27 March 2019
EMA/267129/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: atorvastatin / ezetimibe

Procedure no.: PSUSA/00010385/201807

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ORVATEZ 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3897/001	91315/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
ORVATEZ 10 mg/20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3897/002	91316/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
ORVATEZ 10 mg/40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3897/003	91317/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
ORVATEZ 10 mg/80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3897/004	91318/13/4-11-2014	MERCK SHARP & DOHME LTD.	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
Orvatez® 10 mg/10 mg Filmtabletten	DE/H/3897/001	90231.00.00	MERCK SHARP & DOHME LTD.	DE
Orvatez® 10 mg/20 mg Filmtabletten	DE/H/3897/002	90231.01.00	MERCK SHARP & DOHME LTD.	DE
Orvatez® 10 mg/40 mg Filmtabletten	DE/H/3897/003	90231.02.00	MERCK SHARP & DOHME LTD.	DE
Orvatez® 10 mg/80 mg Filmtabletten	DE/H/3897/004	90231.03.00	MERCK SHARP & DOHME LTD.	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
TIOBLIS® 10 mg/10 mg, comprimés pelliculés	DE/H/3897/001	BE465831	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/10 mg, filmomhulde tabletten	DE/H/3897/001	BE465831	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/10 mg Filmtabletten	DE/H/3897/001	BE465831	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/20 mg, comprimés pelliculés	DE/H/3897/002	BE465840	MERCK SHARP & DOHME BV	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
TIOBLIS® 10 mg/20 mg, filmomhulde tabletten	DE/H/3897/002	BE465840	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/20 mg Filmtabletten	DE/H/3897/002	BE465840	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/40 mg, comprimés pelliculés	DE/H/3897/003	BE465857	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/40 mg, filmomhulde tabletten	DE/H/3897/003	BE465857	MERCK SHARP & DOHME BV	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
TIOBLIS® 10 mg/40 mg Filmtabletten	DE/H/3897/003	BE465857	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/80 mg, comprimés pelliculés	DE/H/3897/004	BE465866	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/80 mg, filmomhulde tabletten	DE/H/3897/004	BE465866	MERCK SHARP & DOHME BV	BE
TIOBLIS® 10 mg/80 mg Filmtabletten	DE/H/3897/004	BE465866	MERCK SHARP & DOHME BV	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
ORVATEZ 10 mg/10 mg filmom obalené tablety	DE/H/3897/001	31/0349/14-S	MERCK SHARP & DOHME BV	SK
ORVATEZ 10 mg/20 mg filmom obalené tablety	DE/H/3897/002	31/0350/14-S	MERCK SHARP & DOHME BV	SK
ORVATEZ 10 mg/40 mg filmom obalené tablety	DE/H/3897/003	31/0351/14-S	MERCK SHARP & DOHME BV	SK
ORVATEZ 10 mg/80 mg filmom obalené tablety	DE/H/3897/004	31/0352/14-S	MERCK SHARP & DOHME BV	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
ORVATEZ 10 mg/10 mg comprimidos recubiertos con película	DE/H/3897/001	79.166	MERCK SHARP & DOHME BV	ES
ORVATEZ 10 mg/20 mg comprimidos recubiertos con película	DE/H/3897/002	79.167	MERCK SHARP & DOHME BV	ES
ORVATEZ 10 mg/40 mg comprimidos recubiertos con película	DE/H/3897/003	79.168	MERCK SHARP & DOHME BV	ES
ORVATEZ 10 mg/80 mg comprimidos recubiertos con película	DE/H/3897/004	79.169	MERCK SHARP & DOHME BV	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
TIOBLIS® 10 mg/10 mg, comprimés pelliculés	DE/H/3897/001	2015040096	MERCK SHARP & DOHME BV	LU
TIOBLIS® 10 mg/20 mg, comprimés pelliculés	DE/H/3897/002	2015040097	MERCK SHARP & DOHME BV	LU
TIOBLIS® 10 mg/40 mg, comprimés pelliculés	DE/H/3897/003	2015040098	MERCK SHARP & DOHME BV	LU
TIOBLIS® 10 mg/80 mg, comprimés pelliculés	DE/H/3897/004	2015040099	MERCK SHARP & DOHME BV	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
Orvatez 10 mg/40 mg comprimidos revestidos por película	DE/H/3897/003	5628748	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT
Orvatez 10 mg/40 mg comprimidos revestidos por película	DE/H/3897/003	5628730	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT
Orvatez 10 mg/20 mg comprimidos revestidos por película	DE/H/3897/002	5628672	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT
Orvatez 10 mg/20 mg comprimidos revestidos por película	DE/H/3897/002	5628664	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Orvatez 10 mg/10 mg comprimidos revestidos por película	DE/H/3897/001	5628466	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT
Orvatez 10 mg/10 mg comprimidos revestidos por película	DE/H/3897/001	5628458	INTERBIAL PRODUTOS FARMACEUTICOS, S.A.	PT
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249010	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249022	MSD ITALIA 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
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249034	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249212	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249135	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/10 mg compresse rivestite con film	DE/H/3897/001	043249174	MSD ITALIA 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
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249046	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249059	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249061	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249147	MSD ITALIA 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
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249186	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/20 mg compresse rivestite con film	DE/H/3897/002	043249224	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249073	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249085	MSD ITALIA 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
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249097	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249150	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249198	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/40 mg compresse rivestite con film	DE/H/3897/003	043249236	MSD ITALIA 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
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249109	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249111	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249123	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249162	MSD ITALIA 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
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249200	MSD ITALIA S.R.L.	IT
ORVATEZ 10 mg/80 mg compresse rivestite con film	DE/H/3897/004	043249248	MSD ITALIA S.R.L.	IT
LIPTRUZET 10 mg/10 mg, comprimé pelliculé	DE/H/3895/001	34009 280 048 9 5	MSD FRANCE	FR
LIPTRUZET 10 mg/10 mg, comprimé pelliculé	DE/H/3895/001	34009 280 049 5 6	MSD 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
LIPTRUZET 10 mg/10 mg, comprimé pelliculé	DE/H/3895/001	34009 587 184 0 3	MSD FRANCE	FR
LIPTRUZET 10 mg/20 mg, comprimé pelliculé	DE/H/3895/002	34009 280 053 2 8	MSD FRANCE	FR
LIPTRUZET 10 mg/20 mg, comprimé pelliculé	DE/H/3895/002	34009 280 054 9 6	MSD FRANCE	FR
LIPTRUZET 10 mg/20 mg, comprimé pelliculé	DE/H/3895/002	34009 587 188 6 1	MSD 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
LIPTRUZET 10 mg/40 mg, comprimé pelliculé	DE/H/3895/003	34009 280 055 5 7	MSD FRANCE	FR
LIPTRUZET 10 mg/40 mg, comprimé pelliculé	DE/H/3895/003	34009 280 056 1 8	MSD FRANCE	FR
LIPTRUZET 10 mg/40 mg, comprimé pelliculé	DE/H/3895/003	34009 587 189 2 2	MSD FRANCE	FR
LIPTRUZET 10 mg/80 mg, comprimé pelliculé	DE/H/3895/004	34009 280 057 8 6	MSD 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
LIPTRUZET 10 mg/80 mg, comprimé pelliculé	DE/H/3895/004	34009 280 058 4 7	MSD FRANCE	FR
LIPTRUZET 10 mg/80 mg, comprimé pelliculé	DE/H/3895/004	34009 587 190 0 4	MSD FRANCE	FR
Liptruzet 10 mg/10 mg filmtabletta	DE/H/3895/001	OGYI-T-22732/01	MSD PHARMA HUNGARY KFT.	HU
Liptruzet 10 mg/20 mg filmtabletta	DE/H/3895/002	OGYI-T-22732/02	MSD PHARMA HUNGARY KFT.	HU

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Liptruzet 10 mg/40 mg filmtabletta	DE/H/3895/003	OGYI-T-22732/03	MSD PHARMA HUNGARY KFT.	HU
Liptruzet 10 mg/80 mg filmtabletta	DE/H/3895/004	OGYI-T-22732/04	MSD PHARMA HUNGARY KFT.	HU
ATOZET 10 mg/10 mg filmom obalené tablety	DE/H/3895/001	31/0329/14-S	MERCK SHARP & DOHME BV	SK
ATOZET 10 mg/20 mg filmom obalené tablety	DE/H/3895/002	31/0330/14-S	MERCK SHARP & DOHME BV	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
ATOZET 10 mg/40 mg filmom obalené tablety	DE/H/3895/003	31/0331/14-S	MERCK SHARP & DOHME BV	SK
ATOZET 10 mg/80 mg filmom obalené tablety	DE/H/3895/004	31/0332/14-S	MERCK SHARP & DOHME BV	SK
ATOZET 10 mg/10 mg filmuhúðaðar töflur	DE/H/3895/001	IS/1/14/086/01	MERCK SHARP & DOHME BV	IS
ATOZET 10 mg/20 mg filmuhúðaðar töflur	DE/H/3895/002	IS/1/14/086/02	MERCK SHARP & DOHME BV	IS

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ATOZET 10 mg/40 mg filmuhúðaðar töflur	DE/H/3895/003	IS/1/14/086/03	MERCK SHARP & DOHME BV	IS
ATOZET 10 mg/80 mg filmuhúðaðar töflur	DE/H/3895/004	IS/1/14/086/04	MERCK SHARP & DOHME BV	IS
Atozet	DE/H/3895/001	53288	MERCK SHARP & DOHME BV	DK
Atozet	DE/H/3895/002	53289	MERCK SHARP & DOHME BV	DK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atozet	DE/H/3895/003	53290	MERCK SHARP & DOHME BV	DK
Atozet	DE/H/3895/004	53291	MERCK SHARP & DOHME BV	DK
Atozet 10 mg/10 mg filmdrasjerte tabletter	DE/H/3895/001	13-9807	MERCK SHARP & DOHME BV	NO
Atozet 10 mg/20 mg filmdrasjerte tabletter	DE/H/3895/002	13-9808	MERCK SHARP & DOHME BV	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
Atozet 10 mg/40 mg filmdrasjerte tabletter	DE/H/3895/003	13-9809	MERCK SHARP & DOHME BV	NO
Atozet 10 mg/80 mg filmdrasjerte tabletter	DE/H/3895/004	13-9810	MERCK SHARP & DOHME BV	NO
LIPTRUZET 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/001	75277/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
LIPTRUZET 10 mg/20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/002	75278/13/4-11-2014	MERCK SHARP & DOHME LTD.	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
LIPTRUZET 10 mg/40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/003	75279/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
LIPTRUZET 10 mg/80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/004	75280/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
Zoletorv® 10 mg/10 mg potahované tablety	DE/H/3895/001	31/403/14-C	MERCK SHARP & DOHME BV	CZ
Zoletorv® 10 mg/20 mg potahované tablety	DE/H/3895/002	31/404/14-C	MERCK SHARP & DOHME BV	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
Zoletorv® 10 mg/40 mg potahované tablety	DE/H/3895/003	31/405/14-C	MERCK SHARP & DOHME BV	CZ
Atozet 10 mg/10 mg Filmtabletten	DE/H/3895/001	135866	MERCK SHARP & DOHME BV	AT
Atozet 10 mg/20 mg Filmtabletten	DE/H/3895/002	135867	MERCK SHARP & DOHME BV	AT
Atozet 10 mg/40 mg Filmtabletten	DE/H/3895/003	135868	MERCK SHARP & DOHME BV	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
Atozet 10 mg/80 mg Filmtabletten	DE/H/3895/004	135869	MERCK SHARP & DOHME BV	AT
Atozet 10 mg/10 mg filmdragerade tabletter	DE/H/3895/001	50194	MERCK SHARP & DOHME BV	SE
Atozet 10 mg/20 mg filmdragerade tabletter	DE/H/3895/002	50195	MERCK SHARP & DOHME BV	SE
Atozet 10 mg/40 mg filmdragerade tabletter	DE/H/3895/003	50196	MERCK SHARP & DOHME BV	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
Atozet 10 mg/80 mg filmdragerade tabletter	DE/H/3895/004	50197	MERCK SHARP & DOHME BV	SE
Atozet® 10 mg/10 mg Filmtabletten	DE/H/3895/001	90223.00.00	MERCK SHARP & DOHME LTD.	DE
Atozet® 10 mg/20 mg Filmtabletten	DE/H/3895/002	90223.01.00	MERCK SHARP & DOHME LTD.	DE
Atozet® 10 mg/40 mg Filmtabletten	DE/H/3895/003	90223.02.00	MERCK SHARP & DOHME LTD.	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
Atozet® 10 mg/80 mg Filmtabletten	DE/H/3895/004	90223.03.00	MERCK SHARP & DOHME LTD.	DE
ATOZET® 10 mg/10 mg, comprimés pelliculés	DE/H/3895/001	BE465795	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/10 mg, filmomhulde tabletten	DE/H/3895/001	BE465795	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/10 mg Filmtabletten	DE/H/3895/001	BE465795	MERCK SHARP & DOHME BV	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
ATOZET® 10 mg/20 mg, comprimés pelliculés	DE/H/3895/002	BE465804	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/20 mg, filmomhulde tabletten	DE/H/3895/002	BE465804	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/20 mg Filmtabletten	DE/H/3895/002	BE465804	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/40 mg, comprimés pelliculés	DE/H/3895/003	BE465813	MERCK SHARP & DOHME BV	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
ATOZET® 10 mg/40 mg, filmomhulde tabletten	DE/H/3895/003	BE465813	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/40 mg Filmtabletten	DE/H/3895/003	BE465813	MERCK SHARP & DOHME BV	BE
Atozet® 10 mg/80 mg, comprimés pelliculés	DE/H/3895/004	BE465822	MERCK SHARP & DOHME BV	BE
ATOZET® 10 mg/80 mg, filmomhulde tabletten	DE/H/3895/004	BE465822	MERCK SHARP & DOHME BV	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
ATOZET® 10 mg/80 mg Filmtabletten	DE/H/3895/004	BE465822	MERCK SHARP & DOHME BV	BE
ATOZET 10 mg/10 mg film-coated tablets	DE/H/3895/001	MA224/00401	MERCK SHARP & DOHME BV	MT
ATOZET 10 mg/20 mg film-coated tablets	DE/H/3895/002	MA224/00402	MERCK SHARP & DOHME BV	MT
ATOZET 10 mg/40 mg film-coated tablets	DE/H/3895/003	MA224/00403	MERCK SHARP & DOHME BV	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
ATOZET 10 mg/80 mg film-coated tablets	DE/H/3895/004	MA224/00404	MERCK SHARP & DOHME BV	MT
ATOZET 10 mg/10 mg Comprimidos revestidos por película	DE/H/3895/001	5628904	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/20 mg Comprimidos revestidos por película	DE/H/3895/002	5629514	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/40 mg Comprimidos revestidos por película	DE/H/3895/003	5629654	MERCK SHARP & DOHME LTD.	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
ATOZET 10 mg/80 mg Comprimidos revestidos por película	DE/H/3895/004	5629811	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/10 mg Comprimidos revestidos por película	DE/H/3895/001	5628912	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/20 mg Comprimidos revestidos por película	DE/H/3895/002	5629506	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/40 mg Comprimidos revestidos por película	DE/H/3895/003	5629662	MERCK SHARP & DOHME LTD.	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
ATOZET 10 mg/80 mg Comprimidos revestidos por película	DE/H/3895/004	5629829	MERCK SHARP & DOHME LTD.	PT
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/06	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/01	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/01	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/04	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/04	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/02	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/02	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/05	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/05	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/05	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/05	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/10 mg comprimate filmate	DE/H/3895/001	7169/2014/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/03	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/03	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/20 mg comprimate filmate	DE/H/3895/002	7170/2014/06	MERCK SHARP & DOHME ROMANIA SRL	RO
ATOZET 10 mg/40 mg comprimate filmate	DE/H/3895/003	7171/2014/06	MERCK SHARP & DOHME ROMANIA SRL	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
ATOZET 10 mg/80 mg comprimate filmate	DE/H/3895/004	7172/2014/06	MERCK SHARP & DOHME ROMANIA SRL	RO
АТОЗЕТ 10mg/10mg филмирани таблетки	DE/H/3895/001	20140381	MERCK SHARP & DOHME BULGARIA EOOD	BG
АТОЗЕТ 10mg/20mg филмирани таблетки	DE/H/3895/002	20140382	MERCK SHARP & DOHME BULGARIA EOOD	BG
АТОЗЕТ 10mg/40mg филмирани таблетки	DE/H/3895/003	20140383	MERCK SHARP & DOHME 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
АТОЗЕТ 10mg/80mg филмирани таблетки	DE/H/3895/004	20140384	MERCK SHARP & DOHME BULGARIA EOOD	BG
LIPTRUZET 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/001	22096	MERCK SHARP & DOHME LTD.	CY
LIPTRUZET 10 mg/20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/002	22097	MERCK SHARP & DOHME LTD.	CY
LIPTRUZET 10 mg/40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/003	22098	MERCK SHARP & DOHME LTD.	CY

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LIPTRUZET 10 mg/80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3895/004	22099	MERCK SHARP & DOHME LTD.	CY
ATOZET 10 mg/10 mg comprimidos recubiertos con película	DE/H/3895/001	79.225	MERCK SHARP & DOHME BV	ES
ATOZET 10 mg/20 mg comprimidos recubiertos con película	DE/H/3895/002	79.224	MERCK SHARP & DOHME BV	ES
ATOZET 10 mg/40 mg comprimidos recubiertos con película	DE/H/3895/003	79.226	MERCK SHARP & DOHME BV	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
ATOZET 10 mg/80 mg comprimidos recubiertos con película	DE/H/3895/004	79.227	MERCK SHARP & DOHME BV	ES
LIPTRUZET 10 mg/80 mg, comprimé pelliculé	DE/H/3895/004	34009 550 045 9 2	MSD FRANCE	FR
ATOZET® 10 mg/10 mg, comprimés pelliculés	DE/H/3895/001	2015040041	MERCK SHARP & DOHME BV	LU
ATOZET® 10 mg/20 mg, comprimés pelliculés	DE/H/3895/002	2015040042	MERCK SHARP & DOHME BV	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
ATOZET® 10 mg/40 mg, comprimés pelliculés	DE/H/3895/003	2015040043	MERCK SHARP & DOHME BV	LU
Atozet® 10 mg/80 mg, comprimés pelliculés	DE/H/3895/004	2015040044	MERCK SHARP & DOHME BV	LU
ATOZET 10 mg/10 mg, filmomhulde tabletten	DE/H/3895/001	RVG 114373	MERCK SHARP & DOHME LTD.	NL
ATOZET 10 mg/20 mg, filmomhulde tabletten	DE/H/3895/002	RVG 114374	MERCK SHARP & DOHME LTD.	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
ATOZET 10 mg/40 mg, filmomhulde tabletten	DE/H/3895/003	RVG 114375	MERCK SHARP & DOHME LTD.	NL
ATOZET 10 mg/80 mg, filmomhulde tabletten	DE/H/3895/004	RVG 114376	MERCK SHARP & DOHME LTD.	NL
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/001	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/007	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/013	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/019	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/002	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/003	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/004	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/005	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/10 mg filmsko obložene tablete	DE/H/3895/001	H/15/02002/006	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/008	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/009	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/010	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/011	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/20 mg filmsko obložene tablete	DE/H/3895/002	H/15/02002/012	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/014	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/015	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/016	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/017	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/40 mg filmsko obložene tablete	DE/H/3895/003	H/15/02002/018	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/020	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/021	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/022	MERCK SHARP & DOHME LTD.	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
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/023	MERCK SHARP & DOHME LTD.	SI
Atozet 10 mg/80 mg filmsko obložene tablete	DE/H/3895/004	H/15/02002/024	MERCK SHARP & DOHME LTD.	SI
ATOZET, 10 mg + 10 mg, tabletki powlekane	DE/H/3895/001	22521	MSD POLSKA SP. Z O.O.	PL
ATOZET, 10 mg + 20 mg, tabletki powlekane	DE/H/3895/002	22522	MSD POLSKA 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
ATOZET, 10 mg + 40 mg, tabletki powlekane	DE/H/3895/003	22523	MSD POLSKA SP. Z O.O.	PL
ATOZET, 10 mg + 80 mg, tabletki powlekane	DE/H/3895/004	22524	MSD POLSKA SP. Z O.O.	PL
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543014	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543026	MSD ITALIA 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
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543038	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543040	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543053	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/10 mg compresse rivestite con film	DE/H/3895/001	043543065	MSD ITALIA 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
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543077	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543089	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543091	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543103	MSD ITALIA 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
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543115	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/20 mg compresse rivestite con film	DE/H/3895/002	043543127	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543139	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543141	MSD ITALIA 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
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543154	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543166	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543178	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/40 mg compresse rivestite con film	DE/H/3895/003	043543180	MSD ITALIA 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
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543192	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543204	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543216	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543228	MSD ITALIA 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
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543230	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/80 mg compresse rivestite con film	DE/H/3895/004	043543242	MSD ITALIA S.R.L.	IT
ATOZET 10 mg/10 mg film-coated tablets	DE/H/3895/001	PL 00025/0618	MERCK SHARP & DOHME LTD.	UK
ATOZET 10 mg/20 mg film-coated tablets	DE/H/3895/002	PL 00025/0619	MERCK SHARP & DOHME LTD.	UK

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ATOZET 10 mg/40 mg film-coated tablets	DE/H/3895/003	PL 00025/0620	MERCK SHARP & DOHME LTD.	UK
ATOZET 10 mg/80 mg film-coated tablets	DE/H/3895/004	PL 00025/0621	MERCK SHARP & DOHME LTD.	UK
ATOZET 10 mg/10 mg filmom obložene tablete	DE/H/3895/001	HR-H-532747774	MERCK SHARP & DOHME D.O.O.	HR
ATOZET 10 mg/20 mg filmom obložene tablete	DE/H/3895/002	HR-H-567039012	MERCK SHARP & DOHME 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
ATOZET 10 mg/40 mg filmom obložene tablete	DE/H/3895/003	HR-H-232001393	MERCK SHARP & DOHME D.O.O.	HR
ATOZET 10 mg/80 mg filmom obložene tablete	DE/H/3895/004	HR-H-602129736	MERCK SHARP & DOHME D.O.O.	HR
ATOZET 10 mg/10 mg film-coated tablets	DE/H/3895/001	PA 1286/060/001	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
ATOZET 10 mg/20 mg film-coated tablets	DE/H/3895/002	PA 1286/060/002	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ATOZET 10 mg/40 mg film-coated tablets	DE/H/3895/003	PA 1286/060/003	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
ATOZET 10 mg/80 mg film-coated tablets	DE/H/3895/004	PA 1286/060/004	MERCK SHARP & DOHME IRELAND (HUMAN HEALTH) LTD	IE
Tioblis® 10 mg/10 mg Filmtabletten	DE/H/3898/001	90235.00.00	MERCK SHARP & DOHME LTD.	DE
Tioblis® 10 mg/20 mg Filmtabletten	DE/H/3898/002	90235.01.00	MERCK SHARP & DOHME LTD.	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
Tioblis® 10 mg/40 mg Filmtabletten	DE/H/3898/003	90235.02.00	MERCK SHARP & DOHME LTD.	DE
Tioblis® 10 mg/80 mg Filmtabletten	DE/H/3898/004	90235.03.00	MERCK SHARP & DOHME LTD.	DE
Ezetimib/Atorvastatin MSD 10 mg/10 mg Filmtabletten	DE/H/3898/001	135870	MERCK SHARP & DOHME BV	AT
Ezetimib/Atorvastatin MSD 10 mg/20 mg Filmtabletten	DE/H/3898/002	135871	MERCK SHARP & DOHME BV	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
Ezetimib/Atorvastatin MSD 10 mg/40 mg Filmtabletten	DE/H/3898/003	135872	MERCK SHARP & DOHME BV	AT
Ezetimib/Atorvastatin MSD 10 mg/80 mg Filmtabletten	DE/H/3898/004	135873	MERCK SHARP & DOHME BV	AT
ZARDIUX 10 mg/10 mg comprimidos recubiertos con película	DE/H/3898/001	79.170	MERCK SHARP & DOHME BV	ES
ZARDIUX 10 mg/20 mg comprimidos recubiertos con película	DE/H/3898/002	79.171	MERCK SHARP & DOHME BV	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
ZARDIUX 10 mg/40 mg comprimidos recubiertos con película	DE/H/3898/003	79.172	MERCK SHARP & DOHME BV	ES
ZARDIUX 10 mg/80 mg comprimidos recubiertos con película	DE/H/3898/004	79.173	MERCK SHARP & DOHME BV	ES
KEXROLT 10 mg/10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3896/001	91322/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
KEXROLT 10 mg/20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3896/002	91323/13/4-11-2014	MERCK SHARP & DOHME LTD.	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
KEXROLT 10 mg/40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3896/003	91324/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
KEXROLT 10 mg/80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3896/004	91325/13/4-11-2014	MERCK SHARP & DOHME LTD.	GR
Kexrolt® 10 mg/10 mg Filmtabletten	DE/H/3896/001	90227.00.00	MERCK SHARP & DOHME LTD.	DE
Kexrolt® 10 mg/20 mg Filmtabletten	DE/H/3896/002	90227.01.00	MERCK SHARP & DOHME LTD.	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
Kexrolt® 10 mg/40 mg Filmtabletten	DE/H/3896/003	90227.02.00	MERCK SHARP & DOHME LTD.	DE
Kexrolt® 10 mg/80 mg Filmtabletten	DE/H/3896/004	90227.03.00	MERCK SHARP & DOHME LTD.	DE
KEXROLT 10 mg/10 mg comprimidos recubiertos con película	DE/H/3896/001	79.162	MERCK SHARP & DOHME BV	ES
KEXROLT 10 mg/20 mg comprimidos recubiertos con película	DE/H/3896/002	79.163	MERCK SHARP & DOHME BV	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
KEXROLT 10 mg/40 mg comprimidos recubiertos con película	DE/H/3896/003	79.164	MERCK SHARP & DOHME BV	ES
KEXROLT 10 mg/80 mg comprimidos recubiertos con película	DE/H/3896/004	79.165	MERCK SHARP & DOHME BV	ES
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247028	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247030	MSD ITALIA 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
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247042	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247016	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247170	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/10 mg compresse rivestite con film	DE/H/3896/001	043247218	MSD ITALIA 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
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247055	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247067	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247079	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247143	MSD ITALIA 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
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247182	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/20 mg compresse rivestite con film	DE/H/3896/002	043247220	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247081	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247093	MSD ITALIA 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
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247105	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247156	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247194	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/40 mg compresse rivestite con film	DE/H/3896/003	043247232	MSD ITALIA 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
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247117	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247129	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247131	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247168	MSD ITALIA 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
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247206	MSD ITALIA S.R.L.	IT
KEXROLT 10 mg/80 mg compresse rivestite con film	DE/H/3896/004	043247244	MSD ITALIA S.R.L.	IT