25 July 2018
EMA/270645/2015
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: rosuvastatin

Procedure no.: PSUSA/00002664/201711

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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
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Coupet 5 mg filmsko obložene tablete	PT/H/0247/001	H/10/00428/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
CRESTOR 10 mg - Filmtabletten	NL/H/0343/001	1-24882	ASTRAZENECA OSTERREICH GMBH	AT
CRESTOR 10 mg apvalkotās tabletes	not available	03-0167	ASTRAZENECA UK LIMITED	LV

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CRESTOR 10 mg apvalkotās tabletes	not available	03-0167	ASTRAZENECA UK LIMITED	LV
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885072	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885146	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885021	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885033	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885019	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885159	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885045	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885084	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885096	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885110	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885108	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885122	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885060	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885058	ASTRAZENECA S.P.A.	IT
Crestor 10 mg compresse rivestite con film	NL/H/0343/001	035885134	ASTRAZENECA S.P.A.	IT
Crestor 10 mg comprimate filmate	not available	5609/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 10 mg comprimate filmate	not available	5609/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 10 mg comprimidos recubiertos con película	NL/H/0343/001	70.243	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357786	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356481	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357687	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356986	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357489	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg	NL/H/0343/001	4356382	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357588	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357380	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357083	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356887	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356689	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357182	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356788	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4356580	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg comprimidos revestidos por película	NL/H/0343/001	4357281	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 10 mg film-coated tablets	not available	19622	ASTRAZENECA UK LIMITED	CY
Crestor 10 mg film-coated tablets	not available	19622	ASTRAZENECA UK LIMITED	CY
Crestor 10 mg film-coated tablets	NL/H/0343/001	PA 970/57/1	ASTRAZENECA UK LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 10 mg film-coated tablets	NL/H/0343/001	MA 044/01502	ASTRAZENECA UK LIMITED	MT
Crestor 10 mg film-coated tablets	NL/H/0343/001	PL 17901/0201	ASTRAZENECA UK LIMITED	UK
Crestor 10 mg filmdragerade tabletter	NL/H/0343/001	19099	ASTRAZENECA AB	SE
Crestor 10 mg filmdrasjerte tabletter	NL/H/0343/001	08-6073	ASTRAZENECA AS	NO
CRESTOR 10 mg filmom obalené tablety	not available	31/0069/03-S	ASTRAZENECA UK LIMITED	SK
CRESTOR 10 mg filmom obalené tablety	not available	31/0069/03-S	ASTRAZENECA UK LIMITED	SK
Crestor 10 mg filmom obložene tablete	not available	HR-H-186159991	ASTRAZENECA D.O.O.	HR
Crestor 10 mg filmom obložene tablete	not available	HR-H-186159991	ASTRAZENECA D.O.O.	HR
CRESTOR 10 mg filmsko obložene tablete	not available	H/04/00430/002	ASTRAZENECA UK LIMITED	SI
CRESTOR 10 mg filmsko obložene tablete	not available	H/04/00430/002	ASTRAZENECA UK LIMITED	SI
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/01	ASTRAZENECA KFT.	HU
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/02	ASTRAZENECA KFT.	HU
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/03	ASTRAZENECA KFT.	HU
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/01	ASTRAZENECA KFT.	HU
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/02	ASTRAZENECA KFT.	HU
Crestor 10 mg filmtabletta	not available	OGYI-T-9574/03	ASTRAZENECA KFT.	HU
Crestor 10 mg filmuhúðaðar töflur	NL/H/0343/001	IS/1/03/008/01	ASTRAZENECA A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 10 mg plėvele dengtos tabletės	not available	LT/1/03/1701/004	ASTRAZENECA UK LIMITED	LT
Crestor 10 mg plėvele dengtos tabletės	not available	LT/1/03/1701/003	ASTRAZENECA UK LIMITED	LT
Crestor 10 mg plėvele dengtos tabletės	not available	LT/1/03/1701/004	ASTRAZENECA UK LIMITED	LT
Crestor 10 mg plėvele dengtos tabletės	not available	LT/1/03/1701/003	ASTRAZENECA UK LIMITED	LT
Crestor 10 mg potahované tablety	not available	31/314/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 10 mg potahované tablety	not available	31/314/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 10 mg tabletti, kalvopäällysteinen	NL/H/0343/001	17885	ASTRAZENECA OY	FI
CRESTOR 10 mg, comprimés pelliculés	NL/H/0343/001	BE250205	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 10 mg, comprimés pelliculés	NL/H/0343/001	BE250187	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 10 mg, filmomhulde tabletten	NL/H/0343/001	BE250205	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 10 mg, filmomhulde tabletten	NL/H/0343/001	BE250187	ASTRAZENECA S.A. / N.V.	BE
Crestor 10, filmomhulde tabletten 10 mg	NL/H/0343/001	RVG 26872	ASTRAZENECA BV	NL
CRESTOR 20 mg - Filmtabletten	NL/H/0343/002	1-24883	ASTRAZENECA OSTERREICH GMBH	AT
CRESTOR 20 mg apvalkotās tabletes	not available	03-0168	ASTRAZENECA UK LIMITED	LV
CRESTOR 20 mg apvalkotās tabletes	not available	03-0168	ASTRAZENECA UK LIMITED	LV
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885247	ASTRAZENECA S.P.A.	IT
Crestor 20 mg	NL/H/0343/002	035885223	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885235	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885286	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885185	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885262	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885173	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885197	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885211	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885209	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885161	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885298	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885274	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885300	ASTRAZENECA S.P.A.	IT
Crestor 20 mg compresse rivestite con film	NL/H/0343/002	035885250	ASTRAZENECA S.P.A.	IT
Crestor 20 mg comprimate filmate	not available	5610/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 20 mg comprimate filmate	not available	5610/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 20 mg comprimidos recubiertos con película	NL/H/0343/002	70.244	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358081	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358487	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358289	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4359188	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4357984	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358784	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4357885	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg	NL/H/0343/002	4359287	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358586	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358883	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358180	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358388	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358982	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4358685	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg comprimidos revestidos por película	NL/H/0343/002	4359089	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 20 mg film-coated tablets.	not available	19623	ASTRAZENECA UK LIMITED	CY
Crestor 20 mg film-coated tablets.	not available	19623	ASTRAZENECA UK LIMITED	CY
Crestor 20 mg film-coated tablets.	NL/H/0343/002	PA 970/57/2	ASTRAZENECA UK LIMITED	IE
Crestor 20 mg film-coated tablets.	NL/H/0343/002	MA 044/01503	ASTRAZENECA UK LIMITED	MT
Crestor 20 mg film-coated tablets.	NL/H/0343/002	PL 17901/0202	ASTRAZENECA UK LIMITED	UK
Crestor 20 mg filmdragerade tabletter	NL/H/0343/002	19100	ASTRAZENECA AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 20 mg filmdrasjerte tabletter	NL/H/0343/002	08-6074	ASTRAZENECA AS	NO
CRESTOR 20 mg filmom obalené tablety	not available	31/0070/03-S	ASTRAZENECA UK LIMITED	SK
CRESTOR 20 mg filmom obalené tablety	not available	31/0070/03-S	ASTRAZENECA UK LIMITED	SK
Crestor 20 mg filmom obložene tablete	not available	HR-H-680317247	ASTRAZENECA D.O.O.	HR
Crestor 20 mg filmom obložene tablete	not available	HR-H-680317247	ASTRAZENECA D.O.O.	HR
CRESTOR 20 mg filmsko obložene tablete	not available	H/04/00430/003	ASTRAZENECA UK LIMITED	SI
CRESTOR 20 mg filmsko obložene tablete	not available	H/04/00430/003	ASTRAZENECA UK LIMITED	SI
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/04	ASTRAZENECA KFT.	HU
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/05	ASTRAZENECA KFT.	HU
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/06	ASTRAZENECA KFT.	HU
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/04	ASTRAZENECA KFT.	HU
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/05	ASTRAZENECA KFT.	HU
Crestor 20 mg filmtabletta	not available	OGYI-T-9574/06	ASTRAZENECA KFT.	HU
Crestor 20 mg filmuhúðaðar töflur	NL/H/0343/002	IS/1/03/008/02	ASTRAZENECA A/S	IS
Crestor 20 mg plėvele dengtos tabletės	not available	LT/1/03/1701/005	ASTRAZENECA UK LIMITED	LT
Crestor 20 mg plėvele dengtos tabletės	not available	LT/1/03/1701/006	ASTRAZENECA UK LIMITED	LT
Crestor 20 mg plėvele dengtos tabletės	not available	LT/1/03/1701/005	ASTRAZENECA UK LIMITED	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
Crestor 20 mg plėvele dengtos tabletės	not available	LT/1/03/1701/006	ASTRAZENECA UK LIMITED	LT
Crestor 20 mg potahované tablety	not available	31/315/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 20 mg potahované tablety	not available	31/315/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 20 mg tabletti, kalvopäällysteinen	NL/H/0343/002	17886	ASTRAZENECA OY	FI
CRESTOR 20 mg, comprimés pelliculés	NL/H/0343/002	BE250223	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 20 mg, comprimés pelliculés	NL/H/0343/002	BE250241	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 20 mg, filmomhulde tabletten	NL/H/0343/002	BE250223	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 20 mg, filmomhulde tabletten	NL/H/0343/002	BE250241	ASTRAZENECA S.A. / N.V.	BE
Crestor 20, filmomhulde tabletten 20 mg	NL/H/0343/002	RVG 26873	ASTRAZENECA BV	NL
CRESTOR 40 mg - Filmtabletten	NL/H/0343/003	1-24884	ASTRAZENECA OSTERREICH GMBH	AT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885363	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885401	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885336	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885399	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885375	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885348	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885449	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885387	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885351	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885413	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885312	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885324	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885437	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885425	ASTRAZENECA S.P.A.	IT
Crestor 40 mg compresse rivestite con film	NL/H/0343/003	035885452	ASTRAZENECA S.P.A.	IT
Crestor 40 mg comprimate filmate	not available	5611/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 40 mg comprimate filmate	not available	5611/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 40 mg	NL/H/0343/003	70.335	ASTRAZENECA	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos recubiertos con película			FARMACÉUTICA SPAIN, S.A.	
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359584	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360087	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360384	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360582	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360780	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359485	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359980	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359683	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359782	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359386	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360285	ASTRAZENECA PRODUTOS FARMACEUTICOS 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
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360681	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360186	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4359881	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg comprimidos revestidos por película	NL/H/0343/003	4360483	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 40 mg film-coated tablets	NL/H/0343/003	PA 970/57/3	ASTRAZENECA UK LIMITED	IE
Crestor 40 mg film-coated tablets	NL/H/0343/003	PL 17901/0203	ASTRAZENECA UK LIMITED	UK
Crestor 40 mg film-coated tablets.	not available	19624	ASTRAZENECA UK LIMITED	CY
Crestor 40 mg film-coated tablets.	not available	19624	ASTRAZENECA UK LIMITED	CY
Crestor 40 mg film-coated tablets.	NL/H/0343/003	MA 044/01504	ASTRAZENECA UK LIMITED	MT
Crestor 40 mg filmdragerade tabletter	NL/H/0343/003	19101	ASTRAZENECA AB	SE
Crestor 40 mg filmdrasjerte tabletter	NL/H/0343/003	08-6075	ASTRAZENECA AS	NO
CRESTOR 40 mg filmom obalené tablety	not available	31/0071/03-S	ASTRAZENECA UK LIMITED	SK
CRESTOR 40 mg filmom obalené tablety	not available	31/0071/03-S	ASTRAZENECA UK LIMITED	SK
Crestor 40 mg filmom obložene tablete	not available	HR-H-857111079	ASTRAZENECA D.O.O.	HR
Crestor 40 mg filmom obložene tablete	not available	HR-H-857111079	ASTRAZENECA D.O.O.	HR
CRESTOR 40 mg filmsko	not available	H/04/00430/004	ASTRAZENECA UK LIMITED	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				
CRESTOR 40 mg filmsko obložene tablete	not available	H/04/00430/004	ASTRAZENECA UK LIMITED	SI
Crestor 40 mg filmuhúðaðar töflur	NL/H/0343/003	IS/1/03/008/03	ASTRAZENECA A/S	IS
Crestor 40 mg potahované tablety	not available	31/316/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 40 mg potahované tablety	not available	31/316/03-C	ASTRAZENECA UK LIMITED	CZ
Crestor 40 mg tabletti, kalvopäällysteinen	NL/H/0343/003	17887	ASTRAZENECA OY	FI
CRESTOR 40 mg, comprimés pelliculés	NL/H/0343/003	BE250266	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 40 mg, comprimés pelliculés	NL/H/0343/003	BE250284	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 40 mg, filmomhulde tabletten	NL/H/0343/003	BE250266	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 40 mg, filmomhulde tabletten	NL/H/0343/003	BE250284	ASTRAZENECA S.A. / N.V.	BE
Crestor 40, filmomhulde tabletten 40 mg	NL/H/0343/003	RVG 26874	ASTRAZENECA BV	NL
CRESTOR 5 mg - Filmtabletten	NL/H/0343/004	1-26023	ASTRAZENECA OSTERREICH GMBH	AT
CRESTOR 5 mg apvalkotās tabletes	not available	09-0351	ASTRAZENECA UK LIMITED	LV
CRESTOR 5 mg apvalkotās tabletes	not available	09-0351	ASTRAZENECA UK LIMITED	LV
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885464	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885540	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885565	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse	NL/H/0343/004	035885490	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rivestite con film				
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885476	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885526	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885589	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885603	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885538	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885514	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885577	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885591	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035883553	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885488	ASTRAZENECA S.P.A.	IT
Crestor 5 mg compresse rivestite con film	NL/H/0343/004	035885502	ASTRAZENECA S.P.A.	IT
Crestor 5 mg comprimate filmate	not available	5608/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 5 mg comprimate filmate	not available	5608/2013/01	ASTRAZENECA UK LIMITED	RO
Crestor 5 mg comprimidos recubiertos con película	NL/H/0343/004	70334	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570585	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg	NL/H/0343/004	5570387	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570288	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570080	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569884	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570684	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570783	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569389	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569488	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569587	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569785	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570189	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569983	ASTRAZENECA PRODUTOS FARMACEUTICOS 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
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5569686	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg comprimidos revestidos por película	NL/H/0343/004	5570486	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Crestor 5 mg film-coated tablets	NL/H/0343/004	PA 970/57/4	ASTRAZENECA UK LIMITED	IE
Crestor 5 mg film-coated tablets	NL/H/0343/004	MA 044/01501	ASTRAZENECA UK LIMITED	MT
Crestor 5 mg film-coated tablets	NL/H/0343/004	PL 17901/0243	ASTRAZENECA UK LIMITED	UK
Crestor 5 mg film-coated tablets.	not available	19728	ASTRAZENECA UK LIMITED	CY
Crestor 5 mg film-coated tablets.	not available	19728	ASTRAZENECA UK LIMITED	CY
Crestor 5 mg filmdragerade tabletter	NL/H/0343/004	21377	ASTRAZENECA AB	SE
Crestor 5 mg filmdrasjerte tabletter	NL/H/0343/004	08-6072	ASTRAZENECA AS	NO
CRESTOR 5 mg filmom obalené tablety	not available	31/0494/09-S	ASTRAZENECA UK LIMITED	SK
CRESTOR 5 mg filmom obalené tablety	not available	31/0494/09-S	ASTRAZENECA UK LIMITED	SK
Crestor 5 mg filmom obložene tablete	not available	HR-H-050912005	ASTRAZENECA D.O.O.	HR
Crestor 5 mg filmom obložene tablete	not available	HR-H-050912005	ASTRAZENECA D.O.O.	HR
CRESTOR 5 mg filmsko obložene tablete	not available	H/04/00430/001	ASTRAZENECA UK LIMITED	SI
CRESTOR 5 mg filmsko obložene tablete	not available	H/04/00430/001	ASTRAZENECA UK LIMITED	SI
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/10	ASTRAZENECA KFT.	HU
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/12	ASTRAZENECA 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
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/11	ASTRAZENECA KFT.	HU
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/10	ASTRAZENECA KFT.	HU
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/12	ASTRAZENECA KFT.	HU
Crestor 5 mg filmtabletta	not available	OGYI-T-9574/11	ASTRAZENECA KFT.	HU
Crestor 5 mg filmuhúðaðar töflur	NL/H/0343/004	IS/1/04/062/01	ASTRAZENECA A/S	IS
Crestor 5 mg plėvele dengtos tabletės	not available	LT/1/03/1701/001	ASTRAZENECA UK LIMITED	LT
Crestor 5 mg plėvele dengtos tabletės	not available	LT/1/03/1701/002	ASTRAZENECA UK LIMITED	LT
Crestor 5 mg plėvele dengtos tabletės	not available	LT/1/03/1701/001	ASTRAZENECA UK LIMITED	LT
Crestor 5 mg plėvele dengtos tabletės	not available	LT/1/03/1701/002	ASTRAZENECA UK LIMITED	LT
Crestor 5 mg potahované tablety	not available	31/472/10-C	ASTRAZENECA UK LIMITED	CZ
Crestor 5 mg potahované tablety	not available	31/472/10-C	ASTRAZENECA UK LIMITED	CZ
Crestor 5 mg tabletti, kalvopäällysteinen	NL/H/0343/004	19816	ASTRAZENECA OY	FI
CRESTOR 5 mg, comprimés pelliculés	NL/H/0343/004	BE276963	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 5 mg, comprimés pelliculés	NL/H/0343/004	BE276945	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 5 mg, filmomhulde tabletten	NL/H/0343/004	BE276963	ASTRAZENECA S.A. / N.V.	BE
CRESTOR 5 mg, filmomhulde tabletten	NL/H/0343/004	BE276945	ASTRAZENECA S.A. / N.V.	BE
Crestor 5, filmomhulde tabletten 5 mg	NL/H/0343/004	RGV 30823	ASTRAZENECA BV	NL
Crestor, 10 mg õhukese polümeerikattega tabletid	not available	416303	ASTRAZENECA UK LIMITED	EE
Crestor, 10 mg õhukese	not available	416303	ASTRAZENECA UK LIMITED	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
polümeerikattega tabletid				
Crestor, 10 mg, tabletki powlekane	NL/H/0343/001	15135	ASTRAZENECA AB	PL
Crestor, 20 mg õhukese polümeerikattega tabletid	not available	416403	ASTRAZENECA UK LIMITED	EE
Crestor, 20 mg õhukese polümeerikattega tabletid	not available	416403	ASTRAZENECA UK LIMITED	EE
Crestor, 20 mg, tabletki powlekane	NL/H/0343/002	15136	ASTRAZENECA AB	PL
Crestor, 40 mg, tabletki powlekane	NL/H/0343/003	15137	ASTRAZENECA AB	PL
Crestor, 5 mg õhukese polümeerikattega tabletid	not available	675110	ASTRAZENECA UK LIMITED	EE
Crestor, 5 mg õhukese polümeerikattega tabletid	not available	675110	ASTRAZENECA UK LIMITED	EE
Crestor, 5 mg, tabletki powlekane	NL/H/0343/004	15134	ASTRAZENECA AB	PL
Crestor, filmovertrukne tabletter	NL/H/0343/004	36993	ASTRAZENECA A/S	DK
Crestor, filmovertrukne tabletter	NL/H/0343/003	34538	ASTRAZENECA A/S	DK
Crestor, filmovertrukne tabletter	NL/H/0343/002	34537	ASTRAZENECA A/S	DK
Crestor, filmovertrukne tabletter	NL/H/0343/001	34536	ASTRAZENECA A/S	DK
CRESTOR® 10 mg Filmtabletten	NL/H/0343/001	58179.00.00	ASTRAZENECA GMBH	DE
CRESTOR® 10 mg, comprimé pelliculé	NL/H/0343/001	NL 28274	ASTRAZENECA S.A.S.	FR
CRESTOR® 20 mg	NL/H/0343/002	58180.00.00	ASTRAZENECA 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				
CRESTOR® 20 mg, comprimé pelliculé	NL/H/0343/002	NL 28275	ASTRAZENECA S.A.S.	FR
Crestor® 40 mg Filmtabletten	NL/H/0343/003	58181.00.00	ASTRAZENECA GMBH	DE
CRESTOR® 5 mg Filmtabletten	NL/H/0343/004	61744.00.00	ASTRAZENECA GMBH	DE
CRESTOR® 5 mg, comprimé pelliculé	NL/H/0343/004	NL 30594	ASTRAZENECA S.A.S.	FR
Epri 15 mg filmom obložene tablete	DK/H/2388/001	HR-H-368649981	PLIVA HRVATSKA D.O.O.	HR
Epri 30 mg filmom obložene tablete	DK/H/2388/002	HR-H-399406571	PLIVA HRVATSKA D.O.O.	HR
Mertenil 15 mg potahované tablety	HU/H/0219/005	31/801/16-C	GEDEON RICHTER PLC.	CZ
Mertenil 30 mg potahované tablety	HU/H/0219/006	31/802/16-C	GEDEON RICHTER PLC.	CZ
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883281	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883279	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883040	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883053	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883065	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883077	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883089	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883091	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883103	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883014	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883115	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883127	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883139	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883026	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883038	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883281	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883279	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con	NL/H/0345/001	035883040	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883053	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883065	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883077	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883089	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883091	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883103	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883014	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883115	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883127	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883139	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg compresse rivestite con film	NL/H/0345/001	035883026	ASTRAZENECA S.P.A.	IT
Provisacor 10 mg	NL/H/0345/001	035883038	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Provisacor 10 mg comprimidos recubiertos con película	NL/H/0345/001	70.242	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 10 mg comprimidos recubiertos con película	NL/H/0345/001	70.242	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 10, filmomhulde tabletten 10 mg	NL/H/0345/001	RGV 27655	ASTRAZENECA BV	NL
Provisacor 10, filmomhulde tabletten 10 mg	NL/H/0345/001	RGV 27655	ASTRAZENECA BV	NL
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883141	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883242	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883255	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883267	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883154	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883178	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883216	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883166	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883180	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883192	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883204	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883228	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883230	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883293	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883305	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883141	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883242	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883255	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con	NL/H/0345/002	035883267	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883154	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883178	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883216	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883166	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883180	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883192	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883204	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883228	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883230	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883293	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg compresse rivestite con film	NL/H/0345/002	035883305	ASTRAZENECA S.P.A.	IT
Provisacor 20 mg	NL/H/0345/002	70.241	ASTRAZENECA	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos recubiertos con película			FARMACÉUTICA SPAIN, S.A.	
Provisacor 20 mg comprimidos recubiertos con película	NL/H/0345/002	70.241	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 20, filmomhulde tabletten 20 mg	NL/H/0345/002	RGV 27656	ASTRAZENECA BV	NL
Provisacor 20, filmomhulde tabletten 20 mg	NL/H/0345/002	RGV 27656	ASTRAZENECA BV	NL
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883329	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883432	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883382	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883457	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883331	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883343	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883356	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883370	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883368	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883444	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883394	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883406	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883317	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883418	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883420	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883329	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883432	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883382	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883457	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con	NL/H/0345/003	035883331	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883343	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883356	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883370	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883368	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883444	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883394	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883406	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883317	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883418	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg compresse rivestite con film	NL/H/0345/003	035883420	ASTRAZENECA S.P.A.	IT
Provisacor 40 mg comprimidos recubiertos con película	NL/H/0345/003	70.336	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 40 mg	NL/H/0345/003	70.336	ASTRAZENECA	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos recubiertos con película			FARMACÉUTICA SPAIN, S.A.	
Provisacor 40, filmomhulde tabletten 40 mg	NL/H/0345/003	RGV 27657	ASTRAZENECA BV	NL
Provisacor 40, filmomhulde tabletten 40 mg	NL/H/0345/003	RGV 27657	ASTRAZENECA BV	NL
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883596	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883608	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883572	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883584	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883471	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883483	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883495	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883507	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883519	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883521	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883533	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883545	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883558	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883469	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883560	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883596	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883608	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883572	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883584	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883471	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con	NL/H/0345/004	035883483	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883495	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883507	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883519	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883521	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883533	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883545	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883558	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883469	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg compresse rivestite con film	NL/H/0345/004	035883560	ASTRAZENECA S.P.A.	IT
Provisacor 5 mg comprimidos recubiertos con película	NL/H/0345/004	70.333	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 5 mg comprimidos recubiertos con película	NL/H/0345/004	70.333	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Provisacor 5,	NL/H/0345/004	RGV 30825	ASTRAZENECA BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmomhulde tabletten 5 mg				
Provisacor 5, filmomhulde tabletten 5 mg	NL/H/0345/004	RGV 30825	ASTRAZENECA BV	NL
Rostat 15 mg comprimate filmate	HU/H/0219/005	10196/2017/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Rostat 15 mg comprimate filmate	HU/H/0219/005	10196/2017/02	GEDEON RICHTER ROMÂNIA S.A.	RO
Rostat 15 mg comprimate filmate	HU/H/0219/005	10196/2017/03	GEDEON RICHTER ROMÂNIA S.A.	RO
Rostat 30 mg comprimate filmate	HU/H/0219/006	10197/2017/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Rostat 30 mg comprimate filmate	HU/H/0219/006	10197/2017/03	GEDEON RICHTER ROMÂNIA S.A.	RO
Rosudia, filmovertrukne tabletter	DK/H/2611/003	57069	KRKA, D.D., NOVO MESTO	DK
Rosudia, filmovertrukne tabletter	DK/H/2611/005	57071	KRKA, D.D., NOVO MESTO	DK
Rosumop 5 mg Potahovan뿱½ tablety	PT/H/0247/001	31/363/10-C	SANDOZ S.R.O.	CZ
Rosuvador® 15 mg Filmtabletten	DK/H/2583/003	96803.00.00	TAD PHARMA GMBH	DE
Rosuvador® 30 mg Filmtabletten	DK/H/2583/005	96805.00.00	TAD PHARMA GMBH	DE
Rosuvastatin "Krka d.d.", filmovertrukne tabletter	DK/H/2583/003	57063	KRKA, D.D., NOVO MESTO	DK
Rosuvastatin "Krka d.d.", filmovertrukne tabletter	DK/H/2583/005	57065	KRKA, D.D., NOVO MESTO	DK
Rosuvastatin "Krka", filmovertrukne tabletter	DK/H/1937/003	46931	KRKA, D.D., NOVO MESTO	DK
Rosuvastatin "Krka", filmovertrukne tabletter	DK/H/1937/005	46933	KRKA, D.D., NOVO MESTO	DK
Rosuvastatin Actavis 15	DK/H/2388/001	32909	TEVA GYÓGYSZERGYÁR ZRT	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
mg tabletti, kalvopäällysteinen				
Rosuvastatin AstraZeneca 10, filmomhulde tabletten 10 mg	NL/H/0346/001	RGV 27659	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 10, filmomhulde tabletten 10 mg	NL/H/0346/001	RGV 27659	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 20, filmomhulde tabletten 20 mg	NL/H/0346/002	RGV 27660	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 20, filmomhulde tabletten 20 mg	NL/H/0346/002	RGV 27660	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 40, filmomhulde tabletten 40 mg	NL/H/0346/003	RGV 27661	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 40, filmomhulde tabletten 40 mg	NL/H/0346/003	RGV 27661	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 5, filmomhulde tabletten 5 mg	NL/H/0346/004	RGV 30826	ASTRAZENECA BV	NL
Rosuvastatin AstraZeneca 5, filmomhulde tabletten 5 mg	NL/H/0346/004	RGV 30826	ASTRAZENECA BV	NL
Rosuvastatin Krka 15 mg	DK/H/1937/003	11-0271	KRKA, D.D., NOVO MESTO	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
apvalkotās tabletes				
Rosuvastatin Krka 15 mg comprimés pelliculés	DK/H/2583/003	BE503555	KRKA, D.D., NOVO MESTO	BE
Rosuvastatin Krka 15 mg filmom obalené tablety	DK/H/1937/003	31/0768/11-S	KRKA, D.D., NOVO MESTO	SK
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/013	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/014	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/015	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/016	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/017	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/018	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/019	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/020	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/021	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/022	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg filmsko obložene tablete	DK/H/1937/003	H/11/01376/012	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 15 mg Filmtabletten	DK/H/2583/003	137420	KRKA, D.D., NOVO MESTO	AT
Rosuvastatin Krka 15 mg kalvopäällysteiset tabletit	DK/H/2583/003	33850	KRKA SVERIGE AB	FI
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/023	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg	DK/H/1937/003	LT/1/11/2621/024	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
plėvele dengtos tabletės				
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/025	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/026	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/027	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/028	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/029	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/030	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/031	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/032	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg plėvele dengtos tabletės	DK/H/1937/003	LT/1/11/2621/033	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 15 mg potahované tablety	DK/H/1937/003	31/081/12-C	KRKA, D.D., NOVO MESTO	CZ
Rosuvastatin Krka 15 mg, õhukese polümeerikattega tabletid	DK/H/1937/003	748311	KRKA, D.D., NOVO MESTO	EE
Rosuvastatin Krka 30 mg apvalkotās tabletes	DK/H/1937/005	11-0273	KRKA, D.D., NOVO MESTO	LV
Rosuvastatin Krka 30 mg comprimés pelliculés	DK/H/2583/005	BE503573	KRKA, D.D., NOVO MESTO	BE
Rosuvastatin Krka 30 mg filmom obalené tablety	DK/H/1937/005	31/0770/11-S	KRKA, D.D., NOVO MESTO	SK
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/035	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg	DK/H/1937/005	H/11/01376/036	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmsko obložene tablete				
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/037	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/038	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/039	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/040	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/041	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/042	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/043	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/044	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg filmsko obložene tablete	DK/H/1937/005	H/11/01376/034	KRKA, D.D., NOVO MESTO	SI
Rosuvastatin Krka 30 mg Filmtabletten	DK/H/2583/005	137422	KRKA, D.D., NOVO MESTO	AT
Rosuvastatin Krka 30 mg kalvopäällysteiset tabletit	DK/H/2583/005	33852	KRKA SVERIGE AB	FI
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/045	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/046	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/047	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/048	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/049	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg	DK/H/1937/005	LT/1/11/2621/050	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
plėvele dengtos tabletės				
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/051	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/052	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/053	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/054	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg plėvele dengtos tabletės	DK/H/1937/005	LT/1/11/2621/055	KRKA, D.D., NOVO MESTO	LT
Rosuvastatin Krka 30 mg potahované tablety	DK/H/1937/005	31/083/12-C	KRKA, D.D., NOVO MESTO	CZ
Rosuvastatin Krka 30 mg, õhukese polümeerikattega tabletid	DK/H/1937/005	748511	KRKA, D.D., NOVO MESTO	EE
Rosuvastatin Krka, 15 mg, tabletki powlekane	DK/H/1937/003	18806	KRKA, D.D., NOVO MESTO	PL
Rosuvastatin Krka, 30 mg, tabletki powlekane	DK/H/1937/005	18808	KRKA, D.D., NOVO MESTO	PL
Rosuvastatin ratiopharm 15 mg filmdragerad tablett	DK/H/2388/001	32909	TEVA GYÓGYSZERGYÁR ZRT	FI
Rosuvastatin ratiopharm 15 mg Filmtabletten	DK/H/2388/001	137569	TEVA B.V	AT
Rosuvastatin ratiopharm 30 mg Filmtabletten	DK/H/2388/002	137568	TEVA B.V	AT
Rosuvastatin ratiopharm 30 mg tabletti, kalvopäällysteinen	DK/H/2388/002	32910	RATIOPHARM GMBH	FI
Rosuvastatin ratiopharm 30 mg tabletti, kalvopäällysteinen	DK/H/2388/002	32910	RATIOPHARM GMBH	FI
Rosuvastatin Sandoz 5	PT/H/0247/001	OGYI-T-21423/02	SANDOZ HUNGÁRIA 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
mg filmtabletta				
Rosuvastatin Sandoz 5 mg filmtabletta	PT/H/0247/001	OGYI-T-21423/01	SANDOZ HUNGÁRIA KFT	HU
Rosuvastatin Sandoz 5 mg filmtabletta	PT/H/0247/001	OGYI-T-21423/03	SANDOZ HUNGÁRIA KFT	HU
Rosuvastatin Sandoz 5 mg filmtabletta	PT/H/0247/001	OGYI-T-21423/04	SANDOZ HUNGÁRIA KFT	HU
Rosuvastatin Teva, 15 mg, tabletki powlekane	DK/H/2388/001	23346	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Rosuvastatin Teva, 30 mg, tabletki powlekane	DK/H/2388/002	23347	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Rosuvastatina HCS 15 mg compresse rivestite con film	DK/H/2611/003	044711036	HCS BVBA	IT
Rosuvastatina HCS 15 mg compresse rivestite con film	DK/H/2611/003	044711099	HCS BVBA	IT
Rosuvastatina HCS 30 mg compresse rivestite con film	DK/H/2611/005	044711051	HCS BVBA	IT
Rosuvastatina HCS 30 mg compresse rivestite con film	DK/H/2611/005	044711113	HCS BVBA	IT
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/01	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/02	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/03	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/04	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/05	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/06	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
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/07	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/08	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/09	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/10	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 15 mg comprimate filmate	DK/H/1937/003	9166/2016/11	KRKA, D.D., NOVO MESTO	RO
Rosuvastatina Krka 15 mg comprimidos revestidos por película	DK/H/2583/003	5712138	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina Krka 15 mg comprimidos revestidos por película	DK/H/2583/003	5712146	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina Krka 20 mg comprimidos recubiertos con película EFG	DK/H/2583/003	81845	KRKA, D.D., NOVO MESTO	ES
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/01	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/02	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/03	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/04	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/05	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/06	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/07	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30	DK/H/1937/005	9168/2016/08	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
mg comprimate filmate				
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/09	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/10	KRKA, D.D., NOVO MESTO	RO
Rosuvastatină Krka 30 mg comprimate filmate	DK/H/1937/005	9168/2016/11	KRKA, D.D., NOVO MESTO	RO
Rosuvastatina Krka 30 mg comprimidos recubiertos con película	DK/H/2583/005	81846	KRKA, D.D., NOVO MESTO	ES
Rosuvastatina Krka 30 mg comprimidos revestidos por película	DK/H/2583/005	5712161	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina ratiopharm 15 mg comprimidos revestidos por película	DK/H/2388/001	5672035	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Rosuvastatina ratiopharm 15 mg comprimidos revestidos por película	DK/H/2388/001	5672027	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Rosuvastatina ratiopharm 15 mg comprimidos revestidos por película	DK/H/2388/001	5672050	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Rosuvastatina ratiopharm 15 mg comprimidos revestidos por película	DK/H/2388/001	5672043	RATIOPHARM LDA	PT
Rosuvastatina ratiopharm 30 mg comprimidos revestidos por película	DK/H/2388/002	5672100	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Rosuvastatina ratiopharm 30 mg	DK/H/2388/002	5672068	RATIOPHARM 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				
Rosuvastatina ratiopharm 30 mg comprimidos revestidos por película	DK/H/2388/002	5672076	RATIOPHARM LDA	PT
Rosuvastatina ratiopharm 30 mg comprimidos revestidos por película	DK/H/2388/002	5672118	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Rosuvastatine Teva 15 mg comprimés pelliculés	DK/H/2388/001	BE488835	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg comprimés pelliculés	DK/H/2388/001	BE488844	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg comprimés pelliculés	DK/H/2388/001	BE508915	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg filmomhulde tabletten	DK/H/2388/001	BE488835	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg filmomhulde tabletten	DK/H/2388/001	BE488844	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg filmomhulde tabletten	DK/H/2388/001	BE508915	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg Filmtabletten	DK/H/2388/001	BE488835	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg Filmtabletten	DK/H/2388/001	BE488844	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 15 mg Filmtabletten	DK/H/2388/001	BE508915	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg comprimés pelliculés	DK/H/2388/002	BE488853	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg comprimés pelliculés	DK/H/2388/002	BE488862	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosuvastatine Teva 30 mg comprimés pelliculés	DK/H/2388/002	BE508924	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg filmomhulde tabletten	DK/H/2388/002	BE488853	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg filmomhulde tabletten	DK/H/2388/002	BE488862	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg filmomhulde tabletten	DK/H/2388/002	BE508924	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg Filmtabletten	DK/H/2388/002	BE488853	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg Filmtabletten	DK/H/2388/002	BE488862	TEVA PHARMA BELGIUM N.V./S.A	BE
Rosuvastatine Teva 30 mg Filmtabletten	DK/H/2388/002	BE508924	TEVA PHARMA BELGIUM N.V./S.A	BE
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/01	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/02	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/03	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/04	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/05	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/06	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/07	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/08	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/09	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
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/10	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg comprimate filmate	DK/H/1938/003	9172/2016/11	KRKA, D.D., NOVO MESTO	RO
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/023	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/024	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/025	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/026	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/027	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/028	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/029	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/030	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/031	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/032	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg plėvele dengtos tabletės	DK/H/1938/003	LT/1/11/2623/033	KRKA, D.D., NOVO MESTO	LT
Roswera 15 mg, õhukese polümeerikattega tabletid	DK/H/1938/003	748911	KRKA, D.D., NOVO MESTO	EE
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/01	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/02	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg	DK/H/1938/005	9174/2016/03	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
comprimate filmate				
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/04	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/05	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/06	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/07	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/08	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/09	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/10	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg comprimate filmate	DK/H/1938/005	9174/2016/11	KRKA, D.D., NOVO MESTO	RO
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/045	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/046	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/047	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/048	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/049	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/050	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/051	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/052	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele	DK/H/1938/005	LT/1/11/2623/053	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				
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/054	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg plėvele dengtos tabletės	DK/H/1938/005	LT/1/11/2623/055	KRKA, D.D., NOVO MESTO	LT
Roswera 30 mg, õhukese polümeerikattega tabletid	DK/H/1938/005	749211	KRKA, D.D., NOVO MESTO	EE
Roswera, 15 mg, tabletki powlekane	DK/H/1938/003	18579	KRKA, D.D., NOVO MESTO	PL
Roswera, 30 mg, tabletki powlekane	DK/H/1938/005	18581	KRKA, D.D., NOVO MESTO	PL
Roswera, filmovertrukne tabletter	DK/H/1938/003	46937	TAD PHARMA GMBH	DK
Roswera, filmovertrukne tabletter	DK/H/1938/005	46939	TAD PHARMA GMBH	DK
Rovanta	DK/H/2388/002	55211	RATIOPHARM GMBH	DK
Rovanta, filmovertrukne tabletter	DK/H/2388/001	55210	RATIOPHARM GMBH	DK
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/23	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/24	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/25	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/26	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/27	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/28	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/29	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg	not available	OGYI-T-21686/30	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmtabletta				
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/31	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/32	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg filmtabletta	not available	OGYI-T-21686/33	KRKA, D.D., NOVO MESTO	HU
Roxera 15 mg õhukese polümeerikattega tabletid	not available	778812	KRKA, D.D., NOVO MESTO	EE
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/45	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/46	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/47	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/48	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/49	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/50	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/51	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/52	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/53	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/54	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg filmtabletta	not available	OGYI-T-21686/55	KRKA, D.D., NOVO MESTO	HU
Roxera 30 mg õhukese polümeerikattega	not available	778912	KRKA, D.D., NOVO MESTO	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletid				
Rozuva-Teva 15 mg filmtabletta	DK/H/2388/001	OGYI-T-22934/17	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 15 mg filmtabletta	DK/H/2388/001	OGYI-T-22934/23	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 15 mg filmtabletta	DK/H/2388/001	OGYI-T-22934/28	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 15 mg filmtabletta	DK/H/2388/001	OGYI-T-22934/29	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 30 mg filmtabletta	DK/H/2388/002	OGYI-T-22934/18	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 30 mg filmtabletta	DK/H/2388/002	OGYI-T-22934/24	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 30 mg filmtabletta	DK/H/2388/002	OGYI-T-22934/30	TEVA GYÓGYSZERGYÁR ZRT	HU
Rozuva-Teva 30 mg filmtabletta	DK/H/2388/002	OGYI-T-22934/31	TEVA GYÓGYSZERGYÁR ZRT	HU
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884143	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884156	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884067	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884042	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884055	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884079	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884081	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884093	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884105	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884016	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884117	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884129	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884131	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884028	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884030	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884143	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884156	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con	NL/H/0346/001	035884067	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884042	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884055	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884079	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884081	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884093	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884105	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884016	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884117	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884129	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884131	SIMESA S.P.A.	IT
Simestat 10 mg compresse rivestite con film	NL/H/0346/001	035884028	SIMESA S.P.A.	IT
Simestat 10 mg	NL/H/0346/001	035884030	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884295	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884194	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884206	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884220	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884232	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884244	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884257	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884218	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884170	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884182	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884283	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884168	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884271	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884269	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884295	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884194	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884206	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884220	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884232	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884244	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884257	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884218	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con	NL/H/0346/002	035884170	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884182	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884283	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884168	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884271	SIMESA S.P.A.	IT
Simestat 20 mg compresse rivestite con film	NL/H/0346/002	035884269	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884446	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884459	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884333	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884345	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884358	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884372	SIMESA S.P.A.	IT
Simestat 40 mg	NL/H/0346/003	035884422	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884384	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884396	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884408	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884319	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884410	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884321	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884434	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884360	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884446	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884459	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884333	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884345	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884358	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884372	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884422	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884384	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884396	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884408	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884319	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884410	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884321	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con film	NL/H/0346/003	035884434	SIMESA S.P.A.	IT
Simestat 40 mg compresse rivestite con	NL/H/0346/003	035884360	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
film				
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884509	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884511	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884523	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884535	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884547	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884550	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884461	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884562	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884574	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884600	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884598	SIMESA S.P.A.	IT
Simestat 5 mg	NL/H/0346/004	035884586	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884473	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884485	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884497	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884509	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884511	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884523	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884535	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884547	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884550	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884461	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884562	SIMESA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884574	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884600	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884598	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884586	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884473	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884485	SIMESA S.P.A.	IT
Simestat 5 mg compresse rivestite con film	NL/H/0346/004	035884497	SIMESA S.P.A.	IT
Sorvasta 15 mg apvalkotās tabletes	not available	11-0028	KRKA, D.D., NOVO MESTO	LV
Sorvasta 15 mg filmom obalené tablety	not available	31/0700/10-S	KRKA, D.D., NOVO MESTO	SK
Sorvasta 15 mg filmsko obložene tablete	not available	H/10/01448/005	KRKA, D.D., NOVO MESTO	SI
Sorvasta 15 mg filmsko obložene tablete	not available	H/10/01448/006	KRKA, D.D., NOVO MESTO	SI
Sorvasta 15 mg potahované tablety	not available	31/165/11-C	KRKA, D.D., NOVO MESTO	CZ
Sorvasta 30 mg apvalkotās tabletes	not available	11-0030	KRKA, D.D., NOVO MESTO	LV
Sorvasta 30 mg filmom obalené tablety	not available	31/0702/10-S	KRKA, D.D., NOVO MESTO	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sorvasta 30 mg filmsko obložene tablete	not available	H/10/01448/009	KRKA, D.D., NOVO MESTO	SI
Sorvasta 30 mg filmsko obložene tablete	not available	H/10/01448/010	KRKA, D.D., NOVO MESTO	SI
Sorvasta 30 mg potahované tablety	not available	31/167/11-C	KRKA, D.D., NOVO MESTO	CZ
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361283	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361382	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361481	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361580	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362083	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4360988	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361085	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361184	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362281	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos	NL/H/0345/001	4361986	ASTRAZENECA PRODUTOS FARMACEUTICOS 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				
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361689	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361788	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4360889	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361887	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362182	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361283	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361382	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361481	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361580	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362083	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4360988	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg	NL/H/0345/001	4361085	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361184	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362281	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361986	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361689	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361788	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4360889	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4361887	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 10 mg comprimidos revestidos por película	NL/H/0345/001	4362182	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362687	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362588	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363487	ASTRAZENECA PRODUTOS FARMACEUTICOS 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
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362984	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363180	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363289	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362380	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363388	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362489	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363586	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362885	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363685	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363784	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363081	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos	NL/H/0345/002	4362786	ASTRAZENECA PRODUTOS FARMACEUTICOS 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				
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362687	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362588	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363487	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362984	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363180	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363289	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362380	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363388	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362489	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363586	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362885	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg	NL/H/0345/002	4363685	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363784	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4363081	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 20 mg comprimidos revestidos por película	NL/H/0345/002	4362786	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365086	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4363982	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364089	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364287	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364188	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364386	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364485	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364584	ASTRAZENECA PRODUTOS FARMACEUTICOS 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
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364683	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364782	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4363883	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364881	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364980	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365086	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4363982	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364089	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364287	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos	NL/H/0345/003	4364188	ASTRAZENECA PRODUTOS FARMACEUTICOS 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				
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364386	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364485	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364584	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364683	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364782	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4363883	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364881	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4364980	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 40 mg comprimidos revestidos por película	NL/H/0345/003	4365185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5572284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg	NL/H/0345/004	5572086	ASTRAZENECA PRODUTOS	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			FARMACEUTICOS LDA	
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5570981	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571286	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571385	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571989	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571583	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571781	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571682	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5570882	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571880	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571187	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571088	ASTRAZENECA PRODUTOS FARMACEUTICOS 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
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5572185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571484	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5572284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5572086	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5570981	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571286	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571385	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571989	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571583	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571781	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571682	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos	NL/H/0345/004	5570882	ASTRAZENECA PRODUTOS FARMACEUTICOS 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				
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571880	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571187	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571088	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5572185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Visacor 5 mg comprimidos revestidos por película	NL/H/0345/004	5571484	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Xeter 15 mg filmtabletta	HU/H/0219/005	OGYI-T-21173/10	GEDEON RICHTER PLC.	HU
Xeter 15 mg filmtabletta	HU/H/0219/005	OGYI-T-21173/11	GEDEON RICHTER PLC.	HU
Xeter 30 mg filmtabletta	HU/H/0219/006	OGYI-T-21173/12	GEDEON RICHTER PLC.	HU
Xeter 30 mg filmtabletta	HU/H/0219/006	OGYI-T-21173/13	GEDEON RICHTER PLC.	HU
Xeter 5 mg filmtabletta	HU/H/0219/001	OGYI-T-21173/09	GEDEON RICHTER PLC.	HU
Xeter 5 mg filmtabletta	HU/H/0219/001	OGYI-T-21173/02	GEDEON RICHTER PLC.	HU
Xeter 5 mg filmtabletta	HU/H/0219/001	OGYI-T-21173/01	GEDEON RICHTER PLC.	HU
ZAHRON 15 mg filmom obalené tablety	PL/H/0422/001	31/0150/17-S	ZAKŁAD FARMACEUTYCZNY ADAMED PHARMA S.A.	SK
Zahron 15 mg potahované tablety	PL/H/0422/001	31/521/16-C	ZAKŁAD FARMACEUTYCZNY ADAMED PHARMA S.A.	CZ
ZAHRON 30 mg filmom obalené tablety	PL/H/0422/002	31/0151/17-S	ZAKŁAD FARMACEUTYCZNY ADAMED PHARMA S.A.	SK
Zahron 30 mg potahované tablety	PL/H/0422/002	31/522/16-C	ZAKŁAD FARMACEUTYCZNY ADAMED PHARMA S.A.	CZ
Zahron, 15 mg, tabletki powlekane	PL/H/0422/001	24048	ZAKŁAD FARMACEUTYCZNY ADAMED PHARMA S.A.	PL
Zahron, 30 mg, tabletki	PL/H/0422/002	24049	ZAKŁAD FARMACEUTYCZNY	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			ADAMED PHARMA S.A.	
Zaranta 15 mg apvalkotās tabletes	HU/H/0219/005	17-0148	GEDEON RICHTER PLC.	LV
Zaranta 30 mg apvalkotās tabletes	HU/H/0219/006	17-0149	GEDEON RICHTER PLC.	LV
ZARANTA, 15 mg õhukese polümeerikattega tabletid	HU/H/0219/005	950217	GEDEON RICHTER PLC.	EE
Zaranta, 15 mg, tabletki powlekane	HU/H/0219/005	24280	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
ZARANTA, 30 mg õhukese polümeerikattega tabletid	HU/H/0219/006	950317	GEDEON RICHTER PLC.	EE
Zaranta, 30 mg, tabletki powlekane	HU/H/0219/006	24281	GEDEON RICHTER POLSKA SP. Z. O.O.	PL
ЗАРАНТА 15 mg филмирани таблетки	HU/H/0219/005	20170287	GEDEON RICHTER PLC.	BG
ЗАРАНТА 30 mg филмирани таблетки	HU/H/0219/006	20170288	GEDEON RICHTER PLC.	BG
ЗАРАНТА 40 mg филмирани таблетки	HU/H/0219/004	20100158	GEDEON RICHTER PLC.	BG
ЗАРАНТА 5 mg филмирани таблетки	HU/H/0219/001	20100155	GEDEON RICHTER PLC.	BG
Крестор 10 mg филмирани таблетки	not available	20040104	ASTRAZENECA UK LIMITED	BG
Крестор 10 mg филмирани таблетки	not available	20040104	ASTRAZENECA UK LIMITED	BG
Крестор 20 mg филмирани таблетки	not available	20040105	ASTRAZENECA UK LIMITED	BG
Крестор 20 mg филмирани таблетки	not available	20040105	ASTRAZENECA UK LIMITED	BG
Крестор 5 mg филмирани таблетки	not available	20090346	ASTRAZENECA UK LIMITED	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
Крестор 5 mg филмирани таблетки	not available	20090346	ASTRAZENECA UK LIMITED	BG
Розвера 15 mg филмирани таблетки	DK/H/1938/003	20110458	KRKA, D.D., NOVO MESTO	BG
Розвера 30 mg филмирани таблетки	DK/H/1938/005	20110460	KRKA, D.D., NOVO MESTO	BG
Розувастатин Крка 15 mg филмирани таблетки	DK/H/1937/003	20110452	KRKA, D.D., NOVO MESTO	BG
Розувастатин Крка 30 mg филмирани таблетки	DK/H/1937/005	20110454	KRKA, D.D., NOVO MESTO	BG
СУЗАСТОР 5 MG ФИЛМИРАНИ ТАБЛЕТКИ	PT/H/0247/001	20100591	SANDOZ PHARMACEUTICALS D.D.	BG