29 November 2018
EMA/837411/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine besilate / ramipril

Procedure no.: PSUSA/00000181/201803

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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
Егирамлон 10 mg/10 mg твърди капсули	HU/H/0303/001-005/DC	20120049	EGIS PHARMACEUTICALS PLC	BG
Егирамлон 5 mg/5 mg твърди капсули	HU/H/0303/001-005/DC	20120046	EGIS PHARMACEUTICALS PLC	BG
Egiramlon 5 mg/5 mg tvrdé tobolky	HU/H/0303/001-005/DC	58/838/11-C	EGIS PHARMACEUTICALS PLC	CZ
Egiramlon 5 mg/5 mg kemény kapszula	HU/H/0303/002/DC	OGYI-T-22016/04-06, 16	EGIS PHARMACEUTICALS PLC	HU
Ramlon 5 mg/5 mg cietās kapsulas	HU/H/0303/001-005/DC	12-0064	EGIS PHARMACEUTICALS PLC	LV
Егирамлон 2,5 mg/2,5 mg твърди капсули	HU/H/0303/001-005/DC	20120045	EGIS PHARMACEUTICALS PLC	BG
Egiramlon 2,5 mg/2,5 mg kemény kapszula	HU/H/0303/001/DC	OGYI-T-22016/01-03	EGIS PHARMACEUTICALS PLC	HU
Ramlon 2,5 mg/2,5 mg cietās kapsulas	HU/H/0303/001-005/DC	12-0063	EGIS PHARMACEUTICALS PLC	LV
Ramlon 2,5 mg/2,5 mg kietosios kapsulės	HU/H/0303/001/DC	LT/1/12/2892/001-005	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 2,5 mg/2,5 mg capsule	HU/H/0303/001/DC	4495/2012/01-05	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 2,5 mg/2,5 mg Tvrdé kapsuly	HU/H/0303/001/DC	58/0743/11-S	EGIS PHARMACEUTICALS PLC	SK
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/002/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 5 mg/5 mg capsule	HU/H/0303/002/DC	4496/2012/01-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg Tvrdé kapsuly	HU/H/0303/002/DC	58/0744/11-S	EGIS PHARMACEUTICALS PLC	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
Егирамлон 10 mg/5 mg твърди капсули	HU/H/0303/003	20120048	EGIS PHARMACEUTICALS PLC	BG
Egiramlon 10 mg/5 mg tvrdé tobolky	HU/H/0303/001-005/DC	58/840/11-C	EGIS PHARMACEUTICALS PLC	CZ
Egiramlon 10 mg/5 mg kemény kapszula	HU/H/0303/004/DC	OGYI-T-22016/10-12, 18	EGIS PHARMACEUTICALS PLC	HU
Ramlon 10 mg/5 mg cietās kapsulas	HU/H/0303/001-005/DC	12-0066	EGIS PHARMACEUTICALS PLC	LV
Ramlon 10 mg/5 mg kietosios kapsulės	HU/H/0303/004/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/5 mg capsule	HU/H/0303/004/DC	4498/2012/01-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/5 mg Tvrdé kapsuly	HU/H/0303/003/DC	58/0745/11-S	EGIS PHARMACEUTICALS PLC	SK
Егирамлон 5 mg/10 mg твърди капсули	HU/H/0303/001-005/DC	20120047	EGIS PHARMACEUTICALS PLC	BG
Egiramlon 5 mg/10 mg tvrdé tobolky	HU/H/0303/001-005/DC	58/839/11-C	EGIS PHARMACEUTICALS PLC	CZ
Egiramlon 5 mg/10 mg kemény kapszula	HU/H/0303/003/DC	OGYI-T-22016/07-09, 17	EGIS PHARMACEUTICALS PLC	HU
Ramlon 5 mg/10 mg cietās kapsulas	HU/H/0303/001-005/DC	12-0065	EGIS PHARMACEUTICALS PLC	LV
Ramlon 5 mg/10 mg kietosios kapsulės	HU/H/0303/003/DC	LT/1/12/2892/011-015	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 5 mg/10 mg capsule	HU/H/0303/003/DC	4497/2012/01-06	EGIS PHARMACEUTICALS PLC	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
Egiramlon 5 mg/10 mg Tvrdé kapsuly	HU/H/0303/004/DC	58/0746/11-S	EGIS PHARMACEUTICALS PLC	SK
Egiramlon 10 mg/10 mg tvrdé tobolky	HU/H/0303/001-005/DC	58/841/11-C	EGIS PHARMACEUTICALS PLC	CZ
Egiramlon 10 mg/10 mg kemény kapszula	HU/H/0303/005/DC	OGYI-T-22016/13-15, 19	EGIS PHARMACEUTICALS PLC	HU
Ramlon 10 mg/10 mg cietās kapsulas	HU/H/0303/001-005/DC	12-0067	EGIS PHARMACEUTICALS PLC	LV
Ramlon 10 mg/10 mg kietosios kapsulės	HU/H/0303/005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/10 mg capsule	HU/H/0303/005/DC	4499/2012/01-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg Tvrdé kapsuly	HU/H/0303/005/DC	58/0747/11- S	EGIS PHARMACEUTICALS PLC	SK
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/01	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/02	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/03	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/06	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/04	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/05	S.C. SANDOZ S.R.L.	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
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/08	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/09	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/01	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule	SE/H/1397/001	6801/2014/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/06	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/03	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/07	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/01	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/09	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/02	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/04	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/05	S.C. SANDOZ S.R.L.	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
Prylar 5 mg/5 mg capsule	SE/H/1397/002	6802/2014/08	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/01	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/02	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/03	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/05	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/04	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/07	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/06	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/09	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/08	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/003	6803/2014/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/02	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/04	S.C. SANDOZ S.R.L.	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
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/03	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/05	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/06	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/07	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/08	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/09	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule	SE/H/1397/004	6804/2014/10	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/02	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/04	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/05	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/03	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/06	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/07	S.C. SANDOZ S.R.L.	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
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/09	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/08	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/10	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule	SE/H/1397/005	6805/2014/01	S.C. SANDOZ S.R.L.	RO
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Ramelso 2,5 mg/5 mg trde kapsule	SE/H/1397/001	H/14/01314/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/032	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/033	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/034	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/035	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/037	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/036	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/038	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/039	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/040	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/025	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/026	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/027	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/028	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/029	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/030	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/042	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/043	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/044	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/045	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/046	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/047	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/048	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/049	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/050	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
PRYLAR, (5+2,5) mg, καψάκιο, σκληρό	SE/H/1397/001	62012/28-12-2015	SANDOZ PHARMACEUTICALS D.D.	GR
PRYLAR, (5+5) mg, καψάκιο, σκληρό	SE/H/1397/002	82910/28-12-2015	SANDOZ PHARMACEUTICALS D.D.	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
PRYLAR, (10+5) mg, καψάκιο σκληρό	SE/H/1397/004	78871/28-12-2015	SANDOZ PHARMACEUTICALS D.D.	GR
PRYLAR, (5+10) mg, καψάκιο, σκληρό	SE/H/1397/003	78872/28-12-2015	SANDOZ PHARMACEUTICALS D.D.	GR
PRYLAR, (10+10) mg, καψάκιο, σκληρό	SE/H/1397/005	78873/28-12-2015	SANDOZ PHARMACEUTICALS D.D.	GR
Ramelso 5 mg/5 mg trde kapsule	SE/H/1397/002	H/14/01314/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 5 mg/10 mg trde kapsule	SE/H/1397/004	H/14/01314/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/5 mg trde kapsule	SE/H/1397/003	H/14/01314/031	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramelso 10 mg/10 mg trde kapsule	SE/H/1397/005	H/14/01314/041	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Ramipril/Amlodipine Sandoz	SE/H/1397/001	50479	SANDOZ A/S	SE
Ramipril/Amlodipine Sandoz	SE/H/1397/002	50480	SANDOZ A/S	SE
Ramipril/Amlodipine Sandoz	SE/H/1397/003	50481	SANDOZ A/S	SE
Ramipril/Amlodipine Sandoz	SE/H/1397/004	50482	SANDOZ A/S	SE
Ramipril/Amlodipine Sandoz	SE/H/1397/005	50483	SANDOZ A/S	SE
Amlodipino/Ramipril Codramol 10 mg/10 mg cápsulas duras	not available	80383	FARMALIDER, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlodipino/Ramipril Codramol 10 mg/5 mg cápsulas duras	not available	80380	FARMALIDER, S.A.	ES
Amlodipino/Ramipril Codramol 5 mg/10 mg cápsulas duras	not available	80381	FARMALIDER, S.A.	ES
Amlodipino/Ramipril Codramol 5 mg/2,5 mg cápsulas duras	not available	80382	FARMALIDER, S.A.	ES
Amlodipino/Ramipril Codramol 5 mg/5 mg cápsulas duras	not available	80384	FARMALIDER, S.A.	ES
Parvati 2,5 mg + 5 mg capsule rigide	not available	043322015	NEOPHARMED GENTILI S.R.L.	IT
Parvati 5 mg + 10 mg capsule rigide	not available	043322041	NEOPHARMED GENTILI S.R.L.	IT
Parvati 5 mg + 5 mg capsule rigide	not available	043322027	NEOPHARMED GENTILI S.R.L.	IT
Parvati 10 mg + 5 mg capsule rigide	not available	043322039	NEOPHARMED GENTILI S.R.L.	IT
Parvati 10 mg + 10 mg capsule rigide	not available	043322054	NEOPHARMED GENTILI S.R.L.	IT
RAMIPRIL/AMLODIPIN Pfizer 5 mg/10 mg Hartkapseln	AT/H/0469/003	135107	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
RAMIPRIL/AMLODIPIN Pfizer 10 mg/5 mg Hartkapseln	AT/H/0469/004	135108	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	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
RAMIPRIL/AMLODIPIN Pfizer 2,5 mg/2,5 mg Hartkapseln	AT/H/0469/001	135105	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
RAMIPRIL/AMLODIPIN Pfizer 10 mg/10 mg Hartkapseln	AT/H/0469/005	135109	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
RAMIPRIL/AMLODIPIN Pfizer 5 mg/5 mg Hartkapseln	AT/H/0469/002	135106	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Icomb 5 mg/5 mg capsule rigide	AT/H/0469/002	042384038	PFIZER ITALIA S.R.L.	IT
Icomb 5 mg/10 mg capsule rigide	AT/H/0469/003	042384065	PFIZER ITALIA S.R.L.	IT
Icomb 10 mg/5 mg capsule rigide	AT/H/0469/004	042384089	PFIZER ITALIA S.R.L.	IT
Icomb 2,5 mg/2,5 mg capsule rigide	AT/H/0469/001	042384026	PFIZER ITALIA S.R.L.	IT
Icomb 10 mg/10 mg capsule rigide	AT/H/0469/005	042384103	PFIZER ITALIA S.R.L.	IT
Icomb 10 mg/5 mg capsule rigide	AT/H/0469/004	042384077	PFIZER ITALIA S.R.L.	IT
Icomb 5 mg/5 mg capsule rigide	AT/H/0469/002	042384040	PFIZER ITALIA S.R.L.	IT
Icomb 2,5 mg/2,5 mg capsule rigide	AT/H/0469/001	042384014	PFIZER ITALIA S.R.L.	IT
Icomb 5 mg/10 mg capsule rigide	AT/H/0469/003	042384053	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Icomb 10 mg/10 mg capsule rigide	AT/H/0469/005	042384091	PFIZER ITALIA S.R.L.	IT
Ramipril Aristo plus Amlodipin 5 mg/5 mg Hartkapseln	SE/H/1340/002/E/001	2200100.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 5 mg/10 mg Hartkapseln	SE/H/1340/003/E/001	2200102.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 10 mg/5 mg Hartkapseln	SE/H/1340/004/E/001	2200101.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 10 mg/10 mg Hartkapseln	SE/H/1340/005/E/001	2200103.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 2,5 mg/5 mg Hartkapseln	SE/H/1340/001/E/001	2200099.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Amlodipino/ramipril Aristo 5 mg/2,5 mg cápsulas duras	SE/H/1340/001/E/001	82686	ARISTO PHARMA IBERIA, S.L.	ES
Amlodipino/ramipril Aristo 5 mg/ 5 mg cápsulas duras	SE/H/1340/002/E/001	82687	ARISTO PHARMA IBERIA, S.L.	ES
Amlodipino/ramipril Aristo 5 mg / 10 mg cápsulas duras	SE/H/1340/003/E/001	82688	ARISTO PHARMA IBERIA, S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlodipino/ramipril Aristo 10 mg/ 5 mg cápsulas duras	SE/H/1340/004/E/001	82689	ARISTO PHARMA IBERIA, S.L.	ES
Amlodipino/ramipril Aristo 10 mg /10 mg cápsulas duras	SE/H/1340/005/E/001	82690	ARISTO PHARMA IBERIA, S.L.	ES
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606011	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606023	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606100	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606050	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606062	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606047	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606098	ARISTO PHARMA GMBH (ART 57)	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
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606035	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606086	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 2,5 mg/5 mg, capsula rigida	SE/H/1340/001/E/001	045606074	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606112	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606124	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606201	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606151	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606163	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606148	ARISTO PHARMA GMBH (ART 57)	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
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606199	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606136	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606187	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/5 mg, capsula rigida	SE/H/1340/002/E/001	045606175	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606213	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606225	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606302	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606252	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606264	ARISTO PHARMA GMBH (ART 57)	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
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606249	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606290	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606237	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606288	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 5 mg/10 mg, capsula rigida	SE/H/1340/003/E/001	045606276	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606314	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606326	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606403	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606353	ARISTO PHARMA GMBH (ART 57)	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
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606365	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606340	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606391	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606338	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606389	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/5 mg, capsula rigida	SE/H/1340/004/E/001	045606377	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606415	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606427	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606504	ARISTO PHARMA GMBH (ART 57)	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
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606454	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606466	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606441	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606492	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606439	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606480	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril e Amlodipina Aristo, 10 mg/10 mg, capsula rigida	SE/H/1340/005/E/001	045606478	ARISTO PHARMA GMBH (ART 57)	IT
Ramipril/Amlodipine Aristo, 2,5 mg/5 mg, kapsel, hård	SE/H/1340/001	49217	ARISTO PHARMA GMBH (ART 57)	SE
Ramipril/Amlodipine Aristo, 5 mg/5 mg, kapsel, hård	SE/H/1340/002	49218	ARISTO PHARMA GMBH (ART 57)	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipine Aristo, 5 mg/10 mg, kapsel, hård	SE/H/1340/003	49220	ARISTO PHARMA GMBH (ART 57)	SE
Ramipril/Amlodipine Aristo, 10 mg/5 mg, kapsel, hård	SE/H/1340/004	49219	ARISTO PHARMA GMBH (ART 57)	SE
Ramipril/Amlodipine Aristo, 10 mg/10 mg, kapsel, hård	SE/H/1340/005	49221	ARISTO PHARMA GMBH (ART 57)	SE
RamiDipin® 5 mg/5 mg Hartkapseln	DE/H/4696/001	97297.00.00	TAD PHARMA GMBH	DE
RamiDipin® 5 mg/10 mg Hartkapseln	DE/H/4696/002	97298.00.00	TAD PHARMA GMBH	DE
RamiDipin® 10 mg/5 mg Hartkapseln	DE/H/4696/003	97299.00.00	TAD PHARMA GMBH	DE
RamiDipin® 10 mg/10 mg Hartkapseln	DE/H/4696/004	97300.00.00	TAD PHARMA GMBH	DE
Ramipril/Amlodipin Hexal 2,5 mg/2,5 mg - Hartkapseln	AT/H/0402/001	1-31215	HEXAL PHARMA GMBH	AT
Ramdacordia	AT/H/0402/001	786012	SANDOZ PHARMACEUTICALS D.D.	EE
Ramipril HEXAL plus Amlodipin 2,5 mg/2,5 mg Hartkapseln	AT/H/0402/001	84284.00.00	HEXAL AG	DE
SAMBETAN 2,5 mg + 5 mg capsule rigide	not available	044544017	CIPROS S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SAMBETAN 5 mg + 5 mg capsule rigide	not available	044544029	CIPROS S.R.L.	IT
SAMBETAN 10 mg + 5 mg capsule rigide	not available	044544031	CIPROS S.R.L.	IT
SAMBETAN 5 mg + 10 mg capsule rigide	not available	044544043	CIPROS S.R.L.	IT
SAMBETAN 10 mg + 10 mg capsule rigide	not available	044544056	CIPROS S.R.L.	IT
Ramantal 2,5 mg + 5 mg capsule rigide	not available	043313016	ERREKAPPA EUROTERAPICI SPA	IT
Ramantal 5 mg + 5 mg capsule rigide	not available	043313028	ERREKAPPA EUROTERAPICI SPA	IT
Ramantal 10 mg + 5 mg capsule rigide	not available	043313030	ERREKAPPA EUROTERAPICI SPA	IT
Ramantal 5 mg + 10 mg capsule rigide	not available	043313042	ERREKAPPA EUROTERAPICI SPA	IT
Ramantal 10 mg + 10 mg capsule rigide	not available	043313055	ERREKAPPA EUROTERAPICI SPA	IT
Ramipril/Amlodipin-ratiopharm® 10 mg/5 mg Hartkapseln	DE/H/5157/004	99471.00.00	RATIOPHARM GMBH	DE
Ramipril/Amlodipin-ratiopharm® 5 mg/5 mg Hartkapseln	DE/H/5157/002	99469.00.00	RATIOPHARM GMBH	DE
Ramipril/Amlodipin-ratiopharm® 10 mg/10 mg Hartkapseln	DE/H/5157/005	99472.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipin-ratiopharm® 5 mg/10 mg Hartkapseln	DE/H/5157/003	99470.00.00	RATIOPHARM GMBH	DE
Ramipril/Amlodipin-ratiopharm® 2,5 mg/5 mg Hartkapseln	DE/H/5157/001	99468.00.00	RATIOPHARM GMBH	DE
RILAMPIN 2,5 mg/5 mg	SE/H/1468/001	58/306/15-C	ACTAVIS GROUP PTC EHF.	CZ
RILAMPIN 5 mg/5 mg	SE/H/1468/002	58/307/15-C	ACTAVIS GROUP PTC EHF.	CZ
RILAMPIN 5 mg/10 mg	SE/H/1468/003	58/308/15-C	ACTAVIS GROUP PTC EHF.	CZ
RILAMPIN 10 mg/5 mg	SE/H/1468/004	58/309/15-C	ACTAVIS GROUP PTC EHF.	CZ
RILAMPIN 10 mg/10 mg	SE/H/1468/005	58/310/15-C	ACTAVIS GROUP PTC EHF.	CZ
Ramipril/Amlodipine Actavis 2.5 mg/5 mg, capsule, hard	SE/H/1468/001	51821	ACTAVIS GROUP PTC EHF.	SE
Ramipril/Amlodipine Actavis 5 mg/5 mg, capsule, hard	SE/H/1468/002	51822	ACTAVIS GROUP PTC EHF.	SE
Ramipril/Amlodipine Actavis 10 mg/5 mg, capsule, hard	SE/H/1468/004	51824	ACTAVIS GROUP PTC EHF.	SE
Ramipril/Amlodipine Actavis 5 mg/10 mg, capsule, hard	SE/H/1468/003	51823	ACTAVIS GROUP PTC EHF.	SE
Ramipril/Amlodipine Actavis 10 mg/10 mg, capsule, hard	SE/H/1468/005	51825	ACTAVIS GROUP PTC EHF.	SE
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/01	ACTAVIS GROUP PTC EHF.	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
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/01	ACTAVIS GROUP PTC EHF.	RO
Вивейс Дуо 5 mg/5 mg твърди капсули	SE/H/1468/002	20150266	ACTAVIS GROUP PTC EHF.	BG
Вивейс Дуо 10 mg/5 mg твърди капсули	SE/H/1468/004	20150268	ACTAVIS GROUP PTC EHF.	BG
Вивейс Дуо 10 mg/10 mg твърди капсули	SE/H/1468/005	20150269	ACTAVIS GROUP PTC EHF.	BG
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/09	ACTAVIS GROUP PTC EHF.	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
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/02	ACTAVIS GROUP PTC EHF.	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
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	8208/2015/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	8209/2015/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/08	ACTAVIS GROUP PTC EHF.	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
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	8207/2015/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	8210/2015/10	ACTAVIS GROUP PTC EHF.	RO
Ramipril/amlodipin Pliva 5 mg/5 mg tvrde kapsule	DE/H/5152/001	HR-H-023212604	PLIVA HRVATSKA D.O.O.	HR
Ramipril/amlodipin Pliva 10 mg/5 mg tvrde kapsule	DE/H/5152/004	HR-H-750926919	PLIVA HRVATSKA D.O.O.	HR
Ramipril/amlodipin Pliva 5 mg/10 mg tvrde kapsule	DE/H/5152/003	HR-H-620376275	PLIVA HRVATSKA D.O.O.	HR
Ramipril/amlodipin Pliva 10 mg/10 mg tvrde kapsule	DE/H/5152/005	HR-H-994428704	PLIVA HRVATSKA D.O.O.	HR
Rameam 5 mg/5 mg tvrde kapsule	DE/H/4683/001	HR-H-128016330	KRKA-FARMA D.O.O.	HR
Rameam 10 mg/5 mg tvrde kapsule	DE/H/4683/003	HR-H-232568101	KRKA-FARMA D.O.O.	HR
Rameam 5 mg/10 mg tvrde kapsule	DE/H/4683/002	HR-H-459256324	KRKA-FARMA D.O.O.	HR
Rameam 10 mg/10 mg tvrde kapsule	DE/H/4683/004	HR-H-830032291	KRKA-FARMA D.O.O.	HR
Ramladio 5 mg/5 mg tvrdé kapsuly	DE/H/4683/001	58/0096/17-S	KRKA, D.D., NOVO MESTO	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramladio 5 mg/10 mg tvrdé kapsuly	DE/H/4683/002	58/0097/17-S	KRKA, D.D., NOVO MESTO	SK
Ramladio 10 mg/5 mg tvrdé kapsuly	DE/H/4683/003	58/0098/17-S	KRKA, D.D., NOVO MESTO	SK
Ramladio 10 mg/10 mg tvrdé kapsuly	DE/H/4683/004	58/0099/17-S	KRKA, D.D., NOVO MESTO	SK
Rameam, 5 mg/5 mg kõvakapslid	DE/H/4683/001	939717	KRKA, D.D., NOVO MESTO	EE
Rameam, 5 mg/10 mg kõvakapslid	DE/H/4683/002	939817	KRKA, D.D., NOVO MESTO	EE
Rameam, 10 mg/5 mg kõvakapslid	DE/H/4683/003	939917	KRKA, D.D., NOVO MESTO	EE
Rameam, 10 mg/10 mg kõvakapslid	DE/H/4683/004	940017	KRKA, D.D., NOVO MESTO	EE
Ramladio 10 mg + 10 mg cietās kapsulas	DE/H/4683/004	17-0096	KRKA, D.D., NOVO MESTO	LV
Ramladio 10 mg + 5 mg cietās kapsulas	DE/H/4683/003	17-0097	KRKA, D.D., NOVO MESTO	LV
Ramladio 5 mg + 10 mg cietās kapsulas	DE/H/4683/002	17-0098	KRKA, D.D., NOVO MESTO	LV
Ramladio 5 mg + 5 mg cietās kapsulas	DE/H/4683/001	17-0099	KRKA, D.D., NOVO MESTO	LV
Ramipril + Amlodipina Krka 5 mg + 5 mg cápsulas	DE/H/4683/001	5715057	KRKA, D.D., NOVO MESTO	PT
Ramipril + Amlodipina Krka 5 mg + 10 mg cápsulas	DE/H/4683/002	5715065	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
Ramipril + Amlodipina Krka 10 mg + 5 mg cápsulas	DE/H/4683/003	5715073	KRKA, D.D., NOVO MESTO	PT
Ramipril + Amlodipina Krka 10 mg + 10 mg cápsulas	DE/H/4683/004	5715107	KRKA, D.D., NOVO MESTO	PT
Ramipril/amlodipine Krka 5 mg/5 mg hard capsules	DE/H/4683/001	PA1347/072/001	KRKA, D.D., NOVO MESTO	IE
Ramipril/amlodipine Krka 5 mg/10 mg hard capsules	DE/H/4683/002	PA1347/072/002	KRKA, D.D., NOVO MESTO	IE
Ramipril/amlodipine Krka 10 mg/5 mg hard capsules	DE/H/4683/003	PA1347/072/003	KRKA, D.D., NOVO MESTO	IE
Ramipril/amlodipine Krka 10 mg/10 mg hard capsules	DE/H/4683/004	PA1347/072/004	KRKA, D.D., NOVO MESTO	IE
Rameam 5 mg/5 mg kovat kapselit	DE/H/4683/001	34105	KRKA, D.D., NOVO MESTO	FI
Rameam 5 mg/10 mg kovat kapselit	DE/H/4683/002	34106	KRKA, D.D., NOVO MESTO	FI
Rameam 10 mg/5 mg kovat kapselit	DE/H/4683/003	34107	KRKA, D.D., NOVO MESTO	FI
Rameam 10 mg/10 mg kovat kapselit	DE/H/4683/004	34108	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
Ramipril/Amlodipin Krka 10 mg/10 mg Hartkapseln	DE/H/4683/004	137712	KRKA, D.D., NOVO MESTO	AT
Ramipril/Amlodipin Krka 10 mg/5 mg Hartkapseln	DE/H/4683/003	137713	KRKA, D.D., NOVO MESTO	AT
Ramipril/Amlodipin Krka 5 mg/10 mg Hartkapseln	DE/H/4683/002	137714	KRKA, D.D., NOVO MESTO	AT
Ramipril/Amlodipin Krka 5 mg/5 mg Hartkapseln	DE/H/4683/001	137715	KRKA, D.D., NOVO MESTO	AT
Рамеам 5 mg/5 mg капсули, твърди	DE/H/4683/001	20170197	KRKA, D.D., NOVO MESTO	BG
Рамеам 5 mg/10 mg капсули, твърди	DE/H/4683/002	20170198	KRKA, D.D., NOVO MESTO	BG
Рамеам 10 mg/5 mg капсули, твърди	DE/H/4683/003	20170199	KRKA, D.D., NOVO MESTO	BG
Рамеам 10 mg/10 mg капсули, твърди	DE/H/4683/004	20170200	KRKA, D.D., NOVO MESTO	BG
Ramipril/Amlodipine Krka 10 mg/10 mg harde capsules	DE/H/4683/004	BE509315	KRKA, D.D., NOVO MESTO	BE
Ramipril/Amlodipine Krka 10 mg/5 mg harde capsules	DE/H/4683/003	BE509324	KRKA, D.D., NOVO MESTO	BE
Ramipril/Amlodipine Krka 5 mg/10 mg harde capsules	DE/H/4683/002	BE509333	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
Ramipril/Amlodipine Krka 5 mg/5 mg harde capsules	DE/H/4683/001	BE509342	KRKA, D.D., NOVO MESTO	BE
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/001	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/002	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/003	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/004	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/005	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/006	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/007	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/008	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/009	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/004	H/17/02378/010	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/011	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/012	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
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/013	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/014	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/015	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/016	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/017	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/018	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/019	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/003	H/17/02378/020	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/021	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/022	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/023	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/024	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/025	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
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/026	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/027	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/028	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/029	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/002	H/17/02378/030	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/031	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/032	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/033	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/034	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/035	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/036	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/037	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/038	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
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/039	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/001	H/17/02378/040	KRKA, D.D., NOVO MESTO	SI
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/004	10071/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/004	10071/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	10071/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/01	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
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/003	10072/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/04	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
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/002	10073/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/07	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
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/001	10074/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramladio 5 mg/5 mg tvrdé tobolky	DE/H/4683/001	58/569/16-C	KRKA, D.D., NOVO MESTO	CZ
Ramladio 5 mg/10 mg tvrdé tobolky	DE/H/4683/002	58/570/16-C	KRKA, D.D., NOVO MESTO	CZ
Ramladio 10 mg/5 mg tvrdé tobolky	DE/H/4683/003	58/571/16-C	KRKA, D.D., NOVO MESTO	CZ
Ramladio 10 mg/10 mg tvrdé tobolky	DE/H/4683/004	58/572/16-C	KRKA, D.D., NOVO MESTO	CZ
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268012	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268024	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268036	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268048	KRKA, D.D., NOVO MESTO	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268051	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268063	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268075	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268087	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268099	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/5 mg capsule rigide	DE/H/4683/004	045268101	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268113	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268125	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268137	KRKA, D.D., NOVO MESTO	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268149	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268152	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268164	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268176	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268188	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268190	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 5 mg/10 mg capsule rigide	DE/H/4683/003	045268202	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268214	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268226	KRKA, D.D., NOVO MESTO	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268238	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268240	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268253	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268265	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268277	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268289	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268291	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/003	045268303	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268315	KRKA, D.D., NOVO MESTO	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268327	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268339	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268341	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268354	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268366	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268378	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268380	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268392	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/004	045268404	KRKA, D.D., NOVO MESTO	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rameam 10 mg/10 mg Hartkapseln	DE/H/4683/004	97123.00.00	TAD PHARMA GMBH	DE
Rameam 10 mg/5 mg Hartkapseln	DE/H/4683/003	97124.00.00	TAD PHARMA GMBH	DE
Rameam 5 mg/10 mg Hartkapseln	DE/H/4683/002	97125.00.00	TAD PHARMA GMBH	DE
Rameam 5 mg/5 mg Hartkapseln	DE/H/4683/001	97126.00.00	TAD PHARMA GMBH	DE
Ramladio, 10 mg + 10 mg, kapsułki, twarde	DE/H/4683/004	24021	KRKA, D.D., NOVO MESTO	PL
Ramladio, 10 mg + 5 mg, kapsułki, twarde	DE/H/4683/003	24022	KRKA, D.D., NOVO MESTO	PL
Ramladio, 5 mg + 10 mg, kapsułki, twarde	DE/H/4683/002	24023	KRKA, D.D., NOVO MESTO	PL
Ramladio, 5 mg + 5 mg, kapsułki, twarde	DE/H/4683/001	24024	KRKA, D.D., NOVO MESTO	PL
Polpram 5 mg/5 mg tvrdé kapsuly	CZ/H/0513/001	58/0201/16-S	MEDANA PHARMA SPOLKA AKCYJNA	SK
Polpram 5 mg/10 mg tvrdé kapsuly	CZ/H/0513/002	58/0202/16-S	MEDANA PHARMA SPOLKA AKCYJNA	SK
Polpram 10 mg/5 mg tvrdé kapsuly	CZ/H/0513/003	58/0203/16-S	MEDANA PHARMA SPOLKA AKCYJNA	SK
Polpram 10 mg/10 mg tvrdé kapsuly	CZ/H/0513/004	58/0204/16-S	MEDANA PHARMA SPOLKA AKCYJNA	SK
Rimal 5 mg / 5 mg cietās kapsulas	CZ/H/0513/001	16-0104	MEDANA PHARMA SPOLKA AKCYJNA	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
Rimal 5 mg / 10 mg cietās kapsulas	CZ/H/0513/002	16-0105	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 10 mg / 5 mg cietās kapsulas	CZ/H/0513/003	16-0106	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 10 mg / 10 mg cietās kapsulas	CZ/H/0513/004	16-0107	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 5 mg/5 mg, tvrdé tobolky	CZ/H/0513/001	58/227/16-C	MEDANA PHARMA SPOLKA AKCYJNA	CZ
Rimal 5 mg/10 mg, tvrdé tobolky	CZ/H/0513/002	58/228/16-C	MEDANA PHARMA SPOLKA AKCYJNA	CZ
Rimal 10 mg/5 mg, tvrdé tobolky	CZ/H/0513/003	58/229/16-C	MEDANA PHARMA SPOLKA AKCYJNA	CZ
Rimal 10 mg/10 mg, tvrdé tobolky	CZ/H/0513/004	58/230/16-C	MEDANA PHARMA SPOLKA AKCYJNA	CZ
Полпрам 5 mg/5 mg капсули, твърди	CZ/H/0513/001	20160220	MEDANA PHARMA SPOLKA AKCYJNA	BG
Полпрам 5 mg/10 mg капсули, твърди	CZ/H/0513/002	20160221	MEDANA PHARMA SPOLKA AKCYJNA	BG
Полпрам 10 mg/5 mg капсули, твърди	CZ/H/0513/003	20160222	MEDANA PHARMA SPOLKA AKCYJNA	BG
Полпрам 10 mg/10 mg капсули, твърди	CZ/H/0513/004	20160223	MEDANA PHARMA SPOLKA AKCYJNA	BG
Rimal, 5 mg + 5 mg, kapsułki, twarde	not available	23832	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Rimal, 5 mg + 10 mg, kapsułki, twarde	not available	23833	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	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
Rimal, 10 mg + 5 mg, kapsułki, twarde	not available	23834	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Rimal, 10 mg + 10 mg, kapsułki, twarde	not available	23835	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Piramil Combi 10 mg/5 mg tvrdé tobolky	AT/H/0402/003	58/362/12-C	SANDOZ S.R.O.	CZ
Sumilar, 10 mg + 5 mg, kapsułki twarde	AT/H/0402/003	20265	SANDOZ GMBH	PL
Ramipril/Amlodipin Hexal 10 mg/5 mg - Hartkapseln	AT/H/0402/003	1-31217	HEXAL PHARMA GMBH	AT
AMIRAP 10 mg/5 mg tvrdé kapsuly	AT/H/0402/003	58/0183/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Ramdacordia 10 mg/5 mg cietās kapsulas	AT/H/0402/003	12-0138	SANDOZ PHARMACEUTICALS D.D.	LV
Ramdacordia, 10 mg/5 mg, kõvakapslid	AT/H/0402/003	785812	SANDOZ PHARMACEUTICALS D.D.	EE
Ramipril HEXAL plus Amlodipin 10 mg/5 mg Hartkapseln	AT/H/0402/003	84286.00.00	HEXAL AG	DE
Ramdacordia, 5 mg/5 mg, kõvakapslid	AT/H/0402/002	785912	SANDOZ PHARMACEUTICALS D.D.	EE
Piramil Combi 5 mg/5 mg tvrdé tobolky	AT/H/0402/002	58/361/12-C	SANDOZ S.R.O.	CZ
Sumilar, 5 mg + 5 mg, kapsułki twarde	AT/H/0402/002	20264	SANDOZ GMBH	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
Ramipril/Amlodipin Hexal 5 mg/5 mg - Hartkapseln	AT/H/0402/002	1-31216	HEXAL PHARMA GMBH	AT
AMIRAP 5 mg/5 mg tvrdé kapsuly	AT/H/0402/002	58/0182/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Ramdacordia 5 mg/5 mg cietās kapsulas	AT/H/0402/002	12-0136	SANDOZ PHARMACEUTICALS D.D.	LV
Amlopin DUO 5mg / 5 mg capsules, hard	AT/H/0402/002	20120330	SANDOZ PHARMACEUTICALS D.D.	BG
Ramipril HEXAL plus Amlodipin 5 mg/5 mg Hartkapseln	AT/H/0402/002	84285.00.00	HEXAL AG	DE
Piramil Combi 10 mg/10 mg tvrdé tobolky	AT/H/0402/005	58/364/12-C	SANDOZ S.R.O.	CZ
Ramipril/Amlodipin Hexal 10 mg/10 mg - Hartkapseln	AT/H/0402/005	1-31219	HEXAL PHARMA GMBH	AT
AMIRAP 10 mg/10 mg tvrdé kapsuly	AT/H/0402/005	58/0185/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Ramdacordia 10 mg/10 mg cietās kapsulas	AT/H/0402/005	12-0139	SANDOZ PHARMACEUTICALS D.D.	LV
Ramdacordia, 10 mg/10 mg, kõvakapslid	AT/H/0402/005	785712	SANDOZ PHARMACEUTICALS D.D.	EE
Sumilar, 10 mg + 10 mg, kapsułki twarde	AT/H/0402/005	20267	SANDOZ GMBH	PL
Ramipril HEXAL plus Amlodipin 10 mg/10 mg Hartkapseln	AT/H/0402/005	84288.00.00	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
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 2,5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/001	22122	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 5 mg, kapsułki, twarde	CZ/H/0674/002	22123	SANOFI-AVENTIS SP Z.O.O.	PL
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	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
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 2,5 mg/5 mg tvrdé tobolky	CZ/H/0674/001	58/333/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	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
Tritace Combi 5 mg/5 mg tvrdé tobolky	CZ/H/0674/002	58/334/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/5 mg tvrdé tobolky	CZ/H/0674/003	58/335/14-C	SANOFI-AVENTIS SRO	CZ
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 10 mg, kapsułki, twarde	CZ/H/0674/005	22126	SANOFI-AVENTIS SP Z.O.O.	PL
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	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
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 5 mg/10 mg tvrdé tobolky	CZ/H/0674/004	58/336/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Tritace Combi 10 mg/10 mg tvrdé tobolky	CZ/H/0674/005	58/337/14-C	SANOFI-AVENTIS SRO	CZ
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	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
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (2,5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/001	022103	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	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
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/002	022104	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	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
TRIAMLO, (5+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/003	022105	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+5) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/004	022106	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	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
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
TRIAMLO, (10+10) MG, ΚΑΨΆΚΙΟ, ΣΚΛΗΡΌ	CZ/H/0674/005	022107	SANOFI-AVENTIS CYPRUS LTD	CY
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 2,5 mg/5 mg tvrdé kapsuly	CZ/H/0674/001	58/0284/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/5 mg tvrdé kapsuly	CZ/H/0674/002	58/0285/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 10 mg/5 mg tvrdé kapsuly	CZ/H/0674/003	58/0287/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 5 mg/10 mg tvrdé kapsuly	CZ/H/0674/004	58/0286/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	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
Ramipril/Amlodipin Zentiva 10 mg/10 mg tvrdé kapsuly	CZ/H/0674/005	58/0288/14-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	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
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+5) mg, καψάκιο, σκληρό	CZ/H/0674/003	91050/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (10+10) mg, καψάκιο, σκληρό	CZ/H/0674/005	91052/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	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
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (2,5+5) mg, καψάκιο, σκληρό	CZ/H/0674/001	91048/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	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
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+5) mg, καψάκιο, σκληρό	CZ/H/0674/002	91049/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Triamlo, (5+10) mg, καψάκιο, σκληρό	CZ/H/0674/004	91051/21-12-2015	SANOFI-AVENTIS AEBE	GR
Tonotec 5 mg/5 mg Hartkapseln	DE/H/3919/002	88599.00.00	UCB INNERE MEDIZIN GMBH & CO. KG	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
Tonotec 5 mg/10 mg Hartkapseln	DE/H/3919/003	88600.00.00	UCB INNERE MEDIZIN GMBH & CO. KG	DE
Tonotec 10 mg/5 mg Hartkapseln	DE/H/3919/004	88601.00.00	UCB INNERE MEDIZIN GMBH & CO. KG	DE
Tonotec 10 mg/10 mg Hartkapseln	DE/H/3919/005	88602.00.00	UCB INNERE MEDIZIN GMBH & CO. KG	DE
Ramizek Combi, 2,5 mg + 5 mg, kapsułki, twarde	SE/H/1336/001	21904	ADAMED	PL
Ramizek Combi, 5 mg + 5 mg, kapsułki, twarde	SE/H/1336/002	21905	ADAMED	PL
Ramizek Combi, 5 mg + 10 mg, kapsułki, twarde	SE/H/1336/003	21906	ADAMED	PL
Ramizek Combi, 10 mg + 5 mg, kapsułki, twarde	SE/H/1336/004	21907	ADAMED	PL
Ramizek Combi, 10 mg + 10 mg, kapsułki, twarde	SE/H/1336/005	21908	ADAMED	PL
Ramipril/Amlodipine Adamed, 2,5 mg/5 mg, hårda kapslar	SE/H/1336/001	49203	ADAMED	SE
Ramipril/Amlodipine Adamed, 5 mg/5 mg, hårda kapslar	SE/H/1336/002	49204	ADAMED	SE
Ramipril/Amlodipine Adamed, 5 mg/10 mg, hårda kapslar	SE/H/1336/004	49206	ADAMED	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipine Adamed, 10 mg/5 mg, hårda kapslar	SE/H/1336/003	49205	ADAMED	SE
Ramipril/Amlodipine Adamed, 10 mg/10 mg, hårda kapslar	SE/H/1336/005	49207	ADAMED	SE
Ramipril/Amlodipin Genericon 2,5 mg/5 mg Hartkapseln	SE/H/1336/001	135662	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin Genericon 5 mg/5 mg Hartkapseln	SE/H/1336/002	135663	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin Genericon 5 mg/10 mg Hartkapseln	SE/H/1336/003	135666	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin Genericon 10 mg/5 mg Hartkapseln	SE/H/1336/004	135664	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin Genericon 10 mg/10 mg Hartkapseln	SE/H/1336/005	135665	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
RAMIZEK 5 mg/5 mg tvrdé tobolky	SE/H/1336/002	58/378/14-C	ADAMED	CZ
RAMIZEK 10 mg/5 mg tvrdé tobolky	SE/H/1336/003	58/379/14-C	ADAMED	CZ
RAMIZEK 5 mg/10 mg tvrdé tobolky	SE/H/1336/004	58/380/14-C	ADAMED	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
RAMIZEK 10 mg/10 mg tvrdé tobolky	SE/H/1336/005	58/381/14-C	ADAMED	CZ
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240011	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240023	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240035	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240047	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240213	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240225	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240237	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240249	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240252	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 2,5 mg/5 mg, capsule rigide	SE/H/1336/001	043240264	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240050	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240062	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240074	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240086	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240276	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240288	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240290	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240302	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240314	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/5 mg, capsule rigide	SE/H/1336/002	043240326	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240136	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240148	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240151	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240163	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240391	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240403	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240415	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240427	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240439	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/5 mg, capsule rigide	SE/H/1336/003	043240441	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240098	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240100	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240112	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240124	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240338	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240340	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240353	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240365	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240377	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 5 mg/10 mg, capsule rigide	SE/H/1336/004	043240389	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240175	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240187	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240199	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240201	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240454	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240466	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240478	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240480	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240492	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Duotens 10 mg/10 mg, capsule rigide	SE/H/1336/005	043240504	SPA – SOCIETÀ PRODOTTI ANTIBIOTICI S.P.A.	IT
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/01	ADAMED	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
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/02	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/03	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/04	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/05	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/06	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/07	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/08	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/09	ADAMED	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/10	ADAMED	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
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/01	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/02	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/03	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/04	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/05	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/06	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/07	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/08	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/09	ADAMED	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
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/10	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/01	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/02	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/03	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/04	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/05	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/06	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/07	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/08	ADAMED	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
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/09	ADAMED	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/10	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/01	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/02	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/03	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/04	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/05	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/06	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/07	ADAMED	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
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/001-005/DC	7202/2014/08	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/09	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/10	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/01	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/02	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/03	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/04	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/05	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/06	ADAMED	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
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/07	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/08	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/09	ADAMED	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/10	ADAMED	RO
RAMI-AMLO, 2,5 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/001	16622/5-3-2015	IASIS PHARMA	GR
RAMI-AMLO, 5 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/002	16623/5-3-2015	IASIS PHARMA	GR
RAMI-AMLO, 5 mg/10 mg , καψάκιο, σκληρό	SE/H/1336/003	16625/5-3-2015	IASIS PHARMA	GR
RAMI-AMLO, 10 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/004	16624/5-3-2015	IASIS PHARMA	GR
RAMI-AMLO, 10 mg/10 mg , καψάκιο, σκληρό	SE/H/1336/005	16626/5-3-2015	IASIS PHARMA	GR
RAMI-AMLO, 2,5 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/001	022123	IASIS PHARMA	CY
RAMI-AMLO, 5 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/002	022124	IASIS PHARMA	CY
RAMI-AMLO, 10 mg/5 mg , καψάκιο, σκληρό	SE/H/1336/004	022125	IASIS PHARMA	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
RAMI-AMLO, 5 mg/10 mg , καψάκιο, σκληρό	SE/H/1336/003	022126	IASIS PHARMA	CY
RAMI-AMLO, 10 mg/10 mg , καψάκιο, σκληρό	SE/H/1336/005	022127	IASIS PHARMA	CY
Импактин Дуо 5 mg/5 mg твърди капсули	SE/H/1336/002	20140180	SOPHARMA AD	BG
Импактин Дуо 5mg/10 mg твърди капсули	SE/H/1336/004	20140182	SOPHARMA AD	BG
Импактин Дуо 10mg/5 mg твърди капсули	SE/H/1336/003	20140181	SOPHARMA AD	BG
Импактин Дуо 10mg/10 mg твърди капсули	SE/H/1336/005	20140183	SOPHARMA AD	BG
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A.	GR
ALAMUT 2,5 mg + 5 mg capsule rigide	not available	044036010	SO.SE.PHARM S.R.L.	IT
ALAMUT 5 mg + 5 mg capsule rigide	not available	044036022	SO.SE.PHARM S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ALAMUT 5 mg + 10 mg capsule rigide	not available	044036034	SO.SE.PHARM S.R.L.	IT
ALAMUT 10 mg + 5 mg capsule rigide	not available	044036046	SO.SE.PHARM S.R.L.	IT
ALAMUT 10 mg + 10 mg capsule rigide	not available	044036059	SO.SE.PHARM S.R.L.	IT
Piramil Combi 5 mg/10 mg tvrdé tobolky	AT/H/0402/004	58/363/12-C	SANDOZ S.R.O.	CZ
Sumilar, 5 mg + 10 mg, kapsułki twarde	AT/H/0402/004	20266	SANDOZ GMBH	PL
Ramipril/Amlodipin Hexal 5 mg/10 mg - Hartkapseln	AT/H/0402/004	1-31218	HEXAL PHARMA GMBH	AT
AMIRAP 5 mg/10 mg tvrdé kapsuly	AT/H/0402/004	58/0184/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Ramdacordia 5 mg/10 mg cietās kapsulas	AT/H/0402/004	12-0137	SANDOZ PHARMACEUTICALS D.D.	LV
Ramdacordia, 5 mg/10 mg, kõvakapslid	AT/H/0402/004	785612	SANDOZ PHARMACEUTICALS D.D.	EE
Ramipril HEXAL plus Amlodipin 5 mg/10 mg Hartkapseln	AT/H/0402/004	84287.00.00	HEXAL AG	DE
RAMIPRIL E AMLODIPINA DOC Generici 2,5 mg + 5 mg capsule rigide	not available	044802015	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 2,5 mg + 5 mg capsule rigide	not available	044802027	DOC GENERICI S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
RAMIPRIL E AMLODIPINA DOC Generici 5 mg + 5 mg capsule rigide	not available	044802039	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 5 mg + 5 mg capsule rigide	not available	044802041	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 10 mg + 5 mg capsule rigide	not available	044802078	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 10 mg + 5 mg capsule rigide	not available	044802080	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 5 mg + 10 mg capsule rigide	not available	044802054	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 5 mg + 10 mg capsule rigide	not available	044802066	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 10 mg + 10 mg capsule rigide	not available	044802092	DOC GENERICI S.R.L.	IT
RAMIPRIL E AMLODIPINA DOC Generici 10 mg + 10 mg capsule rigide	not available	044802104	DOC GENERICI S.R.L.	IT
Ramomark 5 mg/5 mg tvrdé tobolky	SE/H/1337/001	58/281/14-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipine Glenmark, 10 mg/10 mg, hårda kapslar	SE/H/1337/004	49211	GLENMARK PHARMACEUTICALS S.R.O.	SE
Ramipril/Amlodipine Glenmark, 5 mg/5 mg, hårda kapslar	SE/H/1337/001	49208	GLENMARK PHARMACEUTICALS S.R.O.	SE
Ramipril/Amlodipine Glenmark, 10 mg/5 mg, hårda kapslar	SE/H/1337/002	49209	GLENMARK PHARMACEUTICALS S.R.O.	SE
Ramipril/Amlodipine Glenmark, 5 mg/10 mg, hårda kapslar	SE/H/1337/003	49210	GLENMARK PHARMACEUTICALS S.R.O.	SE
Ramomark 5 mg/10 mg tvrdé kapsuly	SE/H/1337/003	58/0205/14-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Ramomark 10 mg/5 mg tvrdé tobolky	SE/H/1337/002	58/282/14-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Ramomark 10 mg/10 mg tvrdé tobolky	SE/H/1337/004	58/284/14-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Ramomark 5 mg/10 mg tvrdé tobolky	SE/H/1337/003	58/283/14-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Ramomark 10 mg/5 mg tvrdé kapsuly	SE/H/1337/002	58/0206/14-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Ramomark 5 mg/5 mg tvrdé kapsuly	SE/H/1337/001	58/0204/14-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Ramomark 10 mg/10 mg tvrdé kapsuly	SE/H/1337/004	58/0207/14-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/01	PHARMA-REGIST LTD	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
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/02	PHARMA-REGIST LTD	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/03	PHARMA-REGIST LTD	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/04	PHARMA-REGIST LTD	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/05	PHARMA-REGIST LTD	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/06	PHARMA-REGIST LTD	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/07	PHARMA-REGIST LTD	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/08	PHARMA-REGIST LTD	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/09	PHARMA-REGIST LTD	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/10	PHARMA-REGIST LTD	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/11	PHARMA-REGIST LTD	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/12	PHARMA-REGIST LTD	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/13	PHARMA-REGIST LTD	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/14	PHARMA-REGIST LTD	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
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/15	PHARMA-REGIST LTD	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/16	PHARMA-REGIST LTD	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/17	PHARMA-REGIST LTD	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/18	PHARMA-REGIST LTD	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/19	PHARMA-REGIST LTD	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/20	PHARMA-REGIST LTD	HU