1 December 2022
EMA/934243/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / candesartan

Procedure no.: PSUSA/00010191/202204

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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
LODICAND 8 mg/5 mg capsule rigide	not available	045122013	BRUNO FARMACEUTICI	IT
LODICAND 8 mg/10 mg capsule rigide	not available	045122037	BRUNO FARMACEUTICI	IT
LODICAND 16 mg/5 mg capsule rigide	not available	045122025	BRUNO FARMACEUTICI	IT
LODICAND 16 mg/10 mg capsule rigide	not available	045122049	BRUNO FARMACEUTICI	IT
Candezek Combi, 8 mg + 5 mg, kapsułki, twarde	PL/H/0379/001	23187	ADAMED PHARMA S.A.	PL
Candezek Combi, 8 mg + 10 mg, kapsułki, twarde	PL/H/0379/002	23188	ADAMED PHARMA S.A.	PL
Candezek Combi, 16 mg + 5 mg, kapsułki, twarde	PL/H/0379/003	23189	ADAMED PHARMA S.A.	PL
Candezek Combi, 16 mg + 10 mg, kapsułki, twarde	PL/H/0379/004	23190	ADAMED PHARMA S.A.	PL
Candezek 8 mg/5 mg tvrdé kapsuly	PL/H/0379/001	58/0315/16-S	ADAMED PHARMA S.A.	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
Candezek 8 mg/10 mg tvrdé kapsuly	PL/H/0379/002	58/0316/16-S	ADAMED PHARMA S.A.	SK
Candezek 16 mg/5 mg tvrdé kapsuly	PL/H/0379/003	58/0317/16-S	ADAMED PHARMA S.A.	SK
Candezek 16 mg/10 mg tvrdé kapsuly	PL/H/0379/004	58/0318/16-S	ADAMED PHARMA S.A.	SK
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/01	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/09	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/17	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/25	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/18	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/20	ADAMED PHARMA S.A.	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
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/19	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/21	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/22	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/23	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/5 mg kemény kapszula	PL/H/0379/003	OGYI-T-23045/24	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/26	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/28	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/27	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/29	ADAMED PHARMA S.A.	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
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/30	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/31	ADAMED PHARMA S.A.	HU
Candezek Plus 16 mg/10 mg kemény kapszula	PL/H/0379/004	OGYI-T-23045/32	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/10	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/15	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/14	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/12	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/11	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/13	ADAMED PHARMA S.A.	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
Candezek Plus 8 mg/10 mg kemény kapszula	PL/H/0379/002	OGYI-T-23045/16	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/02	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/05	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/04	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/03	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/06	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/07	ADAMED PHARMA S.A.	HU
Candezek Plus 8 mg/5 mg kemény kapszula	PL/H/0379/001	OGYI-T-23045/08	ADAMED PHARMA S.A.	HU
Candezek 8 mg/5 mg tvrdé tobolky	PL/H/0379/001	58/382/16-C	ADAMED PHARMA S.A.	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
Candezek 16 mg/5 mg tvrdé tobolky	PL/H/0379/003	58/384/16-C	ADAMED PHARMA S.A.	CZ
Candezek 16 mg/10 mg tvrdé tobolky	PL/H/0379/004	58/385/16-C	ADAMED PHARMA S.A.	CZ
Кандезек Комби 8 mg/ 5 mg твърди капсули	PL/H/0379/001	20160373	ADAMED PHARMA S.A.	BG
Кандезек Комби 8 mg/ 10 mg твърди капсули	PL/H/0379/002	20160374	ADAMED PHARMA S.A.	BG
Кандезек Комби 16 mg/ 5 mg твърди капсули	PL/H/0379/003	20160375	ADAMED PHARMA S.A.	BG
Кандезек Комби 16 mg/ 10 mg твърди капсули	PL/H/0379/004	20160376	ADAMED PHARMA S.A.	BG
Caramlo 16 mg/5 mg Tabletten	DE/H/5736/001	2202564.00.00	ZENTIVA PHARMA GMBH	DE
Candeblo® Amlo 8 mg/5 mg-Tabletten	DE/H/6798/001	141202	G.L. PHARMA GMBH	AT
Candeblo® Amlo 16 mg/5 mg-Tabletten	DE/H/6798/002	141203	G.L. PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Candeblo® Amlo 16 mg/10 mg-Tabletten	DE/H/6798/003	141204	G.L. PHARMA GMBH	AT
Bilamcar	BG/H/0111/001/DC	20190005	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Bilamcar 8 mg/10 mg hard capsules	BG/H/0111/002/DC	20190006	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Bilamcar 16 mg/5 mg hard capsules	BG/H/0111/003/DC	20190007	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Bilamcar 16 mg/10 mg hard capsules	BG/H/0111/004/DC	20190008	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
CANDETENS 8 mg/5 mg capsule rigide	not available	045214018	ERREKAPPA EUROTERAPICI SPA	IT
CANDETENS 8 mg/10 mg capsule rigide	not available	045214032	ERREKAPPA EUROTERAPICI SPA	IT
CANDETENS 16 mg/5 mg capsule rigide	not available	045214020	ERREKAPPA EUROTERAPICI SPA	IT
CANDETENS 16 mg/10 mg capsule rigide	not available	045214044	ERREKAPPA EUROTERAPICI SPA	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
DESARPIN 8 mg + 5 mg, καψάκιο σκληρό	PL/H/0379/001	79415/31-08-2021	RAFARM SA.	GR
DESARPIN 16 mg + 5 mg, καψάκιο σκληρό	PL/H/0379/003	79415/31-08-2021	RAFARM SA.	GR
DESARPIN 8 mg + 10 mg, καψάκιο σκληρό	PL/H/0379/002	79415/31-08-2021	RAFARM SA.	GR
DESARPIN 16 mg + 10 mg, καψάκιο σκληρό	PL/H/0379/004	79415/31-08-2021	RAFARM SA.	GR
Camlostar 16 mg/10 mg Hartkapseln	DE/H/4527/004	95574.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Camlostar 16 mg/5 mg Hartkapseln	DE/H/4527/002	95573.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Camlostar 8 mg/10 mg Hartkapseln	DE/H/4527/003	95572.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Camlostar 8 mg/5 mg Hartkapseln	DE/H/4527/001	95571.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Amlodipina + Candesartan Krka 5 mg + 8 mg comprimidos	DE/H/5108/001	5737366	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
Amlodipina + Candesartan Krka 5 mg + 8 mg comprimidos	DE/H/5108/001	5737374	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 8 mg comprimidos	DE/H/5108/001	5737408	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 8 mg comprimidos	DE/H/5108/001	5737416	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 16 mg comprimidos	DE/H/5108/002	5737424	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 16 mg comprimidos	DE/H/5108/002	5737432	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 16 mg comprimidos	DE/H/5108/002	5737440	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 5 mg + 16 mg comprimidos	DE/H/5108/002	5737457	KRKA, D.D., NOVO MESTO	PT
Camdero, 8 mg/5 mg tabletid	DE/H/5108/001	958918	KRKA, D.D., NOVO MESTO	EE
Camdero, 16 mg/5 mg tabletid	DE/H/5108/002	959018	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
Candesartan/Amlodipine Krka 8 mg/5 mg tabletten	DE/H/5108/001	BE525502	KRKA, D.D., NOVO MESTO	BE
Candesartan/Amlodipine Krka 16 mg/5 mg tabletten	DE/H/5108/002	BE525511	KRKA, D.D., NOVO MESTO	BE
Camdero 8 mg/5 mg tablety	DE/H/5108/001	58/951/16-C	KRKA, D.D., NOVO MESTO	CZ
Camdero 16 mg/5 mg tablety	DE/H/5108/002	58/952/16-C	KRKA, D.D., NOVO MESTO	CZ
Camlocor 8 mg/5 mg tabletes	DE/H/5108/001	18-0043	KRKA, D.D., NOVO MESTO	LV
Camlocor 16 mg/5 mg tabletes	DE/H/5108/002	18-0044	KRKA, D.D., NOVO MESTO	LV
Kandoset 8 mg/5 mg tablety	DE/H/5108/001	58/0150/18-S	KRKA, D.D., NOVO MESTO	SK
Kandoset 16 mg/5 mg tablety	DE/H/5108/002	58/0151/18-S	KRKA, D.D., NOVO MESTO	SK
Camlocor, 8 mg + 5 mg, tabletki	DE/H/5108/001	24633	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
Camlocor, 16 mg + 5 mg, tabletki	DE/H/5108/002	24634	KRKA, D.D., NOVO MESTO	PL
Candecam 8 mg/5 mg tabletit	DE/H/5108/001	34770	KRKA, D.D., NOVO MESTO	FI
Candecam 16 mg/5 mg tabletit	DE/H/5108/002	34771	KRKA, D.D., NOVO MESTO	FI
Кандосет 8 mg/5 mg таблетки	DE/H/5108/001	20180074	KRKA, D.D., NOVO MESTO	BG
Кандосет 16 mg/5 mg таблетки	DE/H/5108/002	20180075	KRKA, D.D., NOVO MESTO	BG
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/001	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/002	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/003	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/004	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
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/005	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/006	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/007	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/008	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/009	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/010	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/011	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/012	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/013	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
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/014	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/015	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/016	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/017	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/018	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/019	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/020	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/021	KRKA, D.D., NOVO MESTO	SI
Camlocor 8 mg/5 mg tablete	DE/H/5108/001	H/18/02440/022	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
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/023	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/024	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/025	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/026	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/027	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/028	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/029	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/030	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/031	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
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/032	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/033	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/034	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/035	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/036	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/037	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/038	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/039	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/040	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
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/041	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/042	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/043	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/5 mg tablete	DE/H/5108/002	H/18/02440/044	KRKA, D.D., NOVO MESTO	SI
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/01	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/02	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/03	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/04	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/05	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
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/06	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/07	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/08	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/09	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/10	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/11	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/12	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/13	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/14	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
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/15	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/16	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/17	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/18	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/19	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/20	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/21	KRKA, D.D., NOVO MESTO	RO
Kandoset 8 mg/5 mg comprimate	DE/H/5108/001	10673/2018/22	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/01	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
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/02	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/03	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/04	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/05	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/06	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/07	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/08	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/09	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/10	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
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/11	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/12	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/13	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/14	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/15	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/16	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/17	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/18	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/19	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
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/20	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/21	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimate	DE/H/5108/002	10674/2018/22	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/5 mg comprimidos	DE/H/5108/002	83016	KRKA, D.D., NOVO MESTO	ES
Kandoset 8 mg/5 mg comprimidos	DE/H/5108/001	83017	KRKA, D.D., NOVO MESTO	ES
Candesartan/Amlodipin TAD 8 mg/5 mg Tabletten	DE/H/5108/001	99020.00.00	TAD PHARMA GMBH	DE
Candesartan/Amlodipin TAD 16 mg/5 mg Tabletten	DE/H/5108/002	99021.00.00	TAD PHARMA GMBH	DE
Candecam® 8 mg/5 mg Tabletten	DE/H/5108/001	138112	HCS BVBA	AT
Candecam® 16 mg/5 mg Tabletten	DE/H/5108/002	138113	HCS BVBA	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kandoset 16 mg/10 mg comprimidos	DE/H/5108/005	84728	KRKA, D.D., NOVO MESTO	ES
Candecam® 16 mg/10 mg Tabletten	DE/H/5108/005	139309	HCS BVBA	AT
Camlocor, 16 mg + 10 mg, tabletki	DE/H/5108/005	25684	KRKA, D.D., NOVO MESTO	PL
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/045	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/046	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/047	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/048	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/049	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/050	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
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/051	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/052	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/053	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/054	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/055	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/056	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/057	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/058	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/059	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
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/060	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/061	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/062	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/063	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/064	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/065	KRKA, D.D., NOVO MESTO	SI
Camlocor 16 mg/10 mg tablete	DE/H/5108/005	H/18/02440/066	KRKA, D.D., NOVO MESTO	SI
Candesartan/Amlodipin TAD 16 mg/ 10 mg Tabletten	DE/H/5108/005	2203030.00.00	TAD PHARMA GMBH	DE
Candecam 16 mg/10 mg tabletit	DE/H/5108/005	36420	KRKA, D.D., NOVO MESTO	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
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/01	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/02	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/03	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/04	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/05	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/06	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/07	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/08	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/09	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
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/10	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/11	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/12	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/13	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/14	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/15	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/16	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/17	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/18	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
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/19	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/20	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/21	KRKA, D.D., NOVO MESTO	RO
Kandoset 16 mg/10 mg comprimate	DE/H/5108/005	12900/2020/22	KRKA, D.D., NOVO MESTO	RO
Camdero 16 mg/10 mg tablety	DE/H/5108/005	58/431/18-C	KRKA, D.D., NOVO MESTO	CZ
Kandoset 16 mg/10 mg tablety	DE/H/5108/005	58/0032/20-S	KRKA, D.D., NOVO MESTO	SK
Amlodipina + Candesartan Krka 10 mg + 16 mg comprimidos	DE/H/5108/005	5811369	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 10 mg + 16 mg comprimidos	DE/H/5108/005	5811377	KRKA, D.D., NOVO MESTO	PT
Amlodipina + Candesartan Krka 10 mg + 16 mg comprimidos	DE/H/5108/005	5811401	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
Amlodipina + Candesartan Krka 10 mg + 16 mg comprimidos	DE/H/5108/005	5811419	KRKA, D.D., NOVO MESTO	PT
Caramlo, 8 mg+5mg, tabletki	DE/H/3677/001	22096	ZENTIVA, K.S.	PL
Zenicamo 8 mg/ 5 mg tabletes	DE/H/3677/001	14-0200	ZENTIVA, K.S.	LV
Carzap AM 8 mg + 5 mg comprimidos	DE/H/3677/001	5610639	ZENTIVA PORTUGAL, LDA	PT
Carzap AM 8 mg + 5 mg comprimidos	DE/H/3677/001	5610654	ZENTIVA PORTUGAL, LDA	PT
Carzap AM 8 mg + 5 mg comprimidos	DE/H/3677/001	5610647	ZENTIVA PORTUGAL, LDA	PT
Карамло 8 mg/5 mg таблетки	DE/H/3677/001	20140354	ZENTIVA, K.S.	BG
Caramlo 8 mg/5 mg tablety	DE/H/3677/001	58/246/14-C	ZENTIVA, K.S.	CZ
Caramlo (8 + 5) mg δισκία	DE/H/3677/001	22063	ZENTIVA, K.S.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/03	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/06	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/07	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/05	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/04	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/01	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg comprimate	DE/H/3677/001	12225/2019/02	ZENTIVA, K.S.	RO
Caramlo 8 mg/5 mg Tabletten	DE/H/3677/001	88639.00.00	ZENTIVA PHARMA GMBH	DE
Caramlo (8 + 5) mg δισκία	DE/H/3677/001	2926/20-01-2021	ZENTIVA, K.S.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Bilamcar 8 mg/5 mg tvrdé kapsuly	BG/H/0111/001	58/0016/19-S	SWYSSI AG	SK
Bilamcar 8 mg/10 mg tvrdé kapsuly	BG/H/0111/002	58/0017/19-S	SWYSSI AG	SK
Bilamcar 16 mg/5 mg tvrdé kapsuly	BG/H/0111/003	58/0018/19-S	SWYSSI AG	SK
Bilamcar 16 mg/10 mg tvrdé kapsuly	BG/H/0111/004	58/0019/19-S	SWYSSI AG	SK
Bilamcar 8 mg/5 mg tvrdé tobolky	BG/H/0111/001	58/388/17-C	SWYSSI AG	CZ
Bilamcar 8 mg/10 mg tvrdé tobolky	BG/H/0111/002	58/389/17-C	SWYSSI AG	CZ
Bilamcar 16 mg/5 mg tvrdé tobolky	BG/H/0111/003	58/390/17-C	SWYSSI AG	CZ
Bilamcar 16 mg/10 mg tvrdé tobolky	BG/H/0111/004	58/391/17-C	SWYSSI AG	CZ
Bilamcar 5 mg + 8 mg cápsulas	BG/H/0111/001	5761549	SWYSSI AG	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
Bilamcar 10 mg + 8 mg cápsulas	BG/H/0111/002	5761648	SWYSSI AG	PT
Bilamcar 5 mg + 16 mg cápsulas	BG/H/0111/003	5761663	SWYSSI AG	PT
Bilamcar 10 mg + 16 mg cápsulas	BG/H/0111/004	5761705	SWYSSI AG	PT
Tilamcar 8 mg/5 mg Hartkapseln	BG/H/0111/001	138621	SWYSSI AG	AT
Tilamcar 8 mg/10 mg Hartkapseln	BG/H/0111/002	138622	SWYSSI AG	AT
Tilamcar 16 mg/5 mg Hartkapseln	BG/H/0111/003	138623	SWYSSI AG	AT
Tilamcar 16 mg/10 mg Hartkapseln	BG/H/0111/004	138624	SWYSSI AG	AT
Tilamcar 8 mg/5 mg capsule	BG/H/0111/001	11344/2019/01	SWYSSI AG	RO
Tilamcar 8 mg/10 mg capsule	BG/H/0111/002	11345/2019/01	SWYSSI AG	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
Tilamcar 16 mg/5 mg capsule	BG/H/0111/003	11346/2019/01	SWYSSI AG	RO
Tilamcar 16mg/10 mg capsule	BG/H/0111/004	11347/2019/01	SWYSSI AG	RO
Bilamcar 8 mg/5 mg, καψάκια, σκληρά	BG/H/0111/001	317410101	SWYSSI AG	GR
Bilamcar 8 mg/10 mg ??????? ??????	BG/H/0111/002/DC	317410201	SWYSSI AG	GR
Bilamcar 16 mg/5 mg ??????? ??????	BG/H/0111/003/DC	317410301	SWYSSI AG	GR
Bilamcar 16 mg/10 mg, καψάκια, σκληρά	BG/H/0111/004	317410401	SWYSSI AG	GR
CandeAmlo HEXAL 8 mg/5 mg Hartkapseln	DE/H/4526/001	95494.00.00	HEXAL AG	DE
CandeAmlo HEXAL 8 mg/10 mg Hartkapseln	DE/H/4526/003	95496.00.00	HEXAL AG	DE
CandeAmlo HEXAL 16 mg/5 mg Hartkapseln	DE/H/4526/002	95495.00.00	HEXAL AG	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
CandeAmlo HEXAL 16 mg/10 mg Hartkapseln	DE/H/4526/004	95497.00.00	HEXAL AG	DE
Candesartan/Amlodipin Sandoz 8 mg/5 mg – Hartkapseln	DE/H/4526/001	137212	SANDOZ GMBH	AT
Candesartan/Amlodipin Sandoz 8 mg/10 mg – Hartkapseln	DE/H/4526/003	137213	SANDOZ GMBH	AT
Candesartan/Amlodipin Sandoz 16 mg/10 mg – Hartkapseln	DE/H/4526/004	137215	SANDOZ GMBH	AT
Candesartan/Amlodipin Sandoz 16 mg/5 mg – Hartkapseln	DE/H/4526/002	137214	SANDOZ GMBH	AT
Framsyl, 8 mg/5 mg, kõvakapslid	DE/H/4526/001	915616	SANDOZ PHARMACEUTICALS D.D.	EE
Framsyl, 8 mg/10 mg, kõvakapslid	DE/H/4526/003	915716	SANDOZ PHARMACEUTICALS D.D.	EE
Framsyl, 16 mg/5 mg, kõvakapslid	DE/H/4526/002	915816	SANDOZ PHARMACEUTICALS D.D.	EE
Framsyl, 16 mg/10 mg, kõvakapslid	DE/H/4526/004	915916	SANDOZ PHARMACEUTICALS D.D.	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
Amcandin 8 mg/5 mg trde kapsule	DE/H/4526/001	H/16/02250/003	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 8 mg/5 mg trde kapsule	DE/H/4526/001	H/16/02250/002	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 8 mg/5 mg trde kapsule	DE/H/4526/001	H/16/02250/001	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 8 mg/10 mg trde kapsule	DE/H/4526/003	H/16/02250/005	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 8 mg/10 mg trde kapsule	DE/H/4526/003	H/16/02250/006	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 8 mg/10 mg trde kapsule	DE/H/4526/003	H/16/02250/004	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 16 mg/5 mg trde kapsule	DE/H/4526/002	H/16/02250/007	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 16 mg/5 mg trde kapsule	DE/H/4526/002	H/16/02250/008	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 16 mg/10 mg trde kapsule	DE/H/4526/004	H/16/02250/011	SANDOZ PHARMACEUTICALS D.D.	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
Amcandin 16 mg/5 mg trde kapsule	DE/H/4526/002	H/16/02250/009	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 16 mg/10 mg trde kapsule	DE/H/4526/004	H/16/02250/010	SANDOZ PHARMACEUTICALS D.D.	SI
Amcandin 16 mg/10 mg trde kapsule	DE/H/4526/004	H/16/02250/012	SANDOZ PHARMACEUTICALS D.D.	SI
Clodipan 8 mg/5 mg hard capsules	not available	MA1243/00501	ASPEN HEALTHCARE MALTA LIMITED	ES
Clodipan 8 mg/10 mg hard capsules	not available	MA1243/00502	ASPEN HEALTHCARE MALTA LIMITED	ES
Clodipan 16 mg/5 mg hard capsules	not available	MA1243/00503	ASPEN HEALTHCARE MALTA LIMITED	ES
Clodipan 16 mg/10 mg hard capsules	not available	MA1243/00504	ASPEN HEALTHCARE MALTA LIMITED	ES
Zenicamo, 16 mg/10 mg tabletid	DE/H/3677/003	847914	ZENTIVA, K.S.	EE
Zenicamo 16 mg/ 10 mg tabletes	DE/H/3677/003	14-0199	ZENTIVA, K.S.	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
Carzap AM 16 mg + 10 mg comprimidos	DE/H/3677/003	5610670	ZENTIVA PORTUGAL, LDA	PT
Carzap AM 16 mg + 10 mg comprimidos	DE/H/3677/003	5610704	ZENTIVA PORTUGAL, LDA	PT
Carzap AM 16 mg + 10 mg comprimidos	DE/H/3677/003	5610662	ZENTIVA PORTUGAL, LDA	PT
Карамло 16 mg/10 mg таблетки	DE/H/3677/003	20140355	ZENTIVA, K.S.	BG
Caramlo 16 mg/10 mg tablety	DE/H/3677/003	58/247/14-C	ZENTIVA, K.S.	CZ
Caramlo (16 + 10) mg δισκία	DE/H/3677/003	22064	ZENTIVA, K.S.	CY
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/03	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/06	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/05	ZENTIVA, K.S.	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
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/04	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/01	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/07	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg comprimate	DE/H/3677/003	12226/2019/02	ZENTIVA, K.S.	RO
Caramlo 16 mg/10 mg Tabletten	DE/H/3677/003	88641.00.00	ZENTIVA PHARMA GMBH	DE
Caramlo, 16 mg+10 mg, tabletki	DE/H/3677/003	22097	ZENTIVA, K.S.	PL
Caramlo (16 + 10) mg δισκία	DE/H/3677/003	2927/20-01-2021	ZENTIVA, K.S.	GR
Altensil 5 mg + 8 mg, cテ。psulas	PT/H/2428/001	5763859	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Altensil 5 mg + 8 mg, cテ。psulas	PT/H/2428/001	5763867	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	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
Altensil 5 mg + 16 mg, cápsulas	PT/H/2428/003	5763909	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Altensil 10 mg + 8 mg, cápsulas	PT/H/2428/002	5763875	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Altensil 10 mg + 16 mg, cápsulas	PT/H/2428/004	5763917	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Candesartan/Amlodipin TAD 8 mg/5 mg Tabletten	DE/H/5107/ 001	99016.00.00	TAD PHARMA GMBH	DE
Candesartan/Amlodipin TAD 16 mg/5 mg Tabletten	DE/H/5107/002	99017.00.00	TAD PHARMA GMBH	DE
Candecor-Amlo 16 mg/10 mg Tabletten	DE/H/5107/003	2203031.00.00	TAD PHARMA GMBH	DE
Caramlo 16 mg/5 mg tablety	DE/H/3677/004	58/283/18-C	ZENTIVA, K.S.	CZ
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/01	ZENTIVA, K.S.	RO
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/02	ZENTIVA, K.S.	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
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/03	ZENTIVA, K.S.	RO
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/04	ZENTIVA, K.S.	RO
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/05	ZENTIVA, K.S.	RO
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/06	ZENTIVA, K.S.	RO
Caramlo 16 mg/5 mg comprimate	DE/H/3677/004	12557/2019/07	ZENTIVA, K.S.	RO
Caramlo, 16 mg + 5 mg, tabletki	DE/H/3677/004	25620	ZENTIVA, K.S.	PL
Carzap Am 16 mg + 5 mg comprimidos	DE/H/3677/004	5783345	ZENTIVA PORTUGAL, LDA	PT
Carzap Am 16 mg + 5 mg comprimidos	DE/H/3677/004	5783352	ZENTIVA PORTUGAL, LDA	PT
Carzap Am 16 mg + 5 mg comprimidos	DE/H/3677/004	5783360	ZENTIVA PORTUGAL, 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
Carzap Am 16 mg + 5 mg comprimidos	DE/H/3677/004	5783923	ZENTIVA PORTUGAL, LDA	PT
Caramlo 16 mg/5 mg Tabletten	DE/H/3677/004	2202567.00.00	ZENTIVA PHARMA GMBH	DE
Zenicamo 16 mg/ 5 mg tabletes	DE/H/3677/004	19-0190	ZENTIVA, K.S.	LV
CandAm 8 mg/5 mg Hartkapseln	AT/H/0958/001	137187	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
CandAm 8 mg/10 mg Hartkapseln	AT/H/0958/002	137186	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
CandAm 16 mg/5 mg Hartkapseln	AT/H/0958/003	137185	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
CandAm 16 mg/10 mg Hartkapseln	AT/H/0958/004	137184	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/01	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/02	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
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/03	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/04	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/05	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/06	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/07	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/08	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/09	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/10	KRKA, D.D., NOVO MESTO	HU
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/11	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
Kandoset 8 mg/5 mg tabletta	not available	OGYI-T-23260/12	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/13	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/14	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/15	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/16	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/17	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/18	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/19	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/20	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
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/21	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/22	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/23	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/5 mg tabletta	not available	OGYI-T-23260/24	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/25	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/26	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/27	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/28	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/29	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
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/30	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/31	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/32	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/33	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/34	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/35	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/5 mg tabletta	not available	OGYI-T-23260/36	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/37	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/38	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
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/39	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/40	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/41	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/42	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/43	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/44	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/45	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/46	KRKA, D.D., NOVO MESTO	HU
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/47	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
Kandoset 32 mg/10 mg tabletta	not available	OGYI-T-23260/48	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/49	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/50	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/51	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/52	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/53	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/54	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/55	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/56	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
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/57	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/58	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/59	KRKA, D.D., NOVO MESTO	HU
Kandoset 16 mg/10 mg tabletta	not available	OGYI-T-23260/60	KRKA, D.D., NOVO MESTO	HU