H/6007/001	34009 361 554 1 1	JANSSEN-CILAG	FR
CONCERTA LP 36 mg, comprimé à libération prolongée	DE/H/6007/002	34009 361 555 8 9	JANSSEN-CILAG	FR
CONCERTA LP 54 mg, comprimé à libération prolongée	DE/H/6007/003	34009 361 556 4 0	JANSSEN-CILAG	FR
Concerta XL 18 mg prolonged-release tablets.	DE/H/6007/001	PA22612/002/001	JANSSEN SCIENCES IRELAND UC	IE
Concerta XL 18 mg prolonged-release tablets.	DE/H/6007/001	PL 00242/0372	JANSSEN-CILAG LIMITED	XI
Concerta XL 27 mg prolonged-release tablets.	DE/H/6007/004	PA22612/002/004	JANSSEN SCIENCES IRELAND UC	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
Concerta XL 27 mg prolonged-release tablets.	DE/H/6007/004	PL 00242/0400	JANSSEN-CILAG LIMITED	XI
Concerta XL 36 mg prolonged-release tablets.	DE/H/6007/002	PA22612/002/002	JANSSEN SCIENCES IRELAND UC	IE
Concerta XL 36 mg prolonged-release tablets.	DE/H/6007/002	PL 00242/0373	JANSSEN-CILAG LIMITED	XI
Concerta XL 54 mg prolonged-release tablets.	DE/H/6007/003	PL 00242/0374	JANSSEN-CILAG LIMITED	XI
Concerta, 18 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/6007/001	583608	UAB "JOHNSON & JOHNSON"	EE
Concerta, 18 mg, tabletki o przedłużonym uwalnianiu	DE/H/6007/001	14751	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta, 36 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/6007/002	583708	UAB "JOHNSON & JOHNSON"	EE
Concerta, 36 mg, tabletki o przedłużonym uwalnianiu	DE/H/6007/002	14752	JANSSEN-CILAG INTERNATIONAL NV	PL
Concerta, 54 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/6007/003	583808	UAB "JOHNSON & JOHNSON"	EE
Concerta, depottabletter	DE/H/6007/001	42318	JANSSEN-CILAG A/S	DK
Concerta, depottabletter	DE/H/6007/002	42319	JANSSEN-CILAG A/S	DK
Concerta, depottabletter	DE/H/6007/003	42320	JANSSEN-CILAG A/S	DK
Equasym 10 mg cápsulas duras de liberación modificada	DK/H/2888/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas	DK/H/2888/001	76061	SHIRE PHARMACEUTICALS	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
duras de liberación modificada			IRELAND LIMITED	
Equasym 10 mg cápsulas duras de liberación modificada	DK/H/2888/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas duras de liberación modificada	DK/H/2888/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg cápsulas duras de liberación modificada	DK/H/2888/001	76061	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 10 mg capsule rigide a rilascio modificato	DK/H/2888/001	041889015	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	DK/H/2888/001	041889039	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	DK/H/2888/001	041889041	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	DK/H/2888/001	041889054	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 10 mg capsule rigide a rilascio modificato	DK/H/2888/001	041889066	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg cápsulas duras de liberación modificada	DK/H/2888/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg cápsulas duras de liberación modificada	DK/H/2888/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg cápsulas duras de liberación modificada	DK/H/2888/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg cápsulas duras de liberación	DK/H/2888/002	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
modificada				
Equasym 20 mg cápsulas duras de liberación modificada	DK/H/2888/002	76076	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 20 mg capsule rigide a rilascio modificato	DK/H/2888/002	041889080	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	DK/H/2888/002	041889092	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	DK/H/2888/002	041889104	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	DK/H/2888/002	041889116	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 20 mg capsule rigide a rilascio modificato	DK/H/2888/002	041889128	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg cápsulas duras de liberación modificada	DK/H/2888/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	DK/H/2888/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	DK/H/2888/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg cápsulas duras de liberación modificada	DK/H/2888/003	76060	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 30 mg capsule rigide a rilascio modificato	DK/H/2888/003	041889130	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg capsule rigide a rilascio modificato	DK/H/2888/003	041889142	SHIRE PHARMACEUTICALS IRELAND LIMITED	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
Equasym 30 mg capsule rigide a rilascio modificato	DK/H/2888/003	041889155	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 30 mg capsule rigide a rilascio modificato	DK/H/2888/003	041889167	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 40 mg cápsulas duras de liberación modificada	DK/H/2888/004	77163	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 40 mg cápsulas duras de liberación modificada	DK/H/2888/004	77163	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 40 mg capsule rigide a rilascio modificato	DK/H/2888/004	041889179	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 40 mg capsule rigide a rilascio modificato	DK/H/2888/004	041889181	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 50 mg cápsulas duras de liberación modificada	DK/H/2888/005	77164	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 50 mg cápsulas duras de liberación modificada	DK/H/2888/005	77164	SHIRE PHARMACEUTICALS IRELAND LIMITED	ES
Equasym 50 mg capsule rigide a rilascio modificato	DK/H/2888/005	041889193	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 50 mg capsule rigide a rilascio modificato	DK/H/2888/005	041889205	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 60 mg capsule rigide a rilascio modificato	DK/H/2888/006	041889217	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym 60 mg capsule rigide a rilascio modificato	DK/H/2888/006	041889229	SHIRE PHARMACEUTICALS IRELAND LIMITED	IT
Equasym Depot 10 mg	DK/H/2888/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS	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
hörð hylki með breyttan losunarhraða.			IRELAND LIMITED	
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/001	IS/1/06/003/01	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 10 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/001	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	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 10 mg kapsler med modifisert frisetting, harde	UK/H/0819/001	05-3711	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan	DK/H/2888/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	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.				
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/002	IS/1/06/003/02	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 20 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/002	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	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 20 mg kapsler med modifisert frisetting, harde	UK/H/0819/002	05-3712	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/003	IS/1/06/003/03	SHIRE PHARMACEUTICALS IRELAND LIMITED	IS
Equasym Depot 30 mg hörð hylki með breyttan losunarhraða.	DK/H/2888/003	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	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	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 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 30 mg kapsler med modifisert frisetting, harde	UK/H/0819/003	05-3713	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 40 mg kapsler med modifisert frisetting, harde	UK/H/0819/004	11-8780	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 40 mg kapsler med modifisert frisetting, harde	UK/H/0819/004	11-8780	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 50 mg kapsler med modifisert frisetting, harde	UK/H/0819/005	11-8781	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 50 mg kapsler med modifisert frisetting, harde	UK/H/0819/005	11-8781	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 60 mg kapsler med modifisert frisetting, harde	UK/H/0819/006	11-8782	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot 60 mg kapsler med modifisert frisetting, harde	UK/H/0819/006	11-8782	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NO
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/001	38534	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/001	38534	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/001	38534	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde	UK/H/0819/001	38534	TAKEDA PHARMACEUTICALS	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			INTERNATIONAL AG IRELAND BRANCH	
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/001	38534	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/002	38535	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/002	38535	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/002	38535	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/002	38535	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/002	38535	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/003	38536	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/003	38536	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/003	38536	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/003	38536	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/005	50038	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret	UK/H/0819/006	50039	TAKEDA PHARMACEUTICALS INTERNATIONAL AG	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
udløsning			IRELAND BRANCH	
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/004	50037	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/004	50037	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/005	50038	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Depot, hårde kapsler med modificeret udløsning	UK/H/0819/006	50039	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	DK
Equasym Retard 10 mg depotkapseli, kova	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg depotkapseli, kova	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad frisättning, hårda	DK/H/2888/001	19257	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 10 mg kapslar med modifierad	DK/H/2888/001	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
frisättning, hårda				
Equasym Retard 20 mg depotkapseli, kova	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg depotkapseli, kova	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 20 mg kapslar med modifierad frisättning, hårda	DK/H/2888/002	19258	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg depotkapseli, kova	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	DK/H/2888/003	19259	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 30 mg kapslar med modifierad frisättning, hårda	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 30 mg kapslar med modifierad frisättning, hårda	DK/H/2888/003	19259	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg depotkapseli, kova	DK/H/2888/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg depotkapseli, kova	DK/H/2888/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/004	7002498.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/004	7002498.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 40 mg kapslar med modifierad frisättning, hårda	DK/H/2888/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 40 mg kapslar med modifierad frisättning, hårda	DK/H/2888/004	30420	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg depotkapseli, kova	DK/H/2888/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg depotkapseli, kova	DK/H/2888/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/005	7002499.00.00	SHIRE PHARMACEUTICALS IRELAND LIMITED	DE
Equasym Retard 50 mg Hartkapseln mit	DK/H/2888/005	7002499.00.00	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 Retard 50 mg kapslar med modifierad frisättning, hårda	DK/H/2888/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 50 mg kapslar med modifierad frisättning, hårda	DK/H/2888/005	30421	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg depotkapseli, kova	DK/H/2888/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg depotkapseli, kova	DK/H/2888/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg kapslar med modifierad frisättning, hårda	DK/H/2888/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym Retard 60 mg kapslar med modifierad frisättning, hårda	DK/H/2888/006	30422	SHIRE PHARMACEUTICALS IRELAND LIMITED	FI
Equasym XL 10 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/001	RVG 33227	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/001	RVG 33227	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/001	RVG 33227	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 10 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/001	RVG 33227	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 10 mg capsule met gereguleerde afgifte,	DK/H/2888/001	RVG 33227	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	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**
hard.				
Equasym XL 10 mg modified-release capsules, hard	DK/H/2888/001	PA 23211/001/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 10 mg modified-release capsules, hard	DK/H/2888/001	PA 23211/001/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 10 mg modified-release capsules, hard	DK/H/2888/001	PA 23211/001/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 10 mg modified-release capsules, hard	DK/H/2888/001	PA 23211/001/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 10 mg modified-release capsules, hard	DK/H/2888/001	PA 23211/001/001	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 20 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/002	RVG 33228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 20 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/002	RVG 33228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 20 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/002	RVG 33228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 20 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/002	RVG 33228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 20 mg modified-release capsules, hard	DK/H/2888/002	PA 23211/001/002	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 20 mg modified-release	DK/H/2888/002	PA 23211/001/002	TAKEDA PHARMACEUTICALS INTERNATIONAL AG	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			IRELAND BRANCH	
Equasym XL 20 mg modified-release capsules, hard	DK/H/2888/002	PA 23211/001/002	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 20 mg modified-release capsules, hard	DK/H/2888/002	PA 23211/001/002	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 20 mg modified-release capsules, hard	DK/H/2888/002	PA 23211/001/002	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 20 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/002	RVG 33228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/003	RVG 33229	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/003	RVG 33229	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/003	RVG 33229	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 30 mg capsule met gereguleerde afgifte, hard.	DK/H/2888/003	RVG 33229	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 30 mg modified-release capsules, hard	DK/H/2888/003	PA 23211/001/003	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 30 mg modified-release capsules, hard	DK/H/2888/003	PA 23211/001/003	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 30 mg	DK/H/2888/003	PA 23211/001/003	TAKEDA PHARMACEUTICALS	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
modified-release capsules, hard			INTERNATIONAL AG IRELAND BRANCH	
Equasym XL 30 mg modified-release capsules, hard	DK/H/2888/003	PA 23211/001/003	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 40 mg modified-release capsules, hard	DK/H/2888/004	PA 1575/001/007	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 40 mg modified-release capsules, hard	DK/H/2888/004	PA 23211/001/004	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 40 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/004	RVG 111228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 40 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/004	RVG 111228	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 50 mg modified-release capsules, hard	DK/H/2888/005	PA 23211/001/005	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 50 mg modified-release capsules, hard	DK/H/2888/005	PA 23211/001/005	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 50 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/005	RVG 111232	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 50 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/005	RVG 111232	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 60 mg modified-release capsules, hard	DK/H/2888/006	PA 23211/001/006	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	IE
Equasym XL 60 mg	DK/H/2888/006	PA 23211/001/006	TAKEDA PHARMACEUTICALS	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
modified-release capsules, hard			INTERNATIONAL AG IRELAND BRANCH	
Equasym XL 60 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/006	RVG 111233	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XL 60 mg, capsule met gereguleerde afgifte, hard.	DK/H/2888/006	RVG 111233	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	NL
Equasym XR 10 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg	DK/H/2888/001	BE423586	TAKEDA 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			INTERNATIONAL AG IRELAND BRANCH	
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	2013120615	SHIRE PHARMACEUTICALS IRELAND LIMITED	LU
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 10 mg gélules à libération modifiée	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 20 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg capsules met	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG	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.			IRELAND BRANCH	
Equasym XR 20 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 20 mg gélules à libération modifiée	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 30 mg	DK/H/2888/003	BE423604	TAKEDA 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
capsules met gereguleerde afgifte, hard.			INTERNATIONAL AG IRELAND BRANCH	
Equasym XR 30 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 30 mg gélules à libération modifiée	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 30 mg	DK/H/2888/003	2013120617	TAKEDA 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			INTERNATIONAL AG IRELAND BRANCH	
Equasym XR 40 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 40 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 40 mg gélules à libération modifiée	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 40 mg gélules à libération modifiée	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 40 mg gélules à libération modifiée	DK/H/2888/004	2013120618	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 40 mg gélules à libération modifiée	DK/H/2888/004	2013120618	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 50 mg capsules met gereguleerde afgifte, hard	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 50 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 50 mg gélules à libération modifiée	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 50 mg gélules à libération modifiée	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 50 mg	DK/H/2888/005	2013120619	TAKEDA 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			INTERNATIONAL AG IRELAND BRANCH	
Equasym XR 50 mg gélules à libération modifiée	DK/H/2888/005	2013120619	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 60 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/006	BE437613	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 60 mg capsules met gereguleerde afgifte, hard.	DK/H/2888/006	BE437613	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 60 mg gélules à libération modifiée	DK/H/2888/006	BE437613	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 60 mg gélules à libération modifiée	DK/H/2888/006	BE437613	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym XR 60 mg gélules à libération modifiée	DK/H/2888/006	2013120620	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym XR 60 mg gélules à libération modifiée	DK/H/2888/006	2013120620	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 10 mg Hartkapseln mit veränderter	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 10 mg Hartkapseln mit veränderter	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	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 10 mg Hartkapseln mit veränderter	DK/H/2888/001	BE423586	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/001	2013120615	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 20 mg Hartkapseln mit veränderter	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 20 mg Hartkapseln mit veränderter	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 20 mg Hartkapseln mit	DK/H/2888/002	BE423595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG	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			IRELAND BRANCH	
Equasym® XR 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/002	2013120616	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 30 mg Hartkapseln mit veränderter	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 30 mg Hartkapseln mit veränderter	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 30 mg Hartkapseln mit veränderter	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 30 mg Hartkapseln mit veränderter	DK/H/2888/003	BE423604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	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 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/003	2013120617	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 40 mg Hartkapseln mit veränderter	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 40 mg Hartkapseln mit veränderter	DK/H/2888/004	BE437595	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/004	2013120618	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 50 mg Hartkapseln mit veränderter	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 50 mg Hartkapseln mit veränderter	DK/H/2888/005	BE437604	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/005	2013120619	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/005	2013120619	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 60 mg	DK/H/2888/006	BE437613	TAKEDA 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
Hartkapseln mit veränderter			INTERNATIONAL AG IRELAND BRANCH	
Equasym® XR 60 mg Hartkapseln mit veränderter	DK/H/2888/006	BE437613	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	BE
Equasym® XR 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/006	2013120620	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Equasym® XR 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DK/H/2888/006	2013120620	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	LU
Medanef 10 mg tabletter	SE/H/1872/002	14-10263	MYLAN AB	NO
Medanef 10 mg tabletter	SE/H/1872/002	51606	MYLAN AB	SE
Medanef 20 mg tabletter	SE/H/1872/003	14-10264	MYLAN AB	NO
Medanef 20 mg tabletter	SE/H/1872/003	51607	MYLAN AB	SE
Medanef 5 mg tabletter	SE/H/1872/001	14-10262	MYLAN AB	NO
Medanef 5 mg tabletter	SE/H/1872/001	51605	MYLAN AB	SE
Medanef, tabletter	SE/H/1872/001	54633	MYLAN AB	DK
Medanef, tabletter	SE/H/1872/002	54634	MYLAN AB	DK
Medanef, tabletter	SE/H/1872/003	54635	MYLAN AB	DK
Medicebran 20 mg comprimidos	DE/H/2222/003	5397757	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medicebran 5 mg comprimidos	DE/H/2222/001	5397732	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
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	AIC N. 041438033	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 10 mg capsule	DE/H/2223/002	AIC N. 041438045	MEDICE ARZNEIMITTEL	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			PÜTTER GMBH & CO. KG	
Medikinet 10 mg kapsler med modifisert frisetting, harde	DE/H/0690/004	06-4119	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
MEDIKINET 10 mg, gélule 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	AIC N. 041438058	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 20 mg capsule rigide a rilascio modificato	DE/H/2223/003	AIC N. 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 tabletes	DE/H/2222/003	10-0460	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet 20 mg tabletės	DE/H/2222/003	LT/1/10/2267/006	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet 20 mg tabletės	DE/H/2222/003	LT/1/10/2267/005	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet 20 mg tablets	DE/H/2222/003	PA1555/001/008	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet 20 mg Tabletten	DE/H/2222/003	75793.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
MEDIKINET 20 mg,	DE/H/2223/003	CIS: 6 365 782 8	MEDICE ARZNEIMITTEL	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 modifiée			PÜTTER GMBH & CO. KG	
Medikinet 20 mg, kapsel med modifierad frisättning, hård	DE/H/0690/005	23841	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Medikinet 20 mg, tabletid	DE/H/2222/003	705410	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet 20 mg, tabletten	DE/H/2222/003	BE 381552	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
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 modificato	DE/H/2223/004	AIC N. 041438072	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 30 mg capsule rigide a rilascio modificato	DE/H/2223/004	AIC N. 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, gélule 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	DE/H/2223/005	AIC N. 041438096	MEDICE ARZNEIMITTEL	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			PÜTTER GMBH & CO. KG	
Medikinet 40 mg capsule rigide a rilascio modificato	DE/H/2223/005	AIC N. 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, gélule 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 libertação modificada	DE/H/2223/001	5361548	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Medikinet 5 mg cápsulas duras de liberación modificada	DE/H/0690/008	73308	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 5 mg capsule rigide a rilascio modificato	DE/H/2223/001	AIC N. 041438019	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IT
Medikinet 5 mg capsule rigide a rilascio modificato	DE/H/2223/001	AIC N. 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 tabletes	DE/H/2222/001	10-0447	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LV
Medikinet 5 mg tabletės	DE/H/2222/001	LT/1/10/2267/002	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet 5 mg tabletės	DE/H/2222/001	LT/1/10/2267/001	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet 5 mg tablets	DE/H/2222/001	PA1555/001/006	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IE
Medikinet 5 mg Tabletten	DE/H/2222/001	75791.00.00	MEDICE ARZNEIMITTEL	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
			PÜTTER GMBH & CO. KG	
MEDIKINET 5 mg, gélule 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 5 mg, tabletid	DE/H/2222/001	705210	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Medikinet 5 mg, tabletten	DE/H/2222/001	BE 381534	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet 50 mg cápsulas duras de liberación modificada	DE/H/0690/009	78453	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	78454	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Medikinet 60 mg kapsler med modifisert frisetting, harde	DE/H/0690/010	12-9268	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
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	63890.00.00	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 ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 40 mg Hartkapseln, retardiert	not available	65965.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 5 mg	not available	66747.00.00	MEDICE 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
Hartkapseln, retardiert			CO. KG	
Medikinet adult 50 mg Hartkapseln, retardiert	not available	88862.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet adult 60 mg Hartkapseln, retardiert	not available	88863.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Medikinet CR 10 mg hörð hylki með breyttan losunarhraða.	DE/H/0690/004	IS/1/14/087/05	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 10 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/004	12844	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 10 mg säädellysti vapauttava kapseli, kova	DE/H/0690/004	22212	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
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	IS/1/14/087/06	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	IS/1/14/087/07	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 30 mg	DE/H/0690/006	12846	MEDICE ARZNEIMITTEL	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
kapsułki o zmodyfikowanym uwalnianiu, twarde			PÜTTER GMBH & CO. KG	
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 losunarhraða.	DE/H/0690/007	IS/1/14/087/08	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
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	IS/1/14/087/04	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,	DE/H/0690/008	RVG 108026	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
hard				
Medikinet CR 50 mg hörð hylki með breyttan losunarhraða	DE/H/0690/009	IS/1/14/087/09	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 50 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/009	21934	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 50 mg säädellysti vapauttava kapseli, kova	DE/H/0690/009	31130	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
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	IS/1/14/087/10	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Medikinet CR 60 mg kapsułki o zmodyfikowanym uwalnianiu, twarde	DE/H/0690/010	21935	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Medikinet CR 60 mg säädellysti vapauttava kapseli, kova	DE/H/0690/010	31131	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
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/0690/008	47393	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	DE/H/0690/009	51661	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
udløsning				
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/0690/010	51662	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 MR 10 mg modified-release capsules, hard	DE/H/2223/002	PA1555/001/002	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
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	DE/H/0690/004	54569.00.01	MEDICE ARZNEIMITTEL	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
Hartkapseln mit veränderter Wirkstofffreisetzung			PÜTTER GMBH & CO. KG	
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 Wirkstofffreisetzung	DE/H/0690/005	1363/07020043	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Medikinet retard 20 mg, Capsules met gereguleerde afgifte, hard	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	DE/H/0690/006	62567.00.00	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 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	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/0690/007	62568.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
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 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	BE 381604	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	BE
Medikinet retard 5 mg Hartkapseln mit veränderter	DE/H/0690/008	1-30056	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
Wirkstofffreisetzung				
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 veränderter Wirkstofffreisetzung	DE/H/0690/009	88767.00.00	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	DE/H/0690/010	88768.00.00	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
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	XI
Medikinet XL 10 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/002	LT/1/10/2274/003	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 10 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/002	705110	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
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	XI
Medikinet XL 20 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/003	LT/1/10/2274/005	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 20 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/003	705010	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
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	DE/H/0690/006	PL 11243/0007	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI

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				
Medikinet XL 30 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/004	LT/1/10/2274/007	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 30 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/004	705310	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
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	XI
Medikinet XL 40 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/005	LT/1/10/2274/009	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 40 mg, toimeainet modifitseeritult vabastavad kõvakapslid	DE/H/2223/005	705710	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
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 modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/001	LT/1/10/2274/001	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 5 mg modifikuoto atpalaidavimo kietosios kapsulės	DE/H/2223/001	LT/1/10/2274/002	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LT
Medikinet XL 5 mg, toimeainet modifitseeritult	DE/H/2223/001	705610	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
vabastavad kõvakapslid				
Medikinet XL 50 mg modified-release capsules, hard	DE/H/0690/009	PL 11243/0011	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI
Medikinet XL 60 mg modified-release capsules, hard	DE/H/0690/010	PL 11243/0012	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI
Methylfenidaat HCl Mylan 10 mg, tabletten	SE/H/1872/002	RVG 116233	MYLAN B.V.	NL
Methylfenidaat HCl Mylan 5 mg, tabletten	SE/H/1872/001	RVG 116232	MYLAN B.V.	NL
Methylfenidaat HCl retard Aristo 10 mg, harde capsules met gereguleerde afgifte	DE/H/6353/001/DC	RVG 125055	ARISTO PHARMA GMBH (ART 57)	NL
Methylfenidaat HCl retard Aristo 20 mg, harde capsules met gereguleerde afgifte	DE/H/6353/002/DC	RVG 125056	ARISTO PHARMA GMBH (ART 57)	NL
Methylfenidaat HCl retard Aristo 30 mg, harde capsules met gereguleerde afgifte	DE/H/6353/003/DC	RVG 125057	ARISTO PHARMA GMBH (ART 57)	NL
Methylfenidaat HCl retard Aristo 40 mg, harde capsules met	DE/H/6353/004/DC	RVG 125058	ARISTO PHARMA GMBH (ART 57)	NL
Methylfenidaat HCl retard Aristo 50 mg, harde capsules met gereguleerde afgifte	DE/H/6353/005/DC	RVG 125059	ARISTO PHARMA GMBH (ART 57)	NL
Methylfenidaat XR EG 10 mg gélules à libération modifiée	DE/H/6338/001	BE571760	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 10 mg gélules à libération modifiée	DE/H/6338/001	2020120334	EUROGENERICS N.V./S.A.	LU
Methylfenidaat XR EG 10	DE/H/6338/001	BE571760	EUROGENERICS N.V./S.A.	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
mg harde capsules met gereguleerde afgifte				
Methylfenidaat XR EG 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/001	BE571760	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 20 mg gélules à libération modifiée	DE/H/6338/002	BE571777	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 20 mg gélules à libération modifiée	DE/H/6338/002	2020120335	EUROGENERICS N.V./S.A.	LU
Methylfenidaat XR EG 20 mg harde capsules met gereguleerde afgifte	DE/H/6338/002	BE571777	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/002	BE571777	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 30 mg gélules à libération modifiée	DE/H/6338/003	BE571786	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 30 mg gélules à libération modifiée	DE/H/6338/003	2020120336	EUROGENERICS N.V./S.A.	LU
Methylfenidaat XR EG 30 mg harde capsules met gereguleerde afgifte	DE/H/6338/003	BE571786	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/003	BE571786	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 40 mg gélules à libération modifiée	DE/H/6338/004	BE571795	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 40 mg gélules à libération	DE/H/6338/004	2020120337	EUROGENERICS N.V./S.A.	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
modifiée				
Methylfenidaat XR EG 40 mg harde capsules met gereguleerde afgifte	DE/H/6338/004	BE571795	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/004	BE571795	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 50 mg gélules à libération modifiée	DE/H/6338/005	BE571804	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 50 mg gélules à libération modifiée	DE/H/6338/005	2020120338	EUROGENERICS N.V./S.A.	LU
Methylfenidaat XR EG 50 mg harde capsules met gereguleerde afgifte	DE/H/6338/005	BE571804	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/005	BE571804	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 60 mg gélules à libération modifiée	DE/H/6338/006	BE571813	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 60 mg gélules à libération modifiée	DE/H/6338/006	2020120339	EUROGENERICS N.V./S.A.	LU
Methylfenidaat XR EG 60 mg harde capsules met gereguleerde afgifte	DE/H/6338/006	BE571813	EUROGENERICS N.V./S.A.	BE
Methylfenidaat XR EG 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/006	BE571813	EUROGENERICS N.V./S.A.	BE
Methylphenidat "Alternova", tabletter	not available	54924	ALTERNOVA A/S	DK
Methylphenidat	not available	54926	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				
Methylphenidate "Sandoz", depottabletter	DK/H/2138/001	49666	SANDOZ A/S	DK
Methylphenidate "Sandoz", depottabletter	DK/H/2138/001	49666	SANDOZ A/S	DK
Methylphenidate "Aristo" 10 mg kapsler med modificeret udløsning, hårde	DE/H/6353/001	62796	ARISTO PHARMA GMBH (ART 57)	DK
Methylphenidate "Aristo" 20 mg hårde kapsler med modificeret udløsning	DE/H/6353/002	62797	ARISTO PHARMA GMBH (ART 57)	DK
Methylphenidate "Aristo" 30 mg hårde kapsler med modificeret udløsning	DE/H/6353/003	62798	ARISTO PHARMA GMBH (ART 57)	DK
Methylphenidate Alternova 20 mg tabletter	not available	51911	ALTERNOVA A/S	SE
Methylphenidate Aristo 10 mg kapsel med modifisert frisetting, hard	DE/H/6353/001/DC	19-12898	ARISTO PHARMA GMBH (ART 57)	NO
Methylphenidate Aristo 20 mg kapsel med modifisert frisetting, hard	DE/H/6353/002/DC	19-12899	ARISTO PHARMA GMBH (ART 57)	NO
Methylphenidate Aristo 30 mg kapsler med modifisert frisetting, harde	DE/H/6353/003/DC	19-12900	ARISTO PHARMA GMBH (ART 57)	NO
Methylphenidate Hydrochloride 10 mg Tablets	SE/H/1872/002	PL 04569/1497	GENERICS [UK] LIMITED	XI
Methylphenidate Hydrochloride 20 mg	SE/H/1872/003	PL 04569/1498	GENERICS [UK] LIMITED	XI

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
Tablets				
Methylphenidate Hydrochloride 5 mg Tablets	SE/H/1872/001	PL 04569/1496	GENERICS [UK] LIMITED	XI
Methylphenidate Mylan 10 mg comprimés	SE/H/1872/002	BE489404	MYLAN BV/SRL	BE
Methylphenidate Mylan 10 mg tabletten	SE/H/1872/002	BE489404	MYLAN BV/SRL	BE
Methylphenidate Mylan 10 mg Tabletten	SE/H/1872/002	BE489404	MYLAN BV/SRL	BE
Methylphenidate Mylan 20 mg comprimés	SE/H/1872/003	BE489413	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 20 mg tabletten	SE/H/1872/003	BE489413	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 20 mg Tabletten	SE/H/1872/003	BE489413	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 5 mg comprimés	SE/H/1872/001	BE489395	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 5 mg tabletten	SE/H/1872/001	BE489395	MYLAN BVBA/SPRL	BE
Methylphenidate Mylan 5 mg Tabletten	SE/H/1872/001	BE489395	MYLAN BVBA/SPRL	BE
Methylphenidat-HCl MEDICE 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2203637.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methylphenidat-HCl MEDICE 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2203638.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methylphenidat-HCl MEDICE 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2203639.00.00	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
Methylphenidat-HCl MEDICE 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2203640.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methylphenidat-HCl MEDICE 5 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	2203636.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methylphenidat-HCl MEDICE 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	97387.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methylphenidat-HCl MEDICE 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	97388.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Methysym 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6353/001/DC	2204196.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Methysym 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6353/002/DC	2204197.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Methysym 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6353/003/DC	2204198.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Methysym 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6353/004/DC	2204199.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Methysym 50 mg	DE/H/6353/005/DC	2204200.00.00	ARISTO 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
Hartkapseln mit veränderter Wirkstofffreisetzung			(ART 57)	
Methysym 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6353/006/DC	2204201.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Methysym Retard 10 mg cápsulas duras de liberación modificada	DE/H/6353/001	85.523	ARISTO PHARMA GMBH (ART 57)	ES
Methysym Retard 20 mg cápsulas duras de liberación modificada	DE/H/6353/002	85.524	ARISTO PHARMA GMBH (ART 57)	ES
Methysym Retard 30 mg cápsulas duras de liberación modificada	DE/H/6353/003	85.525	ARISTO PHARMA GMBH (ART 57)	ES
Methysym Retard 40 mg cápsulas duras de liberación modificada	DE/H/6353/004	85.526	ARISTO PHARMA GMBH (ART 57)	ES
Methysym Retard 50 mg cápsulas duras de liberación modificada	DE/H/6353/005	85.527	ARISTO PHARMA GMBH (ART 57)	ES
Methysym Retard 60 mg cápsulas duras de liberación modificada	DE/H/6353/006	85.528	ARISTO PHARMA GMBH (ART 57)	ES
Metiphenal 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/001	2204112.00.00	ALIUD PHARMA GMBH	DE
Metiphenal 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/002	2204113.00.00	ALIUD PHARMA GMBH	DE
Metiphenal 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/003	2204114.00.00	ALIUD 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
Metiphenal 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/004	2204115.00.00	ALIUD PHARMA GMBH	DE
Metiphenal 50 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/005	2204116.00.00	ALIUD PHARMA GMBH	DE
Metiphenal 60 mg Hartkapseln mit veränderter Wirkstofffreisetzung	DE/H/6338/006	2204117.00.00	ALIUD PHARMA GMBH	DE
Metynor 10 mg hårda kapslar med modifierad frisättning.	DE/H/6338/001	59232	STADA ARZNEIMITTEL AG	SE
Metynor 20 mg hårda kapslar med modifierad frisättning.	DE/H/6338/002	59233	STADA ARZNEIMITTEL AG	SE
Metynor 30 mg hårda kapslar med modifierad frisättning.	DE/H/6338/003	59234	STADA ARZNEIMITTEL AG	SE
Metynor 40 mg hårda kapslar med modifierad frisättning.	DE/H/6338/004	59235	STADA ARZNEIMITTEL AG	SE
Metynor 50 mg hårda kapslar med modifierad frisättning.	DE/H/6338/005	59236	STADA ARZNEIMITTEL AG	SE
Metynor 60 mg hårda kapslar med modifierad frisättning.	DE/H/6338/006	59237	STADA ARZNEIMITTEL AG	SE
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 10 mg cápsulas	UK/H/0819/001	5436563	TAKEDA PHARMACEUTICALS	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			INTERNATIONAL AG IRELAND BRANCH	
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 10 mg cápsulas de libertação modificada	UK/H/0819/001	5436563	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 20 mg cápsulas de libertação modificada	UK/H/0819/002	5436571	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 30 mg cápsulas de libertação modificada	UK/H/0819/003	5436605	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 40 mg cápsulas de libertação modificada	UK/H/0819/004	5550462	TAKEDA PHARMACEUTICALS INTERNATIONAL AG	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
			IRELAND BRANCH	
Quasym 40 mg cápsulas de libertação modificada	UK/H/0819/004	5550462	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 50 mg cápsulas de libertação modificada	UK/H/0819/005	5550454	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 50 mg cápsulas de libertação modificada	UK/H/0819/005	5550454	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 60 mg cápsulas de libertação modificada	UK/H/0819/006	5550470	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
Quasym 60 mg cápsulas de libertação modificada	UK/H/0819/006	5550470	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	PT
QUASYM L.P. 10 mg, gélule à libération modifiée	DK/H/2888/001	34009 377 617 8 9	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	DK/H/2888/001	34009 497 261 6 5	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	DK/H/2888/001	34009 377 618 4 0	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	DK/H/2888/001	34009 570 268 1 3	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 10 mg, gélule à libération modifiée	DK/H/2888/001	34009 570 269 8 1	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	DK/H/2888/002	34009 377 620 9 0	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	DK/H/2888/002	34009 377 619 0 1	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	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. 20 mg, gélule à libération modifiée	DK/H/2888/002	497 262-2	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	DK/H/2888/002	34009 378 313 2 1	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 20 mg, gélule à libération modifiée	DK/H/2888/002	34009 570 271 2 4	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	DK/H/2888/003	34009 377 621 5 1	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	DK/H/2888/003	34009 497 267 4 5	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	DK/H/2888/003	34009 377 622 1 2	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
QUASYM L.P. 30 mg, gélule à libération modifiée	DK/H/2888/003	34009 378 314 9 9	TAKEDA PHARMACEUTICALS INTERNATIONAL AG IRELAND BRANCH	FR
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	BE051597	NOVARTIS PHARMA N.V.	BE
RILATINE 10 mg tabletten	not available	BE051597	NOVARTIS PHARMA N.V.	BE
RILATINE 10 mg, comprimés	not available	BE051597	NOVARTIS PHARMA N.V.	BE
RILATINE 10 mg, comprimés	not available	BE051597	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 10 mg Hartkapseln mit veränderter	not available	BE426413	NOVARTIS PHARMA N.V.	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
Wirkstofffreisetzung				
RILATINE Modified Release 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE426413	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 10 mg, capsules met gereguleerde afgifte, hard	not available	BE426413	NOVARTIS PHARMA N.V.	BE
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	BE426413	NOVARTIS PHARMA N.V.	BE
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	BE241534	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 20 mg, capsules met gereguleerde afgifte, hard	not available	BE241534	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 20 mg, capsules met gereguleerde afgifte,	not available	BE241534	NOVARTIS PHARMA N.V.	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				
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	BE241534	NOVARTIS PHARMA N.V.	BE
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	BE241543	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 30 mg, capsules met gereguleerde afgifte, hard	not available	BE241543	NOVARTIS PHARMA N.V.	BE
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	BE241543	NOVARTIS PHARMA N.V.	BE
RILATINE Modified Release 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	BE241552	NOVARTIS PHARMA N.V.	BE
RILATINE Modified	not available	BE241552	NOVARTIS PHARMA N.V.	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
Release 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung				
RILATINE Modified Release 40 mg, capsules met gereguleerde afgifte, hard	not available	BE241552	NOVARTIS PHARMA N.V.	BE
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	BE241552	NOVARTIS PHARMA N.V.	BE
Ritalin 10 mg hårda kapslar med modifierad frisättning	not available	41556	NOVARTIS SVERIGE AB	SE
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 kapsler med modifisert frisetting, harde	not available	09-6501	NOVARTIS NORGE AS	NO
Ritalin 10 mg tabletes	not available	00-0118	SIA NOVARTIS BALTICS	LV
RITALIN 10 mg Tablets	not available	MA1249/01801	NOVARTIS IRELAND LIMITED	MT
RITALIN 10 mg Tablets	not available	MA1249/01801	NOVARTIS IRELAND LIMITED	MT
Ritalin 10 mg tabletta	not available	OGYI-T-6954/01	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin 10 mg tabletta	not available	OGYI-T-6954/01	NOVARTIS HUNGÁRIA KFT.	HU

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 10 mg tabletter	not available	3449	NOVARTIS NORGE AS	NO
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 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 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 10 mg töflur	not available	640044	NOVARTIS HEALTHCARE A/S	IS
Ritalin 20 mg hårda kapslar med modifierad frisättning	not available	20607	NOVARTIS SVERIGE AB	SE
Ritalin 20 mg hårda kapslar med modifierad frisättning	not available	20607	NOVARTIS SVERIGE AB	SE
Ritalin 20 mg kapsler med modifisert frisetting, harde	not available	04-2876	NOVARTIS NORGE AS	NO
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 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 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 hårda	not available	20609	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
kapslar med modifierad frisättning				
Ritalin 40 mg kapsler med modifisert frisetting, harde	not available	05-3198	NOVARTIS NORGE AS	NO
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 hårda kapslar med modifierad frisättning	not available	50408	NOVARTIS SVERIGE AB	SE
Ritalin 60 mg kapsler med modifisert frisetting, harde	not available	13-9867	NOVARTIS NORGE AS	NO
Ritalin 60 mg kapsler med modifisert frisetting, harde	not available	13-9867	NOVARTIS NORGE AS	NO
Ritalin LA 10 mg, harde capsules met gereguleerde afgifte	not available	RVG 116377	NOVARTIS PHARMA B.V.	NL
Ritalin LA 10 mg, harde capsules met gereguleerde afgifte	not available	RVG 116377	NOVARTIS PHARMA B.V.	NL
Ritalin LA 20 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/02	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 20 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/02	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 20 mg, harde capsules met gereguleerde afgifte	not available	RVG 116379	NOVARTIS PHARMA B.V.	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
Ritalin LA 20 mg, harde capsules met gereguleerde afgifte	not available	RVG 116379	NOVARTIS PHARMA B.V.	NL
Ritalin LA 30 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/03	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 30 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/03	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 30 mg, harde capsules met gereguleerde afgifte	not available	RVG 116380	NOVARTIS PHARMA B.V.	NL
Ritalin LA 30 mg, harde capsules met gereguleerde afgifte	not available	RVG 116380	NOVARTIS PHARMA B.V.	NL
Ritalin LA 40 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/04	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 40 mg módosított hatóanyagleadású kemény kapszula	not available	OGYI-T-6954/04	NOVARTIS HUNGÁRIA KFT.	HU
Ritalin LA 40 mg, harde capsules met gereguleerde afgifte	not available	RVG 116381	NOVARTIS PHARMA B.V.	NL
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.	HU
Ritalin LA 60 mg módosított	not available	OGYI-T-6954/05	NOVARTIS HUNGÁRIA KFT.	HU

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
hatóanyagleadású kemény kapszula				
Ritalin LA 60 mg, harde capsules met gereguleerde afgifte	not available	RVG 116382	NOVARTIS PHARMA B.V.	NL
Ritalin LA 60 mg, harde capsules met gereguleerde afgifte	not available	RVG 116382	NOVARTIS PHARMA B.V.	NL
Ritalin Uno 10 mg hart hylki með breyttan losunarhraða	not available	IS/1/10/133/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 10 mg hart hylki með breyttan losunarhraða	not available	IS/1/10/133/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 20 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 20 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 30 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/02	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 30 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/02	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 40 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/03	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 40 mg hart hylki með breyttan losunarhraða	not available	IS/1/02/127/03	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 60 mg hart hylki með breyttan losunarhraða	not available	IS/1/15/020/01	NOVARTIS HEALTHCARE A/S	IS
Ritalin Uno 60 mg hart hylki með breyttan	not available	IS/1/15/020/01	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
losunarhraða				
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	53335	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34355	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34356	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34357	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	44662	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	53335	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34355	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34356	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	34357	NOVARTIS HEALTHCARE A/S	DK
Ritalin Uno, hårde kapsler med modificeret udløsning	not available	44662	NOVARTIS HEALTHCARE A/S	DK
Ritalin, tabletten 10 mg	not available	RVG 03957	NOVARTIS PHARMA B.V.	NL
Ritalin, tabletten 10 mg	not available	RVG 03957	NOVARTIS PHARMA B.V.	NL
Ritalin, tabletter 10 mg	not available	1870	NOVARTIS HEALTHCARE A/S	DK
Ritalin, tabletter 10 mg	not available	1870	NOVARTIS HEALTHCARE A/S	DK
Ritalin® 10 mg - Tabletten	not available	1-22028	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
Ritalin® 10 mg - Tabletten	not available	1-22028	NOVARTIS PHARMA GMBH	AT
Ritalin® 10 mg Tabletten	not available	6094573.00.00	NOVARTIS PHARMA GMBH	DE
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 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 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 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
Ritalin® Adult 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67254.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 10 mg -	not available	1-31212	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 10 mg - Kapseln	not available	1-31212	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 10 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	77239.00.00	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 - Kapseln	not available	1-24270	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 20 mg - Kapseln	not available	1-24270	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67249.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 20 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67249.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 20mg, prolonged-release capsules.	not available	PA0896/026/002	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 20mg, prolonged-release capsules.	not available	PA0896/026/002	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 30 mg - Kapseln	not available	1-24271	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 30 mg - Kapseln	not available	1-24271	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67250.00.00	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® LA 30 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67250.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 30mg, prolonged-release capsules.	not available	PA0896/026/003	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 30mg, prolonged-release capsules.	not available	PA0896/026/003	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 40 mg - Kapseln	not available	1-24272	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 40 mg - Kapseln	not available	1-24272	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67251.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 40 mg Hartkapseln mit veränderter Wirkstofffreisetzung	not available	67251.00.00	NOVARTIS PHARMA GMBH	DE
Ritalin® LA 40mg, prolonged-release capsules.	not available	PA0896/026/004	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 40mg, prolonged-release capsules.	not available	PA0896/026/004	NOVARTIS IRELAND LIMITED	IE
Ritalin® LA 60mg - Kapseln	not available	138502	NOVARTIS PHARMA GMBH	AT
Ritalin® LA 60mg - Kapseln	not available	138502	NOVARTIS PHARMA GMBH	AT
RITALINE 10 mg, comprimé sécable	not available	34009 339 294 0 4	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 424 1 0	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg,	not available	34009 339 426 4 9	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
comprimé sécable				
RITALINE 10 mg, comprimé sécable	not available	34009 339 294 0 4	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 424 1 0	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 426 4 9	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 294 0 4	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 424 1 0	NOVARTIS PHARMA S.A.S.	FR
RITALINE 10 mg, comprimé sécable	not available	34009 339 426 4 9	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 865 3 5	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 867 6 4	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 579 745 7 2	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 865 3 5	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 867 6 4	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 579 745 7 2	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 865 3 5	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 416 867 6 4	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
RITALINE L.P. 10 mg, gélule à libération prolongée	not available	34009 579 745 7 2	NOVARTIS PHARMA S.A.S.	FR
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	3400936534933	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	3400936231382	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. 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	3400936231382	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. 20 mg, gélule à libération prolongée	not available	3400956496914	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	3400936535015	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 30 mg,	not available	3400956497164	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. 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	3400936535015	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. 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	3400936535015	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	3400936231733	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	3400956497225	NOVARTIS PHARMA S.A.S.	FR
RITALINE L.P. 40 mg, gélule à libération prolongée	not available	3400936231733	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	not available	3400956497225	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
prolongée				
RITALINE L.P. 40 mg, gélule à libération prolongée	not available	3400936231733	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	3400956497225	NOVARTIS PHARMA S.A.S.	FR
Концерта 18 mg таблетки с удължено освобождаване	DE/H/6007 001	20060458	JOHNSON & JOHNSON D.O.O.	BG