25 March 2021
EMA/259778/2021
Pharmacovigilance Risk Assessment Committee (PRAC)

List of nationally authorised medicinal products

Active substance(s): ezetimibe / rosuvastatin

Procedure No.: PSUSA/00010271/202007

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zenon 10 mg/20 mg filmom obalené tablety	CZ/H/0455/002	31/0009/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/20 mg filmom obalené tablety	CZ/H/0455/002	31/0009/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/20 mg filmom obalené tablety	CZ/H/0455/002	31/0009/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/20 mg filmom obalené tablety	CZ/H/0455/002	31/0009/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/20 mg potahované tablety	CZ/H/0455/002	31/322/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/20 mg potahované tablety	CZ/H/0455/002	31/322/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/20 mg potahované tablety	CZ/H/0455/002	31/322/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/20 mg potahované tablety	CZ/H/0455/002	31/322/14-C	SANOFI-AVENTIS SRO	CZ
Зенон 10 mg/20 mg филмирани таблетки	CZ/H/0455/002	20140378	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/20 mg филмирани таблетки	CZ/H/0455/002	20140378	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/20 mg филмирани таблетки	CZ/H/0455/002	20140378	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/20 mg филмирани таблетки	CZ/H/0455/002	20140378	SANOFI BULGARIA EOOD	BG
Suvezen, (10 mg + 20 mg), tabletki powlekane	CZ/H/0455/002	22278	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 20 mg), tabletki powlekane	CZ/H/0455/002	22278	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 20 mg), tabletki powlekane	CZ/H/0455/002	22278	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 20 mg), tabletki powlekane	CZ/H/0455/002	22278	SANOFI-AVENTIS SP Z.O.O.	PL
Ridutrin 10 mg/10 mg capsule, hard	NL/H/3017/001	RVG 114008	EGIS PHARMACEUTICALS PLC	NL
Ridutrin 20 mg/10 mg capsule, hard	NL/H/3017/002	RVG 114009	EGIS PHARMACEUTICALS PLC	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
Ridutrin 40 mg/10 mg capsule, hard	NL/H/3017/003	RVG 114010	EGIS PHARMACEUTICALS PLC	NL
Lipocomb 10 mg/10 mg cápsulas duras	NL/H/3017/001-003/DC	79053	EGIS PHARMACEUTICALS PLC	ES
Lipocomb 20 mg/10 mg cápsulas duras	NL/H/3017/001-003/DC	79051	EGIS PHARMACEUTICALS PLC	ES
Lipocomb 40 mg/10 mg cápsulas duras	NL/H/3017/001-003/DC	79052	EGIS PHARMACEUTICALS PLC	ES
TWICOR 10 mg/10 mg, comprimé pelliculé	NL/H/4126/001	34009 301 638 7 0	MYLAN MEDICAL SAS	FR
TWICOR 10 mg/10 mg, comprimé pelliculé	NL/H/4126/001	34009 301 638 8 7	MYLAN MEDICAL SAS	FR
TWICOR 10 mg/10 mg, comprimé pelliculé	NL/H/4126/001	34009 301 638 9 4	MYLAN MEDICAL SAS	FR
TWICOR 10 mg/10 mg, comprimé pelliculé	NL/H/4126/001	34009 301 639 1 7	MYLAN MEDICAL SAS	FR
TWICOR 20 mg/10 mg, comprimé pelliculé	NL/H/4126/002	34009 301 639 4 8	MYLAN MEDICAL SAS	FR
TWICOR 20 mg/10 mg, comprimé pelliculé	NL/H/4126/002	34009 301 639 5 5	MYLAN MEDICAL SAS	FR
TWICOR 20 mg/10 mg, comprimé pelliculé	NL/H/4126/002	34009 301 639 6 2	MYLAN MEDICAL SAS	FR
TWICOR 20 mg/10 mg, comprimé pelliculé	NL/H/4126/002	34009 301 639 7 9	MYLAN MEDICAL SAS	FR
ROZOR 10 mg/10 mg filmom obalené tablety	NL/H/4126/001	31/0360/18-S	MYLAN IRE HEALTHCARE LIMITED	SK
ROZOR 20 mg/10 mg filmom obalené tablety	NL/H/4126/002	31/0361/18-S	MYLAN IRE HEALTHCARE LIMITED	SK
Twicor 10 mg/10 mg film-coated tablets	NL/H/4126/001	MA 1187/01401	MYLAN IRE HEALTHCARE LIMITED	MT
Twicor 20 mg/10 mg film-coated tablets	NL/H/4126/002	MA 1187/01402	MYLAN IRE HEALTHCARE LIMITED	MT
ROZOR 10 mg/10 mg filmom obložene tablete	NL/H/4126/001	HR-H-900214203	MYLAN HRVATSKA D.O.O.	HR
ROZOR 20 mg/10 mg filmom obložene tablete	NL/H/4126/002	HR-H-978548703	MYLAN HRVATSKA D.O.O.	HR

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
Twicor, filmovertrukne tabletter	NL/H/4126/001	60056	MYLAN DENMARK APS	DK
Twicor, filmovertrukne tabletter	NL/H/4126/002	60057	MYLAN DENMARK APS	DK
TWICOR 10 mg + 10 mg comprimidos revestidos por película	NL/H/4126/001	5760822	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 10 mg + 10 mg comprimidos revestidos por película	NL/H/4126/001	5760830	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 10 mg + 10 mg comprimidos revestidos por película	NL/H/4126/001	5760848	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 10 mg + 10 mg comprimidos revestidos por película	NL/H/4126/001	5760855	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 20 mg + 10 mg comprimidos revestidos por película	NL/H/4126/002	5760863	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 20 mg + 10 mg comprimidos revestidos por película	NL/H/4126/002	5760871	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 20 mg + 10 mg comprimidos revestidos por película	NL/H/4126/002	5760905	BGP PRODUCTS UNIPESSOAL, LDA.	PT
TWICOR 20 mg + 10 mg comprimidos revestidos por película	NL/H/4126/002	5760913	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Myrosor 10 mg / 10 mg comprimés pelliculés	NL/H/4126/001	BE536257	MYLAN EPD BVBA/SPRL	BE
Myrosor 10 mg/10 mg filmomhulde tabletten	NL/H/4126/001	BE536257	MYLAN EPD BVBA/SPRL	BE
Myrosor 10 mg/10 mg Filmtabletten	NL/H/4126/001	BE536257	MYLAN EPD BVBA/SPRL	BE
Myrosor 20 mg / 10 mg comprimés pelliculés	NL/H/4126/002	BE536266	MYLAN EPD BVBA/SPRL	BE
Myrosor 20 mg/10 mg	NL/H/4126/002	BE536266	MYLAN EPD BVBA/SPRL	BE

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filmomhulde tabletten				
Myrosor 20 mg/10 mg Filmtabletten	NL/H/4126/002	BE536266	MYLAN EPD BVBA/SPRL	BE
Rosuvastatin/Ezetimibe Mylan 10 mg/10 mg potahované tablety	NL/H/4126/001	31/364/17-C	MYLAN IRE HEALTHCARE LIMITED	CZ
Twicor 20 mg/10 mg potahované tablety	NL/H/4126/002	31/365/17-C	MYLAN IRE HEALTHCARE LIMITED	CZ
Twicor 10 mg/10 mg film-coated tablets	NL/H/4126/001	PL 46302/0060	MYLAN PRODUCTS LIMITED	UK
Twicor 20 mg/10 mg film-coated tablets	NL/H/4126/002	PL 46302/0061	MYLAN PRODUCTS LIMITED	UK
РОЗОР 10 mg/10 mg филмирани таблетки	NL/H/4126/001	20180338	MYLAN EOOD	BG
РОЗОР 20 mg/10 mg филмирани таблетки	NL/H/4126/002	20180339	MYLAN EOOD	BG
Rozor 10 mg/10 mg filmtabletta	NL/H/4126/001	OGYI-T-23469/01	MYLAN EPD KFT.	HU
Rozor 20 mg/10 mg filmtabletta	NL/H/4126/002	OGYI-T-23469/02	MYLAN EPD KFT.	HU
Rosuvastatine/Ezetimibe Mylan Healthcare 20 mg/10 mg filmomhulde tabletten	NL/H/4126/002	RVG 121836	MYLAN HEALTHCARE B.V.	NL
Rosuvastatine/Ezetimibe Mylan Healthcare 10 mg/10 mg filmomhulde tabletten	NL/H/4126/001	RVG 121835	MYLAN HEALTHCARE B.V.	NL
ROZOR 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/4126/001	022904	MYLAN IRE HEALTHCARE LIMITED	CY
ROZOR 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/4126/002	022905	MYLAN IRE HEALTHCARE LIMITED	CY
Sorento, 10 mg + 10 mg, tabletki powlekane	NL/H/4126/001	25118	MYLAN IRE HEALTHCARE LIMITED	PL
Sorento, 20 mg + 10	NL/H/4126/002	25119	MYLAN IRE HEALTHCARE	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
mg, tabletki powlekane			LIMITED	
Rozor 10 mg/10 mg filmsko obložene tablete	NL/H/4126/001	H/19/02551/002	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 10 mg/10 mg filmsko obložene tablete	NL/H/4126/001	H/19/02551/001	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 10 mg/10 mg filmsko obložene tablete	NL/H/4126/001	H/19/02551/003	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 10 mg/10 mg filmsko obložene tablete	NL/H/4126/001	H/19/02551/004	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 20 mg/10 mg filmsko obložene tablete	NL/H/4126/002	H/19/02551/005	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 20 mg/10 mg filmsko obložene tablete	NL/H/4126/002	H/19/02551/006	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 20 mg/10 mg filmsko obložene tablete	NL/H/4126/002	H/19/02551/007	MYLAN IRE HEALTHCARE LIMITED	SI
Rozor 20 mg/10 mg filmsko obložene tablete	NL/H/4126/002	H/19/02551/008	MYLAN IRE HEALTHCARE LIMITED	SI
Twicor 20 mg/10 mg comprimidos recubiertos con película	NL/H/4126/002	83660	MYLAN IRE HEALTHCARE LIMITED	ES
TWICOR 10 mg/10 mg comprimate filmate	NL/H/4126/001	11316/2018/01	MYLAN HEALTHCARE GMBH	RO
TWICOR 10 mg/10 mg comprimate filmate	NL/H/4126/001	11316/2018/02	MYLAN HEALTHCARE GMBH	RO
TWICOR 10 mg/10 mg comprimate filmate	NL/H/4126/001	11316/2018/03	MYLAN HEALTHCARE GMBH	RO
TWICOR 10 mg/10 mg comprimate filmate	NL/H/4126/001	11316/2018/04	MYLAN HEALTHCARE GMBH	RO
TWICOR 20 mg/10 mg comprimate filmate	NL/H/4126/002	11317/2018/02	MYLAN HEALTHCARE GMBH	RO
TWICOR 20 mg/10 mg comprimate filmate	NL/H/4126/002	11317/2018/01	MYLAN HEALTHCARE GMBH	RO
TWICOR 20 mg/10 mg comprimate filmate	NL/H/4126/002	11317/2018/03	MYLAN HEALTHCARE GMBH	RO
TWICOR 20 mg/10 mg comprimate filmate	NL/H/4126/002	11317/2018/04	MYLAN HEALTHCARE GMBH	RO

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ROZEIOND 10 mg/10 mg compresse rivestite con film	NL/H/4126/001	046030019	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 10 mg/10 mg compresse rivestite con film	NL/H/4126/001	046030021	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 10 mg/10 mg compresse rivestite con film	NL/H/4126/001	046030033	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 10 mg/10 mg compresse rivestite con film	NL/H/4126/001	046030045	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 20 mg/10 mg compresse rivestite con film	NL/H/4126/002	046030058	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 20 mg/10 mg compresse rivestite con film	NL/H/4126/002	046030060	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 20 mg/10 mg compresse rivestite con film	NL/H/4126/002	046030072	MYLAN IRE HEALTHCARE LIMITED	IT
ROZEIOND 20 mg/10 mg compresse rivestite con film	NL/H/4126/002	046030084	MYLAN IRE HEALTHCARE LIMITED	IT
TWICOR 10 mg/10 mg film-coated tablets	NL/H/4126/001	PA 2010/19/1	MYLAN IRE HEALTHCARE LIMITED	IE
TWICOR 20 mg/10 mg film-coated tablets	NL/H/4126/002	PA 2010/19/2	MYLAN IRE HEALTHCARE LIMITED	IE
Twicor 10 mg/10 mg kalvopäällysteiset tabletit	NL/H/4126/001	35495	MYLAN FINLAND OY	FI
Twicor 10 mg/10 mg filmdragerade tabletter	NL/H/4126/001	35495	MYLAN FINLAND OY	FI
Twicor 20 mg/10 mg kalvopäällysteiset tabletit	NL/H/4126/002	35494	MYLAN FINLAND OY	FI
Twicor 20 mg/10 mg filmdragerade tabletter	NL/H/4126/002	35494	MYLAN FINLAND OY	FI
Myrosor 10 mg / 10 mg	NL/H/4126/001	2019050096	MYLAN EPD BVBA/SPRL	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimés pelliculés				
Myrosor 20 mg / 10 mg comprimés pelliculés	NL/H/4126/002	2019050097	MYLAN EPD BVBA/SPRL	LU
Myrosor 10 mg/10 mg Filmtabletten	NL/H/4126/001	2019050096	MYLAN EPD BVBA/SPRL	LU
Myrosor 20 mg/10 mg Filmtabletten	NL/H/4126/002	2019050097	MYLAN EPD BVBA/SPRL	LU
ROZOR 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/4126/001	62683/16-5-2019	BGP PRODUCTS S.LTD	GR
ROZOR 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/4126/002	62684/16-5-2019	BGP PRODUCTS S.LTD	GR
COMPUNA 5 mg/10 mg compresse	not available	045351018	BRUNO FARMACEUTICI	IT
COMPUNA 10 mg/10 mg compresse	not available	045351020	BRUNO FARMACEUTICI	IT
COMPUNA 20 mg/10 mg compresse	not available	045351032	BRUNO FARMACEUTICI	IT
Rosuzet 10 mg/10 mg Filmtabletten	NL/H/3679/001	97016.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Rosuzet 20 mg/10 mg Filmtabletten	NL/H/3679/002	2202348.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Rosuvastatine/Ezetimibe Althera 10 mg/10 mg filmomhulde tabletten	NL/H/3679/001	RVG 118790	ALTHERA LABORATORIES LIMITED	NL
Rosuvastatine/Ezetimibe Althera 20 mg/10 mg filmomhulde tabletten	NL/H/3679/002	RVG 123044	ALTHERA LABORATORIES LIMITED	NL
ROSUVASTATINA + EZETIMIBA BGP 10 mg + 10 mg comprimidos revestidos por película	NL/H/4175/001	5761218	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 10 mg + 10 mg comprimidos revestidos por película	NL/H/4175/001	5761226	BGP PRODUCTS UNIPESSOAL, LDA.	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
ROSUVASTATINA + EZETIMIBA BGP 10 mg + 10 mg comprimidos revestidos por película	NL/H/4175/001	5761234	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 10 mg + 10 mg comprimidos revestidos por película	NL/H/4175/001	5761242	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 20 mg + 10 mg comprimidos revestidos por película	NL/H/4175/002	5761259	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 20 mg + 10 mg comprimidos revestidos por película	NL/H/4175/002	5761267	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 20 mg + 10 mg comprimidos revestidos por película	NL/H/4175/002	5761275	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINA + EZETIMIBA BGP 20 mg + 10 mg comprimidos revestidos por película	NL/H/4175/002	5761309	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROSUVASTATINE/EZETIMIBE MYLAN 10 mg/10 mg, comprimé pelliculé	NL/H/4175/001	34009 301 640 7 5	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETIMIBE MYLAN 10 mg/10 mg, comprimé pelliculé	NL/H/4175/001	34009 301 640 8 2	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETIMIBE MYLAN 10 mg/10 mg, comprimé pelliculé	NL/H/4175/001	34009 301 641 0 5	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETIMIBE MYLAN 10 mg/10 mg, comprimé pelliculé	NL/H/4175/001	34009 301 641 1 2	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETI	NL/H/4175/002	34009 301 640 0 6	MYLAN MEDICAL SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MIBE MYLAN 20 mg/10 mg, comprimé pelliculé				
ROSUVASTATINE/EZETIMIBE MYLAN 20 mg/10 mg, comprimé pelliculé	NL/H/4175/002	34009 301 640 1 3	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETIMIBE MYLAN 20 mg/10 mg, comprimé pelliculé	NL/H/4175/002	34009 301 640 2 0	MYLAN MEDICAL SAS	FR
ROSUVASTATINE/EZETIMIBE MYLAN 20 mg/10 mg, comprimé pelliculé	NL/H/4175/002	34009 301 640 3 7	MYLAN MEDICAL SAS	FR
Rosuvastatine/Ezetimibe Mylan Healthcare 10 mg/10 mg filmomhulde tabletten	NL/H/4175/001	RVG 121849	MYLAN HEALTHCARE B.V.	NL
Rosuvastatine/Ezetimibe Mylan Healthcare 20 mg/10 mg filmomhulde tabletten	NL/H/4175/002	RVG 121853	MYLAN HEALTHCARE B.V.	NL
Ateroger 10 mg/10 mg comprimidos recubiertos con película	NL/H/4175/001	83657	MYLAN IRE HEALTHCARE LIMITED	ES
Ateroger 20 mg/10 mg comprimidos recubiertos con película	NL/H/4175/002	83658	MYLAN IRE HEALTHCARE LIMITED	ES
ROSUVASTATINA E EZETIMIBE MYLAN 10 mg/10 mg compresse rivestite con film	NL/H/4175/001	047294018	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 10 mg/10 mg compresse rivestite con film	NL/H/4175/001	047294020	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 10 mg/10 mg compresse rivestite con film	NL/H/4175/001	047294032	MYLAN IRE HEALTHCARE LIMITED	IT

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ROSUVASTATINA E EZETIMIBE MYLAN 10 mg/10 mg compresse rivestite con film	NL/H/4175/001	047294044	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 20 mg/10 mg compresse rivestite con film	NL/H/4175/002	047294057	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 20 mg/10 mg compresse rivestite con film	NL/H/4175/002	047294069	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 20 mg/10 mg compresse rivestite con film	NL/H/4175/002	047294071	MYLAN IRE HEALTHCARE LIMITED	IT
ROSUVASTATINA E EZETIMIBE MYLAN 20 mg/10 mg compresse rivestite con film	NL/H/4175/002	047294083	MYLAN IRE HEALTHCARE LIMITED	IT
Rozor 10 mg/10 mg Filmtabletten	NL/H/4175/001	138996	MYLAN ÖSTERREICH GMBH	AT
Rozor 20 mg/10 mg Filmtabletten	NL/H/4175/002	138997	MYLAN ÖSTERREICH GMBH	AT
Zuvatek, 5 mg + 10 mg, cápsulas	not available	5796339	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Zuvatek, 5 mg + 10 mg, cápsulas	not available	5796347	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Zuvatek, 10 mg + 10 mg, cápsulas	not available	5795810	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Zuvatek, 20 mg + 10 mg, cápsulas	not available	5796354	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Ezotera 5 mg/10 mg	HR-H-236009772	HR-H-236009772	JADRAN-GALENSKI LABORATORIJ	HR

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
tablete			D.D.	
Ezotera 10 mg/10 mg tablete	HR-H-782405252	HR-H-782405252	JADRAN-GALENSKI LABORATORIJ D.D.	HR
Ezotera 20 mg/10 mg tablete	HR-H-977343942	HR-H-977343942	JADRAN-GALENSKI LABORATORIJ D.D.	HR
ROSUVASTATINA E EZETIMIBE DOC 5 mg/10 mg compresse	not available	045827019	DOC GENERICI S.R.L.	IT
ROSUVASTATINA E EZETIMIBE DOC 5 mg/10 mg compresse	not available	045827021	DOC GENERICI S.R.L.	IT
ROSUVASTATINA E EZETIMIBE DOC 10 mg/10 mg compresse	not available	045827033	DOC GENERICI S.R.L.	IT
ROSUVASTATINA E EZETIMIBE DOC 10 mg/10 mg compresse	not available	045827045	DOC GENERICI S.R.L.	IT
ROSUVASTATINA E EZETIMIBE DOC 20 mg/10 mg compresse	not available	045827058	DOC GENERICI S.R.L.	IT
ROSUVASTATINA E EZETIMIBE DOC 20 mg/10 mg compresse	not available	045827060	DOC GENERICI S.R.L.	IT
Rosuvastatina + Ezetimiba Krka 5 mg + 10 mg comprimidos revestidos por película	CZ/H/0797/001	5761044	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + Ezetimiba Krka 10 mg + 10 mg comprimidos revestidos por película	CZ/H/0797/002	5761051	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + Ezetimiba Krka 20 mg + 10 mg comprimidos revestidos por película	CZ/H/0797/004	5761069	KRKA, D.D., NOVO MESTO	PT
Rosazimib 5 mg/10 mg apvalkotās tabletes	CZ/H/0797/001	18-0169	KRKA, D.D., NOVO MESTO	LV

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
Rosazimib 10 mg/10 mg apvalkotās tabletes	CZ/H/0797/002	18-0170	KRKA, D.D., NOVO MESTO	LV
Rosazimib 15 mg/10 mg apvalkotās tabletes	CZ/H/0797/003	18-0171	KRKA, D.D., NOVO MESTO	LV
Rosazimib 20 mg/10 mg apvalkotās tabletes	CZ/H/0797/004	18-0172	KRKA, D.D., NOVO MESTO	LV
Rosazimib 40 mg/10 mg apvalkotās tabletes	CZ/H/0797/005	18-0173	KRKA, D.D., NOVO MESTO	LV
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/001	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/002	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/003	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/004	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/005	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/006	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/007	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/008	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 5 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/001	LT/1/18/4296/009	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/001	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/002	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/003	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/004	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/005	KRKA, D.D., NOVO MESTO	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
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/006	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/007	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/008	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 10 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/002	LT/1/18/4297/009	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/001	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/002	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/003	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/004	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/005	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/006	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/007	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/008	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 15 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/003	LT/1/18/4298/009	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/001	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/002	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/003	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/004	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/005	KRKA, D.D., NOVO MESTO	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
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/006	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/007	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/008	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 20 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/004	LT/1/18/4299/009	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/001	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/002	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/003	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/004	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/005	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/006	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/007	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/008	KRKA, D.D., NOVO MESTO	LT
Sorvitimb 40 mg/10 mg plėvele dengtos tabletės	CZ/H/0797/005	LT/1/18/4300/009	KRKA, D.D., NOVO MESTO	LT
Rosazimib, 5 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0797/001	976618	KRKA, D.D., NOVO MESTO	EE
Rosazimib, 10 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0797/002	976718	KRKA, D.D., NOVO MESTO	EE
Rosazimib, 15 mg/10 mg õhukese polümeerikattega	CZ/H/0797/003	976818	KRKA, D.D., NOVO MESTO	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
tabletid				
Rosazimib, 20 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0797/004	976918	KRKA, D.D., NOVO MESTO	EE
Rosazimib, 40 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0797/005	977018	KRKA, D.D., NOVO MESTO	EE
Sorvasta Plus 5 mg/10 mg potahované tablety	CZ/H/0797/001	31/547/17-C	KRKA, D.D., NOVO MESTO	CZ
Sorvasta Plus 10 mg/10 mg potahované tablety	CZ/H/0797/002	31/548/17-C	KRKA, D.D., NOVO MESTO	CZ
Sorvasta Plus 15 mg/10 mg potahované tablety	CZ/H/0797/003	31/549/17-C	KRKA, D.D., NOVO MESTO	CZ
Sorvasta Plus 20 mg/10 mg potahované tablety	CZ/H/0797/004	31/550/17-C	KRKA, D.D., NOVO MESTO	CZ
Sorvasta Plus 40 mg/10 mg potahované tablety	CZ/H/0797/005	31/551/17-C	KRKA, D.D., NOVO MESTO	CZ
Co-Roswera 40 mg/10 mg filmom obložene tablete	CZ/H/0797/005	HR-H-145186580	KRKA-FARMA D.O.O.	HR
Co-Roswera 15 mg/10 mg filmom obložene tablete	CZ/H/0797/003	HR-H-319752844	KRKA-FARMA D.O.O.	HR
Co-Roswera 5 mg/10 mg filmom obložene tablete	CZ/H/0797/001	HR-H-578445810	KRKA-FARMA D.O.O.	HR
Co-Roswera 10 mg/10 mg filmom obložene tablete	CZ/H/0797/002	HR-H-775674698	KRKA-FARMA D.O.O.	HR
Co-Roswera 20 mg/10 mg filmom obložene tablete	CZ/H/0797/004	HR-H-919934131	KRKA-FARMA D.O.O.	HR
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/02	KRKA, D.D., NOVO MESTO	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
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 5 mg/10 mg comprimate filmate	CZ/H/0797/001	11331/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 10 mg/10 mg comprimate filmate	CZ/H/0797/002	11332/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/02	KRKA, D.D., NOVO MESTO	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
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 15 mg/10 mg comprimate filmate	CZ/H/0797/003	11333/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 20 mg/10 mg comprimate filmate	CZ/H/0797/004	11334/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/02	KRKA, D.D., NOVO MESTO	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
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Roswera 40 mg/10 mg comprimate filmate	CZ/H/0797/005	11335/2019/09	KRKA, D.D., NOVO MESTO	RO
Rosazimib 5 mg/10 mg filmom obalené tablety	CZ/H/0797/001	31/0011/19-S	KRKA, D.D., NOVO MESTO	SK
Rosazimib 10 mg/10 mg filmom obalené tablety	CZ/H/0797/002	31/0012/19-S	KRKA, D.D., NOVO MESTO	SK
Rosazimib 15 mg/10 mg filmom obalené tablety	CZ/H/0797/003	31/0013/19-S	KRKA, D.D., NOVO MESTO	SK
Rosazimib 20 mg/10 mg filmom obalené tablety	CZ/H/0797/004	31/0014/19-S	KRKA, D.D., NOVO MESTO	SK
Rosazimib 40 mg/10 mg filmom obalené tablety	CZ/H/0797/005	31/0015/19-S	KRKA, D.D., NOVO MESTO	SK
Ко-Розвера 5 mg/10 mg филмирани таблетки	CZ/H/0797/001	20190014	KRKA, D.D., NOVO MESTO	BG
Ко-Розвера 10 mg/10 mg филмирани таблетки	CZ/H/0797/002	20190015	KRKA, D.D., NOVO MESTO	BG
Ко-Розвера 15 mg/10 mg филмирани таблетки	CZ/H/0797/003	20190016	KRKA, D.D., NOVO MESTO	BG
Ко-Розвера 20 mg/10 mg филмирани таблетки	CZ/H/0797/004	20190017	KRKA, D.D., NOVO MESTO	BG
Ко-Розвера 40 mg/10 mg филмирани	CZ/H/0797/005	20190018	KRKA, D.D., NOVO MESTO	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
таблетки				
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/001	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/002	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/003	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/004	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/005	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/006	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/007	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/008	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/009	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/010	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 5 mg/10 mg filmsko obložene tablete	CZ/H/0797/001	H/19/02549/011	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/012	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/013	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/014	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/015	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/016	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/017	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg	CZ/H/0797/002	H/19/02549/018	KRKA, D.D., NOVO MESTO	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
filmsko obložene tablete				
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/019	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/020	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/021	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 10 mg/10 mg filmsko obložene tablete	CZ/H/0797/002	H/19/02549/022	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/023	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/024	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/025	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/026	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/027	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/028	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/029	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/030	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/031	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/032	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 15 mg/10 mg filmsko obložene tablete	CZ/H/0797/003	H/19/02549/033	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/034	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/035	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg	CZ/H/0797/004	H/19/02549/036	KRKA, D.D., NOVO MESTO	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
filmsko obložene tablete				
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/037	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/038	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/039	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/040	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/041	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/042	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/043	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 20 mg/10 mg filmsko obložene tablete	CZ/H/0797/004	H/19/02549/044	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/045	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/046	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/047	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/048	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/049	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/050	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/051	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/052	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/053	KRKA, D.D., NOVO MESTO	SI
Sorvitimb 40 mg/10 mg	CZ/H/0797/005	H/19/02549/054	KRKA, D.D., NOVO MESTO	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
filmsko obložene tablete				
Sorvitimb 40 mg/10 mg filmsko obložene tablete	CZ/H/0797/005	H/19/02549/055	KRKA, D.D., NOVO MESTO	SI
Coroswera, 5 mg + 10 mg, tabletki powlekane	CZ/H/0797/001	25209	KRKA, D.D., NOVO MESTO	PL
Coroswera, 10 mg + 10 mg, tabletki powlekane	CZ/H/0797/002	25210	KRKA, D.D., NOVO MESTO	PL
Coroswera, 15 mg + 10 mg, tabletki powlekane	CZ/H/0797/003	25211	KRKA, D.D., NOVO MESTO	PL
Coroswera, 20 mg + 10 mg, tabletki powlekane	CZ/H/0797/004	25212	KRKA, D.D., NOVO MESTO	PL
Coroswera, 40 mg + 10 mg, tabletki powlekane	CZ/H/0797/005	25213	KRKA, D.D., NOVO MESTO	PL
Rosuvastatina + Ezetimiba Krka 10 mg + 10 mg comprimidos revestidos por película	CZ/H/0797/002	5765540	KRKA, D.D., NOVO MESTO	PT
Rosu - 1 A Pharma plus Ezetimib 5 mg/10 mg Tabletten	not available	2202486.00.00	1 A PHARMA GMBH	DE
Rosu - 1 A Pharma plus Ezetimib 20 mg/10 mg Tabletten	not available	2202488.00.00	1 A PHARMA GMBH	DE
Rosu - 1 A Pharma plus Ezetimib 10 mg/10 mg Tabletten	not available	2202487.00.00	1 A PHARMA GMBH	DE
Rosuvastatine/Ezetimibe Ferrer 5 mg/10 mg film-coated tablets	not available	125896	FERRER INTERNACIONAL, S.A.	NL
Rosuvastatine/Ezetimibe Ferrer 10 mg/10 mg film-coated tablets	not available	125897	FERRER INTERNACIONAL, S.A.	NL
Rosuvastatine/Ezetimibe Ferrer 20 mg/10 mg film-coated tablets	not available	125898	FERRER INTERNACIONAL, S.A.	NL
Rosuvastatine/Ezetimibe Ferrer 40 mg/10 mg	not available	125899	FERRER INTERNACIONAL, 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
film-coated tablets				
Alzil plus 5 mg/10 mg comprimidos recubiertos con película	NL/H/5090/001-004/MR	85379	FERRER INTERNACIONAL, S.A.	ES
Alzil plus 10 mg/10 mg comprimidos recubiertos con película	NL/H/5090/002/MR	85376	FERRER INTERNACIONAL, S.A.	ES
Alzil plus 20 mg/10 mg comprimidos recubiertos con película	NL/H/5090/003/MR	85377	FERRER INTERNACIONAL, S.A.	ES
Alzil plus 40 mg/10 mg comprimidos recubiertos con película	NL/H/5090/004/MR	85378	FERRER INTERNACIONAL, S.A.	ES
Rosuvastatin/Ezetimib Genericon 20 mg/10 mg tablete	AT/H/1079/003	HR-H-870499295	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Rosuvastatin/Ezetimib Genericon 10 mg/10 mg tablete	AT/H/1079/002	HR-H-561909678	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Rosuvastatin/Ezetimib Genericon 5 mg/10 mg tablete	AT/H/1079/001	HR-H-853800213	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Rosamib 5 mg/10 mg Tabletten	AT/H/1079/001	138932	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Rosamib 10 mg/10 mg Tabletten	AT/H/1079/002	138933	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Rosamib 20 mg/10 mg Tabletten	AT/H/1079/003	138934	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Delipid Plus 10 mg/10 mg kemény kapszula	NL/H/3016/001/DC	OGYI-T-22700/01	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 20 mg/10 mg kemény kapszula	NL/H/3016/002/DC	OGYI-T-22700/04	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 40 mg/10 mg kemény kapszula	NL/H/3016/003/DC	OGYI-T-22700/07	EGIS PHARMACEUTICALS PLC	HU
ROSUVASTATINE/EZETIMIBE EGIS 20 mg/10 mg, gélule	NL/H/3016/002	6 263 525 9	EGIS PHARMACEUTICALS PLC	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ROSUVASTATINE/EZETIMIBE EGIS 10 mg/10 mg, gélule	NL/H/3016/001	6 048 151 7	EGIS PHARMACEUTICALS PLC	FR
Rosuvastatine/ezetimibe EGIS 10 mg/10 mg harde capsules	NL/H/3016/001	RVG 113986	EGIS PHARMACEUTICALS PLC	NL
Rosuvastatine/ezetimibe EGIS 20 mg/10 mg harde capsules	NL/H/3016/002	RVG 113991	EGIS PHARMACEUTICALS PLC	NL
Rosuvastatine/ezetimibe EGIS 40 mg/10 mg capsule, hard	NL/H/3016/003	RVG 113992	EGIS PHARMACEUTICALS PLC	NL
Rosuvastatine/ezetimibe EGIS 5 mg/10 mg harde capsules	NL/H/3016/004	RVG 119682	EGIS PHARMACEUTICALS PLC	NL
Delipid Kombi 10mg/10mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/01-03	EGIS PHARMACEUTICALS PLC	HU
Delipid Kombi 20 mg/10 mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/04-06	EGIS PHARMACEUTICALS PLC	HU
Delipid Kombi 10mg/10mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/01-03	EGIS PHARMACEUTICALS PLC	HU
Delipid Kombi 10mg/10mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/01-03	EGIS PHARMACEUTICALS PLC	HU
Delipid Kombi 20 mg/10 mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/04-06	EGIS PHARMACEUTICALS PLC	HU
Delipid Kombi 20 mg/10 mg kemény kapszula	NL/H/3016/005-006/DC:	OGYI-T-23379/04-06	EGIS PHARMACEUTICALS PLC	HU
Rosuvastatine calcium/ezetimibe Egis 10 mg/10 mg harde capsules	NL/H/3016/005	RVG 120480	EGIS PHARMACEUTICALS PLC	NL
Rosuvastatine calcium/ezetimibe Egis 20 mg/10 mg harde	NL/H/3016/006	RVG 120481	EGIS PHARMACEUTICALS PLC	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
capsules				
Delipid Plus 5 mg/10 mg kemény kapszula	NL/H/3016/004	OGYI-T-22700/10	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 5 mg/10 mg kemény kapszula	NL/H/3016/004	OGYI-T-22700/11	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 5 mg/10 mg kemény kapszula	NL/H/3016/004	OGYI-T-22700/12	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 10 mg/10 mg kemény kapszula	NL/H/3016/001	OGYI-T-22700/02	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 10 mg/10 mg kemény kapszula	NL/H/3016/001	OGYI-T-22700/03	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 20 mg/10 mg kemény kapszula	NL/H/3016/002	OGYI-T-22700/05	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 20 mg/10 mg kemény kapszula	NL/H/3016/002	OGYI-T-22700/06	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 40 mg/10 mg kemény kapszula	NL/H/3016/003	OGYI-T-22700/08	EGIS PHARMACEUTICALS PLC	HU
Delipid Plus 40 mg/10 mg kemény kapszula	NL/H/3016/003	OGYI-T-22700/09	EGIS PHARMACEUTICALS PLC	HU
Rosuvastatin/Ezetimib Aristo 5 mg + 10 mg tabletki	PL/H/0499/001	24672	ARISTO PHARMA GMBH (ART 57)	PL
Rosuvastatin/Ezetimib Aristo 10 mg + 10 mg tabletki	PL/H/0499/002	24673	ARISTO PHARMA GMBH (ART 57)	PL
Rosuvastatin/Ezetimib Aristo 20 mg + 10 mg tabletki	PL/H/0499/003	24674	ARISTO PHARMA GMBH (ART 57)	PL
Quiloga 5 mg/10 mg compresse	PL/H/0499/001	046503013	ARISTO PHARMA GMBH (ART 57)	IT
Quiloga 5 mg/10 mg compresse	PL/H/0499/001	046503025	ARISTO PHARMA GMBH (ART 57)	IT
Quiloga 10 mg/10 mg compresse	PL/H/0499/002	046503037	ARISTO PHARMA GMBH (ART 57)	IT
Quiloga 10 mg/10 mg compresse	PL/H/0499/002	046503049	ARISTO PHARMA GMBH (ART 57)	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
Quiloga 20 mg/10 mg compresse	PL/H/0499/003	046503052	ARISTO PHARMA GMBH (ART 57)	IT
Quiloga 20 mg/10 mg compresse	PL/H/0499/003	046503064	ARISTO PHARMA GMBH (ART 57)	IT
Ezerosu 10 mg/5 mg-Filmtabletten	CZ/H/0855/001	139051	G.L. PHARMA GMBH	AT
Ezerosu 10 mg/10 mg-Filmtabletten	CZ/H/0855/002	139052	G.L. PHARMA GMBH	AT
Ezerosu 10 mg/20 mg-Filmtabletten	CZ/H/0855/003	139053	G.L. PHARMA GMBH	AT
Ezerosu 10 mg/40 mg-Filmtabletten	CZ/H/0855/004	139054	G.L. PHARMA GMBH	AT
Rosuvastatin/Ezetimibe Elpen 5 mg/10 mg potahované tablety	CZ/H/0855/001	31/250/18-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
Rosuvastatin/Ezetimibe Elpen 10 mg/10 mg potahované tablety	CZ/H/0855/002	31/251/18-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
Rosuvastatin/Ezetimibe Elpen 20 mg/10 mg potahované tablety	CZ/H/0855/003	31/252/18-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
Rosuvastatin/Ezetimibe Elpen 40 mg/10 mg potahované tablety	CZ/H/0855/004	31/253/18-C	ELPEN PHARMACEUTICAL CO. INC.	CZ
Rosuvastatin/Ezetimib Elpen 5 mg/10 mg Filmtabletten	CZ/H/0855/001	2202637.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Rosuvastatin/Ezetimib Elpen 10 mg/10 mg Filmtabletten	CZ/H/0855/002	2202638.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Rosuvastatin/Ezetimib Elpen 20 mg/10 mg Filmtabletten	CZ/H/0855/003	2202639.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Rosuvastatin/Ezetimib Elpen 40 mg/10 mg Filmtabletten	CZ/H/0855/004	2202640.00.00	ELPEN PHARMACEUTICAL CO. INC.	DE
Rosuvastatina/Ezetimiba	CZ/H/0855/001	84.658	LABORATORIO STADA, S.L.	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
Stadagen 5 mg/10 mg comprimidos recubiertos con película				
Rosuvastatina/Ezetimiba Stadagen 40 mg/10 mg comprimidos recubiertos con película	CZ/H/0855/004	84.657	LABORATORIO STADA, S.L.	ES
Rosuvastatina/Ezetimiba Stadagen 10 mg/10 mg comprimidos recubiertos con película	CZ/H/0855/002	84.659	LABORATORIO STADA, S.L.	ES
Rosuvastatina/Ezetimiba Stadagen 20 mg/10 mg comprimidos recubiertos con película	CZ/H/0855/003	84.660	LABORATORIO STADA, S.L.	ES
Lipopen 5 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0855/001	35568/20-03-2020	ELPEN PHARMACEUTICAL CO. INC.	GR
Lipopen 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0855/002	35569/20-03-2020	ELPEN PHARMACEUTICAL CO. INC.	GR
Lipopen 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0855/003	35570/20-03-2020	ELPEN PHARMACEUTICAL CO. INC.	GR
Lipopen 40 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0855/004	35571/20-03-2020	ELPEN PHARMACEUTICAL CO. INC.	GR
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/01	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/02	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/03	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/04	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/05	KRKA, D.D., NOVO MESTO	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
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/06	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/07	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/08	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 5 mg/10 mg filmtabletta	not available	OGYI-T-23542/09	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/10	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/11	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/12	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/13	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/14	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/15	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/16	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/17	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 10 mg/10 mg filmtabletta	not available	OGYI-T-23542/18	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/19	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/20	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/21	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/22	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/23	KRKA, D.D., NOVO MESTO	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
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/24	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/25	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/26	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 15 mg/10 mg filmtabletta	not available	OGYI-T-23542/27	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/28	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/29	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/30	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/31	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/32	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/33	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/34	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/35	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 20 mg/10 mg filmtabletta	not available	OGYI-T-23542/36	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/37	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/38	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/39	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/40	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/41	KRKA, D.D., NOVO MESTO	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
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/42	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/43	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/44	KRKA, D.D., NOVO MESTO	HU
Roxera Plus 40 mg/10 mg filmtabletta	not available	OGYI-T-23542/45	KRKA, D.D., NOVO MESTO	HU
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	5796677	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	5796701	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 5 mg	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
+ 10 mg, cápsulas				
Rosuvastatina + Ezetimiba Strami, 5 mg + 10 mg, cápsulas	not available	20/H/0023/001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	5796719	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 10 mg + 10 mg, cápsulas	not available	20/H/0023/002	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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**
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	5796727	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Rosuvastatina + Ezetimiba Strami, 20 mg + 10 mg, cápsulas	not available	20/H/0023/003	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
EZATEROS 20 mg/10 mg compresse	not available	045432034	SO.SE.PHARM S.R.L.	IT
EZATEROS 10 mg/10 mg compresse	not available	045432022	SO.SE.PHARM S.R.L.	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
EZATEROS 5mg/10mg compresse	not available	045432010	SO.SE.PHARM S.R.L.	IT
Rosuvastatin+Ezetimibe/ RAFARM 20 mg/10 mg δισκία	PL/H/0556/003	83363/26-08-2020	RAFARM SA.	GR
Rosuvastatin+Ezetimibe/ RAFARM 10 mg/10 mg δισκία	PL/H/0556/002	83362/26-08-2020	RAFARM SA.	GR
Rosuvastatin+Ezetimibe/ RAFARM 5 mg/10 mg δισκία	PL/H/0556/001	83361/26-08-2020	RAFARM SA.	GR
Rosumibe 5 mg + 10 mg comprimidos revestidos por película	not available	5784111	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 5 mg + 10 mg comprimidos revestidos por película	not available	5784129	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 10 mg + 10 mg comprimidos revestidos por película	not available	5784012	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 10 mg + 10 mg comprimidos revestidos por película	not available	5784020	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 10 mg + 10 mg comprimidos revestidos por película	not available	5784004	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 20 mg + 10 mg comprimidos revestidos por película	not available	5784046	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 20 mg + 10 mg comprimidos revestidos por película	not available	5784038	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Rosumibe 40 mg + 10 mg comprimidos revestidos por película	not available	5784053	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Lipocomb 10 mg/10 mg Hartkapseln	NL/H/3006/001-003/DC	135783	EGIS PHARMACEUTICALS PLC	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
Lipocomb 20 mg/10 mg Hartkapseln	NL/H/3006/001-003/DC	135782	EGIS PHARMACEUTICALS PLC	AT
Lipocomb 40 mg/10 mg Hartkapseln	NL/H/3006/001-003/DC	135784	EGIS PHARMACEUTICALS PLC	AT
Delipid Plus 10 mg/10 mg tvrdé tobolky	NL/H/3006/001-003/DC	31/354/14-C	EGIS PHARMACEUTICALS PLC	CZ
Delipid Plus 20 mg/10 mg tvrdé tobolky	NL/H/3006/001-003/DC	31/355/14-C	EGIS PHARMACEUTICALS PLC	CZ
Delipid Plus 40 mg/10 mg tvrdé tobolky	NL/H/3006/001-003/DC	31/356/14-C	EGIS PHARMACEUTICALS PLC	CZ
Lipocomb 10 mg/10 mg hard capsules	NL/H/3006/001-003/DC	PA1470/004/001	EGIS PHARMACEUTICALS PLC	IE
Lipocomb 20 mg/10 mg hard capsules	NL/H/3006/001-003/DC	PA1470/004/002	EGIS PHARMACEUTICALS PLC	IE
Lipocomb 40 mg/10 mg hard capsules	NL/H/3006/001-003/DC	PA1470/004/003	EGIS PHARMACEUTICALS PLC	IE
Lipocomb 10 mg/10 mg capsule	NL/H/3006/001-003/DC	11753/2019/01-07	EGIS PHARMACEUTICALS PLC	RO
Lipocomb 20 mg/10 mg capsule	NL/H/3006/001-003/DC	11754/2019/01-07	EGIS PHARMACEUTICALS PLC	RO
Lipocomb 40 mg/10 mg capsule	NL/H/3006/001-003/DC	11755/2019/01-07	EGIS PHARMACEUTICALS PLC	RO
Lipocomb 10 mg/10 mg tvrdé kapsuly	NL/H/3006/001-003/DC	31/0289/14-S	EGIS PHARMACEUTICALS PLC	SK
Lipocomb 20 mg/10 mg tvrdé kapsuly	NL/H/3006/001-003/DC	31/0290/14-S	EGIS PHARMACEUTICALS PLC	SK
Lipocomb 40 mg/10 mg tvrdé kapsuly	NL/H/3006/001-003/DC	31/0291/14-S	EGIS PHARMACEUTICALS PLC	SK
Lipocomb Neo 10 mg/10 mg tvrdé kapsuly	NL/H/3006/004	31/0128/18-S	EGIS PHARMACEUTICALS PLC	SK
Lipocomb Neo 20 mg/10 mg tvrdé kapsuly	NL/H/3006/005	31/0129/18-S	EGIS PHARMACEUTICALS PLC	SK
Cholecomb calcium 10 mg/10 mg harde capsules	NL/H/3006/004	RVG 120478	EGIS PHARMACEUTICALS PLC	NL
Cholecomb calcium 20	NL/H/3006/005	RVG 120484	EGIS PHARMACEUTICALS PLC	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
mg/10 mg harde capsules				
Delipid Duo 10 mg/10 mg tvrdé tobolky	NL/H/3006/004-005/DC	31/1028/16-C	EGIS PHARMACEUTICALS PLC	CZ
Delipid Duo 20 mg/10 mg tvrdé tobolky	NL/H/3006/004-005/DC	31/1029/16-C	EGIS PHARMACEUTICALS PLC	CZ
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Suvezen 10 mg/10 mg comprimés pelliculés	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Suvezen 10 mg/10 mg filmomhulde tabletten	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Suvezen 20 mg/10 mg comprimés pelliculés	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 20 mg/10 mg filmomhulde tabletten	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 40 mg/10 mg comprimés pelliculés	CZ/H/0696/003	BE539946	SANOFI BELGIUM	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
Suvezen 40 mg/10 mg filmomhulde tabletten	CZ/H/0696/003	BE539946	SANOFI BELGIUM	BE
Suvezen 10 mg/10 mg comprimate filmate	CZ/H/0696/001	11631/2019/02	SANOFI ROMANIA SRL	RO
Suvezen 40 mg/10 mg comprimate filmate	CZ/H/0696/003	11633/2019/02	SANOFI ROMANIA SRL	RO
Suvezen 10 mg/10 mg comprimate filmate	CZ/H/0696/001	11631/2019/01	SANOFI ROMANIA SRL	RO
Suvezen 40 mg/10 mg comprimate filmate	CZ/H/0696/003	11633/2019/01	SANOFI ROMANIA SRL	RO
Suvezen 20 mg/10 mg comprimate filmate	CZ/H/0696/002	11632/2019/02	SANOFI ROMANIA SRL	RO
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon Novum 10 mg/10 mg filmom obalené tablety	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Suvezen 10 mg/10 mg filmsko obložene tablete	CZ/H/0696/001	H/19/02591/001	SANOFI-AVENTIS D.O.O.	SI
Suvezen 20 mg/10 mg filmsko obložene tablete	CZ/H/0696/002	H/19/02591/003	SANOFI-AVENTIS D.O.O.	SI
Suvezen 40 mg/10 mg filmsko obložene tablete	CZ/H/0696/003	H/19/02591/005	SANOFI-AVENTIS D.O.O.	SI
Suvezen 10 mg/10 mg filmsko obložene tablete	CZ/H/0696/001	H/19/02591/002	SANOFI-AVENTIS D.O.O.	SI
Suvezen 20 mg/10 mg	CZ/H/0696/002	H/19/02591/004	SANOFI-AVENTIS D.O.O.	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
filmsko obložene tablete				
Suvezen 40 mg/10 mg filmsko obložene tablete	CZ/H/0696/003	H/19/02591/006	SANOFI-AVENTIS D.O.O.	SI
Suvezen 10 mg/10 mg filmomhulde tabletten	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Suvezen 10 mg/10 mg comprimés pelliculés	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Zenon Novum 10 mg/10 mg filmom obalené tablety	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Suvezen 20 mg/10 mg filmomhulde tabletten	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 20 mg/10 mg comprimés pelliculés	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 40 mg/10 mg comprimés pelliculés	CZ/H/0696/003	BE539946	SANOFI BELGIUM	BE
Suvezen 40 mg/10 mg filmomhulde tabletten	CZ/H/0696/003	BE539946	SANOFI BELGIUM	BE
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg+10 mg comprimidos revestidos por película	CZ/H/0696/001	5772843	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Zenon 40 mg+10 mg comprimidos revestidos por película	CZ/H/0696/003	CZ/H/0696/003	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Zenon 40 mg+10 mg comprimidos revestidos por película	CZ/H/0696/003	5772868	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Zenon 20 mg+10 mg comprimidos revestidos por película	CZ/H/0696/002	5772850	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Zenon 20 mg+10 mg comprimidos revestidos por película	CZ/H/0696/002	CZ/H/0696/002	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Suvreza 10 mg/10 mg	CZ/H/0696/001	046072029	SANOFI S.P.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
compresse rivestite con film				
Suvreza 10 mg/10 mg compresse rivestite con film	CZ/H/0696/001	046072017	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072031	SANOFI S.P.A	IT
Suvreza 40 mg/10 mg compresse rivestite con film	CZ/H/0696/003	046072068	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072043	SANOFI S.P.A	IT
Suvreza 40 mg/10 mg compresse rivestite con film	CZ/H/0696/003	046072056	SANOFI S.P.A	IT
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	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
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 20 mg + 10 mg, tabletki powlekane	CZ/H/0696/002	25499	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 40 mg + 10 mg, tabletki powlekane	CZ/H/0696/003	25500	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen 10 mg/10 mg film-coated tablets	CZ/H/0696/001	PA 540/193/001	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Suvezen 10 mg/10 mg film-coated tablets	CZ/H/0696/001	PA 540/193/001	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Zenon 10 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO
Zenon 10 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO
Zenon 10 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO
Zenon 10 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO
Zenon 10 mg/10 mg	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
filmdrasjerte tabletter				
Zenon 10 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/001	17-11975	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon 40 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/003	17-11977	SANOFI-AVENTIS NORGE AS	NO
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 20 mg/10 mg potahované tablety	CZ/H/0696/002	31/479/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 40 mg/10 mg potahované tablety	CZ/H/0696/003	31/480/17-C	SANOFI-AVENTIS SRO	CZ
Zenon 20 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO
Zenon 20 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO
Zenon 20 mg/10 mg	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmdrasjerte tabletter				
Zenon 20 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO
Zenon 20 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO
Zenon 20 mg/10 mg filmdrasjerte tabletter	CZ/H/0696/002	17-11976	SANOFI-AVENTIS NORGE AS	NO
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
SUVEZEN NEO, 10 mg + 10 mg, tabletki powlekane	CZ/H/0696/001	25498	SANOFI-AVENTIS SP Z.O.O.	PL
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ
Zenon Neo 10 mg/10 mg potahované tablety	CZ/H/0696/001	31/478/17-C	SANOFI-AVENTIS SRO	CZ

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
Zenon Neo 40 mg/10 mg film-coated tablets	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Zenon Neo 40 mg/10 mg film-coated tablets	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Zenon Neo 10 mg/10 mg film-coated tablets	CZ/H/0696/001	20190200	SANOFI BULGARIA EOOD	BG
Zenon Neo 10 mg/10 mg film-coated tablets	CZ/H/0696/001	20190200	SANOFI BULGARIA EOOD	BG
Zenon Neo 20 mg/10 mg film-coated tablets	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	BG
Zenon Neo 20 mg/10 mg film-coated tablets	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	BG
Suvreza 10 mg/10 mg compresse rivestite con film	CZ/H/0696/001	046072070	SANOFI S.P.A	IT
Suvreza 10 mg/10 mg compresse rivestite con film	CZ/H/0696/001	046072094	SANOFI S.P.A	IT
Suvreza 10 mg/10 mg compresse rivestite con film	CZ/H/0696/001	046072082	SANOFI S.P.A	IT
Suvreza 10 mg/10 mg compresse rivestite con film	CZ/H/0696/001	046072106	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072118	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072144	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072120	SANOFI S.P.A	IT
Suvreza 20 mg/10 mg compresse rivestite con film	CZ/H/0696/002	046072132	SANOFI S.P.A	IT
Suvreza 40 mg/10 mg	CZ/H/0696/003	046072171	SANOFI S.P.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
compresse rivestite con film				
Suvreza 40 mg/10 mg compresse rivestite con film	CZ/H/0696/003	046072169	SANOFI S.P.A	IT
Suvreza 40 mg/10 mg compresse rivestite con film	CZ/H/0696/003	046072183	SANOFI S.P.A	IT
Suvreza 40 mg/10 mg compresse rivestite con film	CZ/H/0696/003	046072157	SANOFI S.P.A	IT
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 10 mg/10 mg filmdragerad tablett	CZ/H/0696/001	57180	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE
Suvezen, 40 mg/10 mg filmdragerad tablett	CZ/H/0696/003	57182	SANOFI AB	SE
Suvezen, 20 mg/10 mg filmdragerad tablett	CZ/H/0696/002	57181	SANOFI AB	SE
Suvezen 10 mg/10 mg Filmtabletten	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Suvezen 20 mg/10 mg Filmtabletten	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 10 mg/10 mg Filmtabletten	CZ/H/0696/001	BE539920	SANOFI BELGIUM	BE
Suvezen 40 mg/10 mg Filmtabletten	CZ/H/0696/003	BE539946	SANOFI BELGIUM	BE
Suvezen 20 mg/10 mg Filmtabletten	CZ/H/0696/002	BE539937	SANOFI BELGIUM	BE
Suvezen 40 mg/10 mg Filmtabletten	CZ/H/0696/003	BE539946	SANOFI BELGIUM	BE
Zenon 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/001	13304/18-2-2020	SANOFI-AVENTIS AEBE	GR
Zenon 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/001	13304/18-2-2020	SANOFI-AVENTIS AEBE	GR
Zenon 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/002	13305/18-2-2020	SANOFI-AVENTIS AEBE	GR
Zenon 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/002	13305/18-2-2020	SANOFI-AVENTIS AEBE	GR
Zenon 40 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/003	13306/18-2-2020	SANOFI-AVENTIS AEBE	GR

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
Zenon 40 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/003	13306/18-2-2020	SANOFI-AVENTIS AEBE	GR
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 40 mg/10 mg filmdragerade tabletter	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg/10 mg filmdragerade tabletter	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/001	023096	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/002	023097	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/001	023096	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 40 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/003	023098	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 40 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/003	023098	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 20 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	CZ/H/0696/002	023097	SANOFI-AVENTIS CYPRUS LTD	CY
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-05	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-06	SANOFI-AVENTIS CROATIA D.O.O.	HR

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
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-03	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 40 mg/10 mg filmom obložene tablete	CZ/H/0696/003	HR-H-083490962-04	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-03	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-04	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-05	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 10 mg/10 mg filmom obložene tablete	CZ/H/0696/001	HR-H-523491391-06	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-03	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-04	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-05	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg/10 mg filmom obložene tablete	CZ/H/0696/002	HR-H-348105514-06	SANOFI-AVENTIS CROATIA D.O.O.	HR
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/001	35652	SANOFI OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 20 mg/10 mg filmdragerade tabletter	CZ/H/0696/002	35653	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg/10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg/10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg/10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 40 mg / 10 mg kalvopäällysteiset tabletit	CZ/H/0696/003	35654	SANOFI OY	FI
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon 10 mg/10 mg filmdragerade tabletter	CZ/H/0696/001	35652	SANOFI OY	FI
Zenon Novum 10 mg/10 mg filmom obalené tablety	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 10 mg/10	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg filmom obalené tablety				
Zenon Novum 10 mg/10 mg filmom obalené tablety	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 10 mg/10 mg filmom obalené tablety	CZ/H/0696/001	31/0075/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 20 mg/10 mg filmom obalené tablety	CZ/H/0696/002	31/0076/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon Novum 40 mg/10 mg filmom obalené tablety	CZ/H/0696/003	31/0077/19-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Зенон Нео 20 mg/10 mg филмирани таблетки	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	BG
Зенон Нео 20 mg/10 mg филмирани таблетки	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	BG
Зенон Нео 20 mg/10 mg	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	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
филмирани таблетки				
Зенон Нео 20 mg/10 mg филмирани таблетки	CZ/H/0696/002	20190201	SANOFI BULGARIA EOOD	BG
Зенон Нео 40 mg/10 mg филмирани таблетки	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Зенон Нео 40 mg/10 mg филмирани таблетки	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Зенон Нео 40 mg/10 mg филмирани таблетки	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Зенон Нео 40 mg/10 mg филмирани таблетки	CZ/H/0696/003	20190202	SANOFI BULGARIA EOOD	BG
Росулип плюс 10 mg/10 mg твърди капсули	NL/H/3007/001-003/DC	20140350	EGIS PHARMACEUTICALS PLC	BG
Росулип плюс 20 mg/10 mg твърди капсули	NL/H/3007/001-003/DC	20140351	EGIS PHARMACEUTICALS PLC	BG
Росулип плюс 40 mg/10 mg твърди капсули	NL/H/3007/001-003/DC	20140352	EGIS PHARMACEUTICALS PLC	BG
Rosulip 10 mg/10 mg cietās kapsulas	NL/H/3007/001-003/DC	14-0194	EGIS PHARMACEUTICALS PLC	LV
Rosulip 20 mg/10 mg cietās kapsulas	NL/H/3007/001-003/DC	14-0195	EGIS PHARMACEUTICALS PLC	LV
Rosulip 40 mg/10 mg cietās kapsulas	NL/H/3007/001-003/DC	14-0196	EGIS PHARMACEUTICALS PLC	LV
Rosulip Plus, 10 mg + 10 mg, kapsułki, twarde	NL/H/3007/001-003/DC	22082	EGIS PHARMACEUTICALS PLC	PL
Rosulip Plus, 20 mg + 10 mg, kapsułki, twarde	NL/H/3007/001-003/DC	22083	EGIS PHARMACEUTICALS PLC	PL
Rosulip Plus, 40 mg + 10 mg, kapsułki, twarde	NL/H/3007/001-003/DC	22084	EGIS PHARMACEUTICALS PLC	PL
Cholecomb 10 mg/10 mg, capsule hard	NL/H/3007/001-003/DC	RVG 114002	EGIS PHARMACEUTICALS PLC	NL
Cholecomb 20 mg/10 mg, capsule hard	NL/H/3007/001-003/DC	RVG 114003	EGIS PHARMACEUTICALS PLC	NL
Cholecomb 40 mg/10 mg, capsule hard	NL/H/3007/001-003/DC	RVG 114004	EGIS PHARMACEUTICALS PLC	NL
Ayadont 40 mg/10 mg	NL/H/3007/003	RVG 114004	EGIS PHARMACEUTICALS PLC	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
harde capsule				
Ayadont 10 mg/10 mg, harde capsules	NL/H/3007/001	RVG 114002	EGIS PHARMACEUTICALS PLC	NL
Ayadont 20 mg/10 mg, harde capsules	NL/H/3007/002	RVG 114003	EGIS PHARMACEUTICALS PLC	NL
Delipid Plus 10 mg/10 mg kõvakapslid	NL/H/3007/001	850014	EGIS PHARMACEUTICALS PLC	EE
Delipid Plus 20 mg/10 mg kõvakapslid	NL/H/3007/002	849914	EGIS PHARMACEUTICALS PLC	EE
Delipid Plus 40 mg/10 mg kõvakapslid	NL/H/3007/003	849814	EGIS PHARMACEUTICALS PLC	EE
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Lipocomb 10 mg/10 mg σκληρά καψάκια	NL/H/3007/001-003/DC	3162401	EGIS PHARMACEUTICALS PLC	GR
Lipocomb 20 mg/10 mg σκληρά καψάκια	NL/H/3007/001-003/DC	3162402	EGIS PHARMACEUTICALS PLC	GR
Lipocomb 40 mg/10 mg σκληρά καψάκια	NL/H/3007/001-003/DC	3162403	EGIS PHARMACEUTICALS PLC	GR
Ayadont 5 mg/10 mg capsule, hard	NL/H/3007/001-003/DC	RVG 119689	EGIS PHARMACEUTICALS PLC	NL
Rosulip Plus, 5 mg + 10 mg, kapsułki, twarde	NL/H/3007/001-003/DC	24424	EGIS PHARMACEUTICALS PLC	PL
Росулип Пра 1 О mg/1 О mg твърди капсули	NL/H/3007/005	20180227	EGIS PHARMACEUTICALS PLC	BG
Росули п Про 20 mg/10 mg твърд и капсули	NL/H/3007/006	20180228	EGIS PHARMACEUTICALS PLC	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
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620331	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620349	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10mg/10mg cápsulas	NL/H/3007/001	5620356	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10mg/10mg cápsulas	NL/H/3007/001	5620364	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620372	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620406	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620414	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 10 mg/10 mg cápsulas	NL/H/3007/001	5620422	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620430	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620448	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620455	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620463	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620471	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620505	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620513	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 20 mg/10 mg cápsulas	NL/H/3007/002	5620521	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620539	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620647	EGIS PHARMACEUTICALS PLC	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
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620554	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620562	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620570	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620604	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620612	EGIS PHARMACEUTICALS PLC	PT
Lipocomb 40 mg/10 mg cápsulas	NL/H/3007/003	5620620	EGIS PHARMACEUTICALS PLC	PT
Cholecomb 5 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496254, 043496266, 043496278, 043496280	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 5 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496254, 043496266, 043496278, 043496280	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 5 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496254, 043496266, 043496278, 043496280	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 5 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496254, 043496266, 043496278, 043496280	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg	NL/H/3007/001-003/DC	043496227,043496013,04	EGIS PHARMACEUTICALS PLC	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
capsule rigide		3496025,043496037,043496049,043496052,043496064,043496076		
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 10 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496227,043496013,043496025,043496037,043496049,043496052,043496064,043496076	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496239,043496088,043496090,043496102,043496114,043496126,043496138,043496140	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 20 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496254, 043496266, 043496278, 043496280	EGIS PHARMACEUTICALS PLC	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**
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Cholecomb 40 mg/10 mg capsule rigide	NL/H/3007/001-003/DC	043496241,043496153,043496165,043496177,043496189,043496191,043496203,043496215	EGIS PHARMACEUTICALS PLC	IT
Ayadont calcium 10 mg/10 mg harde capsules	NL/H/3007/005	RVG 120475	EGIS PHARMACEUTICALS PLC	NL
Ayadont calcium 20 mg/10 mg harde capsules	NL/H/3007/006	RVG 120479	EGIS PHARMACEUTICALS PLC	NL
Rosulip Pro 10mg/10mg hard capsule	NL/H/3007/005/DC	24973	EGIS PHARMACEUTICALS PLC	PL
Rosulip Pro 20mg/10mg	NL/H/3007/006/DC	24974	EGIS PHARMACEUTICALS PLC	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
hard capsule				
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/05	SANOFI-AVENTIS ZRT	HU
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/04	SANOFI-AVENTIS ZRT	HU
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/01	SANOFI-AVENTIS ZRT	HU
SUVREZA 20 mg/10mg, comprimé pelliculé	CZ/H/0816/002	34009 301 764 0 5	SANOFI-AVENTIS FRANCE	FR
SUVREZA 10 mg/10 mg, comprimé pelliculé	CZ/H/0816/001	34009 301 763 7 5	SANOFI-AVENTIS FRANCE	FR
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/03	SANOFI-AVENTIS ZRT	HU
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/06	SANOFI-AVENTIS ZRT	HU
SUVREZA 10 mg/10 mg, comprimé pelliculé	CZ/H/0816/001	34009 301 763 8 2	SANOFI-AVENTIS FRANCE	FR
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
SUVREZA 20 mg/10mg, comprimé pelliculé	CZ/H/0816/002	34009 301 763 9 9	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/02	SANOFI-AVENTIS ZRT	HU
Zenon 10 mg/10 mg comprimidos recubiertos con película	CZ/H/0816/001	84121	SANOFI-AVENTIS, S.A.	ES
Zenon 20 mg/10 mg comprimidos recubiertos con película	CZ/H/0816/002	84122	SANOFI-AVENTIS, S.A.	ES
Zenon 40 mg/10 mg comprimidos recubiertos con película	CZ/H/0816/003	84123	SANOFI-AVENTIS, S.A.	ES
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 10 mg/10 mg Filmtabletten	CZ/H/0816/001	2201737.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 20 mg/10 mg Filmtabletten	CZ/H/0816/002	2201738.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Zenon 40 mg/10 mg Filmtabletten	CZ/H/0816/003	2201739.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 10 mg/10 mg potahované tablety	CZ/H/0816/001	31/013/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 20 mg/10 mg potahované tablety	CZ/H/0816/002	31/014/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ

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
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ
Rosuvastatin/Ezetimib Sanofi 40 mg/10 mg potahované tablety	CZ/H/0816/003	31/015/18-C	SANOFI-AVENTIS SRO	CZ
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/09	SANOFI-AVENTIS ZRT	HU
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/12	SANOFI-AVENTIS ZRT	HU
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/18	SANOFI-AVENTIS ZRT	HU
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/07	SANOFI-AVENTIS ZRT	HU
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/08	SANOFI-AVENTIS ZRT	HU
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/14	SANOFI-AVENTIS ZRT	HU
Suvezen 10 mg/10 mg filmtabletta	CZ/H/0816/001	OGYI-T-23520/10	SANOFI-AVENTIS ZRT	HU
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/13	SANOFI-AVENTIS ZRT	HU
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/16	SANOFI-AVENTIS ZRT	HU
Suvezen 20 mg/10 mg filmtabletta	CZ/H/0816/002	OGYI-T-23520/11	SANOFI-AVENTIS ZRT	HU
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/17	SANOFI-AVENTIS ZRT	HU
Suvezen 40 mg/10 mg filmtabletta	CZ/H/0816/003	OGYI-T-23520/15	SANOFI-AVENTIS ZRT	HU
Arosuva plus Ezetimib 5 mg/10 mg Filmtabletten	NL/H/4477/001	139214	GEBRO PHARMA GMBH	AT
Arosuva plus Ezetimib 10 mg/10 mg Filmtabletten	NL/H/4477/002	139215	GEBRO PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Arosuva plus Ezetimib 20 mg/10 mg Filmtabletten	NL/H/4477/003	139216	GEBRO PHARMA GMBH	AT
Arosuva plus Ezetimib 40 mg/10 mg Filmtabletten	NL/H/4477/004	139217	GEBRO PHARMA GMBH	AT
Antilia 5 mg/10 mg Tabletten	PL/H/0497/001	2200745.00.00	HEXAL AG	DE
Antilia 10 mg/10 mg Tabletten	PL/H/0497/002	2200746.00.00	HEXAL AG	DE
Antilia 20 mg/10mg Tabletten	PL/H/0497/003	2200747.00.00	HEXAL AG	DE
Suvardio Plus, 5 mg + 10 mg, tabletki	PL/H/0497/001	24594	SANDOZ GMBH	PL
Suvardio Plus, 10 mg + 10 mg, tabletki	PL/H/0497/002	24595	SANDOZ GMBH	PL
Suvardio Plus, 20 mg + 10 mg, tabletki	PL/H/0497/003	24596	SANDOZ GMBH	PL
Rosumop Combi 10 mg/10 mg tablety	PL/H/0497/002	31/333/17-C	SANDOZ S.R.O.	CZ
Rosumop Combi 20 mg/10 mg tablety	PL/H/0497/003	31/334/17-C	SANDOZ S.R.O.	CZ
Rosumop Combi 5 mg/10 mg tablety	PL/H/0497/001	31/332/17-C	SANDOZ S.R.O.	CZ
Co-Xeter 10 mg/10 mg tabletta	not available	OGYI-T-23413/01	GEDEON RICHTER PLC.	HU
Co-Xeter 20 mg/10 mg tabletta	not available	OGYI-T-23413/02	GEDEON RICHTER PLC.	HU
Eprizet 10 mg/10 mg tablete	PL/H/0500/002	HR-H-862844884	PLIVA HRVATSKA D.O.O.	HR
Eprizet 20 mg/10 mg tablete	PL/H/0500/003	HR-H-797006261	PLIVA HRVATSKA D.O.O.	HR
Тинтарос Плюс 5 mg/10 mg таблетки	PL/H/0500/001	20180113	TEVA B.V	BG
Тинтарос Плюс 10 mg/10 mg таблетки	PL/H/0500/002	20180114	TEVA B.V	BG
Тинтарос Плюс 20 mg/10 mg таблетки	PL/H/0500/003	20180115	TEVA B.V	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
Rosuvastatín/Ezetimib Teva 5 mg/10 mg tablety	PL/H/0500/001	31/0081/18-S	TEVA B.V	SK
Rosuvastatín/Ezetimib Teva 10 mg/10 mg tablety	PL/H/0500/002	31/0082/18-S	TEVA B.V	SK
Rosuvastatín/Ezetimib Teva 10 mg/10 mg tablety	PL/H/0500/003	31/0083/18-S	TEVA B.V	SK
Rosuvastatin/Ezetimibe Teva 20 mg/10 mg tablety	PL/H/0500/003	31/331/17-C	TEVA B.V	CZ
Rosuvastatin/Ezetimibe Teva 10 mg/10 mg tablety	PL/H/0500/002	31/330/17-C	TEVA B.V	CZ
Rosuvastatin/Ezetimibe Teva, 5 mg + 10 mg, tabletki	PL/H/0500/001	24851	TEVA B.V	PL
Rosuvastatin/Ezetimibe Teva, 20 mg + 10 mg, tabletki	PL/H/0500/003	24853	TEVA B.V	PL
Rosuvastatin/Ezetimibe Teva, 10 mg + 10 mg, tabletki	PL/H/0500/002	24852	TEVA B.V	PL
Rosuvastatin/Ezetimib ratiopharm 5 mg/10 mg Tabletten	PL/H/0500/001	139124	TEVA B.V	AT
Rosuvastatin/Ezetimib ratiopharm 10 mg/10 mg Tabletten	PL/H/0500/002	139125	TEVA B.V	AT
Rosuvastatin/Ezetimib ratiopharm 20 mg/10 mg Tabletten	PL/H/0500/003	139126	TEVA B.V	AT
Ezetimibe/Rosuvastatine Teva 10 mg/5 mg comprimés	PL/H/0500/001	BE545057	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10/5mg tabletten	PL/H/0500/001	BE545057	TEVA B.V	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ezetimibe/Rosuvastatine Teva 10 mg/10 mg comprimés	PL/H/0500/002	BE545066	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10/10mg Tabletten	PL/H/0500/002	BE545066	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10/10mg tabletten	PL/H/0500/002	BE545066	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10 mg/20 mg comprimés	PL/H/0500/003	BE545075	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10/20mg tabletten	PL/H/0500/003	BE545075	TEVA B.V	BE
Ezetimibe/Rosuvastatine Teva 10/20mg Tabletten	PL/H/0500/003	BE545075	TEVA B.V	BE
Rosuvastatin/Ezetimibe ratiopharm 5 mg/10 mg tabletti	PL/H/0500/001	36977	TEVA B.V	FI
Rosuvastatin/Ezetimibe ratiopharm 5 mg/10 mg tablett	PL/H/0500/001	36977	TEVA B.V	FI
Rosuvastatin/Ezetimibe ratiopharm 10 mg/10 mg tabletti	PL/H/0500/002	36978	TEVA B.V	FI
Rosuvastatin/Ezetimibe ratiopharm 10 mg/10 mg tablet	PL/H/0500/002	36978	TEVA B.V	FI
Rosuvastatin/Ezetimibe ratiopharm 20 mg/10 mg tabletti	PL/H/0500/003	36979	TEVA B.V	FI
Rosuvastatin/Ezetimibe ratiopharm 20 mg/10 mg tablett	PL/H/0500/003	36979	TEVA B.V	FI
Ezetimibe/Rosuvastatine Teva 10/5mg Tabletten	PL/H/0500/001	BE545057	TEVA B.V	BE
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876014	TEVA B.V	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
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876053	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876091	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876026	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876040	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876038	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876065	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876077	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876103	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876115	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876089	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876127	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876139	TEVA B.V	IT
Rosuvastatina e	PL/H/0500/002	047876166	TEVA B.V	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
Ezetimibe Teva 10 mg/10 mg compresse				
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876192	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876141	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 5 mg/10 mg compresse	PL/H/0500/001	047876154	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876178	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876204	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 20 mg/10 mg compresse	PL/H/0500/003	047876216	TEVA B.V	IT
Rosuvastatina e Ezetimibe Teva 10 mg/10 mg compresse	PL/H/0500/002	047876180	TEVA B.V	IT
ROSETEM 5 mg/10 mg compresse	not available	045310012	ERREKAPPA EUROTERAPICI SPA	IT
ROSETEM 10 mg/10 mg compresse	not available	045310024	ERREKAPPA EUROTERAPICI SPA	IT
ROSETEM 20 mg/10 mg compresse	not available	045310036	ERREKAPPA EUROTERAPICI SPA	IT
ROZOR 10 mg/10 mg comprimidos revestidos por película	NL/H/3647/001	5718739	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROZOR 10 mg/10 mg comprimidos revestidos por película	NL/H/3647/001	5718747	BGP PRODUCTS UNIPESSOAL, LDA.	PT
ROZOR 10 mg/10 mg comprimidos revestidos	NL/H/3647/001	5718754	BGP PRODUCTS UNIPESSOAL, LDA.	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
por película				
ROZOR 10 mg/10 mg comprimidos revestidos por película	NL/H/3647/001	5718762	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Rozor 10 mg/10 mg filmdragerade tabletter	NL/H/3647/001	54350	BGP PRODUCTS AB	SE
Twicor 10 mg/10 mg potahované tablety	NL/H/3647/001	31/511/16-C	MYLAN IRE HEALTHCARE LIMITED	CZ
Rozor, 10 mg + 10 mg, tabletki powlekane	NL/H/3647/001	24220	MYLAN IRE HEALTHCARE LIMITED	PL
Twicor 10 mg/10 mg filmomhulde tabletten	NL/H/3647/001	RVG 118789	MYLAN HEALTHCARE B.V.	NL
Twicor 10 mg/10 mg comprimidos recubiertos con película	NL/H/3647/001	82426	MYLAN IRE HEALTHCARE LIMITED	ES
Ezehron Duo, 5 mg + 10 mg, tabletka	PL/H/0468/001	24646	ADAMED PHARMA S.A.	PL
Ezehron Duo, 10 mg + 10 mg, tabletka	PL/H/0468/002	24647	ADAMED PHARMA S.A.	PL
Ezehron Duo, 20 mg + 10 mg, tabletka	PL/H/0468/003	24648	ADAMED PHARMA S.A.	PL
Ezehron Duo 5 mg/10 mg Tabletten	PL/H/0468/001	99089.00.00	ADAMED PHARMA S.A.	DE
Ezehron Duo 10 mg/10 mg Tabletten	PL/H/0468/002	99090.00.00	ADAMED PHARMA S.A.	DE
Ezehron Duo 20 mg/10 mg Tabletten	PL/H/0468/003	99091.00.00	ADAMED PHARMA S.A.	DE
RUZEB 5 mg/10 mg tablety	PL/H/0468/001	31/0130/18-S	ADAMED PHARMA S.A.	SK
RUZEB 10 mg/10 mg tablety	PL/H/0468/002	31/0131/18-S	ADAMED PHARMA S.A.	SK
RUZEB 20 mg/10 mg tablety	PL/H/0468/003	31/0132/18-S	ADAMED PHARMA S.A.	SK
Ezehron Duo 5 mg/10 mg, tablet	PL/H/0468/001	MA948/00401	ADAMED	MT
Ezehron Duo 10 mg/10	PL/H/0468/002	MA948/00402	ADAMED	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
mg, tablet				
Ezehron Duo 20 mg/10 mg, tablet	PL/H/0468/003	MA948/00403	ADAMED	MT
Ruzeb 5 mg/10 mg tablety	PL/H/0468/001	31/935/16-C	ADAMED PHARMA S.A.	CZ
Ruzeb 10 mg/10 mg tablety	PL/H/0468/002	31/936/16-C	ADAMED PHARMA S.A.	CZ
Ruzeb 20 mg/10 mg tablety	PL/H/0468/003	31/937/16-C	ADAMED PHARMA S.A.	CZ
Colroset 5 mg/10 mg comprimidos	PL/H/0468/001	5741814	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Colroset 5 mg/10 mg comprimidos	PL/H/0468/001	5741822	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Colroset 10 mg/10 mg comprimidos	PL/H/0468/002	5741830	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Colroset 20 mg/10 mg comprimidos	PL/H/0468/003	5741848	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Езехрон Дуо 5 mg/10 mg таблетки	PL/H/0468/001	20180121	ADAMED PHARMA S.A.	BG
Езехрон Дуо 10 mg/10 mg таблетки	PL/H/0468/002	20180122	ADAMED PHARMA S.A.	BG
Езехрон Дуо 20 mg/10 mg таблетки	PL/H/0468/003	20180123	ADAMED PHARMA S.A.	BG
Rosix Combi 5 mg/10 mg tablete	PL/H/0468/001	HR-H-191795318	BELUPO D.D.	HR
Rosix Combi 10 mg/10 mg tablete	PL/H/0468/002	HR-H-660495522	BELUPO D.D.	HR
Rosix Combi 20 mg/10 mg tablete	PL/H/0468/003	HR-H-791429746	BELUPO D.D.	HR
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350016	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350028	PIAM FARMACEUTICI SPA	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
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350030	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350042	PIAM FARMACEUTICI S.P.A.	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350055	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350067	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350079	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350081	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350093	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350105	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 5 mg/10 mg compresse	PL/H/0468/001	045350319	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350129	PIAM FARMACEUTICI S.P.A.	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350131	PIAM FARMACEUTICI S.P.A.	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350117	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350143	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350156	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350168	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350170	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350321	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350182	PIAM FARMACEUTICI SPA	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
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350194	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 10 mg/10 mg compresse	PL/H/0468/002	045350206	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350218	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350220	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350232	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350244	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350257	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350269	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350271	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350333	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350283	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350295	PIAM FARMACEUTICI SPA	IT
ROSUMIBE 20 mg/10 mg compresse	PL/H/0468/003	045350307	PIAM FARMACEUTICI SPA	IT
Compichol 20 mg/10 mg, δισκία	PL/H/0468/003	125446/24-12-2018	WIN MEDICA SA	GR
Compichol 5 mg/10 mg, δισκία	PL/H/0468/001	83422/16/24-12-2018	WIN MEDICA SA	GR
Compichol 10 mg/10 mg, δισκία	PL/H/0468/002	125445/24-12-2018	WIN MEDICA SA	GR
Rozetimad 5 mg/10 mg compresse	not available	045424013	ADAMED S.R.L.	IT
Rozetimad 10 mg/10 mg compresse	not available	045424025	ADAMED S.R.L.	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
Rozetimad 20 mg/10 mg compresse	not available	045424037	ADAMED S.R.L.	IT
Rozetimad 5 mg/10 mg compresse	not available	045424049	ADAMED S.R.L.	IT
Rozetimad 20 mg/10 mg compresse	not available	045424064	ADAMED S.R.L.	IT
Rozetimad 10 mg/10 mg compresse	not available	045424052	ADAMED S.R.L.	IT
Zenon 10 mg/40 mg filmom obalené tablety	CZ/H/0455/003	31/0010/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/40 mg filmom obalené tablety	CZ/H/0455/003	31/0010/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/40 mg filmom obalené tablety	CZ/H/0455/003	31/0010/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/40 mg filmom obalené tablety	CZ/H/0455/003	31/0010/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/40 mg potahované tablety	CZ/H/0455/003	31/323/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/40 mg potahované tablety	CZ/H/0455/003	31/323/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/40 mg potahované tablety	CZ/H/0455/003	31/323/14-C	SANOFI-AVENTIS SRO	CZ
Зенон 10 mg/40 mg филмирани таблетки	CZ/H/0455/003	20140379	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/40 mg филмирани таблетки	CZ/H/0455/003	20140379	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/40 mg филмирани таблетки	CZ/H/0455/003	20140379	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/40 mg филмирани таблетки	CZ/H/0455/003	20140379	SANOFI BULGARIA EOOD	BG
Zenon 10 mg/40 mg potahované tablety	CZ/H/0455/003	31/323/14-C	SANOFI-AVENTIS SRO	CZ
Suvezen, (10 mg + 40 mg), tabletki powlekane	CZ/H/0455/003	22279	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 40 mg), tabletki powlekane	CZ/H/0455/003	22279	SANOFI-AVENTIS SP Z.O.O.	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
Suvezen, (10 mg + 40 mg), tabletki powlekane	CZ/H/0455/003	22279	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 40 mg), tabletki powlekane	CZ/H/0455/003	22279	SANOFI-AVENTIS SP Z.O.O.	PL
Zenon 10 mg/10 mg filmom obalené tablety	CZ/H/0455/001	31/0008/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/10 mg filmom obalené tablety	CZ/H/0455/001	31/0008/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/10 mg filmom obalené tablety	CZ/H/0455/001	31/0008/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/10 mg filmom obalené tablety	CZ/H/0455/001	31/0008/15-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Zenon 10 mg/10 mg potahované tablety	CZ/H/0455/001	31/321/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/10 mg potahované tablety	CZ/H/0455/001	31/321/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/10 mg potahované tablety	CZ/H/0455/001	31/321/14-C	SANOFI-AVENTIS SRO	CZ
Zenon 10 mg/10 mg potahované tablety	CZ/H/0455/001	31/321/14-C	SANOFI-AVENTIS SRO	CZ
Зенон 10 mg/10 mg филмирани таблетки	CZ/H/0455/001	20140377	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/10 mg филмирани таблетки	CZ/H/0455/001	20140377	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/10 mg филмирани таблетки	CZ/H/0455/001	20140377	SANOFI BULGARIA EOOD	BG
Зенон 10 mg/10 mg филмирани таблетки	CZ/H/0455/001	20140377	SANOFI BULGARIA EOOD	BG
Suvezen, (10 mg + 10 mg), tabletki powlekane	CZ/H/0455/001	22277	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 10 mg), tabletki powlekane	CZ/H/0455/001	22277	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 10 mg), tabletki powlekane	CZ/H/0455/001	22277	SANOFI-AVENTIS SP Z.O.O.	PL
Suvezen, (10 mg + 10 mg), tabletki powlekane	CZ/H/0455/001	22277	SANOFI-AVENTIS SP Z.O.O.	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
MAORIS 20 mg/10 mg compresse	not available	045304033	NEOPHARMED GENTILI SPA	IT
MAORIS 5 mg/10 mg compresse	not available	045304019	NEOPHARMED GENTILI SPA	IT
MAORIS 10 mg/10 mg compresse	not available	045304021	NEOPHARMED GENTILI SPA	IT