1 December 2022
EMA/933936/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine besilate / hydrochlorothiazide / olmesartan medoxomil

Procedure no.: PSUSA/00002210/202204

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Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661012	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661024	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661036	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661048	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661051	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661063	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661075	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661087	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661099	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661101	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661113	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661125	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661137	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661149	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 20 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/001	041661152	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661164	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661176	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661188	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661190	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661202	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661214	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661226	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661238	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661240	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661253	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661265	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661277	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661289	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661291	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/12,5 mg compresse rivestite con film	NL/H/1858/002	041661303	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661315	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661327	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661339	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661341	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661354	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661366	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661378	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661380	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661392	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661404	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661416	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661428	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661430	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661442	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/12,5 mg compresse rivestite con film	NL/H/1858/003	041661455	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661467	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661479	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661481	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661493	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661505	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661517	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661529	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661531	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661543	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661556	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661568	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661570	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661582	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661594	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1858/004	041661606	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661618	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661620	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661632	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661644	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661657	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661669	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661671	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661683	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661695	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661707	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661719	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661721	DAIICHI SANKYO ITALIA S.P.A	IT

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SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661733	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661745	DAIICHI SANKYO ITALIA S.P.A	IT
SEVITREX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1858/005	041661758	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR/HCT 20 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/001	BE387466	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/002	BE387475	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/12,5 mg comprimés pelliculées	NL/H/1858/003	BE387484	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/25 mg comprimés pelliculées	NL/H/1858/004	BE387493	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/25 mg comprimés pelliculées	NL/H/1858/005	BE387502	DAIICHI SANKYO BELGIUM S.A	BE
Sevikar Plus 20 mg/5 mg/12.5 mg film-coated tablets	NL/H/1858/001	PA 1595/3/1	DAIICHI SANKYO IRELAND LIMITED	IE

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Sevikar Plus 40 mg/5 mg/12.5 mg film-coated tablets	NL/H/1858/002	PA 1595/3/2	DAIICHI SANKYO IRELAND LIMITED	IE
Sevikar Plus 40 mg/10 mg/12.5 mg film-coated tablets	NL/H/1858/003	PA 1595/3/3	DAIICHI SANKYO IRELAND LIMITED	IE
Sevikar Plus 40 mg/5 mg/25 mg film-coated tablets	NL/H/1858/004	PA 1595/3/4	DAIICHI SANKYO IRELAND LIMITED	IE
Sevikar Plus 40 mg/10 mg/25 mg film-coated tablets	NL/H/1858/005	PA 1595/3/5	DAIICHI SANKYO IRELAND LIMITED	IE
SEVIKAR HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1858/001	1-30068	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1858/002	1-30069	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1858/003	1-30070	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1858/004	1-30071	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1858/005	1-30072	DAIICHI SANKYO AUSTRIA GMBH	AT

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Sevikar Comp, filmovertrukne tabletter	NL/H/1858/001	46260	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar Comp, filmovertrukne tabletter	NL/H/1858/002	46261	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar Comp, filmovertrukne tabletter	NL/H/1858/003	46262	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar Comp, filmovertrukne tabletter	NL/H/1858/004	46263	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar Comp, filmovertrukne tabletter	NL/H/1858/005	46264	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar Comp 20 mg/5 mg/12,5 mg filmuhúðaðar töflur	NL/H/1858/001	IS/1/10/056/01	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar Comp 40 mg/5 mg/12,5 mg filmuhúðaðar töflur	NL/H/1858/002	IS/1/10/056/02	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar Comp 40 mg/10 mg/12,5 mg filmuhúðaðar töflur	NL/H/1858/003	IS/1/10/056/03	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar Comp 40 mg/5 mg/25 mg filmuhúðaðar töflur	NL/H/1858/004	IS/1/10/056/04	DAIICHI SANKYO EUROPE GMBH	IS

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Sevikar Comp 40 mg/10 mg/25 mg filmuhúðaðar töflur	NL/H/1858/005	IS/1/10/056/05	DAIICHI SANKYO EUROPE GMBH	IS
SEVIKAR HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1858/001	79810.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1858/002	79811.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1858/003	79812.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1858/004	79813.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1858/005	79814.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR/HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/001	BE387466	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/002	BE387475	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1858/003	BE387484	DAIICHI SANKYO BELGIUM S.A	BE

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SEVIKAR/HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1858/004	BE387493	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1858/005	BE387502	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 20 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/001	1851/11040064	DAIICHI SANKYO BELGIUM S.A	LU
SEVIKAR/HCT 40 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/002	1851/11040065	DAIICHI SANKYO BELGIUM S.A	LU
SEVIKAR/HCT 40 mg/10 mg/12,5 mg comprimés pelliculées	NL/H/1858/003	1851/11040066	DAIICHI SANKYO BELGIUM S.A	LU
SEVIKAR/HCT 40 mg/5 mg/25 mg comprimés pelliculées	NL/H/1858/004	1851/11040067	DAIICHI SANKYO BELGIUM S.A	LU
SEVIKAR/HCT 40 mg/10 mg/25 mg comprimés pelliculées	NL/H/1858/005	1851/11040068	DAIICHI SANKYO BELGIUM S.A	LU
Sevikar HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/001	RVG 106667	DAIICHI SANKYO NEDERLAND B.V.	NL
Sevikar HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/002	RVG 106671	DAIICHI SANKYO NEDERLAND B.V.	NL

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Sevikar HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1858/003	RVG 106672	DAIICHI SANKYO NEDERLAND B.V.	NL
Sevikar HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1858/004	RVG 106673	DAIICHI SANKYO NEDERLAND B.V.	NL
Sevikar HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1858/005	RVG 106674	DAIICHI SANKYO NEDERLAND B.V.	NL
SEVIKAR HCT 40 mg /5 mg/25 mg comprimidos recubiertos con película	NL/H/1858/004	73.453	DAIICHI SANKYO ESPAÑA, S.A.	ES
SEVIKAR HCT 20 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1858/001	73.450	DAIICHI SANKYO ESPAÑA, S.A.	ES
SEVIKAR HCT 40 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1858/002	73.451	DAIICHI SANKYO ESPAÑA, S.A.	ES
SEVIKAR HCT 40 mg /10 mg/12,5 mg comprimidos recubiertos con película	NL/H/1858/003	73.452	DAIICHI SANKYO ESPAÑA, S.A.	ES
SEVIKAR HCT 40 mg /10 mg/25 mg comprimidos recubiertos con película	NL/H/1858/005	73.454	DAIICHI SANKYO ESPAÑA, S.A.	ES
Sevikar HCT 20 mg/5 mg/12.5 mg film-coated tablets	NL/H/1858/001	PL 08265/0031	DAIICHI SANKYO UK LTD	XI

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Sevikar HCT 40 mg/5 mg/12.5 mg film-coated tablets	NL/H/1858/002	PL 08265/0032	DAIICHI SANKYO UK LTD	XI
Sevikar HCT 40 mg/10 mg/12.5 mg film-coated tablets	NL/H/1858/003	PL 08265/0033	DAIICHI SANKYO UK LTD	XI
Sevikar HCT 40 mg/5 mg/25 mg film-coated tablets	NL/H/1858/004	PL 08265/0034	DAIICHI SANKYO UK LTD	XI
Sevikar HCT 40 mg/10 mg/25 mg film-coated tablets	NL/H/1858/005	PL 08265/0035	DAIICHI SANKYO UK LTD	XI
SEVIKAR/HCT 20 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/001	BE387414	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/12,5 mg comprimés pelliculées	NL/H/1858/002	BE387423	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/12,5 mg comprimés pelliculées	NL/H/1858/003	BE387432	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/25 mg comprimés pelliculées	NL/H/1858/004	BE387441	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/25 mg comprimés pelliculées	NL/H/1858/005	BE387457	DAIICHI SANKYO BELGIUM 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
SEVIKAR/HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/001	BE387414	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1858/002	BE387423	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1858/003	BE387432	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1858/004	BE387441	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR/HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1858/005	BE387457	DAIICHI SANKYO BELGIUM S.A	BE
Capenon HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1860/001	RVG 106682	DAIICHI SANKYO EUROPE GMBH	NL
Capenon HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1860/002	RVG 106683	DAIICHI SANKYO EUROPE GMBH	NL
Capenon HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1860/003	RVG 106684	DAIICHI SANKYO EUROPE GMBH	NL
Capenon HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1860/004	RVG 106685	DAIICHI SANKYO EUROPE GMBH	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
Capenon HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1860/005	RVG 106686	DAIICHI SANKYO EUROPE GMBH	NL
CAPENON HCT 20 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1860/001	73433	DAIICHI SANKYO ESPAÑA, S.A.	ES
CAPENON HCT 40 mg /10 mg/12,5 mg comprimidos recubiertos con película	NL/H/1860/003	73435	DAIICHI SANKYO ESPAÑA, S.A.	ES
CAPENON HCT 40 mg /10 mg/25 mg comprimidos recubiertos con película	NL/H/1860/005	73432	DAIICHI SANKYO ESPAÑA, S.A.	ES
CAPENON HCT 40 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1860/002	73436	DAIICHI SANKYO ESPAÑA, S.A.	ES
CAPENON HCT 40 mg /5 mg/25 mg comprimidos recubiertos con película	NL/H/1860/004	73434	DAIICHI SANKYO ESPAÑA, S.A.	ES
Nistik 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/3862/005	RVG 119986	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Nistik 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/3862/001	RVG 119981	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Nistik 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/3862/002	RVG 119983	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Nistik 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/3862/003	RVG 119984	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Nistik 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/3862/004	RVG 119985	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Nistik 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/001	049732050	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/3862/005	049732252	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/3862/005	049732225	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/3862/005	049732213	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/3862/003	049732136	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/3862/004	049732163	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/002	049732074	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Nistik 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/3862/005	049732249	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/002	049732062	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/001	049732023	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/002	049732100	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/001	049732047	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/3862/005	049732237	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/002	049732098	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/002	049732086	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/3862/004	049732187	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Nistik 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/3862/004	049732175	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/001	049732035	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/3862/004	049732199	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/3862/003	049732124	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/3862/004	049732201	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/3862/003	049732151	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/3862/001	049732011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/3862/003	049732112	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Nistik 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/3862/003	049732148	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
FORZATEN/HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1859/004	BE390503	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1859/004	BE390503	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1859/004	BE390512	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1859/004	BE390512	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/002	BE390564	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1859/005	BE390546	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1859/003	BE390521	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/001	BE390485	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/002	BE390564	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG 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
Forzaten/HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1859/005	BE390555	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/002	BE390494	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/001	BE390485	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/001	BE390573	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/002	BE390494	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1859/005	BE390555	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1859/003	BE390537	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/001	BE390573	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1859/003	BE390537	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG 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
Forzaten/HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1859/003	BE390521	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten/HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1859/005	BE390546	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/038	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/058	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/051	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/040	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/043	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/042	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/055	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/022	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/015	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/054	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/020	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/065	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/035	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/045	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/029	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/039	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/056	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/027	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/044	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/021	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/028	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/072	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/071	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/052	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/025	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/059	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/030	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/057	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/026	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/053	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/050	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/041	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/069	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/066	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/070	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/068	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/060	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/074	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/037	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/067	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/073	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/023	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/013	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/001	8618/2016/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1859/005	8622/2016/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO

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
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1859/002	8619/2016/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1859/004	8621/2016/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1859/003	8620/2016/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Amelior® plus HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/001	1-30073	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Amelior® plus HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1859/003	1-30075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Amelior® plus HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/002	1-30074	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Amelior® plus HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1859/005	1-30077	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
FORZATEN/HCT 40 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/002	BE390494	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/002	BE390564	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 20 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/001	BE390573	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Amelior® plus HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1859/004	1-30076	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
FORZATEN/HCT 40 mg/5 mg/25 mg comprimés pelliculés	NL/H/1859/004	BE390503	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/12,5 mg comprimés pelliculés	NL/H/1859/003	BE390521	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/25 mg comprimés pelliculés	NL/H/1859/005	BE390546	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/5 mg/25 mg comprimés pelliculés	NL/H/1859/004	BE390512	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG 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
FORZATEN/HCT 20 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/001	BE390485	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/25 mg comprimés pelliculés	NL/H/1859/005	BE390555	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
FORZATEN/HCT 40 mg/10 mg/12,5 mg comprimés pelliculés	NL/H/1859/003	BE390537	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
ТЕСПАДАН HCT 20 mg/5 mg/12,5 mg филмирани таблетки	NL/H/1859/001	20110282	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
ТЕСПАДАН HCT 40 mg/10 mg/12,5 mg филмирани таблетки	NL/H/1859/003	20110284	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
ТЕСПАДАН HCT 40 mg/5 mg/25 mg филмирани таблетки	NL/H/1859/004	20110285	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
ТЕСПАДАН HCT 40 mg/5 mg/12,5 mg филмирани таблетки	NL/H/1859/002	20110283	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Sintonyn Combi 20 mg/5 mg/12,5 mg potahované tablety	NL/H/1859/001	58/544/11-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
ТЕСПАДАН HCT 40 mg/10 mg/25 mg филмирани таблетки	NL/H/1859/005	20110286	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG

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
Sintonyn Combi 40 mg/10 mg/12,5 mg potahované tablety	NL/H/1859/003	58/546/11-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sintonyn Combi 40 mg/5 mg/12,5 mg potahované tablety	NL/H/1859/002	58/545/11-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sintonyn Combi 40 mg/5 mg/25 mg potahované tablety	NL/H/1859/004	58/547/11-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sintonyn Combi 40 mg/10 mg/25 mg potahované tablety	NL/H/1859/005	58/548/11-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sanoral HCT, 40 mg / 10 mg / 25 mg õhukese polümeerikattega tabletid	NL/H/1859/005	730111	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Sanoral HCT, 20 mg / 5 mg / 12,5 mg õhukese polümeerikattega tabletid	NL/H/1859/001	730411	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Sanoral HCT, 40 mg / 5 mg / 12,5 mg õhukese polümeerikattega tabletid	NL/H/1859/002	730311	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Sanoral HCT, 40 mg / 10 mg / 12,5 mg õhukese polümeerikattega tabletid	NL/H/1859/003	730211	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Sanoral HCT, 40 mg / 5 mg / 25 mg õhukese polümeerikattega tabletid	NL/H/1859/004	730511	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Vocado® HCT 40 mg/5 mg/25 mg Filmtabletten	NL/H/1859/004	79818.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Duactan HCT 40 mg/5 mg/12,5 mg filmtabletta	NL/H/1859/002	OGYI-T-21745/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Vocado® HCT 40 mg/10 mg/25 mg Filmtabletten	NL/H/1859/005	79819.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Duactan HCT 40 mg/5 mg/12,5 mg filmtabletta	NL/H/1859/002	OGYI-T-21745/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Vocado® HCT 40 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/002	79816.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Duactan HCT 20 mg/5 mg/12,5 mg filmtabletta	NL/H/1859/001	OGYI-T-21745/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan HCT 20 mg/5 mg/12,5 mg filmtabletta	NL/H/1859/001	OGYI-T-21745/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan HCT 40 mg/5 mg/25 mg filmtabletta	NL/H/1859/004	OGYI-T-21745/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan HCT 40 mg/5 mg/25 mg filmtabletta	NL/H/1859/004	OGYI-T-21745/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Vocado® HCT 20 mg/5 mg/12,5 mg Filmtabletten	NL/H/1859/001	79815.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Duactan HCT 40 mg/10 mg/12,5 mg filmtabletta	NL/H/1859/003	OGYI-T-21745/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan HCT 40 mg/10 mg/12,5 mg filmtabletta	NL/H/1859/003	OGYI-T-21745/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan HCT 40 mg/10 mg/25 mg filmtabletta	NL/H/1859/005	OGYI-T-21745/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Konverge Plus 20 mg/5 mg/12.5 mg film-coated tablets	NL/H/1859/001	PA 865/19/1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Konverge Plus 40 mg/5 mg/12.5 mg film-coated tablets	NL/H/1859/002	PA 865/19/2	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Duactan HCT 40 mg/10 mg/25 mg filmtabletta	NL/H/1859/005	OGYI-T-21745/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Konverge Plus 40 mg/10 mg/12.5 mg film-coated tablets	NL/H/1859/003	PA 865/19/3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Konverge Plus 40 mg/10 mg/25 mg film-coated tablets	NL/H/1859/005	PA 865/19/5	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Konverge Plus 40 mg/5 mg/25 mg film-coated tablets	NL/H/1859/004	PA 865/19/4	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Vocado® HCT 40 mg/10 mg/12,5 mg Filmtabletten	NL/H/1859/003	79817.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Sanoral HCT 20 mg/5 mg/12,5 mg apvalkotās tabletes	NL/H/1859/001	11-0034	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Sanoral HCT 40 mg/5 mg/12,5 mg apvalkotās tabletes	NL/H/1859/002	11-0037	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Sanoral HCT 40 mg/10 mg/12,5 mg apvalkotās tabletes	NL/H/1859/003	11-0035	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Sanoral HCT 40 mg/5 mg/25 mg apvalkotās tabletes	NL/H/1859/004	11-0038	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Sanoral HCT 40 mg/10 mg/25 mg apvalkotās tabletes	NL/H/1859/005	11-0036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/031	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/016	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/025	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/026	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/019	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/023	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/027	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/022	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/020	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/029	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/021	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/042	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/041	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/030	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/002	LT/1/11/2460/028	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/013	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/015	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/001	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/040	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/032	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 20 mg/5 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/001	LT/1/11/2460/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/039	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/038	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/033	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/045	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/037	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/034	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/043	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/035	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/12,5 mg plėvele dengtos tabletės	NL/H/1859/003	LT/1/11/2460/044	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/046	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/055	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/048	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/053	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/056	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/057	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/054	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/047	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/050	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/049	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/051	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/052	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/060	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/071	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/059	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/069	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/5 mg/25 mg plėvele dengtos tabletės	NL/H/1859/004	LT/1/11/2460/058	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/072	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/064	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/065	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/066	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/073	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/061	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/062	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/074	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/067	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/070	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral HCT 40 mg/10 mg/25 mg plėvele dengtos tabletės	NL/H/1859/005	LT/1/11/2460/068	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
FORZATEN/HCT 20 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/001	2011060014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
FORZATEN/HCT 40 mg/5 mg/25 mg comprimés pelliculés	NL/H/1859/004	2011060017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
FORZATEN/HCT 40 mg/5 mg/12,5 mg comprimés pelliculés	NL/H/1859/002	2011060015	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
FORZATEN/HCT 40 mg/10 mg/12,5 mg comprimés pelliculés	NL/H/1859/003	2011060016	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Konverge Plus 40 mg/5 mg/25 mg film-coated tablets	NL/H/1859/004	MA204/00604	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
FORZATEN/HCT 40 mg/10 mg/25 mg comprimés pelliculés	NL/H/1859/005	2011060018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Belfor HCT 40 mg/5 mg/25 mg filmomhulde tabletten	NL/H/1859/004	RVG 106679	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Konverge Plus 20 mg/5 mg/12.5 mg film-coated tablets	NL/H/1859/001	MA204/00601	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT

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
Belfor HCT 20 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/001	RVG 106675	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Konverge Plus 40 mg/5 mg/12.5 mg film-coated tablets	NL/H/1859/002	MA204/00602	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Belfor HCT 40 mg/10 mg/25 mg filmomhulde tabletten	NL/H/1859/005	RVG 106680	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Belfor HCT 40 mg/10 mg/12,5 mg filmomhulde tabletten	NL/H/1859/003	RVG 106678	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Konverge Plus 40 mg/10 mg/25 mg film-coated tablets	NL/H/1859/005	MA204/00605	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Belfor HCT 40 mg/5 mg/12,5 mg filmomhulde tabletten	NL/H/1859/002	RVG 106676	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Konverge Plus 40 mg/10 mg/12.5 mg film-coated tablets	NL/H/1859/003	MA204/00603	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Elestar HCT, 20 mg + 5 mg + 12,5 mg, tabletki powlekane	NL/H/1859/001	18947	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Elestar HCT, 40 mg + 5 mg + 25 mg, tabletki powlekane	NL/H/1859/004	18950	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Elestar HCT, 40 mg + 10 mg + 12,5 mg, tabletki powlekane	NL/H/1859/003	18949	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Elestar HCT, 40 mg + 10 mg + 25 mg, tabletki powlekane	NL/H/1859/005	18951	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Elestar HCT, 40 mg + 5 mg + 12,5 mg, tabletki powlekane	NL/H/1859/002	18948	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Zolnor HCT 20 mg+5 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/001	5369327	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+10 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/003	5369350	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 20 mg+5 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/001	5369319	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+5 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/002	5369335	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+5 mg+25 mg, comprimidos revestidos por película	NL/H/1859/004	5369400	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+5 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/002	5369343	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zolnor HCT 40 mg+10 mg+25 mg, comprimidos revestidos por película	NL/H/1859/005	5369418	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+5 mg+25 mg, comprimidos revestidos por película	NL/H/1859/004	5369376	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+10 mg+25 mg, comprimidos revestidos por película	NL/H/1859/005	5369426	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor HCT 40 mg+10 mg+12,5 mg, comprimidos revestidos por película	NL/H/1859/003	5369368	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Folgan HCT 40 mg/10 mg/12,5 mg filmom obalené tablety	NL/H/1859/003	58/0036/11-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Folgan HCT 40 mg/10 mg/25 mg filmom obalené tablety	NL/H/1859/005	58/0038/11-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Folgan HCT 20 mg/5 mg/12,5 mg filmom obalené tablety	NL/H/1859/001	58/0034/11-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Folgan HCT 40 mg/5 mg/12,5 mg filmom obalené tablety	NL/H/1859/002	58/0035/11-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/034	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/001	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 20 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/001	H/11/01161/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Folgan HCT 40 mg/5 mg/25 mg filmom obalené tablety	NL/H/1859/004	58/0037/11-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/032	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/019	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/016	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI

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
Olectan HCT 40 mg/5 mg/12,5 mg filmsko obložene tablete	NL/H/1859/002	H/11/01161/018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/031	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/12,5 mg filmsko obložene tablete	NL/H/1859/003	H/11/01161/033	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/046	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Balzak Plus 20 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1859/001	73.437	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/047	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/048	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/5 mg/25 mg filmsko obložene tablete	NL/H/1859/004	H/11/01161/049	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Balzak Plus 40 mg /5 mg/12,5 mg comprimidos recubiertos con película	NL/H/1859/002	73.438	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Balzak Plus 40 mg /10 mg/12,5 mg comprimidos recubiertos con película	NL/H/1859/003	73.440	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/061	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Balzak Plus 40 mg /5 mg/25 mg comprimidos recubiertos con película	NL/H/1859/004	73.441	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/062	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Balzak Plus 40 mg /10 mg/25 mg comprimidos recubiertos con película	NL/H/1859/005	73.439	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olectan HCT 40 mg/10 mg/25 mg filmsko obložene tablete	NL/H/1859/005	H/11/01161/064	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 20 mg/5 mg/12.5 mg	NL/H/1859/001	21492	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/10 mg/25 mg	NL/H/1859/005	21496	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/5 mg/12.5 mg	NL/H/1859/002	21493	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/5 mg/25 mg	NL/H/1859/004	21495	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/10 mg/12.5 mg	NL/H/1859/003	21494	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 20 mg/5 mg/12.5 mg	NL/H/1859/001	39688	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/10 mg/12.5 mg	NL/H/1859/003	39690	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/5 mg/12.5 mg	NL/H/1859/002	39689	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/10 mg/25 mg	NL/H/1859/005	39692	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal plus 40 mg/5 mg/25 mg	NL/H/1859/004	39691	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679145	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679095	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679133	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679071	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679121	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679032	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679083	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679069	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679044	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679020	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679107	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679119	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679057	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 20 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/001	049679715	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679362	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679158	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679297	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679350	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679335	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679374	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679398	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679677	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679754	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679208	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679689	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679626	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679703	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679210	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679653	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679246	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679576	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679691	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679424	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679259	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679234	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679602	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679196	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679638	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679172	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679222	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679309	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679323	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679614	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679588	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679347	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679739	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679665	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679311	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679436	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679727	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679386	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679640	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679400	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/12.5 mg compresse rivestite con film	NL/H/1859/003	049679412	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679273	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679160	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679475	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679451	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679184	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679463	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679525	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/10 mg/25 mg compresse rivestite con film	NL/H/1859/005	049679590	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679537	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679513	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679261	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/12.5 mg compresse rivestite con film	NL/H/1859/002	049679285	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679499	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679501	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679487	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679552	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679448	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679564	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679741	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANTRIX 40 mg/5 mg/25 mg compresse rivestite con film	NL/H/1859/004	049679549	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/01	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/02	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/03	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/04	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/05	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/12	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/13	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/14	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/15	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/06	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/07	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/08	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/09	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/10	LABORMED PHARMA S.A.	RO
Sevikar HCT 20 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/001	8613/2016/11	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/06	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/07	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/08	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/09	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/10	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/11	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/10	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/09	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/08	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/07	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/15	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/14	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/13	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/12	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/11	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/06	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/05	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/04	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/03	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/02	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/25 mg comprimate filmate	NL/H/1858/004	8616/2016/01	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/11	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/10	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/09	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/08	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/07	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/06	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/05	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/04	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/03	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/02	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/01	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/15	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/14	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/13	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/5 mg/12,5 mg comprimate filmate	NL/H/1858/002	8614/2016/12	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/15	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/14	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/13	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/12	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/05	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/04	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/03	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/02	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/01	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/11	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/10	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/09	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/08	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/07	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/25 mg comprimate filmate	NL/H/1858/005	8617/2016/06	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/01	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/15	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/14	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/13	LABORMED PHARMA S.A.	RO

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
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/12	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/05	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/04	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/03	LABORMED PHARMA S.A.	RO
Sevikar HCT 40 mg/10 mg/12,5 mg comprimate filmate	NL/H/1858/003	8615/2016/02	LABORMED PHARMA S.A.	RO