27 June 2024
EMA/372134/2024
Pharmacovigilance Risk Assessment Committee (PRAC)

List of nationally authorised medicinal products

Active substance(s): atorvastatin

Procedure no.: PSUSA/00010347/202310

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Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atorvastatina Atorvan 30 mg comprimidos revestidos por película	not available	5859434	ALTER S.A.	PT
Atorvastatina Atorvan 30 mg comprimidos revestidos por película	not available	5859442	ALTER S.A.	PT
Atorvastatina Atorvan 30 mg comprimidos revestidos por película	not available	5859459	ALTER S.A.	PT
Atorvastatina Atorvan 30 mg comprimidos revestidos por película	not available	5859467	ALTER S.A.	PT
Atorvastatina Atorvan 60 mg comprimidos revestidos por película	not available	5859475	ALTER S.A.	PT
Atorvastatina Atorvan 60 mg comprimidos revestidos por película	not available	5859509	ALTER S.A.	PT
Atorvastatina Atorvan 60 mg comprimidos revestidos por película	not available	5859517	ALTER S.A.	PT
Atorvastatina Atorvan 60 mg comprimidos revestidos por película	not available	5859525	ALTER S.A.	PT
Atorvastatine Aurobindo 30 mg, filmomhulde tabletten	NL/H/2982/005	RVG 119340	AUROBINDO PHARMA B.V.	NL
Atorvastatine Aurobindo 60 mg, filmomhulde tabletten	NL/H/2982/006	RVG 119342	AUROBINDO PHARMA B.V.	NL
Atorvastatin PUREN 30 mg Filmtabletten	NL/H/2982/005	97801.00.00	PUREN PHARMA GMBH & CO. KG	DE
Atorvastatin PUREN 60 mg Filmtabletten	NL/H/2982/006	97802.00.00	PUREN PHARMA GMBH & CO. KG	DE
Atorvastatin Aurobindo 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2982/003	022702	AUROBINDO PHARMA (MALTA) LIMITED	CY
Apo-Atorva, 30 mg, tabletki powlekane	NL/H/2982/005/DC	25360	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
Apo-Atorva, 60 mg, tabletki powlekane	NL/H/2982/006/DC	25361	AUROVITAS PHARMA POLSKA SP. Z O.O	PL
Liprimar 10 mg Filmtabletten	DE/H/3385/001	39587.00.00	VIATRIS PHARMA GMBH	DE
Liprimar 20 mg Filmtabletten	DE/H/3385/002	39587.01.00	VIATRIS PHARMA GMBH	DE
Liprimar 40 mg Filmtabletten	DE/H/3385/003	39587.02.00	VIATRIS PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Liprimar 80 mg Filmtabletten	DE/H/3385/004	77656.00.00	VIATRIS PHARMA GMBH	DE
Orbeos 10 mg filmdragerade tabletter	DE/H/3385/001	24871	UPJOHN EESV	FI
Orbeos 10 mg kalvopäällysteiset tabletit	DE/H/3385/001	24871	UPJOHN EESV	FI
Orbeos 20 mg filmdragerade tabletter	DE/H/3385/002	24872	UPJOHN EESV	FI
Orbeos 40 mg filmdragerade tabletter	DE/H/3385/003	24873	UPJOHN EESV	FI
Orbeos 80 mg filmdragerade tabletter	DE/H/3385/004	24874	UPJOHN EESV	FI
Orbeos 40 mg kalvopäällysteiset tabletit	DE/H/3385/003	24873	UPJOHN EESV	FI
Orbeos 20 mg kalvopäällysteiset tabletit	DE/H/3385/002	24872	UPJOHN EESV	FI
Orbeos 80 mg kalvopäällysteiset tabletit	DE/H/3385/004	24874	UPJOHN EESV	FI
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007016	VIATRIS PHARMA S.R.L.	IT
TORVAST 20 mg compresse rivestite con film	DE/H/3385/002	033007030	VIATRIS PHARMA S.R.L.	IT
TORVAST 20 mg compresse rivestite con film	DE/H/3385/002	033007081	VIATRIS PHARMA S.R.L.	IT
TORVAST 20 mg compresse rivestite con film	DE/H/3385/002	033007042	VIATRIS PHARMA S.R.L.	IT
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007079	VIATRIS PHARMA S.R.L.	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007055	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007384	VIATRIS PHARMA S.R.L.	IT
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007028	VIATRIS PHARMA S.R.L.	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007067	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse	DE/H/3385/004	033007360	VIATRIS PHARMA S.R.L.	IT

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rivestite con film				
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007257	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007321	VIATRIS PHARMA S.R.L.	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007093	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007244	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007333	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007271	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007269	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007372	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007345	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007319	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007295	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007283	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007358	VIATRIS PHARMA S.R.L.	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007307	VIATRIS PHARMA S.R.L.	IT
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/58	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/54	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/06	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/51	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/53	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/50	UPJOHN EESV	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
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/55	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/52	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/60	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/63	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/57	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/49	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/59	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/64	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/56	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/05	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/65	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/62	UPJOHN EESV	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/61	UPJOHN EESV	HU
Zarator 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/001	023717	VIATRIS HELLAS LTD.	CY
Zarator 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/002	023718	VIATRIS HELLAS LTD.	CY
Zarator 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/003	023719	VIATRIS HELLAS LTD.	CY
Atorvastatin Galenicum Health 30 mg film-coated tablets	MT/H/0404/003	MA255/04303	GALENICUM HEALTH, S.L.	MT
Atorvastatin Galenicum Health 60 mg film-coated tablets	MT/H/0404/006	MA255/04305	GALENICUM HEALTH, S.L.	MT
Atorvastatin AbZ 60 mg Filmtabletten	not available	88207.00.00	ABZ-PHARMA GMBH	DE
Atorvastatin AbZ 30 mg Filmtabletten	not available	88206.00.00	ABZ-PHARMA GMBH	DE
Atorvastatine CF 60 mg, filmomhulde tabletten	NL/H/3345/005	RVG 116325	CENTRAFARM B.V.	NL
Atorvastatine CF 30 mg, filmomhulde tabletten	NL/H/3345/003	RVG 116323	CENTRAFARM B.V.	NL
Atorvastatina Stada 60 mg comprimidos recubiertos con película	NL/H/3345/005	80.732	LABORATORIO STADA, S.L.	ES
Atorvastatina Stada 30 mg	NL/H/3345/003	80.730	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
comprimidos recubiertos con película				
Atorvastatin US Pharmacia, 10 mg, tabletki powlekane	not available	25693	US PHARMACIA SP. Z O.O.	PL
Atoris 30 mg potahované tablety	not available	31/661/11-C	KRKA, D.D., NOVO MESTO	CZ
Atoris 60 mg potahované tablety	not available	31/662/11-C	KRKA, D.D., NOVO MESTO	CZ
Atorvastatina Pharmakern 30 mg comprimidos revestidos por película	MT/H/0404/003	5808902	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 30 mg comprimidos revestidos por película	MT/H/0404/003	5808910	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 30 mg comprimidos revestidos por película	MT/H/0404/003	5808928	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 30 mg comprimidos revestidos por película	MT/H/0404/003	5808936	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 30 mg comprimidos revestidos por película	MT/H/0404/003	5808944	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 60 mg comprimidos revestidos por película	MT/H/0404/005	5809041	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 60 mg comprimidos revestidos por película	MT/H/0404/005	5809066	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 60 mg comprimidos revestidos por película	MT/H/0404/005	5809058	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS,	PT

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			SOCIEDADE UNIPESSOAL, LDA.	
Atorvastatina Pharmakern 60 mg comprimidos revestidos por película	MT/H/0404/005	5809025	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Pharmakern 60 mg comprimidos revestidos por película	MT/H/0404/005	5809033	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Atorvastatina Kern Pharma 30 mg comprimidos recubiertos con película	MT/H/0404/003	85406	KERN PHARMA, S.L.	ES
Atorvastatina Kern Pharma 60 mg comprimidos recubiertos con película	MT/H/0404/005	85408	KERN PHARMA, S.L.	ES
Аторис 30 mg филмирани таблетки	not available	20120175	KRKA, D.D., NOVO MESTO	BG
Аторис 60 mg филмирани таблетки	not available	20120176	KRKA, D.D., NOVO MESTO	BG
Atorvastatina ratio 30 mg comprimidos recubiertos con película	not available	77298	TEVA PHARMA S.L.U.,	ES
Atorvastatina ratio 60 mg comprimidos recubiertos con película	not available	77297	TEVA PHARMA S.L.U.,	ES
Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/005	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/006	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/007	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/008	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/009	KRKA, D.D., NOVO MESTO	SI

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Atoris 30 mg filmsko obložene tablete	not available	H/01/00234/010	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/013	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/014	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/015	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/016	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/017	KRKA, D.D., NOVO MESTO	SI
Atoris 60 mg filmsko obložene tablete	not available	H/01/00234/018	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/019	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/020	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/021	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/022	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/023	KRKA, D.D., NOVO MESTO	SI
Atoris 80 mg filmsko obložene tablete	not available	H/01/00234/024	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg apvalkotās tabletes	not available	10-0605	KRKA, D.D., NOVO MESTO	LV
Atoris 60 mg apvalkotās tabletes	not available	10-0606	KRKA, D.D., NOVO MESTO	LV
Sortis 10 mg filmom obložene tablete	not available	HR-H-943745481	UPJOHN EESV	HR
Sortis 20 mg filmom obložene tablete	not available	HR-H-076787872	UPJOHN EESV	HR
Sortis 40 mg filmom obložene tablete	not available	HR-H-232587549	UPJOHN EESV	HR
Sortis 80 mg filmom obložene tablete	not available	HR-H-611144692	UPJOHN EESV	HR

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Atorvastatin HEXAL 30 mg Filmtabletten	AT/H/0197/003	72009.00.00	HEXAL AG	DE
Atolipidrin 30 mg - Filmtabletten	AT/H/0197/003	1-28677	HEXAL PHARMA GMBH	AT
Prevencor 10 mg comprimidos recubiertos con película	DE/H/3820/001	61742	VIATRIS HEALTHCARE S.L.	ES
Prevencor 20 mg comprimidos recubiertos con película	DE/H/3820/002	61743	VIATRIS HEALTHCARE S.L.	ES
Prevencor 40 mg comprimidos recubiertos con película	DE/H/3820/003	61744	VIATRIS HEALTHCARE S.L.	ES
Prevencor 80 mg comprimidos recubiertos con película	DE/H/3820/004	64642	VIATRIS HEALTHCARE S.L.	ES
ATOR Viatris 10 mg Filmtabletten	DE/H/3820/001	87744.00.00	VIATRIS PHARMA GMBH	DE
ATOR Viatris 20 mg Filmtabletten	DE/H/3820/002	87745.00.00	VIATRIS PHARMA GMBH	DE
ATOR Viatris 40 mg Filmtabletten	DE/H/3820/003	87746.00.00	VIATRIS PHARMA GMBH	DE
ATOR Viatris 80 mg Filmtabletten	DE/H/3820/004	87747.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatine STADA 60 mg, filmomhulde tabletten	NL/H/3346/006	RVG 123397	STADA ARZNEIMITTEL AG	NL
Atorvastatine STADA 30 mg, filmomhulde tabletten	NL/H/3346/005	RVG 123396	STADA ARZNEIMITTEL AG	NL
Atorvastatin AL 60 mg Filmtabletten	NL/H/3346/006	2202621.00.00	ALIUD PHARMA GMBH	DE
Atorvastatin AL 30 mg Filmtabletten	NL/H/3346/005	2202620.00.00	ALIUD PHARMA GMBH	DE
ATORVASTATINA EG 30 mg compresse rivestite con film	NL/H/3346/005	044144133	EG S.P.A.	IT
ATORVASTATINA EG 30 mg compresse rivestite con film	NL/H/3346/005	044144145	EG S.P.A.	IT
ATORVASTATINA EG 60 mg compresse rivestite con film	NL/H/3346/006	044144184	EG S.P.A.	IT
ATORVASTATINA EG 60 mg compresse rivestite con film	NL/H/3346/006	044144160	EG S.P.A.	IT
ATORVASTATINA EG 60 mg compresse rivestite con film	NL/H/3346/006	044144172	EG S.P.A.	IT
ATORVASTATINA EG 30 mg compresse rivestite con film	NL/H/3346/005	044144158	EG S.P.A.	IT
Atorvastatin 60 mg film-coated tablets	CZ/H/0868/005	PL 17780/0833	ZENTIVA PHARMA UK LIMITED	XI

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Atorvastatin 30 mg film-coated tablets	CZ/H/0868/003	PL 17780/0832	ZENTIVA PHARMA UK LIMITED	XI
Torvacard Neo 60 mg potahované tablety	CZ/H/0868/005	31/353/18-C	ZENTIVA, K.S.	CZ
Torvacard Neo 30 mg potahované tablety	CZ/H/0868/003	31/351/18-C	ZENTIVA, K.S.	CZ
Atorvastatin Zentiva 30 mg Filmtabletten	CZ/H/0868/003	2202806.00.00	ZENTIVA PHARMA GMBH	DE
Atorvastatin Zentiva 60 mg Filmtabletten	CZ/H/0868/005	2202808.00.00	ZENTIVA PHARMA GMBH	DE
Atorvastatin-ratiopharm® 30 mg Filmtabletten	not available	88204.00.00	RATIOPHARM GMBH	DE
Atorvastatin-ratiopharm® 60 mg Filmtabletten	not available	88205.00.00	RATIOPHARM GMBH	DE
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/004	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/018	KRKA, D.D., NOVO MESTO	LT
Atoris, 30 mg, tabletki powlekane	LT/H/0114/001	18462	KRKA, D.D., NOVO MESTO	PL
Atoris, 60 mg, tabletki powlekane	LT/H/0114/002	18463	KRKA, D.D., NOVO MESTO	PL
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/005	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/006	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/007	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/008	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/009	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/010	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/011	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/012	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
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/013	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/014	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/015	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/016	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	LT/H/0114/001	LT/1/03/0865/017	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/019	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/020	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/021	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/022	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/023	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/024	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/025	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/026	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/027	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/028	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/029	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/030	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	LT/H/0114/002	LT/1/03/0865/031	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
Atorvastatina Tevagen 60 mg comprimidos recubiertos con película	not available	77291	TEVA PHARMA S.L.U.,	ES
Atorvastatina Tevagen 30 mg comprimidos recubiertos con película	not available	77292	TEVA PHARMA S.L.U.,	ES
Atorvastatine Centrient 30 mg, filmomhulde tabletten	NL/H/3795/003	RVG 119390	CENTRIENT PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine Centrient 60 mg, filmomhulde tabletten	NL/H/3795/005	RVG 119392	CENTRIENT PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatin HEXAL 60 mg Filmtabletten	AT/H/0305/006	81792.00.00	HEXAL AG	DE
Atorvastatin Sandoz 60 mg – Filmtabletten	AT/H/0305/006	1-30705	SANDOZ GMBH	AT
Atorvastatin Sandoz 30 mg – Filmtabletten	AT/H/0305/005	1-30704	SANDOZ GMBH	AT
Atorvastatin Heumann 30 mg Filmtabletten	not available	2202894.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Atorvastatin Heumann 60 mg Filmtabletten	not available	2202896.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Atorvastatina Normon 30 mg comprimidos recubiertos con película	not available	87270	LABORATORIOS NORMON, S.A.	ES
Atorvastatina Normon 60 mg comprimidos recubiertos con película	not available	87271	LABORATORIOS NORMON, S.A.	ES
Atorvasa 20 mg Filmtabletten	DE/H/3882/002	87753.00.00	VIATRIS PHARMA GMBH	DE
Atorvasa 40 mg Filmtabletten	DE/H/3882/003	87754.00.00	VIATRIS PHARMA GMBH	DE
Atorvasa 80 mg Filmtabletten	DE/H/3882/004	87755.00.00	VIATRIS PHARMA GMBH	DE
Atorvasa 10 mg Filmtabletten	DE/H/3882/001	87752.00.00	VIATRIS PHARMA GMBH	DE
Cardyl 10 mg comprimidos recubiertos con película	DE/H/3882/001	61716	VIATRIS HEALTHCARE S.L.	ES
Cardyl 20 mg comprimidos recubiertos con película	DE/H/3882/002	61717	VIATRIS HEALTHCARE 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
Cardyl 40 mg comprimidos recubiertos con película	DE/H/3882/003	61718	VIATRIS HEALTHCARE S.L.	ES
Cardyl 80 mg comprimidos recubiertos con película	DE/H/3882/004	64571	VIATRIS HEALTHCARE S.L.	ES
Atorvastatin 10 mg film-coated tablets	not available	PL 43461/0017	FLAMINGO PHARMA UK LTD	XI
Atorvastatin 20 mg film-coated tablets	not available	PL 43461/0018	FLAMINGO PHARMA UK LTD	XI
Atorvastatin 40 mg film-coated tablets	not available	PL 43461/0019	FLAMINGO PHARMA UK LTD	XI
Atorvastatin 80 mg film-coated tablets	not available	PL 43461/0020	FLAMINGO PHARMA UK LTD	XI
Atorvastatin Bluefish AB, 30 mg, tabletki powlekane	IE/H/1099/003	24905	BLUEFISH PHARMACEUTICALS AB	PL
Atorvastatin Bluefish AB, 60 mg, tabletki powlekane	IE/H/1099/005	24907	BLUEFISH PHARMACEUTICALS AB	PL
Atorvastatin Bluefish 30 mg film-coated tablets	IE/H/1099/003	PA1436/027/003	BLUEFISH PHARMACEUTICALS AB	IE
Atorvastatin Bluefish 60 mg film-coated tablets	IE/H/1099/005	PA1436/027/005	BLUEFISH PHARMACEUTICALS AB	IE
Atorvastatina Bluefish 30 mg comprimidos recubiertos con película	IE/H/1099/003	86211	BLUEFISH PHARMACEUTICALS AB	ES
Atorvastatina Bluefish 60 mg comprimidos recubiertos con película	IE/H/1099/005	86213	BLUEFISH PHARMACEUTICALS AB	ES
Atorvastatin Krka 30 mg Filmtabletten	SE/H/0642/005	1-30807	KRKA, D.D., NOVO MESTO	AT
Atorvastatin Krka 60 mg Filmtabletten	SE/H/0642/006	1-30808	KRKA, D.D., NOVO MESTO	AT
Atorvastatin Krka 30 mg filmdragerade tabletter	SE/H/0642/005	44160	KRKA SVERIGE AB	SE
Atorvastatin Krka 60 mg filmdragerade tabletter	SE/H/0642/006	44161	KRKA SVERIGE AB	SE
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561449	KRKA, D.D., NOVO MESTO	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
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561452	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561553	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561565	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561589	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561603	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561678	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561680	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561437	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561464	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561476	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561490	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561502	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561514	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561526	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561540	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561577	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561591	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561615	KRKA, D.D., NOVO MESTO	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
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561627	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561666	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561704	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561488	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	AIC N. 040561538	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561639	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561641	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561654	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	AIC N. 040561692	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/01	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/02	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/03	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/04	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/05	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/06	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/07	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/08	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/09	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/10	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/11	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/12	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/13	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	10776/2018/14	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/01	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/02	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/03	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
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/04	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/05	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/06	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/07	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/08	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/09	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/10	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/11	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/12	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/13	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	10777/2018/14	KRKA, D.D., NOVO MESTO	RO
Atorvastatin Krka 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	021287	KRKA, D.D., NOVO MESTO	CY
Atorvastatin Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	021288	KRKA, D.D., NOVO MESTO	CY
Atoridor 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	85908/03-09-2020	KRKA, D.D., NOVO MESTO	GR
Atoridor 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	87079/03-09-2020	KRKA, D.D., NOVO MESTO	GR
Atorvastatin HCS 30 mg, filmomhulde tabletten	SE/H/0642/005	BE398036	HCS BVBA	BE
Atorvastatin HCS 60 mg, filmomhulde tabletten	SE/H/0642/006	BE398045	HCS BVBA	BE
Atorvastatina Alter Genéricos 30 mg comprimidos recubiertos con película	not available	85.902	LABORATORIOS ALTER, S.A.	ES
Atorvastatina Alter Genéricos 60 mg comprimidos recubiertos con película	not available	85.907	LABORATORIOS ALTER, S.A.	ES
Atorvastatin 10 mg Film-coated Tablets	not available	PL 36687/0258	TORRENT PHARMA (UK) LTD.	XI
Atorvastatin 20 mg Film-coated Tablets	not available	PL 36687/0259	TORRENT PHARMA (UK) LTD.	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atorvastatin 40 mg Film-coated Tablets	not available	PL 36687/0260	TORRENT PHARMA (UK) LTD.	XI
Atoris 30 filmom obalené tablety	not available	31/0057/11-S	KRKA, D.D., NOVO MESTO	SK
Atoris 60 filmom obalené tablety	not available	31/0058/11-S	KRKA, D.D., NOVO MESTO	SK
Atorvastatin 10 mg film-coated tablets	not available	PL 17907/0547	BRISTOL LABORATORIES LIMITED	XI
Atorvastatin 20 mg film-coated tablets	not available	PL 17907/0548	BRISTOL LABORATORIES LIMITED	XI
Atorvastatin 40 mg film-coated tablets	not available	PL 17907/0549	BRISTOL LABORATORIES LIMITED	XI
Atorvastatin 80 mg film-coated tablets	not available	PL 17907/0550	BRISTOL LABORATORIES LIMITED	XI
Atorvastatin Krka, 30 mg, tabletki powlekane	SI/H/0211/001	18465	KRKA, D.D., NOVO MESTO	PL
Atorvastatin Krka, 60 mg, tabletki powlekane	SI/H/0211/002	18466	KRKA, D.D., NOVO MESTO	PL
Astator 30 mg õhukese polümeerikattega tabletid	SI/H/0211/001	742311	KRKA, D.D., NOVO MESTO	EE
Astator 60 mg õhukese polümeerikattega tabletid	SI/H/0211/002	742511	KRKA, D.D., NOVO MESTO	EE
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/001	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/002	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/003	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/004	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/005	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/006	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/007	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/008	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
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/009	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/010	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/011	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/012	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/013	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SI/H/0211/001	LT/1/11/2450/014	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/015	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/016	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/017	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/018	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/019	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/020	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/021	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/022	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/023	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/024	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/025	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/026	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
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/027	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SI/H/0211/002	LT/1/11/2450/028	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/01	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/02	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/03	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/04	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/05	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SI/H/0211/001	OGYI-T-21813/06	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/07	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/08	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/09	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/10	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/11	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SI/H/0211/002	OGYI-T-21813/12	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmom obalené tablety	SI/H/0211/002	31/0633/11-S	KRKA, D.D., NOVO MESTO	SK
Astator 30 mg filmom obalené tablety	SI/H/0211/001	31/0632/11-S	KRKA, D.D., NOVO MESTO	SK
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/001	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/002	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/003	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/004	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/005	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/006	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/007	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene	SI/H/0211/001	H/11/00226/008	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
tablete				
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/009	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/010	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/011	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/012	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/013	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SI/H/0211/001	H/11/00226/014	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/015	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/016	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/017	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/018	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/019	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/020	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/021	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/022	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/023	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/024	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/025	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene	SI/H/0211/002	H/11/00226/026	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
tablete				
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/027	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SI/H/0211/002	H/11/00226/028	KRKA, D.D., NOVO MESTO	SI
Liprimar® 10 mg Kautabletten	DE/H/3616/002	86866.00.00	VIATRIS PHARMA GMBH	DE
Liprimar® 20 mg Kautabletten	DE/H/3616/003	86867.00.00	VIATRIS PHARMA GMBH	DE
Orbeos 20 mg tuggtabletter	DE/H/3616/003	28937	UPJOHN EESV	FI
Orbeos 10 mg tuggtabletter	DE/H/3616/002	28938	UPJOHN EESV	FI
Orbeos 10 mg purutabletit	DE/H/3616/002	28938	UPJOHN EESV	FI
Orbeos 20 mg purutabletit	DE/H/3616/003	28937	UPJOHN EESV	FI
TORVAST 10 mg compresse masticabili	DE/H/3616/002	033007408	VIATRIS PHARMA S.R.L.	IT
TORVAST 20 mg compresse masticabili	DE/H/3616/003	033007410	VIATRIS PHARMA S.R.L.	IT
OBRADON 10 mg rágótabletta	DE/H/3616/002	OGYI-T-8306/12	UPJOHN EESV	HU
ATORVASTATINA DOCgen 30 mg compresse rivestite con film	not available	045140035	DOC GENERICI S.R.L.	IT
ATORVASTATINA DOCgen 60 mg compresse rivestite con film	not available	045140050	DOC GENERICI S.R.L.	IT
Zarator 10 mg μασώμενα δισκία	DE/H/3616/002	2345905	WIN MEDICA SA	GR
Zarator 20 mg μασώμενα δισκία	DE/H/3616/003	2345906	WIN MEDICA SA	GR
Zarator 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/001	2345901	WIN MEDICA SA	GR
Zarator 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/002	2345902	WIN MEDICA SA	GR
Zarator 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/003	2345903	WIN MEDICA SA	GR
Atorvastatine Liconsa 60 mg filmomhulde tabletten	NL/H/5512/002	RVG 129172	LABORATORIOS LICONSA, S.A.	NL
Atorvastatine Liconsa 30 mg filmomhulde tabletten	NL/H/5512/001	RVG 129171	LABORATORIOS LICONSA, S.A.	NL
Atorvastatina Liconsa 30 mg comprimidos recubiertos con película	NL/H/5512/001	88892	LABORATORIOS LICONSA, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atorvastatina Liconsa 60 mg comprimidos recubiertos con película	NL/H/5512/002	88893	LABORATORIOS LICONSA, S.A.	ES
Atorvastatin Diagonalis 30 mg film-coated tablets	MT/H/0624/003	MA255/06903	GALENICUM HEALTH, S.L.	MT
Atorvastatin Diagonalis 60 mg film-coated tablets	MT/H/0624/006	MA255/06905	GALENICUM HEALTH, S.L.	MT
Atorvastatin/Sandoz 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	AT/H/0196/003	41953/19-6-2015	SANDOZ GMBH	GR
Atorvadivid 30 mg - Filmtabletten	AT/H/0196/003	1-28644	HEXAL PHARMA GMBH	AT
Atorvastatin 30 mg film-coated tablets	not available	PL 16363/0512	MILPHARM LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 16363/0513	MILPHARM LIMITED	XI
Atoris 30 mg comprimidos recubiertos con película	DE/H/6594/005	74684	KRKA, D.D., NOVO MESTO	ES
Atoris® 30 mg Filmtabletten	DE/H/6594/005	81701.00.00	TAD PHARMA GMBH	DE
Atoris® 60 mg Filmtabletten	DE/H/6594/006	81702.00.00	TAD PHARMA GMBH	DE
Atorvastatin Aristo 30 mg Filmtabletten	NL/H/3797/003	97973.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Atorvastatin Aristo 60 mg Filmtabletten	NL/H/3797/005	97975.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Atorvastatine Aristo 30 mg, filmomhulde tabletten	NL/H/3797/003	RVG 119402	ARISTO PHARMA GMBH (ART 57)	NL
Atorvastatine Aristo 60 mg, filmomhulde tabletten	NL/H/3797/005	RVG 119404	ARISTO PHARMA GMBH (ART 57)	NL
Atorvastatina Aristo 30 mg comprimidos recubiertos con película	NL/H/3797/003	85385	ARISTO PHARMA GMBH (ART 57)	ES
Atorvastatina Aristo 60 mg comprimidos recubiertos con película	NL/H/3797/005	85387	ARISTO PHARMA GMBH (ART 57)	ES
Atorvastatina Aristo Pharma 30 mg compresse rivestite con film	NL/H/3797/003	048731145	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 30 mg	NL/H/3797/003	048731158	ARISTO PHARMA GMBH (ART	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			57)	
Atorvastatina Aristo Pharma 30 mg compresse rivestite con film	NL/H/3797/003	048731160	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 30 mg compresse rivestite con film	NL/H/3797/003	048731172	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 60 mg compresse rivestite con film	NL/H/3797/005	048731234	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 60 mg compresse rivestite con film	NL/H/3797/005	048731246	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 60 mg compresse rivestite con film	NL/H/3797/005	048731259	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristo Pharma 60 mg compresse rivestite con film	NL/H/3797/005	048731261	ARISTO PHARMA GMBH (ART 57)	IT
Atorvastatina Aristogen 30 mg comprimidos revestidos por película	NL/H/3797/003	NL/H/3797/001-006/E/001	ARISTO PHARMA GMBH (ART 57)	PT
Atorvastatina Aristogen 60 mg comprimidos revestidos por película	NL/H/3797/005	NL/H/3797/001-006/E/001	ARISTO PHARMA GMBH (ART 57)	PT
Atorvastatin 80 mg film-coated tablets	not available	PL 36687/0261	TORRENT PHARMA (UK) LTD.	XI
Atoris 30 mg õhukese polümeerikattega tabletid	not available	744111	KRKA, D.D., NOVO MESTO	EE
Atoris 60 mg õhukese polümeerikattega tabletid	not available	744211	KRKA, D.D., NOVO MESTO	EE
Atorvastatina pensa pharma 30 mg comprimidos recubiertos con película	MT/H/0624/003	89026	TOWA PHARMACEUTICAL S.A.	ES
Atorvastatina pensa pharma 60 mg comprimidos recubiertos con película	MT/H/0624/005	89028	TOWA PHARMACEUTICAL S.A.	ES
Thervan 30 mg comprimidos recubiertos con película	not available	85.927	LABORATORIOS ALTER, S.A.	ES
Thervan 60 mg comprimidos recubiertos con película	not available	85.929	LABORATORIOS ALTER, S.A.	ES
Atorvastatin 10mg Film-coated Tablets	not available	PL 20075/0551	ACCORD HEALTHCARE LIMITED	XI
Atorvastatin 10mg Film-coated	not available	PL 20075/0551	ACCORD HEALTHCARE	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tablets			LIMITED	
Atorvastatin 20mg Film-coated Tablets	not available	PL 20075/0552	ACCORD HEALTHCARE LIMITED	XI
Atorvastatin 40mg Film-coated Tablets	not available	PL 20075/0553	ACCORD HEALTHCARE LIMITED	XI
Atorvastatin 20mg Film-coated Tablets	not available	PL 20075/0552	ACCORD HEALTHCARE LIMITED	XI
Atorvastatin 40mg Film-coated Tablets	not available	PL 20075/0553	ACCORD HEALTHCARE LIMITED	XI
ATORVASTATIN BASICS 30 mg Filmtabletten	MT/H/0125/005	91318.00.00	BASICS GMBH	DE
ATORVASTATIN BASICS 60 mg Filmtabletten	MT/H/0125/006	91319.00.00	BASICS GMBH	DE
Atorab 30 mg film-coated tablets	MT/H/0125/005	MA1048/00105	BASICS GMBH	MT
Atorab 60 mg film-coated tablets	MT/H/0125/006	MA1048/00106	BASICS GMBH	MT
Atorvastatina SUN 30 mg comprimidos recubiertos con película.	MT/H/0125/005/E/001	84056	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
ATORVASTATINA SUN 60 MG COMPRIMIDOS RECUBIERTOS CON PELICULA	MT/H/0125/006/E/001	84055	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Storvas CRT, 30 mg, tabletki powlekane	MT/H/0125/005	25654	RANBAXY POLAND SP. ZO.O	PL
Storvas CRT, 60 mg, tabletki powlekane	MT/H/0125/006	25655	RANBAXY POLAND SP. ZO.O	PL
TOTALIP 40 mg compresse rivestite con film	IT/H/0299/003	033006091	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 10 mg compresse rivestite con film	IT/H/0299/001	033006077	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse rivestite con film	IT/H/0299/002	033006089	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 10 mg compresse rivestite con film	IT/H/0299/001	033006014	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 10 mg compresse rivestite con film	IT/H/0299/001	033006026	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite	IT/H/0299/004	033006317	LABORATORI GUIDOTTI 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
con film				
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006279	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006356	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006368	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006293	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006329	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006281	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006331	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006343	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006255	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006242	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006370	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 40 mg compresse rivestite con film	IT/H/0299/003	033006053	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006305	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 40 mg compresse rivestite con film	IT/H/0299/003	033006065	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006382	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006267	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse rivestite con film	IT/H/0299/002	033006040	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse rivestite	IT/H/0299/002	033006038	LABORATORI GUIDOTTI 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
con film				
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	2022040086	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg	DE/H/6315/006	2022040087	MICRO LABS GMBH	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
Filmtabletten				
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	2022040087	MICRO LABS GMBH	LU
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg	DE/H/6315/005	7004432.00.00	MICRO LABS 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
Filmtabletten				
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 30 mg Filmtabletten	DE/H/6315/005	7004432.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatin Micro Labs 60 mg Filmtabletten	DE/H/6315/006	7004433.00.00	MICRO LABS GMBH	DE
Atorvastatina Davur 60 mg comprimidos recubiertos con película	not available	77293	TEVA PHARMA S.L.U.,	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
Atorvastatina Davur 30 mg comprimidos recubiertos con película	not available	77296	TEVA PHARMA S.L.U.,	ES
TOTALIP 10 mg compresse masticabili	IT/H/0299/006	033006406	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse masticabili	IT/H/0299/007	033006418	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 40 mg compresse masticabili	IT/H/0299/008	033006420	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 5 mg compresse masticabili	IT/H/0299/005	033006394	LABORATORI GUIDOTTI S.P.A.	IT
Atorvastatina Cinfa 30 mg comprimidos recubiertos con película	not available	85715	LABORATORIOS CINFA, S.A.	ES
Atorvastatina Cinfa 60 mg comprimidos recubiertos con película	not available	85716	LABORATORIOS CINFA, S.A.	ES
Sortis® 10 mg Filmtabletten	DE/H/3384/001	39581.00.00	VIATRIS PHARMA GMBH	DE
Sortis® 20 mg Kautabletten	DE/H/3384/007	82884.00.00	VIATRIS PHARMA GMBH	DE
Sortis® 10 mg Kautabletten	DE/H/3384/006	82883.00.00	VIATRIS PHARMA GMBH	DE
Sortis® 20 mg Filmtabletten	DE/H/3384/002	39581.01.00	VIATRIS PHARMA GMBH	DE
Sortis® 40 mg Filmtabletten	DE/H/3384/003	39581.02.00	VIATRIS PHARMA GMBH	DE
Sortis® 80 mg Filmtabletten	DE/H/3384/004	77655.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatin - 1 A Pharma 30 mg Filmtabletten	AT/H/0198/003	71989.00.00	1 A PHARMA GMBH	DE
Callator 30 mg - Filmtabletten	AT/H/0198/003	1-28687	1A PHARMA GMBH	AT
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 30 mg film-coated tablets	not available	PL 25298/0306	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Atorvastatin 60 mg film-coated tablets	not available	PL 25298/0307	BROWN & BURK UK LIMITED	XI
Sortis 10 mg apvalkotās tabletes	DE/H/0109/001	98-0598	UPJOHN EESV	LV
Sortis 10 mg košļājamās tabletes	DE/H/0109/006	10-0403	UPJOHN EESV	LV
Sortis 20 mg apvalkotās tabletes	DE/H/0109/002	98-0599	UPJOHN EESV	LV
Sortis 20 mg košļājamās tabletes	DE/H/0109/007	10-0404	UPJOHN EESV	LV
Sortis 40 mg apvalkotās tabletes	DE/H/0109/003	98-0600	UPJOHN EESV	LV
Sortis 40 mg košļājamās tabletes	DE/H/0109/008	10-0405	UPJOHN EESV	LV
Sortis 5 mg košļājamās tabletes	DE/H/0109/005	10-0402	UPJOHN EESV	LV
Sortis, 10 mg närimistabletid	DE/H/0109/006	694510	UPJOHN EESV	EE
Sortis, 10 mg õhukese polümeerikattega tabletid	DE/H/0109/001	205698	UPJOHN EESV	EE
Sortis, 40 mg närimistabletid	DE/H/0109/008	694710	UPJOHN EESV	EE
Sortis, 40 mg õhukese polümeerikattega tabletid	DE/H/0109/003	205898	UPJOHN EESV	EE
Lipitor 5 mg chewable tablets	DE/H/0109/005	MA1396/00902	UPJOHN HELLAS L.T.D.	MT
Lipitor 10 mg chewable tablets	DE/H/0109/001	MA1396/00903	UPJOHN HELLAS L.T.D.	MT
Lipitor 20 mg chewable tablets	DE/H/0109/002	MA1396/00904	UPJOHN HELLAS L.T.D.	MT
Lipitor 40 mg chewable tablets	DE/H/0109/003	MA1396/00905	UPJOHN HELLAS L.T.D.	MT
Lipitor 10 mg film-coated tablets	DE/H/0109/001	MA1396/00901	UPJOHN HELLAS L.T.D.	MT
Lipitor 10 mg film-coated tablets	DE/H/0109/001	PA 23055/017/001	UPJOHN EESV	IE
Lipitor 20mg chewable tablets	DE/H/0109/002	PA 23055/017/007	UPJOHN EESV	IE
Lipitor 20mg film-coated tablets	DE/H/0109/002	PA 23055/017/002	UPJOHN EESV	IE
Lipitor 40mg chewable tablets	DE/H/0109/003	PA 23055/017/008	UPJOHN EESV	IE
Lipitor 40mg film-coated tablets	DE/H/0109/003	PA 23055/017/003	UPJOHN EESV	IE
Lipitor 5mg chewable tablets	DE/H/0109/005	PA 23055/017/005	UPJOHN EESV	IE
SORTIS 10 mg filmom obalené tablety	DE/H/0109/001	31/0409/98–S	UPJOHN EESV	SK
SORTIS 20 mg filmom obalené tablety	DE/H/0109/002	31/0413/11–S	UPJOHN EESV	SK
SORTIS 40 mg filmom obalené tablety	DE/H/0109/003	31/0414/11–S	UPJOHN EESV	SK
SORTIS 80 mg filmom obalené tablety	DE/H/0109/004	31/0018/03–S	UPJOHN EESV	SK
SORTIS 10 mg žuvacie tablety	DE/H/0109/001	31/0534/10-S	UPJOHN EESV	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
SORTIS 20 mg žuvacie tablety	DE/H/0109/002	31/0535/10-S	UPJOHN EESV	SK
SORTIS 40 mg žuvacie tablety	DE/H/0109/003	31/0536/10-S	UPJOHN EESV	SK
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/03	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/09	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/13	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/01	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/14	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/19	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/06	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/05	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/12	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/04	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/11	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/07	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/02	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/15	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/17	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/08	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/18	UPJOHN EESV	RO
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/015	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/002	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/016	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/012	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/007	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/006	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/004	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/010	UPJOHN EESV	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
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/001	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/008	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/003	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/017	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/018	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/005	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/009	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/011	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/013	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/019	UPJOHN EESV	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/014	UPJOHN EESV	LT
SORTIS 10 mg kramtomosios tabletės	DE/H/0109/001	LT/1/98/2147/002	UPJOHN EESV	LT
Lipitor 80mg film-coated tablets	DE/H/0109/004	PA 23055/017/004	UPJOHN EESV	IE
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/023	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/037	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/022	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/026	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/035	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos	DE/H/0109/002	LT/1/98/3805/021	UPJOHN EESV	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletės				
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/033	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/029	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/031	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/027	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/038	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/020	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/025	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/032	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/034	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/024	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/028	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/036	UPJOHN EESV	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/030	UPJOHN EESV	LT
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/04	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/16	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/18	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/02	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/14	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/11	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/07	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/10	UPJOHN EESV	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
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/13	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/01	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/19	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/03	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/17	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/12	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/09	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/05	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/06	UPJOHN EESV	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/15	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/03	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/01	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/09	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/11	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/13	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/15	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/05	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/07	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/02	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/06	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/12	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/17	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/04	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/16	UPJOHN EESV	RO
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/076	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/064	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/072	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/062	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/058	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos	DE/H/0109/004	LT/1/98/3805/060	UPJOHN EESV	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletės				
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/066	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/068	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/070	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/074	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/075	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/069	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/061	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/071	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/067	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/063	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/065	UPJOHN EESV	LT
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/19	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/17	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/07	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/15	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/03	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/09	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/13	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/11	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/01	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/16	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/04	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/10	UPJOHN EESV	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
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/02	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/12	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/18	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/08	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/14	UPJOHN EESV	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/06	UPJOHN EESV	RO
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/054	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/056	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/040	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/039	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/044	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/046	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/050	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/052	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/042	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/048	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/055	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/057	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/049	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/053	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos	DE/H/0109/003	LT/1/98/3805/043	UPJOHN EESV	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletės				
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/051	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/041	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/047	UPJOHN EESV	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/045	UPJOHN EESV	LT
Sortis 80 mg comprimate filmate	DE/H/0109 MRP	8195/2015/19	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109 MRP	8195/2015/08	UPJOHN EESV	RO
Sortis 5 mg comprimate masticabile	DE/H/0109/005	8188/2015/01	UPJOHN EESV	RO
Sortis 20 mg comprimate masticabile	DE/H/0109/007	8190/2015/01	UPJOHN EESV	RO
Sortis 40 mg comprimate masticabile	DE/H/0109/008	8191/2015/01	UPJOHN EESV	RO
SORTIS 40 mg kramtomosios tabletės	DE/H/0109/008	LT/1/98/2147/004	UPJOHN EESV	LT
Sortis 10 mg comprimate masticabile	DE/H/0109/006	8189/2015/01	UPJOHN EESV	RO
SORTIS 5 mg kramtomosios tabletes	DE/H/0109/005	LT/1/98/2147/001	UPJOHN EESV	LT
SORTIS 20 mg kramtomosios tabletės	DE/H/0109/007	LT/1/98/2147/003	UPJOHN EESV	LT
Lipitor 10 mg chewable tablets	DE/H/0109/006	PA 23055/017/006	UPJOHN EESV	IE
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051587	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051389	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050985	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051488	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051686	UPJOHN EESV	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
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050886	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	2535680	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050688	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050787	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051280	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051181	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	2535789	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051785	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052981	UPJOHN EESV	PT
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/05	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/14	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/18	UPJOHN EESV	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/10	UPJOHN EESV	RO
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4051884	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052783	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052387	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052585	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052684	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	2535888	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4051983	UPJOHN EESV	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
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052189	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052882	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	3538980	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052486	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052080	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053286	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053484	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053088	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	2535987	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053880	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4054185	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4054284	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053187	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020988	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021184	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020582	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021382	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020681	UPJOHN EESV	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
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020889	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021085	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	5047360	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020186	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020483	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020285	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020780	UPJOHN EESV	PT
Zarator 10 mg comprimidos para mastigar	DE/H/0109/001	5319512	UPJOHN EESV	PT
Zarator 20 mg comprimidos para mastigar	DE/H/0109 MRP	5319520	UPJOHN EESV	PT
Zarator 40 mg comprimidos para mastigar	DE/H/0109/003	5319538	UPJOHN EESV	PT
Zarator 5 mg comprimidos para mastigar	DE/H/0109/005	5319504	UPJOHN EESV	PT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/073	UPJOHN EESV	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/059	UPJOHN EESV	LT
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/16	UPJOHN EESV	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/10	UPJOHN EESV	RO
Sortis, 5 mg närimistabletid	DE/H/0109/005	694410	UPJOHN EESV	EE
Sortis 80 mg, 80 mg õhukese polümeerikattega tabletid	DE/H/0109/004	423003	UPJOHN EESV	EE
Sortis, 20 mg närimistabletid	DE/H/0109/002	694610	UPJOHN EESV	EE
Sortis, 20 mg õhukese polümeerikattega tabletid	DE/H/0109/002	205798	UPJOHN EESV	EE
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/08	UPJOHN EESV	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
Lipitor 20 mg film-coated tablets	DE/H/0109/002	PL 50622/0039	UPJOHN UK LIMITED	XI
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/54	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/59	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/79	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/51	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/60	UPJOHN EESV	HU
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053583	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053781	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053682	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4054086	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053385	UPJOHN EESV	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053989	UPJOHN EESV	PT
SORTIS 10 mg rágótabletta	DE/H/0109/001	OGYI-T-6542/18	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/53	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/61	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/64	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/07	UPJOHN EESV	HU
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021283	UPJOHN EESV	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021481	UPJOHN EESV	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052288	UPJOHN EESV	PT
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/52	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/62	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/57	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/58	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/56	UPJOHN EESV	HU
Lipitor 40 mg film-coated tablets	DE/H/0109/003	PL 50622/0041	UPJOHN UK LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 80 mg film-coated tablets	DE/H/0109/004	PL 50622/0043	UPJOHN UK LIMITED	XI
Lipitor 10 mg film-coated tablets	DE/H/0109/001	PL 50622/0036	UPJOHN UK LIMITED	XI
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/55	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/09	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/65	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/08	UPJOHN EESV	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/63	UPJOHN EESV	HU
Sortis 80 mg apvalkotās tabletes	DE/H/0109/004	03-0502	UPJOHN EESV	LV
SORTIS 5 mg žuvacie tablety	DE/H/0109/005	31/0533/10-S	UPJOHN EESV	SK
Lipitor 10 mg Filmtabletten	DE/H/0109/001	2008019645	UPJOHN SRL	LU
Lipitor 10 mg Kautabletten	DE/H/0109/006	2010110025	UPJOHN SRL	LU
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	2008019645	UPJOHN SRL	LU
Lipitor 20 mg Kautabletten	DE/H/0109/007	2010110026	UPJOHN SRL	LU
Lipitor 10 mg comprimés à croquer	DE/H/0109/006	2010110025	UPJOHN SRL	LU
Lipitor 20 mg Filmtabletten	DE/H/0109/002	2008019646	UPJOHN SRL	LU
Lipitor 40 mg Filmtabletten	DE/H/0109/003	2008019647	UPJOHN SRL	LU
Lipitor 20 mg comprimés à croquer	DE/H/0109/007	2010110026	UPJOHN SRL	LU
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	2008019646	UPJOHN SRL	LU
Lipitor 40 mg Kautabletten	DE/H/0109/008	2010110027	UPJOHN SRL	LU
Lipitor 40 mg comprimés à croquer	DE/H/0109/008	2010110027	UPJOHN SRL	LU
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	2008019647	UPJOHN SRL	LU
Lipitor 5 mg Kautabletten	DE/H/0109/005	2010110024	UPJOHN SRL	LU
Lipitor 5 mg comprimés à croquer	DE/H/0109/005	2010110024	UPJOHN SRL	LU
Lipitor 80 mg Filmtabletten	DE/H/0109/004	2008019648	UPJOHN SRL	LU
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	2008019648	UPJOHN SRL	LU
SORTIS 10 mg potahované tablety	DE/H/0109/001	31/233/99-C	UPJOHN EESV	CZ
SORTIS 20 mg potahované tablety	DE/H/0109/002	31/234/99-C	UPJOHN EESV	CZ
SORTIS 40 mg potahované tablety	DE/H/0109/003	31/235/99-C	UPJOHN EESV	CZ
SORTIS 80 mg potahované tablety	DE/H/0109/004	31/397/03-C	UPJOHN EESV	CZ
Atorvastatin Viatris 10 mg Kautabletten	DE/H/0109/006	82572.00.00	VIATRIS PHARMA GMBH	DE
Lipitor 10 mg tuggtabletter	DE/H/0109/006	43475	UPJOHN EESV	SE
Lipitor 20 mg tuggtabletter	DE/H/0109/007	43476	UPJOHN EESV	SE
Lipitor 40 mg tuggtabletter	DE/H/0109/008	43477	UPJOHN EESV	SE
Lipitor 5 mg tuggtabletter	DE/H/0109/005	43473	UPJOHN EESV	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
Lipitor 10 mg filmdragerade tabletter	DE/H/0109/001	13415	UPJOHN EESV	SE
Lipitor 20 mg filmdragerade tabletter	DE/H/0109/002	13416	UPJOHN EESV	SE
Lipitor 40 mg filmdragerade tabletter	DE/H/0109/003	13417	UPJOHN EESV	SE
Lipitor 80 mg filmdragerade tabletter	DE/H/0109/004	17836	UPJOHN EESV	SE
Atorvastatin Viatris 20 mg Kautabletten	DE/H/0109/007	82573.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 40 mg Kautabletten	DE/H/0109/008	82574.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 5 mg Kautabletten	DE/H/0109/005	82571.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 10 mg Filmtabletten	DE/H/0109/001	38635.00.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 20 mg Filmtabletten	DE/H/0109/002	38635.01.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 40 mg Filmtabletten	DE/H/0109/003	38635.02.00	VIATRIS PHARMA GMBH	DE
Atorvastatin Viatris 80 mg Filmtabletten	DE/H/0109/004	38635.03.00	VIATRIS PHARMA GMBH	DE
Lipitor 10 mg comprimés à croquer	DE/H/0109/006	BE375986	UPJOHN SRL	BE
Lipitor 10 mg kauwtabletten	DE/H/0109/006	BE375986	UPJOHN SRL	BE
Lipitor 20 mg kauwtabletten	DE/H/0109/007	BE375995	UPJOHN SRL	BE
Lipitor 20 mg comprimés à croquer	DE/H/0109/007	BE375995	UPJOHN SRL	BE
Lipitor 40 mg kauwtabletten	DE/H/0109/008	BE376004	UPJOHN SRL	BE
Lipitor 40 mg comprimés à croquer	DE/H/0109/008	BE376004	UPJOHN SRL	BE
Lipitor 40 mg Kautabletten	DE/H/0109/008	BE376004	UPJOHN SRL	BE
Lipitor 5 mg Kautabletten	DE/H/0109/005	BE375977	UPJOHN SRL	BE
Lipitor 5 mg kauwtabletten	DE/H/0109/005	BE375977	UPJOHN SRL	BE
Lipitor 5 mg comprimés à croquer	DE/H/0109/005	BE375977	UPJOHN SRL	BE
Lipitor 10 mg filmomhulde tabletten	DE/H/0109/001	BE184082	UPJOHN SRL	BE
Lipitor 10 mg filmomhulde tabletten	DE/H/0109/001	BE307727	UPJOHN SRL	BE
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	BE307727	UPJOHN SRL	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
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	BE184082	UPJOHN SRL	BE
Lipitor 20 mg filmomhulde tabletten	DE/H/0109/002	BE307736	UPJOHN SRL	BE
Lipitor 20 mg filmomhulde tabletten	DE/H/0109/002	BE184073	UPJOHN SRL	BE
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	BE307736	UPJOHN SRL	BE
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	BE184073	UPJOHN SRL	BE
Lipitor 40 mg Filmtabletten	DE/H/0109/003	BE184064	UPJOHN SRL	BE
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	BE184064	UPJOHN SRL	BE
Lipitor 40 mg filmomhulde tabletten	DE/H/0109/003	BE184064	UPJOHN SRL	BE
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	BE307745	UPJOHN SRL	BE
Lipitor 40 mg filmomhulde tabletten	DE/H/0109/003	BE307745	UPJOHN SRL	BE
Lipitor 40 mg Filmtabletten	DE/H/0109/003	BE307745	UPJOHN SRL	BE
Lipitor 80 mg filmomhulde tabletten	DE/H/0109/004	BE307754	UPJOHN SRL	BE
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	BE307754	UPJOHN SRL	BE
Lipitor 80 mg Filmtabletten	DE/H/0109/004	BE232933	UPJOHN SRL	BE
Lipitor 80 mg filmomhulde tabletten	DE/H/0109/004	BE232933	UPJOHN SRL	BE
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	BE232933	UPJOHN SRL	BE
Lipitor 80 mg Filmtabletten	DE/H/0109/004	BE307754	UPJOHN SRL	BE
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005099	VIATRIS PHARMA S.R.L.	IT
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005063	VIATRIS PHARMA S.R.L.	IT
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005051	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005339	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005315	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005380	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005366	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005303	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005277	VIATRIS PHARMA 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
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005291	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005354	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005240	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005378	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005289	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005253	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005265	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005327	VIATRIS PHARMA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005341	VIATRIS PHARMA S.R.L.	IT
XARATOR 40 mg compresse masticabili	DE/H/0109/008	033005428	VIATRIS PHARMA S.R.L.	IT
XARATOR 5 mg compresse masticabili	DE/H/0109/005	033005392	VIATRIS PHARMA S.R.L.	IT
XARATOR 10 mg compresse masticabili	DE/H/0109/006	033005404	VIATRIS PHARMA S.R.L.	IT
XARATOR 20 mg compresse masticabili	DE/H/0109/007	033005416	VIATRIS PHARMA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005087	VIATRIS PHARMA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005036	VIATRIS PHARMA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005048	VIATRIS PHARMA S.R.L.	IT
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005012	VIATRIS PHARMA S.R.L.	IT
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005024	VIATRIS PHARMA 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
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005075	VIATRIS PHARMA S.R.L.	IT
Lipitor 80 mg filmdrasjerte tabletter	DE/H/0109/004	00-7221	UPJOHN EESV	NO
Lipitor 20 mg filmdrasjerte tabletter	DE/H/0109/002	96-3033	UPJOHN EESV	NO
Lipitor 10 mg filmdrasjerte tabletter	DE/H/0109/001	96-3032	UPJOHN EESV	NO
Lipitor 40 mg filmdrasjerte tabletter	DE/H/0109/003	96-3034	UPJOHN EESV	NO
Lipitor 5 mg tyggetabletter	DE/H/0109/005	10-7961	UPJOHN EESV	NO
Lipitor 40 mg tyggetabletter	DE/H/0109/008	10-7964	UPJOHN EESV	NO
Lipitor 20 mg tyggetabletter	DE/H/0109/007	10-7963	UPJOHN EESV	NO
Lipitor 10 mg tyggetabletter	DE/H/0109/006	10-7962	UPJOHN EESV	NO
Lipitor 10 mg purutabletit	DE/H/0109/006	28933	UPJOHN EESV	FI
Lipitor 20 mg purutabletit	DE/H/0109/007	28934	UPJOHN EESV	FI
Lipitor 40 mg purutabletit	DE/H/0109/008	28935	UPJOHN EESV	FI
Lipitor 5 mg purutabletit	DE/H/0109/005	28932	UPJOHN EESV	FI
Lipitor 10 mg kalvopäällysteiset tabletit	DE/H/0109/001	12612	UPJOHN EESV	FI
Lipitor 20 mg kalvopäällysteiset tabletit	DE/H/0109/002	12613	UPJOHN EESV	FI
Lipitor 40 mg kalvopäällysteiset tabletit	DE/H/0109/003	12614	UPJOHN EESV	FI
Lipitor 80 mg kalvopäällysteiset tabletit	DE/H/0109/004	16881	UPJOHN EESV	FI
Zarator 10 mg tuggutöflur	DE/H/0109/006	IS/1/10/081/02	UPJOHN EESV	IS
Zarator 20 mg tuggutöflur	DE/H/0109/007	IS/1/10/081/03	UPJOHN EESV	IS
Zarator 40 mg tuggutöflur	DE/H/0109/008	IS/1/10/081/04	UPJOHN EESV	IS
Zarator 5 mg tuggutöflur	DE/H/0109/005	IS/1/10/081/01	UPJOHN EESV	IS
Zarator 80 mg filmuhúðaðar töflur	DE/H/0109/004	IS/1/01/059/01	UPJOHN EESV	IS
Sortis® 10 mg Filmtabletten	DE/H/0109/001	1-21927	UPJOHN EESV	AT
Sortis® 20 mg Filmtabletten	DE/H/0109/002	1-21928	UPJOHN EESV	AT
Sortis® 40 mg Filmtabletten	DE/H/0109/003	1-21926	UPJOHN EESV	AT
Sortis® 80 mg Filmtabletten	DE/H/0109/004	1-24525	UPJOHN EESV	AT
Sortis 10 mg Kautabletten	DE/H/0109/006	1-29555	UPJOHN EESV	AT
Sortis 20 mg Kautabletten	DE/H/0109/007	1-29556	UPJOHN EESV	AT
Sortis 5 mg Kautabletten	DE/H/0109/005	1-29554	UPJOHN EESV	AT
Sortis 40 mg Kautabletten	DE/H/0109/008	1-29557	UPJOHN EESV	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
Lipitor 10 mg, kauwtabletten	DE/H/0109/006	RVG 107607	VIATRIS NETHERLANDS B.V.	NL
Lipitor 20 mg, kauwtabletten	DE/H/0109/007	RVG 107608	VIATRIS NETHERLANDS B.V.	NL
Lipitor 40 mg, kauwtabletten	DE/H/0109/008	RVG 107610	VIATRIS NETHERLANDS B.V.	NL
Lipitor 5 mg, kauwtabletten	DE/H/0109/005	RVG 107605	VIATRIS NETHERLANDS B.V.	NL
Lipitor 10, filmomhulde tabletten 10 mg	DE/H/0109/001	RVG 21081	VIATRIS NETHERLANDS B.V.	NL
Lipitor 20, filmomhulde tabletten 20 mg	DE/H/0109/002	RVG 21082	VIATRIS NETHERLANDS B.V.	NL
Lipitor 40, filmomhulde tabletten 40 mg	DE/H/0109/003	RVG 21083	VIATRIS NETHERLANDS B.V.	NL
Lipitor 80, filmomhulde tabletten 80 mg	DE/H/0109/004	RVG 27148	VIATRIS NETHERLANDS B.V.	NL
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/032	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/024	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/026	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/027	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/037	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/036	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/021	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/025	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/020	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/029	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/031	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/035	UPJOHN EESV	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
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/023	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/030	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/028	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/033	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/034	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/038	UPJOHN EESV	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/022	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/067	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/069	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/060	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/073	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/062	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/072	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/058	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/061	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/059	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/063	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/070	UPJOHN EESV	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
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/065	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/066	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/068	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/076	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/075	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/071	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/074	UPJOHN EESV	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/064	UPJOHN EESV	SI
Sortis 40 mg žvečljive tablete	DE/H/0109/008	H/00/01447/080	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/044	UPJOHN EESV	SI
Lipitor 10 mg tuggtabletter	DE/H/0109/006	28933	UPJOHN EESV	FI
Lipitor 20 mg tuggtabletter	DE/H/0109/007	28934	UPJOHN EESV	FI
Lipitor 40 mg tuggtabletter	DE/H/0109/008	28935	UPJOHN EESV	FI
Lipitor 5 mg tuggtabletter	DE/H/0109/005	28932	UPJOHN EESV	FI
Lipitor 10 mg filmdragerade tabletter	DE/H/0109/001	12612	UPJOHN EESV	FI
Lipitor 20 mg filmdragerade tabletter	DE/H/0109/002	12613	UPJOHN EESV	FI
Lipitor 40 mg filmdragerade tabletter	DE/H/0109/003	12614	UPJOHN EESV	FI
Lipitor 80 mg filmdragerade tabletter	DE/H/0109/004	16881	UPJOHN EESV	FI
TAHOR 10 mg, comprimé à croquer	DE/H/0109/006	34009 494 587 8 3	VIATRIS UP	FR
TAHOR 20 mg, comprimé à croquer	DE/H/0109/007	34009 494 588 4 4	VIATRIS UP	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 218 982 3 1	VIATRIS UP	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 343 067 5 4	VIATRIS UP	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
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 560 303 9 2	VIATRIS UP	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 560 302 2 4	VIATRIS UP	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 371 992 1 6	VIATRIS UP	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 560 304 5 3	VIATRIS UP	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 218 984 6 0	VIATRIS UP	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 343 068 1 5	VIATRIS UP	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 371 993 8 4	VIATRIS UP	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 560 305 1 4	VIATRIS UP	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 560 307 4 3	VIATRIS UP	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 371 994 4 5	VIATRIS UP	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 343 069 8 3	VIATRIS UP	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 560 306 8 2	VIATRIS UP	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 218 985 2 1	VIATRIS UP	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 576 7 4	VIATRIS UP	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 575 0 6	VIATRIS UP	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 356 335 3 8	VIATRIS UP	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 371 995 0 6	VIATRIS UP	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 577 3 5	VIATRIS UP	FR
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/051	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/043	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/056	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/047	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/055	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/050	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/041	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/046	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene	DE/H/0109/003	H/00/01447/052	UPJOHN EESV	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
tablete				
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/053	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/040	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/048	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/057	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/054	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/049	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/042	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/045	UPJOHN EESV	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/039	UPJOHN EESV	SI
Sortis 10 mg žvečljive tablete	DE/H/0109/006	H/00/01447/078	UPJOHN EESV	SI
Sortis 5 mg žvečljive tablete	DE/H/0109/005	H/00/01447/077	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/008	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/016	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/015	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/006	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/012	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/007	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/005	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene	DE/H/0109/001	H/00/01447/014	UPJOHN EESV	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
tablete				
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/003	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/001	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/004	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/010	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/011	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/009	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/017	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/002	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/013	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/018	UPJOHN EESV	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/019	UPJOHN EESV	SI
Sortis 20 mg žvečljive tablete	DE/H/0109/007	H/00/01447/079	UPJOHN EESV	SI
СОРТИС 80 mg филмирани таблетки	DE/H/0109/004	20030461	UPJOHN EESV	BG
СОРТИС 10 mg таблетки за дъвчене	DE/H/0109/006	20100549	UPJOHN EESV	BG
СОРТИС 20 mg таблетки за дъвчене	DE/H/0109/007	20100550	UPJOHN EESV	BG
СОРТИС 40 mg таблетки за дъвчене	DE/H/0109/008	20100551	UPJOHN EESV	BG
СОРТИС 5 mg таблетки за дъвчене	DE/H/0109/005	20100548	UPJOHN EESV	BG
СОРТИС 10 mg филмирани таблетки	DE/H/0109/001	9800143	UPJOHN EESV	BG
СОРТИС 20 mg филмирани	DE/H/0109/002	9800142	UPJOHN EESV	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				
СОРТИС 40 mg филмирани таблетки	DE/H/0109/003	9800141	UPJOHN EESV	BG
SORTIS 10, 10 mg, tabletki do rozgryzania i żucia	DE/H/0109/006	17436	UPJOHN EESV	PL
SORTIS 20, 20 mg, tabletki do rozgryzania i żucia	DE/H/0109/007	17435	UPJOHN EESV	PL
SORTIS 10, 10 mg, tabletki powlekane	DE/H/0109/001	7846	UPJOHN EESV	PL
SORTIS 20, 20 mg, tabletki powlekane	DE/H/0109/002	7847	UPJOHN EESV	PL
SORTIS 40, 40 mg, tabletki powlekane	DE/H/0109/003	7848	UPJOHN EESV	PL
SORTIS 80, 80 mg, tabletki powlekane	DE/H/0109/004	10188	UPJOHN EESV	PL
Lipitor 10 mg Kautabletten	DE/H/0109/006	BE375986	UPJOHN SRL	BE
Lipitor 20 mg Filmtabletten	DE/H/0109/002	BE184073	UPJOHN SRL	BE
Lipitor 20 mg Filmtabletten	DE/H/0109/002	BE307736	UPJOHN SRL	BE
Lipitor 20 mg Kautabletten	DE/H/0109/007	BE375995	UPJOHN SRL	BE
Lipitor 10 mg Filmtabletten	DE/H/0109/001	BE307727	UPJOHN SRL	BE
Lipitor 10 mg Filmtabletten	DE/H/0109/001	BE184082	UPJOHN SRL	BE
Lipitor 10 mg μασώμενα δισκία	DE/H/0109/006	20728	VIATRIS HELLAS LTD.	CY
Lipitor 20 mg μασώμενα δισκία	DE/H/0109/007	20729	VIATRIS HELLAS LTD.	CY
Lipitor 40 mg μασώμενα δισκία	DE/H/0109/008	20730	VIATRIS HELLAS LTD.	CY
Lipitor 5 mg μασώμενα δισκία	DE/H/0109/005	20727	VIATRIS HELLAS LTD.	CY
Lipitor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/001	19489	VIATRIS HELLAS LTD.	CY
Lipitor 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/002	19490	VIATRIS HELLAS LTD.	CY
Lipitor 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/003	19491	VIATRIS HELLAS LTD.	CY
Lipitor 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/002	34557/19-04-2016	VIATRIS HELLAS LTD.	GR
Lipitor 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/003	34558/19-04-2016	VIATRIS HELLAS 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
Lipitor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/001	34556/19-04-2016	VIATRIS HELLAS LTD.	GR
Lipitor 20 mg μασώμενα δισκία	DE/H/0109/007	34562/19-04-2016	VIATRIS HELLAS LTD.	GR
Lipitor 40 mg μασώμενα δισκία	DE/H/0109/008	34563/19-04-2016	VIATRIS HELLAS LTD.	GR
Lipitor 80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/004	34559/19-04-2016	VIATRIS HELLAS LTD.	GR
Lipitor 10 mg μασώμενα δισκία	DE/H/0109/006	34561/19-04-2016	VIATRIS HELLAS LTD.	GR
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020384	UPJOHN EESV	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051082	UPJOHN EESV	PT
Lipitor 10 mg chewable tablets.	DE/H/0109/006	PL 50622/0037	UPJOHN UK LIMITED	XI
Lipitor 20 mg chewable tablets.	DE/H/0109/007	PL 50622/0038	UPJOHN UK LIMITED	XI
Lipitor 40 mg chewable tablets.	DE/H/0109/008	PL 50622/0040	UPJOHN UK LIMITED	XI
Zarator 40 mg filmuhúðaðar töflur	DE/H/0109/003	960277 (IS)	UPJOHN EESV	IS
Zarator, 5 mg tyggetabletter	DE/H/0109/005	47143	UPJOHN EESV	DK
Lipitor 5 mg μασώμενα δισκία	DE/H/0109/005	34560/19-04-2016	VIATRIS HELLAS LTD.	GR
Zarator 10 mg filmuhúðaðar töflur	DE/H/0109/001	960275 (IS)	UPJOHN EESV	IS
Zarator, 20 mg filmovertrukne tabletter	DE/H/0109/002	18613	UPJOHN EESV	DK
Zarator, 40 mg tyggetabletter	DE/H/0109/008	47146	UPJOHN EESV	DK
Zarator, 10 mg tyggetabletter	DE/H/0109/006	47144	UPJOHN EESV	DK
Lipitor 5 mg chewable tablets	DE/H/0109/005	PL 50622/0042	UPJOHN UK LIMITED	XI
Zarator, 20 mg tyggetabletter	DE/H/0109/007	47145	UPJOHN EESV	DK
Zarator, 10 mg filmovertrukne tabletter	DE/H/0109/001	18612	UPJOHN EESV	DK
Zarator 20 mg filmuhúðaðar töflur	DE/H/0109/002	960276 (IS)	UPJOHN EESV	IS
Zarator, 40 mg filmovertrukne tabletter	DE/H/0109/003	18614	UPJOHN EESV	DK
Zarator, 80 mg filmovertrukne tabletter	DE/H/0109/004	32955	UPJOHN EESV	DK
Zarator 10 mg comprimidos recubiertos con película	DE/H/0109/001	61654	VIATRIS HEALTHCARE LTD	ES
Zarator 80 mg comprimidos recubiertos con película	DE/H/0109/004	64529	VIATRIS HEALTHCARE LTD	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
Zarator 10 mg comprimidos masticables	DE/H/0109/006	72599	VIATRIS HEALTHCARE LTD	ES
Zarator 20 mg comprimidos masticables	DE/H/0109/007	72601	VIATRIS HEALTHCARE LTD	ES
Zarator 20 mg comprimidos recubiertos con película	DE/H/0109/002	61655	VIATRIS HEALTHCARE LTD	ES
Zarator 40 mg comprimidos recubiertos con película	DE/H/0109/003	61656	VIATRIS HEALTHCARE LTD	ES