EMA/PRAC/384956/2018
Human Medicines Evaluation Division

List of nationally authorised products

Active substance(s): atorvastatin

Procedure No.: PSUSA/00010347/201710

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Аторис 80 mg филмирани таблетки	not available	20120177	KRKA, D.D., NOVO MESTO	BG
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 10 mg compresse masticabili	IT/H/0299/006	033006406	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006317	LABORATORI GUIDOTTI S.P.A.	IT
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	IT/H/0299/004	033006368	LABORATORI GUIDOTTI	IT

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compresse rivestite con film			S.P.A.	
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	IT/H/0299/003	033006065	LABORATORI GUIDOTTI S.P.A.	IT

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film				
TOTALIP 80 mg compresse rivestite con film	IT/H/0299/004	033006382	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse masticabili	IT/H/0299/007	033006418	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 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
TOTALIP 20 mg compresse rivestite con film	IT/H/0299/002	033006040	LABORATORI GUIDOTTI S.P.A.	IT
TOTALIP 20 mg compresse rivestite con film	IT/H/0299/002	033006038	LABORATORI GUIDOTTI S.P.A.	IT
Atoris 30 potahované tablety	not available	31/661/11-C	KRKA, D.D., NOVO MESTO	CZ
Atoris 60 potahované tablety	not available	31/662/11-C	KRKA, D.D., NOVO MESTO	CZ
Atorvastatine DSM Sinochem 30 mg, filmomhulde tabletten	NL/H/3796/003	RVG 119396	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine DSM Sinochem 60 mg, filmomhulde tabletten	NL/H/3796/005	RVG 119398	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/034	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/022	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele	DE/H/0109/002	LT/1/98/3805/024	PFIZER EUROPE MA EEIG	LT

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dengtos tabletės				
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/026	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/028	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/036	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/020	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/037	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/030	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/033	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/038	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/023	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/031	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/029	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/032	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/021	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/035	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/025	PFIZER EUROPE MA EEIG	LT
SORTIS 20 mg plėvele dengtos tabletės	DE/H/0109/002	LT/1/98/3805/027	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele	DE/H/0109/003	LT/1/98/3805/047	PFIZER EUROPE MA EEIG	LT

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dengtos tabletės				
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/043	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/041	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/045	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/039	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/040	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/051	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/055	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/049	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/057	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/053	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/050	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/044	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/042	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/048	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/046	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/056	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg plėvele	DE/H/0109/003	LT/1/98/3805/054	PFIZER EUROPE MA EEIG	LT

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dengtos tabletės				
SORTIS 40 mg plėvele dengtos tabletės	DE/H/0109/003	LT/1/98/3805/052	PFIZER EUROPE MA EEIG	LT
SORTIS 40 mg kramtomosios tabletės	DE/H/0109/008	LT/1/98/2147/004	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/076	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/066	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/064	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/062	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/072	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/074	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/058	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/060	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/068	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/059	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/069	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/070	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/067	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/061	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele	DE/H/0109/004	LT/1/98/3805/073	PFIZER EUROPE MA EEIG	LT

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dengtos tabletės				
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/065	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/075	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/071	PFIZER EUROPE MA EEIG	LT
SORTIS 80 mg plėvele dengtos tabletės	DE/H/0109/004	LT/1/98/3805/063	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/010	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/012	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/008	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/016	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/005	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/007	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/006	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/003	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/004	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/014	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/009	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/015	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele	DE/H/0109/001	LT/1/98/3805/017	PFIZER EUROPE MA EEIG	LT

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dengtos tabletės				
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/001	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/018	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/002	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/013	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/019	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg plėvele dengtos tabletės	DE/H/0109/001	LT/1/98/3805/011	PFIZER EUROPE MA EEIG	LT
SORTIS 10 mg kramtomosios tabletės	DE/H/0109/006	LT/1/98/2147/002	PFIZER EUROPE MA EEIG	LT
SORTIS 5 mg kramtomosios tabletės	DE/H/0109/005	LT/1/98/2147/001	PFIZER EUROPE MA EEIG	LT
Lipitor 10 mg film-coated tablets	DE/H/0109/001	PL 39933/0001	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 10 mg kalvopäällysteiset tabletit	DE/H/0109/001	12612	PFIZER OY	FI
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/01	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/33	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/35	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/27	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/31	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/71	PFIZER KFT.	HU
SORTIS 10 mg	DE/H/0109/001	OGYI-T-6542/28	PFIZER KFT.	HU

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filmtabletta				
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/29	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/02	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/13	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/67	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/14	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/11	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/15	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/12	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/69	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/16	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/66	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/70	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/68	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/80	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/10	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/75	PFIZER KFT.	HU
SORTIS 80 mg	DE/H/0109/004	OGYI-T-6542/73	PFIZER KFT.	HU

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filmtabletta				
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/74	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/76	PFIZER KFT.	HU
SORTIS 80 mg filmtabletta	DE/H/0109/004	OGYI-T-6542/72	PFIZER KFT.	HU
Zarator, filmovertrukne tabletter	DE/H/0109/001	18612	PFIZER APS	DK
Zarator, filmovertrukne tabletter	DE/H/0109/002	18613	PFIZER APS	DK
Zarator, filmovertrukne tabletter	DE/H/0109/003	18614	PFIZER APS	DK
Zarator, tyggetabletter	DE/H/0109/005	47143	PFIZER APS	DK
Zarator, tyggetabletter	DE/H/0109/008	47146	PFIZER APS	DK
Zarator, filmovertrukne tabletter	DE/H/0109/004	32955	PFIZER APS	DK
Zarator, tyggetabletter	DE/H/0109/006	47144	PFIZER APS	DK
Zarator, tyggetabletter	DE/H/0109/007	47145	PFIZER APS	DK
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/64	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/62	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/53	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/56	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/61	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/07	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/79	PFIZER KFT.	HU
SORTIS 40 mg	DE/H/0109/003	OGYI-T-6542/58	PFIZER KFT.	HU

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filmtabletta				
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/52	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/59	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/57	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/60	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/51	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/08	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/54	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/63	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/55	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/65	PFIZER KFT.	HU
SORTIS 40 mg filmtabletta	DE/H/0109/003	OGYI-T-6542/09	PFIZER KFT.	HU
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005265	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005339	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005315	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con	DE/H/0109/004	033005354	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005253	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005277	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005303	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005240	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005327	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005378	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005289	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005380	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005291	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005341	PFIZER ITALIA S.R.L.	IT
XARATOR 80 mg compresse rivestite con film	DE/H/0109/004	033005366	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005012	PFIZER ITALIA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005036	PFIZER ITALIA S.R.L.	IT
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005099	PFIZER ITALIA S.R.L.	IT
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005063	PFIZER ITALIA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005048	PFIZER ITALIA S.R.L.	IT
XARATOR 40 mg compresse rivestite con film	DE/H/0109/003	033005051	PFIZER ITALIA S.R.L.	IT
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005075	PFIZER ITALIA S.R.L.	IT
XARATOR 20 mg compresse rivestite con film	DE/H/0109/002	033005087	PFIZER ITALIA S.R.L.	IT
Lipitor 80 mg kalvopäällysteiset tabletit	DE/H/0109/004	16881	PFIZER OY	FI
XARATOR 10 mg compresse rivestite con film	DE/H/0109/001	033005024	PFIZER ITALIA S.R.L.	IT
LIPITOR 80, filmomhulde tabletten 80 mg	DE/H/0109/004	RVG 27148	PFIZER B.V.	NL
XARATOR 10 mg compresse masticabili	DE/H/0109/006	033005404	PFIZER ITALIA S.R.L.	IT
XARATOR 20 mg	DE/H/0109/007	033005416	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse masticabili				
XARATOR 5 mg compresse masticabili	DE/H/0109/005	033005392	PFIZER ITALIA S.R.L.	IT
XARATOR 40 mg compresse masticabili	DE/H/0109/008	033005428	PFIZER ITALIA S.R.L.	IT
Lipitor 80 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/004	34559/19-04-2016	PFIZER HELLAS, A.E.	GR
Lipitor 80 mg filmdragerade tabletter	DE/H/0109/004	16881	PFIZER OY	FI
Lipitor 80 mg filmdragerade tabletter	DE/H/0109/004	17836	PFIZER AB	SE
Lipitor 80 mg film-coated tablets	DE/H/0109/004	PL 39933/0004	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 5 mg tuggtabletter	DE/H/0109/005	43473	PFIZER AB	SE
Lipitor 5 mg purutabletit	DE/H/0109/005	28932	PFIZER OY	FI
Lipitor 5 mg tuggtabletter	DE/H/0109/005	28932	PFIZER OY	FI
Lipitor 5 mg, kauwtabletten	DE/H/0109/005	RVG 107605	PFIZER B.V.	NL
Lipitor 5 mg μασώμενα δισκία	DE/H/0109/005	34560/19-04-2016	PFIZER HELLAS, A.E.	GR
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/30	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/34	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/21	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/25	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/32	PFIZER KFT.	HU
SORTIS 10 mg	DE/H/0109/001	OGYI-T-6542/22	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmtabletta				
Sortis® 10 mg Kautabletten	DE/H/0109/006	1-29555	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sortis® 20 mg Kautabletten	DE/H/0109/007	1-29556	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sortis® 40 mg Kautabletten	DE/H/0109/008	1-29557	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/03	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/24	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/26	PFIZER KFT.	HU
Sortis® 5 mg Kautabletten	DE/H/0109/005	1-29554	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/23	PFIZER KFT.	HU
SORTIS 10 mg filmtabletta	DE/H/0109/001	OGYI-T-6542/77	PFIZER KFT.	HU
Lipitor 10 mg filmdragerade tabletter	DE/H/0109/001	13415	PFIZER AB	SE
Lipitor 10 mg tuggtabletter	DE/H/0109/006	43475	PFIZER AB	SE
Lipitor 10 mg, kauwtabletten	DE/H/0109/006	RVG 107607	PFIZER B.V.	NL
Lipitor 10 mg tuggtabletter	DE/H/0109/006	28933	PFIZER OY	FI
Lipitor 10 mg purutabletit	DE/H/0109/006	28933	PFIZER OY	FI
SORTIS 10 mg	DE/H/0109/006	OGYI-T-6542/18	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rágótabletta				
Lipitor 20 mg film-coated tablets	DE/H/0109/002	PL 39933/0002	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 20 mg kalvopäällysteiset tabletit	DE/H/0109/002	12613	PFIZER OY	FI
Lipitor 20 mg filmdragerade tabletter	DE/H/0109/002	13416	PFIZER AB	SE
Lipitor 20 mg filmdragerade tabletter	DE/H/0109/002	12613	PFIZER OY	FI
Lipitor 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/002	34557/19-04-2016	PFIZER HELLAS, A.E.	GR
Lipitor 20 mg μασώμενα δισκία	DE/H/0109/007	34562/19-04-2016	PFIZER HELLAS, A.E.	GR
LIPITOR 20, filmomhulde tabletten 20 mg	DE/H/0109/002	RVG 21082	PFIZER B.V.	NL
Lipitor 20 mg, kauwtabletten	DE/H/0109/007	RVG 107608	PFIZER B.V.	NL
Lipitor 20 mg purutabletit	DE/H/0109/007	28934	PFIZER OY	FI
Lipitor 20 mg tuggtabletter	DE/H/0109/007	28934	PFIZER OY	FI
Lipitor 20 mg tuggtabletter	DE/H/0109/007	43476	PFIZER AB	SE
Lipitor 40 mg filmdragerade tabletter	DE/H/0109/003	12614	PFIZER OY	FI
Lipitor 40 mg tuggtabletter	DE/H/0109/008	28935	PFIZER OY	FI
Lipitor 40 mg filmdragerade tabletter	DE/H/0109/003	13417	PFIZER AB	SE
Lipitor 40 mg purutabletit	DE/H/0109/008	28935	PFIZER OY	FI
Lipitor 40 mg kalvopäällysteiset tabletit	DE/H/0109/003	12614	PFIZER OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 40 mg tuggtabletter	DE/H/0109/008	43477	PFIZER AB	SE
Zarator 20 mg filmuhúðaðar töflur	DE/H/0109/002	960276	PFIZER APS	IS
Zarator 40 mg filmuhúðaðar töflur	DE/H/0109/003	960277	PFIZER APS	IS
Zarator 80 mg filmuhúðaðar töflur	DE/H/0109/004	IS/1/01/059/01	PFIZER APS	IS
Zarator 10 mg filmuhúðaðar töflur	DE/H/0109/001	960275	PFIZER APS	IS
Zarator 20 mg tuggutöflur	DE/H/0109/007	IS/1/10/081/03	PFIZER APS	IS
Zarator 40 mg tuggutöflur	DE/H/0109/008	IS/1/10/081/04	PFIZER APS	IS
Zarator 10 mg tuggutöflur	DE/H/0109/006	IS/1/10/081/02	PFIZER APS	IS
Zarator 5 mg tuggutöflur	DE/H/0109/005	IS/1/10/081/01	PFIZER APS	IS
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/032	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/020	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/072	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/035	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/002	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/014	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/024	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/044	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko	DE/H/0109/004	H/00/01447/061	PFIZER LUXEMBOURG SARL	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
obložene tablete				
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/066	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/033	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/071	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/076	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/027	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/062	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/023	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/001	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/042	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/043	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/049	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/019	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/047	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/050	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/007	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg žvečljive tablete	DE/H/0109/007	H/00/01447/079	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko	DE/H/0109/003	H/00/01447/040	PFIZER LUXEMBOURG SARL	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
obložene tablete				
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/012	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/073	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/037	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/015	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/053	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/025	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/029	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/057	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/052	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/036	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/058	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/026	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/060	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/022	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/068	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/038	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko	DE/H/0109/002	H/00/01447/034	PFIZER LUXEMBOURG SARL	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
obložene tablete				
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/003	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg žvečljive tablete	DE/H/0109/008	H/00/01447/080	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/046	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/045	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/008	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/056	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/075	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/009	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/018	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/041	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/004	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/005	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/059	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/048	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/039	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/055	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko	DE/H/0109/003	H/00/01447/051	PFIZER LUXEMBOURG SARL	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
obložene tablete				
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/030	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/013	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/063	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/017	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/064	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/065	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/021	PFIZER LUXEMBOURG SARL	SI
Sortis 5 mg žvečljive tablete	DE/H/0109/005	H/00/01447/077	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg žvečljive tablete	DE/H/0109/006	H/00/01447/078	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/011	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/067	PFIZER LUXEMBOURG SARL	SI
Sortis 40 mg filmsko obložene tablete	DE/H/0109/003	H/00/01447/054	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/028	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/074	PFIZER LUXEMBOURG SARL	SI
Sortis 20 mg filmsko obložene tablete	DE/H/0109/002	H/00/01447/031	PFIZER LUXEMBOURG SARL	SI
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/069	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko	DE/H/0109/001	H/00/01447/010	PFIZER LUXEMBOURG SARL	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
obložene tablete				
Sortis 80 mg filmsko obložene tablete	DE/H/0109/004	H/00/01447/070	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/016	PFIZER LUXEMBOURG SARL	SI
Sortis 10 mg filmsko obložene tablete	DE/H/0109/001	H/00/01447/006	PFIZER LUXEMBOURG SARL	SI
SORTIS 20 mg kramtomosios tabletės	DE/H/0109/007	LT/1/98/2147/003	PFIZER EUROPE MA EEIG	LT
LIPITOR 10, filmomhulde tabletten 10 mg	DE/H/0109/001	RVG 21081	PFIZER B.V.	NL
LIPITOR 40, filmomhulde tabletten 40 mg	DE/H/0109/003	RVG 21083	PFIZER B.V.	NL
Lipitor 40 mg, kauwtabletten	DE/H/0109/008	RVG 107610	PFIZER B.V.	NL
Lipitor 40 mg μασώμενα δισκία	DE/H/0109/008	34563/19-04-2016	PFIZER HELLAS, A.E.	GR
Lipitor 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/003	34558/19-04-2016	PFIZER HELLAS, A.E.	GR
Lipitor 10 mg μασώμενα δισκία	DE/H/0109/006	34561/19-04-2016	PFIZER HELLAS, A.E.	GR
Lipitor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/001	34556/19-04-2016	PFIZER HELLAS, A.E.	GR
СОРТИС 10 mg таблетки за дъвчене	DE/H/0109/006	20100549	PFIZER EUROPE MA EEIG	BG
СОРТИС 80 mg филмирани таблетки	DE/H/0109/004	20030461	PFIZER EUROPE MA EEIG	BG
СОРТИС 20 mg таблетки за дъвчене	DE/H/0109/007	20100550	PFIZER EUROPE MA EEIG	BG
СОРТИС 10 mg филмирани таблетки	DE/H/0109/001	9800143	PFIZER EUROPE MA EEIG	BG
СОРТИС 20 mg	DE/H/0109/002	9800142	PFIZER EUROPE MA EEIG	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
филмирани таблетки				
СОРТИС 40 mg таблетки за дъвчене	DE/H/0109/008	20100551	PFIZER EUROPE MA EEIG	BG
СОРТИС 5 mg таблетки за дъвчене	DE/H/0109/005	20100548	PFIZER EUROPE MA EEIG	BG
СОРТИС 40 mg филмирани таблетки	DE/H/0109/003	9800141	PFIZER EUROPE MA EEIG	BG
Sortis 10 mg comprimate masticabile	DE/H/0109/006	8189/2015/01	PFIZER EUROPE MA EEIG	RO
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/42	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/40	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/44	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/41	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/46	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/36	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/48	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/05	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/39	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/47	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/37	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/49	PFIZER KFT.	HU
SORTIS 20 mg	DE/H/0109/002	OGYI-T-6542/06	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmtabletta				
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/04	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/38	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/45	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/50	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/43	PFIZER KFT.	HU
SORTIS 20 mg filmtabletta	DE/H/0109/002	OGYI-T-6542/78	PFIZER KFT.	HU
Lipitor 40 mg, επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/003	19491	PFIZER HELLAS, A.E.	CY
Lipitor 10 mg, επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/001	19489	PFIZER HELLAS, A.E.	CY
Lipitor 40 mg μασώμενα δισκία	DE/H/0109/008	20730	PFIZER HELLAS, A.E.	CY
Lipitor 5 mg μασώμενα δισκία	DE/H/0109/005	20727	PFIZER HELLAS, A.E.	CY
Lipitor 20 mg, επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0109/002	19490	PFIZER HELLAS, A.E.	CY
Lipitor 20 mg μασώμενα δισκία	DE/H/0109/007	20729	PFIZER HELLAS, A.E.	CY
Lipitor 10 mg μασώμενα δισκία	DE/H/0109/006	20728	PFIZER HELLAS, A.E.	CY
SORTIS 40 mg rágótabletta	DE/H/0109/008	OGYI-T-6542/20	PFIZER KFT.	HU
SORTIS 5 mg rágótabletta	DE/H/0109/005	OGYI-T-6542/17	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SORTIS 20 mg rágótabletta	DE/H/0109/007	OGYI-T-6542/19	PFIZER KFT.	HU
Sortis 40 mg comprimate masticabile	DE/H/0109/008	8191/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/13	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/05	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/11	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/15	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/04	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/16	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/07	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/09	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/03	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/12	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/19	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/10	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/17	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/14	PFIZER EUROPE MA EEIG	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 80 mg comprimate filmate	DE/H/0109/004	8195/2015/08	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/18	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/02	PFIZER EUROPE MA EEIG	RO
Sortis 80 mg comprimate filmate	DE/H/0109/004	8195/2015/06	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate masticabile	DE/H/0109/007	8190/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 5 mg comprimate masticabile	DE/H/0109/005	8188/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/05	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/09	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/04	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/02	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/11	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/19	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/06	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/10	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/15	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/13	PFIZER EUROPE MA EEIG	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	8193/2015/03	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/12	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/18	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/17	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/16	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/07	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/14	PFIZER EUROPE MA EEIG	RO
Sortis 20 mg comprimate filmate	DE/H/0109/002	8193/2015/08	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/19	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/11	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/05	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/12	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/03	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/16	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/09	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/07	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/17	PFIZER EUROPE MA EEIG	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	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/06	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/13	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/15	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/04	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/01	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/10	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/14	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/18	PFIZER EUROPE MA EEIG	RO
Sortis 40 mg comprimate filmate	DE/H/0109/003	8194/2015/08	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/10	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/17	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/18	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/16	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/08	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/02	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/15	PFIZER EUROPE MA EEIG	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 10 mg comprimate filmate	DE/H/0109/001	8192/2015/12	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/11	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/07	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/04	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/14	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/06	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/13	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/19	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/09	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/05	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/03	PFIZER EUROPE MA EEIG	RO
Sortis 10 mg comprimate filmate	DE/H/0109/001	8192/2015/01	PFIZER EUROPE MA EEIG	RO
Zarator 20 mg comprimidos masticables	DE/H/0109/007	72601	PARKE-DAVIS, S.L.	ES
Zarator 10 mg comprimidos masticables	DE/H/0109/006	72599	PARKE-DAVIS, S.L.	ES
Zarator 20 mg comprimidos recubiertos con película	DE/H/0109/002	61.655	PARKE-DAVIS, S.L.	ES
Zarator 10 mg comprimidos recubiertos con película	DE/H/0109/001	61.654	PARKE-DAVIS, 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
Zarator 40 mg comprimidos recubiertos con película	DE/H/0109/003	61.656	PARKE-DAVIS, S.L.	ES
Zarator 80 mg comprimidos recubiertos con película	DE/H/0109/004	64.529	PARKE-DAVIS, S.L.	ES
Lipitor 20 mg chewable tablets	DE/H/0109/007	MA505/04007	PFIZER HELLAS, A.E.	MT
Lipitor 5 mg chewable tablets	DE/H/0109/005	MA505/04005	PFIZER HELLAS, A.E.	MT
Lipitor 10 mg chewable tablets	DE/H/0109/006	MA505/04006	PFIZER HELLAS, A.E.	MT
Lipitor 40 mg chewable tablets	DE/H/0109/008	MA505/04008	PFIZER HELLAS, A.E.	MT
Lipitor 80 mg film-coated tablets	DE/H/0109/004	MA505/04004	PFIZER HELLAS, A.E.	MT
Lipitor 10 mg film-coated tablets	DE/H/0109/001	MA505/04001	PFIZER HELLAS, A.E.	MT
Sortis® 20 mg Filmtabletten	DE/H/0109/002	1-21928	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sortis® 40 mg Filmtabletten	DE/H/0109/003	1-21926	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sortis® 10 mg Filmtabletten	DE/H/0109/001	1-21927	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Sortis® 80 mg Filmtabletten	DE/H/0109/004	1-24525	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
SORTIS 80 mg potahované tablety	DE/H/0109/004	31/397/03-C	PFIZER, SPOL. S R.O.	CZ
SORTIS 10 mg potahované tablety	DE/H/0109/001	31/233/99-C	PFIZER, SPOL. S R.O.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SORTIS 20 mg potahované tablety	DE/H/0109/002	31/234/99-C	PFIZER, SPOL. S R.O.	CZ
SORTIS 40 mg potahované tablety	DE/H/0109/003	31/235/99-C	PFIZER, SPOL. S R.O.	CZ
Lipitor 20 mg tyggetabletter	DE/H/0109/007	10-7963	PFIZER AS	NO
Lipitor 10 mg tyggetabletter	DE/H/0109/006	10-7962	PFIZER AS	NO
Lipitor 40 mg tyggetabletter	DE/H/0109/008	10-7964	PFIZER AS	NO
Lipitor 5 mg tyggetabletter	DE/H/0109/005	10-7961	PFIZER AS	NO
Atorvastatin Pfizer 5 mg Kautabletten	DE/H/0109/005	82571.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 20 mg Kautabletten	DE/H/0109/007	82573.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 10 mg Kautabletten	DE/H/0109/006	82572.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 40 mg Kautabletten	DE/H/0109/008	82574.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 80 mg Filmtabletten	DE/H/0109/004	38635.03.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 40 mg Filmtabletten	DE/H/0109/003	38635.02.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 20 mg Filmtabletten	DE/H/0109/002	38635.01.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin Pfizer 10 mg Filmtabletten	DE/H/0109/001	38635.00.00	PFIZER PHARMA PFE GMBH	DE
Lipitor 10 mg filmdrasjerte tabletter	DE/H/0109/001	96-3032	PFIZER AS	NO
Lipitor 20 mg filmdrasjerte tabletter	DE/H/0109/002	96-3033	PFIZER AS	NO
Lipitor 80 mg filmdrasjerte tabletter	DE/H/0109/004	00-7221	PFIZER AS	NO

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 40 mg filmdrasjerte tabletter	DE/H/0109/003	96-3034	PFIZER AS	NO
SORTIS 40 mg filmom obalené tablety	DE/H/0109/003	31/0414/11–S	PFIZER EUROPE MA EEIG	SK
SORTIS 10 mg filmom obalené tablety	DE/H/0109/001	31/0409/98–S	PFIZER EUROPE MA EEIG	SK
SORTIS 40 mg žuvacie tablety	DE/H/0109/008	31/0536/10-S	PFIZER EUROPE MA EEIG	SK
SORTIS 20 mg žuvacie tablety	DE/H/0109/007	31/0535/10-S	PFIZER EUROPE MA EEIG	SK
SORTIS 5 mg žuvacie tablety	DE/H/0109/005	31/0533/10-S	PFIZER EUROPE MA EEIG	SK
SORTIS 80 mg filmom obalené tablety	DE/H/0109/004	31/0018/03–S	PFIZER EUROPE MA EEIG	SK
SORTIS 20 mg filmom obalené tablety	DE/H/0109/002	31/0413/11–S	PFIZER EUROPE MA EEIG	SK
SORTIS 10 mg žuvacie tablety	DE/H/0109/006	31/0534/10-S	PFIZER EUROPE MA EEIG	SK
Lipitor 10 mg Kautabletten	DE/H/0109/006	BE375986	PFIZER S.A. (BELGIUM)	BE
Lipitor 5 mg Kautabletten	DE/H/0109/005	BE375977	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg kauwtabletten	DE/H/0109/006	BE375986	PFIZER S.A. (BELGIUM)	BE
Lipitor 5 mg kauwtabletten	DE/H/0109/005	BE375977	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg comprimés à croquer	DE/H/0109/006	BE375986	PFIZER S.A. (BELGIUM)	BE
Lipitor 5 mg comprimés à croquer	DE/H/0109/005	BE375977	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg Kautabletten	DE/H/0109/008	BE376004	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg comprimés à croquer	DE/H/0109/007	BE375995	PFIZER S.A. (BELGIUM)	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 20 mg kauwtabletten	DE/H/0109/007	BE375995	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg kauwtabletten	DE/H/0109/008	BE376004	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg comprimés à croquer	DE/H/0109/008	BE376004	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg Kautabletten	DE/H/0109/007	BE375995	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg comprimés à croquer	DE/H/0109/006	2010110025	PFIZER S.A. (BELGIUM)	LU
Lipitor 20 mg Kautabletten	DE/H/0109/007	2010110026	PFIZER S.A. (BELGIUM)	LU
Lipitor 10 mg Kautabletten	DE/H/0109/006	2010110025	PFIZER S.A. (BELGIUM)	LU
Lipitor 5 mg Kautabletten	DE/H/0109/005	2010110024	PFIZER S.A. (BELGIUM)	LU
Lipitor 20 mg comprimés à croquer	DE/H/0109/007	2010110026	PFIZER S.A. (BELGIUM)	LU
Lipitor 5 mg comprimés à croquer	DE/H/0109/005	2010110024	PFIZER S.A. (BELGIUM)	LU
Lipitor 40 mg comprimés à croquer	DE/H/0109/008	2010110027	PFIZER S.A. (BELGIUM)	LU
Lipitor 40 mg Kautabletten	DE/H/0109/008	2010110027	PFIZER S.A. (BELGIUM)	LU
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	2008019645	PFIZER S.A. (BELGIUM)	LU
Lipitor 10 mg Filmtabletten	DE/H/0109/001	2008019645	PFIZER S.A. (BELGIUM)	LU
Lipitor 20 mg Filmtabletten	DE/H/0109/002	2008019646	PFIZER S.A. (BELGIUM)	LU
Lipitor 40 mg filmomhulde tabletten	DE/H/0109/003	BE307745	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	2008019646	PFIZER S.A. (BELGIUM)	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
Lipitor 40 mg Filmtabletten	DE/H/0109/003	2008019647	PFIZER S.A. (BELGIUM)	LU
Lipitor 10 mg filmomhulde tabletten	DE/H/0109/001	BE184082	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg filmomhulde tabletten	DE/H/0109/004	BE232933	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	2008019647	PFIZER S.A. (BELGIUM)	LU
Lipitor 80 mg Filmtabletten	DE/H/0109/004	2008019648	PFIZER S.A. (BELGIUM)	LU
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	2008019648	PFIZER S.A. (BELGIUM)	LU
Lipitor 20 mg filmomhulde tabletten	DE/H/0109/002	BE307736	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg filmomhulde tabletten	DE/H/0109/001	BE307727	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg filmomhulde tabletten	DE/H/0109/004	BE307754	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg filmomhulde tabletten	DE/H/0109/002	BE184073	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg filmomhulde tabletten	DE/H/0109/003	BE184064	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg Filmtabletten	DE/H/0109/001	BE184082	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg Filmtabletten	DE/H/0109/003	BE184064	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	BE307736	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	BE232933	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg Filmtabletten	DE/H/0109/002	BE184073	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg Filmtabletten	DE/H/0109/002	BE307736	PFIZER S.A. (BELGIUM)	BE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 10 mg Filmtabletten	DE/H/0109/001	BE307727	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg Filmtabletten	DE/H/0109/004	BE232933	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	BE184064	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg comprimés pelliculés	DE/H/0109/004	BE307754	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	BE184082	PFIZER S.A. (BELGIUM)	BE
Lipitor 20 mg comprimés pelliculés	DE/H/0109/002	BE184073	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg comprimés pelliculés	DE/H/0109/003	BE307745	PFIZER S.A. (BELGIUM)	BE
Lipitor 10 mg comprimés pelliculés	DE/H/0109/001	BE307727	PFIZER S.A. (BELGIUM)	BE
Lipitor 40 mg Filmtabletten	DE/H/0109/003	BE307745	PFIZER S.A. (BELGIUM)	BE
Lipitor 80 mg Filmtabletten	DE/H/0109/004	BE307754	PFIZER S.A. (BELGIUM)	BE
Zarator 5 mg comprimidos para mastigar	DE/H/0109/005	5319504	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos para mastigar	DE/H/0109/006	5319512	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos para mastigar	DE/H/0109/008	5319538	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos para mastigar	DE/H/0109/007	5319520	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos	DE/H/0109/004	4021481	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película				
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021184	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020285	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020582	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020681	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021085	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020780	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	5047360	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020889	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020483	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021382	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020186	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020384	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4020988	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 80 mg comprimidos revestidos por película	DE/H/0109/004	4021283	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4054185	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4054284	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053989	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052486	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052684	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053187	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052882	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	2535987	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg	DE/H/0109/003	4054086	LABORATÓRIOS PFIZER,	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
comprimidos revestidos por película			LDA.	
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052288	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053781	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053385	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052387	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053088	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053682	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	2535888	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053583	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052585	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053880	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos	DE/H/0109/002	3538980	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
por película				
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053286	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052981	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052783	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 40 mg comprimidos revestidos por película	DE/H/0109/003	4053484	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051686	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050688	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051587	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051280	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051389	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050985	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051488	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051181	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4051983	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052080	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051082	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050886	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4051884	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	2535680	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4050787	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	2535789	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 10 mg comprimidos revestidos por película	DE/H/0109/001	4051785	LABORATÓRIOS PFIZER, LDA.	PT
Zarator 20 mg comprimidos revestidos por película	DE/H/0109/002	4052189	LABORATÓRIOS PFIZER, LDA.	PT
TAHOR 20 mg, comprimé	DE/H/0109/007	34009 494 588 4 4	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
à croquer				
TAHOR 10 mg, comprimé à croquer	DE/H/0109/006	34009 494 587 8 3	PFIZER HOLDING FRANCE	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 343 069 8 3	PFIZER HOLDING FRANCE	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 560 305 1 4	PFIZER HOLDING FRANCE	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 218 984 6 0	PFIZER HOLDING FRANCE	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 560 306 8 2	PFIZER HOLDING FRANCE	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 560 303 9 2	PFIZER HOLDING FRANCE	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 218 985 2 1	PFIZER HOLDING FRANCE	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 343 068 1 5	PFIZER HOLDING FRANCE	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 371 994 4 5	PFIZER HOLDING FRANCE	FR
TAHOR 40 mg, comprimé pelliculé	DE/H/0109/003	34009 560 307 4 3	PFIZER HOLDING FRANCE	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 371 993 8 4	PFIZER HOLDING FRANCE	FR
TAHOR 20 mg, comprimé pelliculé	DE/H/0109/002	34009 560 304 5 3	PFIZER HOLDING FRANCE	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 343 067 5 4	PFIZER HOLDING FRANCE	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 218 982 3 1	PFIZER HOLDING FRANCE	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 560 302 2 4	PFIZER HOLDING FRANCE	FR
TAHOR 10 mg, comprimé pelliculé	DE/H/0109/001	34009 371 992 1 6	PFIZER HOLDING FRANCE	FR
TAHOR 80 mg, comprimé	DE/H/0109/004	34009 371 995 0 6	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pelliculé				
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 575 0 6	PFIZER HOLDING FRANCE	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 356 335 3 8	PFIZER HOLDING FRANCE	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 576 7 4	PFIZER HOLDING FRANCE	FR
TAHOR 80 mg, comprimé pelliculé	DE/H/0109/004	34009 355 577 3 5	PFIZER HOLDING FRANCE	FR
Sortis 10 mg apvalkotās tabletes	DE/H/0109/001	98-0598	PFIZER LIMITED	LV
Sortis 80 mg apvalkotās tabletes	DE/H/0109/004	03-0502	PFIZER LIMITED	LV
Sortis 40 mg apvalkotās tabletes	DE/H/0109/003	98-0600	PFIZER LIMITED	LV
Sortis 20 mg apvalkotās tabletes	DE/H/0109/002	98-0599	PFIZER LIMITED	LV
Sortis 5 mg košļājamās tabletes	DE/H/0109/005	10-0402	PFIZER LIMITED	LV
Sortis 40 mg košļājamās tabletes	DE/H/0109/008	10-0405	PFIZER LIMITED	LV
Sortis 10 mg košļājamās tabletes	DE/H/0109/006	10-0403	PFIZER LIMITED	LV
Sortis 20 mg košļājamās tabletes	DE/H/0109/007	10-0404	PFIZER LIMITED	LV
SORTIS, 10 mg närimistabletid	DE/H/0109/006	694510	PFIZER EUROPE MA EEIG	EE
SORTIS, 5 mg närimistabletid	DE/H/0109/005	694410	PFIZER EUROPE MA EEIG	EE
SORTIS, 20 mg närimistabletid	DE/H/0109/007	694610	PFIZER EUROPE MA EEIG	EE
SORTIS, 40 mg närimistabletid	DE/H/0109/008	694710	PFIZER EUROPE MA EEIG	EE
SORTIS 20, 20 mg,	DE/H/0109/007	17435	PFIZER EUROPE MA EEIG	PL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletki do rozgryzania i żucia				
SORTIS 80, 80 mg, tabletki powlekane	DE/H/0109/004	10188	PFIZER POLSKA SP. Z O.O.	PL
SORTIS 40, 40 mg, tabletki powlekane	DE/H/0109/003	7848	PFIZER EUROPE MA EEIG	PL
SORTIS 10, 10 mg, tabletki do rozgryzania i żucia	DE/H/0109/006	17436	PFIZER EUROPE MA EEIG	PL
SORTIS 10, 10 mg, tabletki powlekane	DE/H/0109/001	7846	PFIZER EUROPE MA EEIG	PL
SORTIS 20, 20 mg, tabletki powlekane	DE/H/0109/002	7847	PFIZER EUROPE MA EEIG	PL
SORTIS, 20 mg õhukese polümeerikattega tabletid	DE/H/0109/002	205798	PFIZER EUROPE MA EEIG	EE
SORTIS, 40 mg õhukese polümeerikattega tabletid	DE/H/0109/003	205898	PFIZER EUROPE MA EEIG	EE
SORTIS, 10 mg õhukese polümeerikattega tabletid	DE/H/0109/001	205698	PFIZER EUROPE MA EEIG	EE
SORTIS 80 mg, 80 mg õhukese polümeerikattega tabletid	DE/H/0109/004	423003	PFIZER EUROPE MA EEIG	EE
Lipitor 10 mg chewable tablets	DE/H/0109/006	PL 39933/0006	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 5 mg chewable tablets	DE/H/0109/005	PL 39933/0005	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 20 mg chewable tablets	DE/H/0109/007	PL 39933/0007	PFIZER IRELAND PHARMACEUTICALS	UK
Lipitor 40 mg film-coated tablets	DE/H/0109/003	PL 39933/0003	PFIZER IRELAND PHARMACEUTICALS	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lipitor 40 mg chewable tablets	DE/H/0109/008	PL 39933/0008	PFIZER IRELAND PHARMACEUTICALS	UK
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 30 mg filmsko obložene tablete	AT/H/0305/005	H/10/00236/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/025	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/034	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/035	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/036	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/038	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/037	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Atorvastatin Lek 60 mg filmsko obložene tablete	AT/H/0305/006	H/10/00236/039	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
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

Product 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 Sandoz 30 mg – Filmtabletten	AT/H/0305/005	1-30704	SANDOZ GMBH	AT
Atorasat 30 mg filmomhulde tabletten	AT/H/0305/005	BE434332	SANDOZ N.V.	BE
Atorasat 60 mg filmomhulde tabletten	AT/H/0305/006	BE434341	SANDOZ N.V.	BE
Atorvastatin 10 mg film-coated tablets	not available	PL 43461/0017	FLAMINGO PHARMA UK LTD	UK
Atorvastatin 20 mg film-coated tablets	not available	PL 43461/0018	FLAMINGO PHARMA UK LTD	UK
Atorvastatin 40 mg film-coated tablets	not available	PL 43461/0019	FLAMINGO PHARMA UK LTD	UK
Atorvastatin 80 mg film-coated tablets	not available	PL 43461/0020	FLAMINGO PHARMA UK LTD	UK
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 επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	14475/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin/Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	14478/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin Krka 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	21287	KRKA, D.D., NOVO MESTO	CY
Atorvastatin Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	21288	KRKA, D.D., NOVO MESTO	CY
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/01	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/01	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
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	040561449	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561452	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561553	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561565	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561589	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561603	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561678	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561680	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561437	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561464	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561476	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561490	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561502	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561514	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561526	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	040561540	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561577	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561591	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561615	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561627	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561666	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561704	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561488	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561538	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561639	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561641	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561654	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561692	KRKA, D.D., NOVO MESTO	IT
Atorvastatin Krka 30 mg, filmomhulde tabletten	SE/H/0642/005	BE398036	KRKA, D.D., NOVO MESTO	BE
Atorvastatin Krka 60 mg, filmomhulde tabletten	SE/H/0642/006	BE398045	KRKA, D.D., NOVO MESTO	BE
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/03	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 30 mg comprimate filmate	SE/H/0642/005	3796/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/07	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/11	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/14	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/03	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/07	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	3797/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/11	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/14	KRKA, D.D., NOVO MESTO	RO
Atorvastatin 30 mg film-coated tablets	NL/H/2982/005	PL 16363/0512	MILPHARM LIMITED	UK
Atorvastatin 60 mg film-coated tablets	NL/H/2982/006	PL 16363/0513	MILPHARM LIMITED	UK
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
Atorvastatina Pharmacia 10 mg comprimidos recubiertos con película EFG	DE/H/3385/001	69.555	PHARMACIA NOSTRUM, S.A.	ES
Atorvastatina Pharmacia 80 mg comprimidos recubiertos con película EFG	DE/H/3385/004	69.566	PHARMACIA NOSTRUM, S.A.	ES
Atorvastatina Pharmacia 40 mg comprimidos	DE/H/3385/003	69.565	PHARMACIA NOSTRUM, 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
recubiertos con película EFG				
Atorvastatina Pharmacia 20 mg comprimidos recubiertos con película EFG	DE/H/3385/002	69.564	PHARMACIA NOSTRUM, S.A.	ES
Orbeos 80 mg filmdragerade tabletter	DE/H/3385/004	24874	PFIZER OY	FI
Orbeos 10 mg filmdragerade tabletter	DE/H/3385/001	24871	PFIZER OY	FI
Orbeos 20 mg filmdragerade tabletter	DE/H/3385/002	24872	PFIZER OY	FI
Orbeos 40 mg filmdragerade tabletter	DE/H/3385/003	24873	PFIZER OY	FI
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/71	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/10	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/26	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/01	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/27	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/77	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/76	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/25	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/15	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/22	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/34	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/73	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/31	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/28	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/08	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/09	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/30	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/33	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/79	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/72	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/69	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/23	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/36	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/80	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/66	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/07	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/74	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/04	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/78	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/81	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/70	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/68	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/32	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/21	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/82	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/20	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/67	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/16	PFIZER KFT.	HU
OBRADON 80 mg filmtabletta	DE/H/3385/004	OGYI-T-8306/75	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/24	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/19	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/29	PFIZER KFT.	HU
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/17	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/35	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OBRADON 10 mg filmtabletta	DE/H/3385/001	OGYI-T-8306/18	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/37	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/38	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/06	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/43	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/51	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/39	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/44	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/58	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/54	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/53	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/50	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/63	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/55	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/40	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/52	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/57	PFIZER KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/60	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/45	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/41	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/49	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/47	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/42	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/64	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/48	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/56	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/65	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/59	PFIZER KFT.	HU
OBRADON 20 mg filmtabletta	DE/H/3385/002	OGYI-T-8306/46	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/05	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/62	PFIZER KFT.	HU
OBRADON 40 mg filmtabletta	DE/H/3385/003	OGYI-T-8306/61	PFIZER KFT.	HU
Liprimar 40 mg Filmtabletten	DE/H/3385/003	39587.02.00	PFIZER PHARMA PFE GMBH	DE
Liprimar 80 mg Filmtabletten	DE/H/3385/004	77656.00.00	PFIZER PHARMA PFE 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 10 mg Filmtabletten	DE/H/3385/001	39587.00.00	PFIZER PHARMA PFE GMBH	DE
Liprimar 20 mg Filmtabletten	DE/H/3385/002	39587.01.00	PFIZER PHARMA PFE GMBH	DE
TORVAST 20 mg compresse rivestite con film	DE/H/3385/002	033007081	PFIZER LIMITED	IT
TORVAST 20 mg compresse rivestite con film	DE/H/3385/002	033007030	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007321	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007257	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007360	PFIZER LIMITED	IT
Orbeos 40 mg kalvopäällysteiset tabletit	DE/H/3385/003	24873	PFIZER OY	FI
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007244	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007333	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007271	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007345	PFIZER LIMITED	IT
TORVAST 80 mg	DE/H/3385/004	033007269	PFIZER LIMITED	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				
Orbeos 80 mg kalvopäällysteiset tabletit	DE/H/3385/004	24874	PFIZER OY	FI
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007283	PFIZER LIMITED	IT
Orbeos 20 mg kalvopäällysteiset tabletit	DE/H/3385/002	24872	PFIZER OY	FI
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007384	PFIZER LIMITED	IT
Orbeos 10 mg kalvopäällysteiset tabletit	DE/H/3385/001	24871	PFIZER OY	FI
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007319	PFIZER LIMITED	IT
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007079	PFIZER LIMITED	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007055	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007307	PFIZER LIMITED	IT
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007028	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007372	PFIZER LIMITED	IT
TORVAST 20 mg compresse rivestite con	DE/H/3385/002	033007042	PFIZER LIMITED	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
film				
TORVAST 10 mg compresse rivestite con film	DE/H/3385/001	033007016	PFIZER LIMITED	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007067	PFIZER LIMITED	IT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007358	PFIZER LIMITED	IT
TORVAST 40 mg compresse rivestite con film	DE/H/3385/003	033007093	PFIZER LIMITED	IT
Atorvastatina Parke-Davis 20 mg comprimidos revestidos por película	DE/H/3385/002	5129630	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 10 mg comprimidos revestidos por película	DE/H/3385/001	5129564	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 40 mg comprimidos revestidos por película	DE/H/3385/003	5129648	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
TORVAST 80 mg compresse rivestite con film	DE/H/3385/004	033007295	PFIZER LIMITED	IT
Atorvastatina Parke-Davis 40 mg comprimidos revestidos por película	DE/H/3385/003	5129655	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 10 mg	DE/H/3385/001	5129556	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película				
Atorvastatina Parke-Davis 20 mg comprimidos revestidos por película	DE/H/3385/002	5129614	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 20 mg comprimidos revestidos por película	DE/H/3385/002	5129622	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 40 mg comprimidos revestidos por película	DE/H/3385/003	5129663	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 10 mg comprimidos revestidos por película	DE/H/3385/001	5129606	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
Atorvastatina Parke-Davis 10 mg comprimidos revestidos por película	DE/H/3385/001	5129572	PARKE-DAVIS, PRODUTOS FARMACÊUTICOS, LDA.	PT
ZARATOR 40 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23960/3-4-2015	WIN MEDICA SA	GR
ZARATOR 20 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23959/3-4-2015	WIN MEDICA SA	GR
ZARATOR 10 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23958/3-4-2015	WIN MEDICA SA	GR
Atoris, 30 mg, tabletki powlekane	SE/H/1000/001	18462	KRKA - POLSKA SP. Z O.O.	PL
Atoris, 60 mg, tabletki	SE/H/1000/002	18463	KRKA - POLSKA SP. Z O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
powlekane				
Tinator 30mg filmdragerade tabletter	SE/H/1000/001	44184	KRKA, D.D., NOVO MESTO	SE
Tinator 60mg filmdragerade tabletter	SE/H/1000/002	44185	KRKA, D.D., NOVO MESTO	SE
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/004	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/018	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/005	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/006	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/007	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/008	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/009	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/010	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/011	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/012	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/013	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/014	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/015	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele dengtos tabletės	SE/H/1000/001	LT/1/03/0865/016	KRKA, D.D., NOVO MESTO	LT
Atoris 30 mg plėvele	SE/H/1000/001	LT/1/03/0865/017	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
dengtos tabletės				
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/019	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/020	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/021	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/022	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/023	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/024	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/025	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/026	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/027	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/028	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/029	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/030	KRKA, D.D., NOVO MESTO	LT
Atoris 60 mg plėvele dengtos tabletės	SE/H/1000/002	LT/1/03/0865/031	KRKA, D.D., NOVO MESTO	LT
Tulip 30 mg filmsko obložene tablete	not available	H/01/01575/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
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
Atorvastatina Tevagen	not available	77291	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
60 mg comprimidos recubiertos con película				
Atorvastatina Tevagen 30 mg comprimidos recubiertos con película	not available	77292	TEVA PHARMA S.L.U	ES
Atorvastatin-CT 30 mg Filmtabletten	not available	88208.00.00	ABZ-PHARMA GMBH	DE
Astator 60 mg apvalkotās tabletes	SE/H/0998/002	11-0206	KRKA, D.D., NOVO MESTO	LV
Atorvastatin Krka, 30 mg, tabletki powlekane	SE/H/0998/001	18465	KRKA, D.D., NOVO MESTO	PL
Atorvastatin Krka, 60 mg, tabletki powlekane	SE/H/0998/002	18466	KRKA, D.D., NOVO MESTO	PL
Астатор 30 mg филмирани таблетки	SE/H/0998/001	20110264	KRKA, D.D., NOVO MESTO	BG
Астатор 60 mg филмирани таблетки	SE/H/0998/002	20110265	KRKA, D.D., NOVO MESTO	BG
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/01	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/02	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/03	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/04	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/05	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/06	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/07	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/08	TAD PHARMA GMBH	RO
Statorva 30 mg	SE/H/0998/001	3799/2011/09	TAD PHARMA GMBH	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
comprimate filmate				
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/10	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/11	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/12	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/13	TAD PHARMA GMBH	RO
Statorva 30 mg comprimate filmate	SE/H/0998/001	3799/2011/14	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/01	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/02	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/03	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/04	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/05	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/06	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/07	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/08	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/09	TAD PHARMA GMBH	RO
Statorva 60mg comprimate filmate	SE/H/0998/002	3800/2011/10	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/11	TAD PHARMA GMBH	RO
Statorva 60 mg	SE/H/0998/002	3800/2011/12	TAD PHARMA GMBH	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
comprimate filmate				
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/13	TAD PHARMA GMBH	RO
Statorva 60 mg comprimate filmate	SE/H/0998/002	3800/2011/14	TAD PHARMA GMBH	RO
Astator 30 mg filmdragerad tablett	SE/H/0998/001	44175	KRKA, D.D., NOVO MESTO	SE
Astator 60 mg filmdragerad tablett	SE/H/0998/002	44176	KRKA, D.D., NOVO MESTO	SE
Astator, 30 mg õhukese polümeerikattega tabletid	SE/H/0998/001	742311	KRKA, D.D., NOVO MESTO	EE
Astator, 60 mg õhukese polümeerikattega tabletid	SE/H/0998/002	742511	KRKA, D.D., NOVO MESTO	EE
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/001	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/002	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/003	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/004	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/005	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/006	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/007	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/008	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/009	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele	SE/H/0998/001	LT/1/11/2450/010	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
dengtos tabletės				
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/011	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/012	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/013	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg plėvele dengtos tabletės	SE/H/0998/001	LT/1/11/2450/014	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/015	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/016	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/017	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/018	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/019	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/020	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/021	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/022	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/023	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/024	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/025	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/026	KRKA, D.D., NOVO MESTO	LT
Astator 60 mg plėvele	SE/H/0998/002	LT/1/11/2450/027	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
dengtos tabletės				
Astator 60 mg plėvele dengtos tabletės	SE/H/0998/002	LT/1/11/2450/028	KRKA, D.D., NOVO MESTO	LT
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/01	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/02	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/03	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/04	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/05	KRKA, D.D., NOVO MESTO	HU
Astator 30 mg filmtabletta	SE/H/0998/001	OGYI-T-21813/06	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/07	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/08	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/09	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/10	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/11	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmtabletta	SE/H/0998/002	OGYI-T-21813/12	KRKA, D.D., NOVO MESTO	HU
Astator 60 mg filmom obalené tablety	SE/H/0998/002	31/0633/11-S	KRKA, D.D., NOVO MESTO	SK
Astator 30 mg filmom obalené tablety	SE/H/0998/001	31/0632/11-S	KRKA, D.D., NOVO MESTO	SK
Astator 30 mg apvalkotās tabletes	SE/H/0998/001	11-0205	KRKA, D.D., NOVO MESTO	LV
Astator 30 mg filmsko	SE/H/0998/001	H/11/00226/001	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
obložene tablete				
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/002	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/003	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/004	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/005	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/006	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/007	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/008	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/009	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/010	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/011	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/012	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/013	KRKA, D.D., NOVO MESTO	SI
Astator 30 mg filmsko obložene tablete	SE/H/0998/001	H/11/00226/014	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/015	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/016	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/017	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko	SE/H/0998/002	H/11/00226/018	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
obložene tablete				
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/019	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/020	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/021	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/022	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/023	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/024	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/025	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/026	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/027	KRKA, D.D., NOVO MESTO	SI
Astator 60 mg filmsko obložene tablete	SE/H/0998/002	H/11/00226/028	KRKA, D.D., NOVO MESTO	SI
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
Atorvastatine DSM Sinochem 30 mg, filmomhulde tabletten	NL/H/3798/003	RVG 119408	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine DSM Sinochem 60 mg, filmomhulde tabletten	NL/H/3798/005	RVG 119410	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatin Krka 30 mg	SE/H/0642/005	1-30807	KRKA, D.D., NOVO MESTO	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
Filmtabletten				
Atorvastatin Krka 60 mg Filmtabletten	SE/H/0642/006	1-30808	KRKA, D.D., NOVO MESTO	AT
Atorvastatin/Krka 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	14475/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin/Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	14478/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin Krka 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	21287	KRKA, D.D., NOVO MESTO	CY
Atorvastatin Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	21288	KRKA, D.D., NOVO MESTO	CY
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/01	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/01	KRKA, D.D., NOVO MESTO	RO
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	040561449	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561452	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561553	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561565	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561589	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse	SE/H/0642/006	040561603	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
rivestite con film				
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561678	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561680	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561437	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561464	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561476	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561490	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561502	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561514	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561526	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561540	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561577	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561591	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561615	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561627	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561666	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561704	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse	SE/H/0642/005	040561488	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
rivestite con film				
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561538	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561639	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561641	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561654	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561692	KRKA, D.D., NOVO MESTO	IT
Atorvastatin Krka 30 mg, filmomhulde tabletten	SE/H/0642/005	BE398036	KRKA, D.D., NOVO MESTO	BE
Atorvastatin Krka 60 mg, filmomhulde tabletten	SE/H/0642/006	BE398045	KRKA, D.D., NOVO MESTO	BE
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/03	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/07	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate	SE/H/0642/005	3796/2011/11	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
filmate				
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/14	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/03	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/07	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/11	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/14	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg filmsko	not available	H/01/00234/013	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
obložene tablete				
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
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
Orbeos 5 mg tuggtabletter	DE/H/3616/001	28939	PFIZER OY	FI
Orbeos 10 mg tuggtabletter	DE/H/3616/002	28938	PFIZER OY	FI
Orbeos 40 mg tuggtabletter	DE/H/3616/004	28936	PFIZER OY	FI
Orbeos 20 mg tuggtabletter	DE/H/3616/003	28937	PFIZER OY	FI
OBRADON 10 mg rágótabletta	DE/H/3616/002	OGYI-T-8306/12	PFIZER KFT.	HU
OBRADON 20 mg rágótabletta	DE/H/3616/003	OGYI-T-8306/13	PFIZER KFT.	HU
OBRADON 5 mg rágótabletta	DE/H/3616/001	OGYI-T-8306/11	PFIZER KFT.	HU
OBRADON 40 mg rágótabletta	DE/H/3616/004	OGYI-T-8306/14	PFIZER KFT.	HU
TORVAST 5 mg compresse masticabili	DE/H/3616/001	033007396	PFIZER LIMITED	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
TORVAST 40 mg compresse masticabili	DE/H/3616/004	033007422	PFIZER LIMITED	IT
TORVAST 20 mg compresse masticabili	DE/H/3616/003	033007410	PFIZER LIMITED	IT
Liprimar® 5 mg Kautabletten	DE/H/3616/001	86865.00.00	PFIZER PHARMA PFE GMBH	DE
Liprimar® 20 mg Kautabletten	DE/H/3616/003	86867.00.00	PFIZER PHARMA PFE GMBH	DE
Liprimar® 10 mg Kautabletten	DE/H/3616/002	86866.00.00	PFIZER PHARMA PFE GMBH	DE
Liprimar® 40 mg Kautabletten	DE/H/3616/004	86868.00.00	PFIZER PHARMA PFE GMBH	DE
Orbeos 10 mg purutabletit	DE/H/3616/002	28938	PFIZER OY	FI
Orbeos 20 mg purutabletit	DE/H/3616/003	28937	PFIZER OY	FI
TORVAST 10 mg compresse masticabili	DE/H/3616/002	033007408	PFIZER LIMITED	IT
Orbeos 5 mg purutabletit	DE/H/3616/001	28939	PFIZER OY	FI
Orbeos 40 mg purutabletit	DE/H/3616/004	28936	PFIZER OY	FI
Atorvastatin 10 mg Film-coated Tablets	not available	PL 36687/0258	TORRENT PHARMA (UK) LTD.	UK
Atorvastatin 20 mg Film-coated Tablets	not available	PL 36687/0259	TORRENT PHARMA (UK) LTD.	UK
Atorvastatin 40 mg Film-coated Tablets	not available	PL 36687/0260	TORRENT PHARMA (UK) LTD.	UK
Zarator 5 mg μασώμενα δισκία	DE/H/3616/001	83315/19-10-2017	WIN MEDICA SA	GR
Zarator 10 mg μασώμενα δισκία	DE/H/3616/002	83316/19-10-2017	WIN MEDICA SA	GR
Zarator 20 mg μασώμενα δισκία	DE/H/3616/003	83317/19-10-2017	WIN MEDICA SA	GR
Zarator 40 mg μασώμενα	DE/H/3616/004	32533/16/19-10-2017	WIN MEDICA SA	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
δισκία				
Zarator 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/001	94839/ 20-10-2017	WIN MEDICA SA	GR
Zarator 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/002	94840/20-10-2017	WIN MEDICA SA	GR
Zarator 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/3385/003	94841/20-10-2017	WIN MEDICA SA	GR
Prevencor 10 mg comprimidos recubiertos con película	DE/H/3820/001	61.742	PFIZER GEP, S.L.	ES
Prevencor 20 mg comprimidos recubiertos con película	DE/H/3820/002	61.743	PFIZER GEP, S.L.	ES
Prevencor 80 mg comprimidos recubiertos con película	DE/H/3820/004	64.642	PFIZER GEP, S.L.	ES
Prevencor 40 mg comprimidos recubiertos con película	DE/H/3820/003	61.744	PFIZER GEP, S.L.	ES
ATOR Pfizer 80 mg Filmtabletten	DE/H/3820/004	87747.00.00	PFIZER PHARMA PFE GMBH	DE
ATOR Pfizer 20 mg Filmtabletten	DE/H/3820/002	87745.00.00	PFIZER PHARMA PFE GMBH	DE
ATOR Pfizer 40 mg Filmtabletten	DE/H/3820/003	87746.00.00	PFIZER PHARMA PFE GMBH	DE
ATOR Pfizer 10 mg Filmtabletten	DE/H/3820/001	87744.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 5 mg Kautabletten	DE/H/3882/005	87756.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 10 mg Kautabletten	DE/H/3882/006	87757.00.00	PFIZER PHARMA PFE 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
Atorvasa 20 mg Filmtabletten	DE/H/3882/002	87753.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 20 mg Kautabletten	DE/H/3882/007	87758.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 40 mg Kautabletten	DE/H/3882/008	87759.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 80 mg Filmtabletten	DE/H/3882/004	87755.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 40 mg Filmtabletten	DE/H/3882/003	87754.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvasa 10 mg Filmtabletten	DE/H/3882/001	87752.00.00	PFIZER PHARMA PFE GMBH	DE
Cardyl 10 mg comprimidos recubiertos con película	DE/H/3882/001	61.716	PFIZER, S.L.	ES
Cardyl 20 mg comprimidos recubiertos con película	DE/H/3882/002	61.717	PFIZER, S.L.	ES
Cardyl 40 mg comprimidos recubiertos con película	DE/H/3882/003	61.718	PFIZER, S.L.	ES
Cardyl 80 mg comprimidos recubiertos con película	DE/H/3882/004	64.571	PFIZER, S.L.	ES
Lambrinex 80 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DK/H/1840/004	21003	PHARMATHEN S.A.	CY
Atorvastatin 10 mg film-coated tablets	not available	PL 17907/0547	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Atorvastatin 20 mg film-coated tablets	not available	PL 17907/0548	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Atorvastatin 40 mg film-coated tablets	not available	PL 17907/0549	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Atorvastatin 80 mg film-	not available	PL 17907/0550	BRISTOL LABORATORIES	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
coated tablets			LTD (BERKHAMSTED)	
ZARATOR 40 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23960/3-4-2015	WIN MEDICA SA	GR
ZARATOR 20 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23959/3-4-2015	WIN MEDICA SA	GR
ZARATOR 10 mg, επικαλυμμένα με λεπτό υμένιο δισκία	not available	23958/3-4-2015	WIN MEDICA SA	GR
SORTIS 40 mg tablete	not available	UP/I-530-09/11-02/300	PFIZER CROATIA D.O.O.	HR
SORTIS 10 mg tablete	not available	UP/I-530-09/11-02/298	PFIZER CROATIA D.O.O.	HR
SORTIS 20 mg tablete	not available	UP/I-530-09/11-02/299	PFIZER CROATIA D.O.O.	HR
SORTIS 80 mg tablete	not available	UP/I-530-09/11-02/301	PFIZER CROATIA D.O.O.	HR
Sortis® 80 mg Filmtabletten	DE/H/3384/004	77655.00.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 20 mg Filmtabletten	DE/H/3384/002	39581.01.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 40 mg Filmtabletten	DE/H/3384/003	39581.02.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 10 mg Filmtabletten	DE/H/3384/001	39581.00.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 10 mg Kautabletten	DE/H/3384/006	82883.00.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 5 mg Kautabletten	DE/H/3384/005	82882.00.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 20 mg Kautabletten	DE/H/3384/007	82884.00.00	PFIZER PHARMA PFE GMBH	DE
Sortis® 40 mg Kautabletten	DE/H/3384/008	82886.00.00	PFIZER PHARMA PFE GMBH	DE
Atorvastatin 80 mg film-coated tablets	not available	PL 36687/0261	TORRENT PHARMA (UK) LTD.	UK
Atorvastatin/Sandoz 30 mg επικαλυμμένα με	AT/H/0196/003	41953/19-6-2015	SANDOZ GMBH	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
λεπτό υμένιο δισκία				
Atorvadivid 30 mg - Filmtabletten	AT/H/0196/003	1-28644	HEXAL PHARMA GMBH	AT
Atorvastatin Sandoz 30mg	AT/H/0196/003	20986	SANDOZ GMBH	CY
Atorvastatin 10mg Film-coated Tablets	not available	PL 20075/0551	ACCORD HEALTHCARE LIMITED	UK
Atorvastatin 20mg Film-coated Tablets	not available	PL 20075/0552	ACCORD HEALTHCARE LIMITED	UK
Atorvastatin 40mg Film-coated Tablets	not available	PL 20075/0553	ACCORD HEALTHCARE LIMITED	UK
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/005	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/006	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/007	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/008	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/009	KRKA, D.D., NOVO MESTO	SI
Atoris 30 mg filmsko obložene tablete	NP/H/0173/004	H/01/00234/010	KRKA, D.D., NOVO MESTO	SI
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 επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/005	14475/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin/Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	14478/24-2-2012	KRKA, D.D., NOVO MESTO	GR
Atorvastatin Krka 30 mg	SE/H/0642/005	21287	KRKA, D.D., NOVO MESTO	CY

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
επικαλυμμένα με λεπτό υμένιο δισκία				
Atorvastatin Krka 60 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0642/006	21288	KRKA, D.D., NOVO MESTO	CY
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/01	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/01	KRKA, D.D., NOVO MESTO	RO
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	040561449	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561452	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561553	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561565	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561589	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561603	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561678	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561680	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561437	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561464	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse	SE/H/0642/005	040561476	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
rivestite con film				
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561490	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561502	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561514	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561526	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561540	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561577	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561591	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561615	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561627	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561666	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561704	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561488	KRKA, D.D., NOVO MESTO	IT
Atoris 30 mg compresse rivestite con film	SE/H/0642/005	040561538	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561639	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561641	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse rivestite con film	SE/H/0642/006	040561654	KRKA, D.D., NOVO MESTO	IT
Atoris 60 mg compresse	SE/H/0642/006	040561692	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
rivestite con film				
Atorvastatin Krka 30 mg, filmomhulde tabletten	SE/H/0642/005	BE398036	KRKA, D.D., NOVO MESTO	BE
Atorvastatin Krka 60 mg, filmomhulde tabletten	SE/H/0642/006	BE398045	KRKA, D.D., NOVO MESTO	BE
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/03	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/07	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/11	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 30 mg comprimate filmate	SE/H/0642/005	3796/2011/14	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/02	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate	SE/H/0642/006	3797/2011/03	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
filmate				
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/04	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/05	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/06	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/07	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/08	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/09	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/10	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/11	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/12	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/13	KRKA, D.D., NOVO MESTO	RO
Atoris 60 mg comprimate filmate	SE/H/0642/006	3797/2011/14	KRKA, D.D., NOVO MESTO	RO
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
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 STADAGEN 60 mg comprimidos recubiertos con película	NL/H/3345/005	80.732	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
Atorvastatina STADAGEN 30 mg comprimidos recubiertos con película	NL/H/3345/003	80.730	LABORATORIO STADA, S.L.	ES
Atoris 30 mg comprimidos recubiertos con película	SE/H/0758/005	74684	KRKA, D.D., NOVO MESTO	ES
Atostin 30mg filmdragerade tabletter	SE/H/0758/005	44166	KRKA, D.D., NOVO MESTO	SE
Atostin 60mg filmdragerade tabletter	SE/H/0758/006	44167	KRKA, D.D., NOVO MESTO	SE
Atoris® 30 mg Filmtabletten	SE/H/0758/005	81701.00.00	TAD PHARMA GMBH	DE
Atoris® 60 mg Filmtabletten	SE/H/0758/006	81702.00.00	TAD PHARMA GMBH	DE
Atorvastatina ratio 30 mg comprimidos recubiertos con película	not available	77298	RATIOPHARM ESPANA SA	ES
Atorvastatina ratio 60 mg comprimidos recubiertos con película	not available	77297	RATIOPHARM ESPANA SA	ES
Atorvastatine DSM Sinochem 30 mg, filmomhulde tabletten	NL/H/3795/003	RVG 119390	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine DSM Sinochem 60 mg, filmomhulde tabletten	NL/H/3795/005	RVG 119392	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine DSM Sinochem 30 mg, filmomhulde tabletten	NL/H/3797/003	RVG 119402	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
Atorvastatine DSM Sinochem 60 mg, filmomhulde tabletten	NL/H/3797/005	RVG 119404	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
ATORVASTATIN BASICS 30 mg Filmtabletten	MT/H/0125/005	91318.00.00	BASICS 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
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
Atorvastatin 30 mg film-coated tablets	NL/H/3794/003	PL 40170/0014	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	UK
Atorvastatin 60 mg film-coated tablets	NL/H/3794/005	PL 40170/0016	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	UK
Atorvastatin DSM Sinochem 30 mg film-coated tablets	NL/H/3794/003	PA1832/005/003	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	IE
Atorvastatin DSM Sinochem 60 mg film-coated tablets	NL/H/3794/005	PA1832/005/005	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	IE
Atorvastatine DSM Sinochem 30 mg, filmomhulde tabletten	NL/H/3794/003	RVG 119384	DSM SINOCHEM PHARMACEUTICALS NETHERLANDS BV	NL
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
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 971 1 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 984 6 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 005 1 1	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 996 4 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 974 0 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 970 5 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 579 966 3 5	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 972 8 6	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 004 5 0	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 006 8 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 003 9 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 973 4 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 982 3 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 985 2 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE	not available	34009 417 981 7 7	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZENTIVA LAB 20 mg, comprimé pelliculé				
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 986 9 6	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 993 5 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 992 9 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 995 8 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 994 1 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 990 6 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 009 7 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 998 7 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 997 0 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 987 5 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg,	not available	34009 417 991 2 9	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimé pelliculé				
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 008 0 1	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 417 999 3 8	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 007 4 0	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 418 010 5 1	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 418 002 2 1	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 979 2 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 978 6 6	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 976 3 7	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 40 mg, comprimé pelliculé	not available	34009 418 001 6 0	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 989 8 6	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 975 7 6	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ATORVASTATINE ZENTIVA LAB 10 mg, comprimé pelliculé	not available	34009 417 980 0 9	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 80 mg, comprimé pelliculé	not available	34009 579 968 6 4	SANOFI-AVENTIS FRANCE	FR
ATORVASTATINE ZENTIVA LAB 20 mg, comprimé pelliculé	not available	34009 417 988 1 8	SANOFI-AVENTIS FRANCE	FR
Atovans 30 mg filmdragerade tabletter	SE/H/0757/005	44163	KRKA, D.D., NOVO MESTO	SE
Atovans 60 mg filmdragerade tabletter	SE/H/0757/006	44164	KRKA, D.D., NOVO MESTO	SE
Atorvastatin 30 mg film-coated tablets	SE/H/0757/005	PL 01656/0178	KRKA, D.D., NOVO MESTO	UK
Atorvastatin 60 mg film-coated tablets	SE/H/0757/006	PL 01565/0179	KRKA, D.D., NOVO MESTO	UK
Atorvastatina Davur 60 mg comprimidos recubiertos con película	not available	77293	LABORATORIOS DAVUR SL	ES
Atorvastatina Davur 30 mg comprimidos recubiertos con película	not available	77296	LABORATORIOS DAVUR SL	ES
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
Atorvastatin AbZ 30 mg Filmtabletten	not available	88206.00.00	ABZ-PHARMA GMBH	DE
Atorvastatin-ratiopharm® 30 mg Filmtabletten	not available	88204.00.00	RATIOPHARM GMBH	DE
Atorvastatin-ratiopharm® 60 mg	not available	88205.00.00	RATIOPHARM 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				