14 March 2019
EMA/181063/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / rosuvastatin

Procedure no.: PSUSA/00010434/201807

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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
Zahron Combi 10 mg/5 mg tvrdé tobolky	PL/H/0462/001	83/899/16-C	ADAMED	CZ
Zahron Combi 10 mg/10 mg tvrdé tobolky	PL/H/0462/002	83/900/16-C	ADAMED	CZ
Zahron Combi 20 mg/5 mg tvrdé tobolky	PL/H/0462/003	83/901/16-C	ADAMED	CZ
Zahron Combi 20 mg/10 mg tvrdé tobolky	PL/H/0462/004	83/902/16-C	ADAMED	CZ
Zahron Combi, 10 mg + 5 mg, kapsułki, twarde	PL/H/0462/001	24661	ADAMED	PL
Zahron Combi, 10 mg + 10 mg, kapsułki, twarde	PL/H/0462/002	24662	ADAMED	PL
Zahron Combi, 20 mg + 5 mg, kapsułki, twarde	PL/H/0462/003	24663	ADAMED	PL
Zahron Combi, 20 mg + 10 mg, kapsułki, twarde	PL/H/0462/004	24664	ADAMED	PL
ZAHRON COMBI 10 mg/5 mg tvrdé kapsuly	PL/H/0462/001	58/0110/18-S	ADAMED	SK
ZAHRON COMBI 10 mg/10 mg tvrdé kapsuly	PL/H/0462/002	58/0111/18-S	ADAMED	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
ZAHRON COMBI 20 mg/5 mg tvrdé kapsuly	PL/H/0462/003	58/0112/18-S	ADAMED	SK
ZAHRON COMBI 20 mg/10 mg tvrdé kapsuly	PL/H/0462/004	58/0113/18-S	ADAMED	SK
Захрон Комби 10 mg/5 mg твърди капсули	PL/H/0462/001	20180190	ADAMED	BG
Захрон Комби 10 mg/10 mg твърди капсули	PL/H/0462/002	20180191	ADAMED	BG
Захрон Комби 20 mg/5 mg твърди капсули	PL/H/0462/003	20180192	ADAMED	BG
Захрон Комби 20 mg/10 mg твърди капсули	PL/H/0462/004	20180193	ADAMED	BG
Zahron Combi 10 mg/5 mg	PL/H/0462/001-004	275391137	ADAMED	HR
Zahron Combi 10 mg/10 mg	PL/H/0462/001-004	463840961	ADAMED	HR
Zahron Combi 20 mg/5 mg	PL/H/0462/001-004	977231010	ADAMED	HR
Zahron Combi 20 mg/10 mg	PL/H/0462/001-004	542074590	ADAMED	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Asutam 10 mg + 5 mg Cápsula	PL/H/0462/001	5744727	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Asutam 10 mg + 5 mg cápsula	PL/H/0462/001	5744735	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Asutam 10 mg + 10 mg cápsula	PL/H/0462/002	5745278	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Asutam 20 mg + 5 mg cápsula	PL/H/0462/003	5745310	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Asutam 20 mg + 10 mg cápsula	PL/H/0462/004	5745336	LUSOMEDICAMENTA – SOCIEDADE TÉCNICA FARMACÊUTICA, SA	PT
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/001	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/002	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/003	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
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/004	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/005	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/006	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/007	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/5 mg filmsko obložene tablete	HU/H/0388/001	H/15/2035/008	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/009	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/010	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/011	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/012	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/013	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/014	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/015	KRKA, D.D., NOVO MESTO	SI
Rosmela 10 mg/10 mg filmsko obložene tablete	HU/H/0388/002	H/15/2035/016	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/017	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/018	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/019	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/020	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/021	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/022	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/023	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
Rosmela 15 mg/5 mg filmsko obložene tablete	HU/H/0388/005	H/15/2035/024	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/025	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/026	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/027	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/028	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/029	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/030	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/031	KRKA, D.D., NOVO MESTO	SI
Rosmela 15 mg/10 mg filmsko obložene tablete	HU/H/0388/006	H/15/2035/032	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/033	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
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/034	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/035	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/036	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/037	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/038	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/039	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/5 mg filmsko obložene tablete	HU/H/0388/003	H/15/2035/040	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/041	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/042	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/043	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
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/044	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/045	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/046	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/047	KRKA, D.D., NOVO MESTO	SI
Rosmela 20 mg/10 mg filmsko obložene tablete	HU/H/0388/004	H/15/2035/048	KRKA, D.D., NOVO MESTO	SI
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/01	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/02	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/03	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/04	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/05	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/06	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/07	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmtabletta	HU/H/0388/001	OGYI-T-22894/08	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/09	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/10	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/11	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/12	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/13	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/14	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/15	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
Rosudapin 10 mg/10 mg filmtabletta	HU/H/0388/002	OGYI-T-22894/16	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/17	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/18	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/19	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/20	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/21	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/22	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/23	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/5 mg filmtabletta	HU/H/0388/003	OGYI-T-22894/24	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/25	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
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/26	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/27	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/28	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/29	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/30	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/31	KRKA, D.D., NOVO MESTO	HU
Rosudapin 15 mg/10 mg filmtabletta	HU/H/0388/004	OGYI-T-22894/32	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/33	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/34	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/35	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
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/36	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/37	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/38	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/39	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/5 mg filmtabletta	HU/H/0388/005	OGYI-T-22894/40	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/41	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/42	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/43	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/44	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/45	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
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/46	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/47	KRKA, D.D., NOVO MESTO	HU
Rosudapin 20 mg/10 mg filmtabletta	HU/H/0388/006	OGYI-T-22894/48	KRKA, D.D., NOVO MESTO	HU
Rosudapin 10 mg/5 mg filmom obalené tablety	HU/H/0388/001	58/0384/15-S	KRKA, D.D., NOVO MESTO	SK
Rosudapin 10 mg/10 mg filmom obalené tablety	HU/H/0388/002	58/0385/15-S	KRKA, D.D., NOVO MESTO	SK
Rosudapin 20 mg/5 mg filmom obalené tablety	HU/H/0388/005	58/0386/15-S	KRKA, D.D., NOVO MESTO	SK
Rosudapin 20 mg/10 mg filmom obalené tablety	HU/H/0388/006	58/0387/15-S	KRKA, D.D., NOVO MESTO	SK
Rosudapin 15 mg/5 mg filmom obalené tablety	HU/H/0388/003	58/0388/15-S	KRKA, D.D., NOVO MESTO	SK
Rosudapin 15 mg/10 mg filmom obalené tablety	HU/H/0388/004	58/0389/15-S	KRKA, D.D., NOVO MESTO	SK
Розудапин 10 mg/5 mg филмирани таблетки	HU/H/0388/001	20150304	KRKA, D.D., NOVO MESTO	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Розудапин 10 mg/10 mg филмирани таблетки	HU/H/0388/002	20150305	KRKA, D.D., NOVO MESTO	BG
Розудапин 20 mg/5 mg филмирани таблетки	HU/H/0388/005	20150306	KRKA, D.D., NOVO MESTO	BG
Розудапин 20 mg/10 mg филмирани таблетки	HU/H/0388/006	20150307	KRKA, D.D., NOVO MESTO	BG
Розудапин 15 mg/5 mg филмирани таблетки	HU/H/0388/003	20150308	KRKA, D.D., NOVO MESTO	BG
Розудапин 15 mg/10 mg филмирани таблетки	HU/H/0388/004	20150309	KRKA, D.D., NOVO MESTO	BG
Rosudapin 10 mg+5 mg apvalkotās tabletes	HU/H/0388/001	15-0244	KRKA, D.D., NOVO MESTO	LV
Rosudapin 10 mg+10 mg apvalkotās tabletes	HU/H/0388/002	15-0245	KRKA, D.D., NOVO MESTO	LV
Rosudapin 15 mg+5 mg apvalkotās tabletes	HU/H/0388/003	15-0246	KRKA, D.D., NOVO MESTO	LV
Rosudapin 15 mg+10 mg apvalkotās tabletes	HU/H/0388/004	15-0247	KRKA, D.D., NOVO MESTO	LV
Rosudapin 20 mg+5 mg apvalkotās tabletes	HU/H/0388/005	15-0248	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
Rosudapin 20 mg+10 mg apvalkotās tabletes	HU/H/0388/006	15-0249	KRKA, D.D., NOVO MESTO	LV
Rosvaden, 10 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0388/001	890115	KRKA, D.D., NOVO MESTO	EE
Rosvaden, 10 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0388/002	890215	KRKA, D.D., NOVO MESTO	EE
Rosvaden, 20 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0388/005	890315	KRKA, D.D., NOVO MESTO	EE
Rosvaden, 20 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0388/006	890415	KRKA, D.D., NOVO MESTO	EE
Rosvaden, 15 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0388/003	890515	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
Rosvaden, 15 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0388/004	890615	KRKA, D.D., NOVO MESTO	EE
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/001	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/002	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/003	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/004	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/005	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/006	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/007	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/5 mg plėvele dengtos tabletės	HU/H/0388/001	LT/1/15/3811/008	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/009	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
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/010	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/011	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/012	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/013	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/014	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/015	KRKA, D.D., NOVO MESTO	LT
Rosvaden 10 mg/10 mg plėvele dengtos tabletės	HU/H/0388/002	LT/1/15/3811/016	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/017	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/018	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/019	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
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/020	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/021	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/022	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/023	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/5 mg plėvele dengtos tabletės	HU/H/0388/003	LT/1/15/3811/024	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/025	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/026	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/027	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/028	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/029	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
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/030	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/031	KRKA, D.D., NOVO MESTO	LT
Rosvaden 15 mg/10 mg plėvele dengtos tabletės	HU/H/0388/004	LT/1/15/3811/032	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/033	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/034	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/035	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/036	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/037	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/038	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/039	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
Rosvaden 20 mg/5 mg plėvele dengtos tabletės	HU/H/0388/005	LT/1/15/3811/040	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/041	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/042	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/043	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/044	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/045	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/046	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/047	KRKA, D.D., NOVO MESTO	LT
Rosvaden 20 mg/10 mg plėvele dengtos tabletės	HU/H/0388/006	LT/1/15/3811/048	KRKA, D.D., NOVO MESTO	LT
Rosudapin, 10 mg + 5 mg, tabletki powlekane	HU/H/0388/001	22723	KRKA, D.D., NOVO MESTO	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
Rosudapin, 10 mg + 10 mg, tabletki powlekane	HU/H/0388/002	22724	KRKA, D.D., NOVO MESTO	PL
Rosudapin, 20 mg + 5 mg, tabletki powlekane	HU/H/0388/005	22725	KRKA, D.D., NOVO MESTO	PL
Rosudapin, 20 mg + 10 mg, tabletki powlekane	HU/H/0388/006	22726	KRKA, D.D., NOVO MESTO	PL
Rosudapin, 15 mg + 5 mg, tabletki powlekane	HU/H/0388/003	22727	KRKA, D.D., NOVO MESTO	PL
Rosudapin, 15 mg + 10 mg, tabletki powlekane	HU/H/0388/004	22728	KRKA, D.D., NOVO MESTO	PL
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/01	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/03	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/05	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
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/07	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/5 mg comprimate filmate	HU/H/0388/001	8265/2015/08	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/01	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/03	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/05	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/07	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosudapin 10 mg/10 mg comprimate filmate	HU/H/0388/002	8266/2015/08	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/01	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/03	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/05	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/07	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/5 mg comprimate filmate	HU/H/0388/005	8267/2015/08	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/01	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/03	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/05	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/07	KRKA, D.D., NOVO MESTO	RO
Rosudapin 15 mg/10 mg comprimate filmate	HU/H/0388/006	8268/2015/08	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/01	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/03	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
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/05	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/07	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/5 mg comprimate filmate	HU/H/0388/003	8269/2015/08	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/01	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/02	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/03	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/04	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/05	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
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/06	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/07	KRKA, D.D., NOVO MESTO	RO
Rosudapin 20 mg/10 mg comprimate filmate	HU/H/0388/004	8270/2015/08	KRKA, D.D., NOVO MESTO	RO
Roxuten 10 mg/10 mg comprimidos recubiertos con película	ES/H/0320/002	80564	KRKA, D.D., NOVO MESTO	ES
Roxuten 10 mg/5 mg comprimidos recubiertos con película	ES/H/0320/001	80565	KRKA, D.D., NOVO MESTO	ES
Roxuten 20 mg/5 mg comprimidos recubiertos con película	ES/H/0320/003	80566	KRKA, D.D., NOVO MESTO	ES
Roxuten 20 mg/10 mg comprimidos recubiertos con película	ES/H/0320/004	80567	KRKA, D.D., NOVO MESTO	ES
Rosuvastatin/Amlodipin Klinge 10 mg/5 mg Hartkapseln	PL/H/0463/001	98885.00.00	KLINGE PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosuvastatin/Amlodipin Klinge 10 mg/10 mg Hartkapseln	PL/H/0463/002	98886.00.00	KLINGE PHARMA GMBH	DE
Rosuvastatin/Amlodipin Klinge 20 mg/5 mg Hartkapseln	PL/H/0463/003	98887.00.00	KLINGE PHARMA GMBH	DE
Rosuvastatin/Amlodipin Klinge 20 mg/10 mg Hartkapseln	PL/H/0463/004	98888.00.00	KLINGE PHARMA GMBH	DE
Rosuvastatina + amlodipina Krka 10 mg + 5 mg comprimidos revestidos por película	HU/H/0396/001	5671425	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 10 mg + 5 mg comprimidos revestidos por película	HU/H/0396/001	5671433	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 10 mg + 10 mg comprimidos revestidos por película	HU/H/0396/002	5671441	KRKA, D.D., NOVO MESTO	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosuvastatina + amlodipina Krka 10 mg + 10 mg comprimidos revestidos por película	HU/H/0396/002	5671458	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 20 mg + 5 mg comprimidos revestidos por película	HU/H/0396/005	5671466	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 20 mg + 5 mg comprimidos revestidos por película	HU/H/0396/005	5671474	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 20 mg + 10 mg comprimidos revestidos por película	HU/H/0396/006	5671508	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 20 mg + 10 mg comprimidos revestidos por película	HU/H/0396/006	5671516	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 15 mg + 5 mg comprimidos revestidos por película	HU/H/0396/003	5671524	KRKA, D.D., NOVO MESTO	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosuvastatina + amlodipina Krka 15 mg + 5 mg comprimidos revestidos por película	HU/H/0396/003	5671532	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 15 mg + 10 mg comprimidos revestidos por película	HU/H/0396/004	5671540	KRKA, D.D., NOVO MESTO	PT
Rosuvastatina + amlodipina Krka 15 mg + 10 mg comprimidos revestidos por película	HU/H/0396/004	5671557	KRKA, D.D., NOVO MESTO	PT
Rosudapin 10 mg/5 mg potahované tablety	HU/H/0396/001	83/039/16-C	KRKA, D.D., NOVO MESTO	CZ
Rosudapin 10 mg/10 mg potahované tablety	HU/H/0396/002	83/040/16-C	KRKA, D.D., NOVO MESTO	CZ
Rosudapin 20 mg/5 mg potahované tablety	HU/H/0396/005	83/041/16-C	KRKA, D.D., NOVO MESTO	CZ
Rosudapin 20 mg/10 mg potahované tablety	HU/H/0396/006	83/042/16-C	KRKA, D.D., NOVO MESTO	CZ
Rosudapin 15 mg/5 mg potahované tablety	HU/H/0396/003	83/043/16-C	KRKA, D.D., NOVO MESTO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosudapin 15 mg/10 mg potahované tablety	HU/H/0396/004	83/044/16-C	KRKA, D.D., NOVO MESTO	CZ
Rosudapin 10 mg/5 mg kalvopäällysteiset tabletit	HU/H/0396/001	32783	KRKA, D.D., NOVO MESTO	FI
Rosudapin 10 mg/10 mg kalvopäällysteiset tabletit	HU/H/0396/002	32784	KRKA, D.D., NOVO MESTO	FI
Rosudapin 20 mg/5 mg kalvopäällysteiset tabletit	HU/H/0396/005	32785	KRKA, D.D., NOVO MESTO	FI
Rosudapin 20 mg/10 mg kalvopäällysteiset tabletit	HU/H/0396/006	32786	KRKA, D.D., NOVO MESTO	FI
Rosudapin 15 mg/5 mg kalvopäällysteiset tabletit	HU/H/0396/003	32787	KRKA, D.D., NOVO MESTO	FI
Rosudapin 15 mg/10 mg kalvopäällysteiset tabletit	HU/H/0396/004	32788	KRKA, D.D., NOVO MESTO	FI
Rosuvastatin/amlodipine Krka 10 mg/5 mg film-coated tablets	HU/H/0396/001	PA1347/053/001	KRKA, D.D., NOVO MESTO	IE
Rosuvastatin/amlodipine Krka 10 mg/10 mg film-coated tablets	HU/H/0396/002	PA1347/053/002	KRKA, D.D., NOVO MESTO	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
Rosuvastatin/amlodipine Krka 20 mg/5 mg film-coated tablets	HU/H/0396/005	PA1347/053/003	KRKA, D.D., NOVO MESTO	IE
Rosuvastatin/amlodipine Krka 20 mg/10 mg film-coated tablets	HU/H/0396/006	PA1347/053/004	KRKA, D.D., NOVO MESTO	IE
Rosuvastatin/amlodipine Krka 15 mg/5 mg film-coated tablets	HU/H/0396/003	PA1347/053/005	KRKA, D.D., NOVO MESTO	IE
Rosuvastatin/amlodipine Krka 15 mg/10 mg film-coated tablets	HU/H/0396/004	PA1347/053/006	KRKA, D.D., NOVO MESTO	IE
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/01	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/02	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/03	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/04	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/05	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/06	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/07	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/5 mg filmtabletta	HU/H/0396/001	OGYI-T-22976/08	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/09	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/10	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/11	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/12	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/13	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/14	KRKA, D.D., NOVO MESTO	HU
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/15	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
Rosvaden 10 mg/10 mg filmtabletta	HU/H/0396/002	OGYI-T-22976/16	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/17	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/18	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/19	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/20	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/21	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/22	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/23	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/5 mg filmtabletta	HU/H/0396/003	OGYI-T-22976/24	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/25	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
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/26	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/27	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/28	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/29	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/30	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/31	KRKA, D.D., NOVO MESTO	HU
Rosvaden 15 mg/10 mg filmtabletta	HU/H/0396/004	OGYI-T-22976/32	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/33	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/34	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/35	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
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/36	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/37	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/38	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/39	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/5 mg filmtabletta	HU/H/0396/005	OGYI-T-22976/40	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/41	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/42	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/43	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/44	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/45	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
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/46	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/47	KRKA, D.D., NOVO MESTO	HU
Rosvaden 20 mg/10 mg filmtabletta	HU/H/0396/006	OGYI-T-22976/48	KRKA, D.D., NOVO MESTO	HU
Rosuvastatine/amlodipine Krka 10 mg/5 mg filmomhulde tabletten	HU/H/0396/001	BE489697	KRKA, D.D., NOVO MESTO	BE
Rosuvastatine/amlodipine Krka 10 mg/10 mg filmomhulde tabletten	HU/H/0396/002	BE489706	KRKA, D.D., NOVO MESTO	BE
Rosuvastatine/amlodipine Krka 15 mg/5 mg filmomhulde tabletten	HU/H/0396/003	BE489715	KRKA, D.D., NOVO MESTO	BE
Rosuvastatine/amlodipine Krka 15 mg/10 mg filmomhulde tabletten	HU/H/0396/004	BE489724	KRKA, D.D., NOVO MESTO	BE
Rosuvastatine/amlodipine Krka 20 mg/5 mg filmomhulde tabletten	HU/H/0396/005	BE489733	KRKA, D.D., NOVO MESTO	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/amlodipine Krka 20 mg/10 mg filmomhulde tabletten	HU/H/0396/006	BE489742	KRKA, D.D., NOVO MESTO	BE
Roxuten 10 mg/5 mg Filmtabletten	HU/H/0396/001	137363	KRKA, D.D., NOVO MESTO	AT
Roxuten 10 mg/10 mg Filmtabletten	HU/H/0396/002	137364	KRKA, D.D., NOVO MESTO	AT
Roxuten 15 mg/5 mg Filmtabletten	HU/H/0396/005	137365	KRKA, D.D., NOVO MESTO	AT
Roxuten 15 mg/10 mg Filmtabletten	HU/H/0396/006	137366	KRKA, D.D., NOVO MESTO	AT
Roxuten 20 mg/5 mg Filmtabletten	HU/H/0396/003	137367	KRKA, D.D., NOVO MESTO	AT
Roxuten 20 mg/10 mg Filmtabletten	HU/H/0396/004	137368	KRKA, D.D., NOVO MESTO	AT