19 June 2025
EMADOC-1700519818-2340477
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): methylphenidate

EURD list No. PSUSA/00002024/202410

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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
5 mg, Medikinet	DE/H/2222/001		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
50 mg, Medikinet adult		88862.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Atenza	SE/H/1912/005	89.260	Exeltis Healthcare S.L.	Kingdom of Spain
Atenza	SE/H/1912/005	28248	Exeltis Poland Sp. z. o.o.	Republic of Poland
Concerta	DE/H/6007003	06/409/08-C	Janssen-Cilag s.r.o.	Czech Republic
Concerta	DE/H/6007002	06/408/08-C	Janssen-Cilag s.r.o.	Czech Republic
Concerta	DE/H/6007001	06/407/08-C	Janssen-Cilag s.r.o.	Czech Republic
Concerta	DE/H/6007003	54861.02.00	Janssen-Cilag GmbH	Federal Republic of Germany
Concerta	DE/H/6007002	54861.01.00	Janssen-Cilag GmbH	Federal Republic of Germany
Concerta	DE/H/6007004	72263.00.00	Janssen-Cilag GmbH	Federal Republic of Germany
Concerta	DE/H/6007001	54861.00.00	Janssen-Cilag GmbH	Federal Republic of Germany
Concerta	DE/H/6007004	2008120051	Janssen-Cilag N.V.	Grand Duchy of Luxembourg
Concerta	DE/H/6007003	2008100012	Janssen-Cilag N.V.	Grand Duchy of Luxembourg
Concerta	DE/H/6007001	2008100010	Janssen-Cilag N.V.	Grand Duchy of Luxembourg
Concerta	DE/H/6007002	2008100011	Janssen-Cilag N.V.	Grand Duchy of Luxembourg
Concerta	DE/H/6007004	12585/19.02.2015	Janssen Cilag Pharmaceutical S.A.C.I.	Hellenic Republic
Concerta	DE/H/6007002	BE242697	Janssen-Cilag N.V.	Kingdom of Belgium
Concerta	DE/H/6007001	BE242681	Janssen-Cilag N.V.	Kingdom of Belgium
Concerta	DE/H/6007003	BE242706	Janssen-Cilag N.V.	Kingdom of Belgium
Concerta	DE/H/6007004	BE324913	Janssen-Cilag N.V.	Kingdom of Belgium
Concerta	DE/H/6007003	42320	Janssen-Cilag A/S	Kingdom of Denmark
Concerta	DE/H/6007001	42318	Janssen-Cilag A/S	Kingdom of Denmark
Concerta	DE/H/6007002	42319	Janssen-Cilag A/S	Kingdom of Denmark
Concerta	DE/H/6007001	02-1221	Janssen-Cilag AS	Kingdom of Norway
Concerta	DE/H/6007004	08-5646	Janssen-Cilag AS	Kingdom of Norway

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	DE/H/6007002	02-1222	Janssen-Cilag AS	Kingdom of Norway
Concerta	DE/H/6007003	02-1223	Janssen-Cilag AS	Kingdom of Norway
Concerta	DE/H/6007004	69.988	Janssen Cilag S.A.	Kingdom of Spain
Concerta	DE/H/6007003	65.171	Janssen Cilag S.A.	Kingdom of Spain
Concerta	DE/H/6007001	65.148	Janssen Cilag S.A.	Kingdom of Spain
Concerta	DE/H/6007002	65.170	Janssen Cilag S.A.	Kingdom of Spain
Concerta	DE/H/6007003	18543	Janssen-Cilag AB	Kingdom of Sweden
Concerta	DE/H/6007002	18542	Janssen-Cilag AB	Kingdom of Sweden
Concerta	DE/H/6007001	18541	Janssen-Cilag AB	Kingdom of Sweden
Concerta	DE/H/6007004	26554	Janssen-Cilag AB	Kingdom of Sweden
Concerta	DE/H/6007002	RVG 28074	Janssen-Cilag B.V.	Kingdom of the Netherlands
Concerta	DE/H/6007001	RVG 28073	Janssen-Cilag B.V.	Kingdom of the Netherlands
Concerta	DE/H/6007004	RVG 101739	Janssen-Cilag B.V.	Kingdom of the Netherlands
Concerta	DE/H/6007003	RVG 28075	Janssen-Cilag B.V.	Kingdom of the Netherlands
Concerta	DE/H/6007004		Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007001	4260782	Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007003	4261186	Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007003	4261285	Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007001	4260881	Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007002		Janssen - Cilag Farmacêutica Lda.	Portuguese Republic
Concerta	DE/H/6007002	1-24813	Janssen-Cilag Pharma GmbH	Republic of Austria
Concerta	DE/H/6007003	1-24814	Janssen-Cilag Pharma GmbH	Republic of Austria
Concerta	DE/H/6007001	1-24812	Janssen-Cilag Pharma GmbH	Republic of Austria
Concerta	DE/H/6007004	1-27727	Janssen-Cilag Pharma GmbH	Republic of Austria
Concerta		HR-H-413338677	Johnson & Johnson S.E.	Republic of Croatia

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
			d.o.o.	
Concerta		HR-H-078951991	Johnson & Johnson S.E. d.o.o.	Republic of Croatia
Concerta	DE/H/6007002	20340	Janssen Cilag International	Republic of Cyprus
Concerta	DE/H/6007001	20339	Janssen Cilag International	Republic of Cyprus
Concerta	DE/H/6007003	20336	Janssen Cilag International	Republic of Cyprus
Concerta	DE/H/6007003	583808	UAB Johnson And Johnson	Republic of Estonia
Concerta	DE/H/6007002	583708	UAB Johnson And Johnson	Republic of Estonia
Concerta	DE/H/6007001	583608	UAB Johnson And Johnson	Republic of Estonia
Concerta	DE/H/6007001	17408	Janssen-Cilag Oy	Republic of Finland
Concerta	DE/H/6007002	17409	Janssen-Cilag Oy	Republic of Finland
Concerta	DE/H/6007003	17410	Janssen-Cilag Oy	Republic of Finland
Concerta	DE/H/6007004	24631	Janssen-Cilag Oy	Republic of Finland
Concerta	DE/H/6007002	IS/1/02/032/02	Janssen-Cilag AB	Republic of Iceland
Concerta	DE/H/6007003	IS/1/02/032/03	Janssen-Cilag AB	Republic of Iceland
Concerta	DE/H/6007004	IS/1/08/032/01	Janssen-Cilag AB	Republic of Iceland
Concerta	DE/H/6007001	IS/1/02/032/01	Janssen-Cilag AB	Republic of Iceland
CONCERTA	DE/H/6007003		UAB Johnson And Johnson	Republic of Lithuania
CONCERTA	DE/H/6007001		UAB Johnson And Johnson	Republic of Lithuania
CONCERTA	DE/H/6007002		UAB Johnson And Johnson	Republic of Lithuania
Concerta	DE/H/6007001	MA018/02601	Janssen Cilag International	Republic of Malta
Concerta	DE/H/6007002	MA018/02602	Janssen Cilag International	Republic of Malta
Concerta	DE/H/6007003	MA018/02603	Janssen Cilag International	Republic of Malta
Concerta	DE/H/6007001	14751	Janssen Cilag International	Republic of Poland
Concerta	DE/H/6007002	14752	Janssen Cilag International	Republic of Poland
Concerta	DE/H/6007001		Johnson & Johnson d.o.o.	Republic of Slovenia
Concerta	DE/H/6007002		Johnson & Johnson d.o.o.	Republic of Slovenia
Concerta	DE/H/6007003		Johnson & Johnson d.o.o.	Republic of Slovenia
Concerta	DE/H/6007001	78/0126/08-S	Johnson & Johnson s.r.o.	Slovak Republic
Concerta	DE/H/6007003	78/0128/08-S	Johnson & Johnson s.r.o.	Slovak Republic
Concerta	DE/H/6007002	78/0127/08-S	Johnson & Johnson s.r.o.	Slovak Republic
CONCERTA LP	DE/H/6007002	34009 361 555 8 9		French Republic
CONCERTA LP	DE/H/6007001	34009 361 554 1 1		French Republic
CONCERTA LP	DE/H/6007003	34009 361 556 4 0		French Republic
Concerta XL	DE/H/6007001	PL 00242/0372	Janssen-Cilag Limited	United Kingdom

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
				(Northern Ireland)
Concerta XL	DE/H/6007002	PL 00242/0373	Janssen-Cilag Limited	United Kingdom (Northern Ireland)
Concerta XL	DE/H/6007004	PL 00242/0400	Janssen-Cilag Limited	United Kingdom (Northern Ireland)
Concerta XL	DE/H/6007003	PL 00242/0374	Janssen-Cilag Limited	United Kingdom (Northern Ireland)
Equasym	DK/H/2888/006		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/001		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/003		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/005		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/002		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/004		Takeda Pharmaceuticals International AG	Italian Republic
Equasym	DK/H/2888/004	77163	Takeda Pharmaceuticals International AG	Kingdom of Spain
Equasym	DK/H/2888/005	77164	Takeda Pharmaceuticals International AG	Kingdom of Spain
Equasym	DK/H/2888/003	76060	Takeda Pharmaceuticals International AG	Kingdom of Spain
Equasym	DK/H/2888/002	76076	Takeda Pharmaceuticals International AG	Kingdom of Spain
Equasym	DK/H/2888/001	76061	Takeda Pharmaceuticals International AG	Kingdom of Spain
Equasym Depot	UK/H/0819/001	38534	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Denmark
Equasym Depot	UK/H/0819/002	38535	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Denmark
Equasym Depot	UK/H/0819/003	38536	[INACTIVE] Takeda	Kingdom of Denmark

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
			Pharmaceuticals International AG	
Equasym Depot	UK/H/0819/005	50038	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Denmark
Equasym Depot	UK/H/0819/004	50037	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Denmark
Equasym Depot	UK/H/0819/006	50039	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Denmark
Equasym Depot	UK/H/0819/001	05-3711	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	UK/H/0819/005	11-8781	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	UK/H/0819/004	11-8780	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	UK/H/0819/006	11-8782	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	UK/H/0819/002	05-3712	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	UK/H/0819/003	05-3713	Takeda Pharmaceuticals International AG	Kingdom of Norway
Equasym Depot	DK/H/2888/001	IS/1/06/003/01	[INACTIVE] Takeda Pharmaceuticals International AG	Republic of Iceland
Equasym Depot	DK/H/2888/002	IS/1/06/003/02	[INACTIVE] Takeda Pharmaceuticals International AG	Republic of Iceland
Equasym Depot	DK/H/2888/003	IS/1/06/003/03	[INACTIVE] Takeda Pharmaceuticals International AG	Republic of Iceland
Equasym Retard	DK/H/2888/001	64003.00.00	Takeda Pharmaceuticals International AG	Federal Republic of Germany
Equasym Retard	DK/H/2888/004	7002498.00.00	Takeda Pharmaceuticals International AG	Federal Republic of Germany

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	DK/H/2888/005	7002499.00.00	Takeda Pharmaceuticals International AG	Federal Republic of Germany
Equasym Retard	DK/H/2888/002	64004.00.00	Takeda Pharmaceuticals International AG	Federal Republic of Germany
Equasym Retard	DK/H/2888/003	64005.00.00	Takeda Pharmaceuticals International AG	Federal Republic of Germany
Equasym Retard	DK/H/2888/005	30421	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/003	19259	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/002	19258	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/001	19257	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/002	19258	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/004	30420	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym Retard	DK/H/2888/006	30422	Takeda Pharmaceuticals International AG	Republic of Finland
Equasym XL	DK/H/2888/002	PA 23211/001/002	[INACTIVE] Takeda Pharmaceuticals International AG	Ireland
Equasym XL	DK/H/2888/003	PA 23211/001/003	[INACTIVE] Takeda Pharmaceuticals International AG	Ireland
Equasym XL	DK/H/2888/005	PA 23211/001/005	[INACTIVE] Takeda Pharmaceuticals International AG	Ireland
Equasym XL	DK/H/2888/006	PA 23211/001/006	[INACTIVE] Takeda Pharmaceuticals International AG	Ireland
Equasym XL	DK/H/2888/004	PA 23211/001/004	[INACTIVE] Takeda Pharmaceuticals International AG	Ireland
Equasym XL	DK/H/2888/001	PA 23211/001/001	[INACTIVE] Takeda	Ireland

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
			Pharmaceuticals International AG	
Equasym XL	DK/H/2888/002	RVG 33228	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/005	RVG 111232	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/001	RVG 33227	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/006	RVG 111233	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/004	RVG 111228	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/002	RVG 33228	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/003	RVG 33229	Takeda Pharmaceuticals International AG	Kingdom of the Netherlands
Equasym XL	DK/H/2888/002	PL 54937/0002	[INACTIVE] Takeda Pharmaceuticals International AG	United Kingdom (Northern Ireland)
Equasym XL	DK/H/2888/003	PL 54937/0003	[INACTIVE] Takeda Pharmaceuticals International AG	United Kingdom (Northern Ireland)
Equasym XL	DK/H/2888/001	PL 54937/0001	[INACTIVE] Takeda Pharmaceuticals International AG	United Kingdom (Northern Ireland)
Equasym XR	DK/H/2888/006	2013120620	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/001	2013120615	[INACTIVE] Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/005	2013120619	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/004	2013120618	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/003	2013120617	[INACTIVE] Takeda Pharmaceuticals	Grand Duchy of Luxembourg

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
			International AG	
Equasym XR	DK/H/2888/004	2013120618	[INACTIVE] Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/002	2013120616	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/003	2013120617	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/001	2013120615	Takeda Pharmaceuticals International AG	Grand Duchy of Luxembourg
Equasym XR	DK/H/2888/001	BE423586	Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/004	BE437595	Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/006	BE437613	Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/006	BE437613	[INACTIVE] Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/005	BE437604	Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/003	BE423604	Takeda Pharmaceuticals International AG	Kingdom of Belgium
Equasym XR	DK/H/2888/002	BE423595	Takeda Pharmaceuticals International AG	Kingdom of Belgium
MEDICEBRAN	DE/H/0690/003	68.548	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medicebran	DE/H/0690/001	68546	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medicebran	DE/H/2222/003	5397757	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medicebran	DE/H/2222/001	5397732	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2222/003		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
MEDIKINET	DE/H/2223/003	CIS: 6 365 782 8	Medice Arzneimittel Puetter	French Republic

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			GmbH & Co. KG	
MEDIKINET	DE/H/2223/001	CIS: 6 114 564 0	Medice Arzneimittel Puetter GmbH & Co. KG	French Republic
MEDIKINET	DE/H/2223/005	CIS: 6 421 119 1	Medice Arzneimittel Puetter GmbH & Co. KG	French Republic
MEDIKINET	DE/H/2223/004	CIS: 6 659 298 3	Medice Arzneimittel Puetter GmbH & Co. KG	French Republic
MEDIKINET	DE/H/2223/002	CIS: 6 300 053 3	Medice Arzneimittel Puetter GmbH & Co. KG	French Republic
Medikinet	DE/H/0690/001	2007020039	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet	DE/H/0690/003	2007020041	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet	DE/H/2222/001	PA1555/001/006	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet	DE/H/2222/003	PA1555/001/008	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet	DE/H/2223/001	AIC N. 041438021	Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2223/001	041438019	Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2223/003		Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2223/002		Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2223/004		Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2223/005		Medice Arzneimittel Puetter GmbH & Co. KG	Italian Republic
Medikinet	DE/H/2222/001	BE 381534	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Belgium
Medikinet	DE/H/2222/003	BE 381552	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Belgium
Medikinet	DE/H/0690/001	39397	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet	DE/H/0690/003	39399	Medice Arzneimittel Puetter	Kingdom of Denmark

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			GmbH & Co. KG	
Medikinet	DE/H/0690/001	06-4110	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/009	12-9267	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/007	06-4122	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/003	06-4118	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/010	12-9268	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/004	06-4119	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/005	06-4120	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/008	10-7785	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/006	06-4121	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Norway
Medikinet	DE/H/0690/006	68544	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/009	78453	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/004	68.542	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/008	73308	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/005	68.543	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/010	78454	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/007	68.545	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Spain
Medikinet	DE/H/0690/001	23835	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/004	23840	Medice Arzneimittel Puetter	Kingdom of Sweden

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			GmbH & Co. KG	
Medikinet	DE/H/0690/003	23837	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/006	23842	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/010	48668	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/008	44810	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/005	23841	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/009	48667	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/007	23843	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Sweden
Medikinet	DE/H/0690/003	RVG 34026	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet	DE/H/0690/001	RVG 34024	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet	DE/H/2223/005	5361605	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2223/003	5361563	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2223/002	5361555	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2223/004	5361571	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2223/001	5361548	Medice Arzneimittel Puetter GmbH & Co. KG	Portuguese Republic
Medikinet	DE/H/2222/001	705210	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet	DE/H/2222/003	705410	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet	DE/H/0690/001	22209	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet	DE/H/0690/003	22211	Medice Arzneimittel Puetter	Republic of Finland

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
			GmbH & Co. KG	
Medikinet	DE/H/0690/003	IS/1/14/087/03	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet	DE/H/0690/001	IS/1/14/087/01	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet	DE/H/2222/001	10-0447	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet	DE/H/2222/003	10-0460	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet	DE/H/2222/001		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet	DE/H/2222/003/DC		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet	DE/H/0690/001	12841	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet	DE/H/0690/003	12843	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet	DE/H/0690/001	PL 11243/0002	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet	DE/H/0690/003	PL 11243/0004	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet adult		63891.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet adult		66747.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet adult		63890.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet adult		65964.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet adult		65965.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet adult		88863.00.00	Medice Pharma GmbH & Co. KG	Federal Republic of Germany
Medikinet CR	DE/H/0690/004	39458	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/007	39461	Medice Arzneimittel Puetter	Kingdom of Denmark

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
			GmbH & Co. KG	
Medikinet CR	DE/H/0690/010	51662	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/005	39459	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/008	47393	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/006	39460	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/009	51661	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of Denmark
Medikinet CR	DE/H/0690/009	RVG 112771	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/008	RVG 108026	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/006	RVG 34029	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/007	RVG 34030	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/005	RVG 34028	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/010	RVG 112772	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/004	RVG 34027	Medice Arzneimittel Puetter GmbH & Co. KG	Kingdom of the Netherlands
Medikinet CR	DE/H/0690/007	22215	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/008	28970	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/010	31131	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/006	22214	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/005	22213	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/004	22212	Medice Arzneimittel Puetter	Republic of Finland

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
			GmbH & Co. KG	
Medikinet CR	DE/H/0690/009	31130	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Finland
Medikinet CR	DE/H/0690/008	IS/1/14/087/04	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/010	IS/1/14/087/10	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/006	IS/1/14/087/07	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/007	IS/1/14/087/08	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/005	IS/1/14/087/06	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/004	IS/1/14/087/05	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/009	IS/1/14/087/09	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Iceland
Medikinet CR	DE/H/0690/010	21935	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/008	18355	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/007	12847	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/005	12845	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/009	21934	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/006	12846	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet CR	DE/H/0690/004	12844	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Poland
Medikinet MR	DE/H/2223/004	PA1555/001/004	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet MR	DE/H/2223/001	PA1555/001/001	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet MR	DE/H/2223/003	PA1555/001/003	Medice Arzneimittel Puetter	Ireland

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
			GmbH & Co. KG	
Medikinet MR	DE/H/2223/005	PA1555/001/005	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet MR	DE/H/2223/002	PA1555/001/002	Medice Arzneimittel Puetter GmbH & Co. KG	Ireland
Medikinet retard	DE/H/0690/010	88768.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/2223/003		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/2223/005		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/0690/009	88767.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/2223/002		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/2223/001		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/2223/004		Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Medikinet retard	DE/H/0690/004	1754/07020042	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/006	1754/07020044	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/005	1754/07020043	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/009	2014060146	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/010	2014060147	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/007	1754/07020045	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/008	1754/11030018	Medice Arzneimittel Puetter GmbH & Co. KG	Grand Duchy of Luxembourg
Medikinet retard	DE/H/0690/009	135358	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet retard	DE/H/0690/008	1-30056	Medice Arzneimittel Puetter	Republic of Austria

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
			GmbH & Co. KG	
Medikinet retard	DE/H/0690/004	1-26725	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet retard	DE/H/0690/010	135389	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet retard	DE/H/0690/005	1-26726	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet retard	DE/H/0690/006	1-26727	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet retard	DE/H/0690/007	1-26728	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Austria
Medikinet XL	DE/H/2223/004	705310	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet XL	DE/H/2223/002	705110	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet XL	DE/H/2223/005	705710	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet XL	DE/H/2223/001	705610	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet XL	DE/H/2223/003	705010	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Estonia
Medikinet XL	DE/H/2223/003	10-0462	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet XL	DE/H/2223/005	10-0464	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet XL	DE/H/2223/004	10-0463	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet XL	DE/H/2223/001	10-0448	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet XL	DE/H/2223/002	10-0461	Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Latvia
Medikinet XL	DE/H/2223/003		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet XL	DE/H/2223/001/DC		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet XL	DE/H/2223/005		Medice Arzneimittel Puetter	Republic of Lithuania

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
			GmbH & Co. KG	
Medikinet XL	DE/H/2223/004		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet XL	DE/H/2223/002		Medice Arzneimittel Puetter GmbH & Co. KG	Republic of Lithuania
Medikinet XL	DE/H/0690/007	PL 11243/0008	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/005	PL 11243/0006	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/006	PL 11243/0007	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/004	PL 11243/0005	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/009	PL 11243/0011	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/010	PL 11243/0012	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Medikinet XL	DE/H/0690/008	PL 11243/0010	Medice Arzneimittel Puetter GmbH & Co. KG	United Kingdom (Northern Ireland)
Methylfenidaat HCl Aristo	DE/H/6353/004	RVG 125058	Aristo Pharma GmbH	Kingdom of the Netherlands
Methylfenidaat HCl Aristo	DE/H/6353/002	RVG 125056	Aristo Pharma GmbH	Kingdom of the Netherlands
Methylfenidaat HCl Aristo	DE/H/6353/003	RVG 125057	Aristo Pharma GmbH	Kingdom of the Netherlands
Methylfenidaat HCl Aristo	DE/H/6353/005	RVG 125059	Aristo Pharma GmbH	Kingdom of the Netherlands
Methylfenidaat HCl Aristo	DE/H/6353/001	RVG 125055	Aristo Pharma GmbH	Kingdom of the Netherlands
Methylfenidaat XR EG	DE/H/6338/004	2020120337	EG	Grand Duchy of Luxembourg
Methylfenidaat XR EG	DE/H/6338/001	2020120334	EG	Grand Duchy of Luxembourg
Methylfenidaat XR EG	DE/H/6338/005	2020120338	EG	Grand Duchy of Luxembourg
Methylfenidaat XR EG	DE/H/6338/002	2020120335	EG	Grand Duchy of

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
				Luxembourg
Methylfenidaat XR EG	DE/H/6338/003	2020120336	EG	Grand Duchy of Luxembourg
Methylfenidaat XR EG	DE/H/6338/006	2020120339	EG	Grand Duchy of Luxembourg
Methylfenidaat XR EG	DE/H/6338/002	BE571777	EG	Kingdom of Belgium
Methylfenidaat XR EG	DE/H/6338/005	BE571804	EG	Kingdom of Belgium
Methylfenidaat XR EG	DE/H/6338/006	BE571813	EG	Kingdom of Belgium
Methylfenidaat XR EG	DE/H/6338/004	BE571795	EG	Kingdom of Belgium
Methylfenidaat XR EG	DE/H/6338/001	BE571760	EG	Kingdom of Belgium
Methylfenidaat XR EG	DE/H/6338/003	BE571786	EG	Kingdom of Belgium
Methylphenidat Alternova		54924	Orifarm Generics A/S	Kingdom of Denmark
Methylphenidat Alternova		54926	Orifarm Generics A/S	Kingdom of Denmark
Methylphenidate	DK/H/3415/001/DC	68574	Consilient Health Limited	Kingdom of Denmark
Methylphenidate	DK/H/3415/001/DC.	63957	Consilient Health Limited	Kingdom of Sweden
Methylphenidate "Sandoz"	DK/H/2138/001	49666	Sandoz A/S	Kingdom of Denmark
Methylphenidate Consilient Health	DK/H/3415/001/DC	22-15044	Consilient Health Limited	Kingdom of Norway
Methylphenidate Hydrochloride Consilient Health		PL 24837/0166	[INACTIVE] Consilient Health Limited	United Kingdom (Northern Ireland)
Methylphenidate Hydrochloride Generics UK	SE/H/1872/003	PL 04569/1498	Generics (U.K.) Limited	United Kingdom (Northern Ireland)
Methylphenidate Hydrochloride Generics UK	SE/H/1872/001	PL 04569/1496	Generics (U.K.) Limited	United Kingdom (Northern Ireland)
Methylphenidate Hydrochloride Generics UK	SE/H/1872/002	PL 04569/1497	Generics (U.K.) Limited	United Kingdom (Northern Ireland)
Methylphenidat-HCl MEDICE		2203637.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany

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		2203637.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methylphenidat-HCl MEDICE		2203638.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methylphenidat-HCl MEDICE		2203636.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methylphenidat-HCl MEDICE		97388.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methylphenidat-HCl MEDICE		97387.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methylphenidat-HCl MEDICE		2203639.00.00	Medice Arzneimittel Puetter GmbH & Co. KG	Federal Republic of Germany
Methysym	DE/H/6353/003	2204198.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym	DE/H/6353/002	2204197.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym	DE/H/6353/005	2204200.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym	DE/H/6353/001	2204196.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym	DE/H/6353/006	2204201.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym	DE/H/6353/004	2204199.00.00	Aristo Pharma GmbH	Federal Republic of Germany
Methysym Retard	DE/H/6353/001	85523	Aristo Pharma GmbH	Kingdom of Spain
Methysym Retard	DE/H/6353/006	85528	Aristo Pharma GmbH	Kingdom of Spain
Methysym Retard	DE/H/6353/002	85524	Aristo Pharma GmbH	Kingdom of Spain
Methysym Retard	DE/H/6353/005	85527	Aristo Pharma GmbH	Kingdom of Spain
Methysym Retard	DE/H/6353/004	85526	Aristo Pharma GmbH	Kingdom of Spain
Methysym Retard	DE/H/6353/003	85525	Aristo Pharma GmbH	Kingdom of Spain
Metiphenal	DE/H/6338/006	2204117.00.00	Aliud Pharma GmbH	Federal Republic of Germany
Metiphenal	DE/H/6338/002	2204113.00.00	Aliud Pharma GmbH	Federal Republic of Germany
Metiphenal	DE/H/6338/003	2204114.00.00	Aliud Pharma GmbH	Federal Republic of

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
				Germany
Metiphenal	DE/H/6338/005	2204116.00.00	Aliud Pharma GmbH	Federal Republic of Germany
Metiphenal	DE/H/6338/004	2204115.00.00	Aliud Pharma GmbH	Federal Republic of Germany
Metiphenal	DE/H/6338/001	2204112.00.00	Aliud Pharma GmbH	Federal Republic of Germany
Metynor hårda	DE/H/6338/001	59232	STADA Arzneimittel AG	Kingdom of Sweden
Metynor med modifierad frisättning.	DE/H/6338/004	59235	STADA Arzneimittel AG	Kingdom of Sweden
Metynor med modifierad frisättning.	DE/H/6338/005	59236	STADA Arzneimittel AG	Kingdom of Sweden
Metynor med modifierad frisättning.	DE/H/6338/006	59237	STADA Arzneimittel AG	Kingdom of Sweden
Metynor med modifierad frisättning.	DE/H/6338/002	59233	STADA Arzneimittel AG	Kingdom of Sweden
Metynor med modifierad frisättning.	DE/H/6338/003	59234	STADA Arzneimittel AG	Kingdom of Sweden
Phenichem	SE/H/1912/005	64389		Kingdom of Sweden
Quasym	UK/H/0819/005	5550454	[INACTIVE] Takeda Pharmaceuticals International AG	Portuguese Republic
Quasym	UK/H/0819/002	5436571	[INACTIVE] Takeda Pharmaceuticals International AG	Portuguese Republic
Quasym	UK/H/0819/001	5436563	[INACTIVE] Takeda Pharmaceuticals International AG	Portuguese Republic
Quasym	UK/H/0819/003	5436605	[INACTIVE] Takeda Pharmaceuticals International AG	Portuguese Republic
Quasym	UK/H/0819/006	5550470	[INACTIVE] Takeda Pharmaceuticals International AG	Portuguese Republic
Quasym	UK/H/0819/004	5550462	[INACTIVE] Takeda Pharmaceuticals	Portuguese Republic

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			International AG	
QUASYM L.P.	DK/H/2888/003	34009 377 622 1 2	Takeda Pharmaceuticals International AG	French Republic
QUASYM L.P.	DK/H/2888/002	34009 570 271 2 4	Takeda Pharmaceuticals International AG	French Republic
QUASYM L.P.	DK/H/2888/001	34009 497 261 6 5	Takeda Pharmaceuticals International AG	French Republic
Ritalin	DE/H/7682/001	7010489.00.00	Padia GmbH	Federal Republic of Germany
Концерта	DE/H/6007001	20060458	Johnson & Johnson Bulgaria EOOD	Republic of Bulgaria