5 September 2024
EMA/483180/2024
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / losartan

Procedure no.: PSUSA/00010512/202401

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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
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	5716477	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/001	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	5716501	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/001	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	5716519	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/001	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/001	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	5716626	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/002	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	5716659	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/002	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	5716667	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/002	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 50 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/002	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	5716816	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/003	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	5716824	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/003	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	5716832	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/003	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 5 mg comprimidos revestidos por película	not available	16/H/0034/003	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	5716840	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/004	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	5716857	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/004	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	5716865	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/004	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Blocan 100 mg + 10 mg comprimidos revestidos por película	not available	16/H/0034/004	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/001	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/002	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/003	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/004	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/005	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/006	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/007	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/5 mg filmsko obložene tablete	HU/H/0375/001	H/16/02205/008	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/009	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/010	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/011	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/012	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/013	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/014	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/015	KRKA, D.D., NOVO MESTO	SI
Alortya 50 mg/10 mg filmsko obložene tablete	HU/H/0375/002	H/16/02205/016	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/017	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
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/018	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/019	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/020	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/021	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/022	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/023	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/5 mg filmsko obložene tablete	HU/H/0375/003	H/16/02205/024	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/025	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/026	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/027	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/028	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/029	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/030	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/031	KRKA, D.D., NOVO MESTO	SI
Alortya 100 mg/10 mg filmsko obložene tablete	HU/H/0375/004	H/16/02205/032	KRKA, D.D., NOVO MESTO	SI
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/01	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/02	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/03	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
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/04	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/05	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/06	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/07	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/5 mg filmtabletta	HU/H/0375/001	OGYI-T-23039/08	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/09	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/10	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/11	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/12	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
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/13	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/14	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/15	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 50 mg/10 mg filmtabletta	HU/H/0375/002	OGYI-T-23039/16	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/17	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/18	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/19	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/20	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/21	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
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/22	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/23	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/5 mg filmtabletta	HU/H/0375/003	OGYI-T-23039/24	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/25	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/26	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/27	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/28	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/29	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/30	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
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/31	KRKA, D.D., NOVO MESTO	HU
Losartan/Amlodipin Krka 100 mg/10 mg filmtabletta	HU/H/0375/004	OGYI-T-23039/32	KRKA, D.D., NOVO MESTO	HU
LosAmlo® 50 mg/5 mg Filmtabletten	HU/H/0375/001	92515.00.00	KRKA, D.D., NOVO MESTO	DE
LosAmlo® 50 mg/10 mg Filmtabletten	HU/H/0375/002	92516.00.00	KRKA, D.D., NOVO MESTO	DE
LosAmlo® 100 mg/5 mg Filmtabletten	HU/H/0375/003	92517.00.00	KRKA, D.D., NOVO MESTO	DE
LosAmlo® 100 mg/10 mg Filmtabletten	HU/H/0375/004	92518.00.00	KRKA, D.D., NOVO MESTO	DE
Tenloris 50 mg/5 mg apvalkotās tabletes	HU/H/0350/001	14-0071	KRKA, D.D., NOVO MESTO	LV
Tenloris 50 mg/10 mg apvalkotās tabletes	HU/H/0350/002	14-0072	KRKA, D.D., NOVO MESTO	LV
Tenloris 100 mg/5 mg apvalkotās tabletes	HU/H/0350/003	14-0073	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
Tenloris 100 mg/10 mg apvalkotās tabletes	HU/H/0350/004	14-0074	KRKA, D.D., NOVO MESTO	LV
Тенлорис 50 mg/5 mg филмирани таблетки	HU/H/0350/001	20140104	KRKA, D.D., NOVO MESTO	BG
Тенлорис 50 mg/10 mg филмирани таблетки	HU/H/0350/002	20140105	KRKA, D.D., NOVO MESTO	BG
Тенлорис 100 mg/5 mg филмирани таблетки	HU/H/0350/003	20140106	KRKA, D.D., NOVO MESTO	BG
Тенлорис 100 mg/10 mg филмирани таблетки	HU/H/0350/004	20140107	KRKA, D.D., NOVO MESTO	BG
Alortia, 50 mg + 5 mg, tabletki powlekane	HU/H/0350/001	21710	KRKA, D.D., NOVO MESTO	PL
Alortia, 50 mg + 10 mg, tabletki powlekane	HU/H/0350/002	21711	KRKA, D.D., NOVO MESTO	PL
Alortia, 100 mg + 5 mg, tabletki powlekane	HU/H/0350/003	21712	KRKA, D.D., NOVO MESTO	PL
Alortia, 100 mg + 10 mg, tabletki powlekane	HU/H/0350/004	21713	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
Tenloris 50 mg/5 mg filmom obalené tablety	HU/H/0350/001	58/0091/14-S	KRKA, D.D., NOVO MESTO	SK
Tenloris 50 mg/10 mg filmom obalené tablety	HU/H/0350/002	58/0092/14-S	KRKA, D.D., NOVO MESTO	SK
Tenloris 100 mg/5 mg filmom obalené tablety	HU/H/0350/003	58/0093/14-S	KRKA, D.D., NOVO MESTO	SK
Tenloris 100 mg/10 mg filmom obalené tablety	HU/H/0350/004	58/0094/14-S	KRKA, D.D., NOVO MESTO	SK
Tenloris 100 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0350/003	837614	KRKA, D.D., NOVO MESTO	EE
Tenloris 100 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0350/004	837714	KRKA, D.D., NOVO MESTO	EE
Tenloris 50 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0350/001	837814	KRKA, D.D., NOVO MESTO	EE
Tenloris 50 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0350/002	837914	KRKA, D.D., NOVO MESTO	EE
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/001	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
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/002	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/003	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/004	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/005	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/006	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/007	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/008	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/009	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/010	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/011	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/012	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/013	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/014	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/015	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/016	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/017	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/018	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/019	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
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/020	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/021	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/022	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/023	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/024	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/5 mg plėvele dengtos tabletės	HU/H/0350/001	LT/1/14/3512/025	KRKA, D.D., NOVO MESTO	LT
Tenloris 50 mg/10 mg plėvele dengtos tabletės	HU/H/0350/002	LT/1/14/3512/026	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/5 mg plėvele dengtos tabletės	HU/H/0350/003	LT/1/14/3512/027	KRKA, D.D., NOVO MESTO	LT
Tenloris 100 mg/10 mg plėvele dengtos tabletės	HU/H/0350/004	LT/1/14/3512/028	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
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/01	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/02	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/03	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/04	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/05	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/06	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/07	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/08	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/09	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
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/10	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/11	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/12	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/13	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/14	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/15	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/16	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/17	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/18	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
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/19	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/20	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/21	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/22	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/23	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/24	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/5 mg filmtabletta	HU/H/0350/001	OGYI-T-22612/25	KRKA, D.D., NOVO MESTO	HU
Tenloris 50 mg/10 mg filmtabletta	HU/H/0350/002	OGYI-T-22612/26	KRKA, D.D., NOVO MESTO	HU
Tenloris 100 mg/5 mg filmtabletta	HU/H/0350/003	OGYI-T-22612/27	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
Tenloris 100 mg/10 mg filmtabletta	HU/H/0350/004	OGYI-T-22612/28	KRKA, D.D., NOVO MESTO	HU