02 December 2021
EMA/731539/202121
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine besilate / ramipril, amlodipine / hydrochlorothiazide / ramipril, hydrochlorothiazide / ramipril

Procedure no.: PSUSA/00010774/202103

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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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
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

Product 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/14/01314/006	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
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
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

Product 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/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
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
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/01	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/02	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/03	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/05	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 ramipril/amlodipină	SI/H/0231/001	13821/2021/06	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/04	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/07	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/08	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/09	S.C. SANDOZ S.R.L.	RO
Prylar 2,5 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/001	13821/2021/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/01	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/02	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/03	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/04	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/05	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 ramipril/amlodipina	SI/H/0231/002	13822/2021/06	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/07	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/08	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/09	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/5 mg capsule ramipril/amlodipina	SI/H/0231/002	13822/2021/10	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/01	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/02	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/03	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/04	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/05	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/06	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/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/08	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/09	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/004	13824/2021/02	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/004	13824/2021/03	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/004	13824/2021/04	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/004	13824/2021/05	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/004	13824/2021/06	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/004	13824/2021/07	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/004	13824/2021/08	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/004	13824/2021/09	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 ramipril/amlodipină	SI/H/0231/004	13824/2021/10	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/5 mg capsule ramipril/amlodipină	SI/H/0231/003	13823/2021/07	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/005	13825/2021/01	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/005	13825/2021/02	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/005	13825/2021/03	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/005	13825/2021/04	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/005	13825/2021/05	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/005	13825/2021/06	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipină	SI/H/0231/005	13825/2021/07	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/005	13825/2021/08	S.C. SANDOZ S.R.L.	RO
Prylar 10 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/005	13825/2021/09	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 ramipril/amlodipina	SI/H/0231/005	13825/2021/10	S.C. SANDOZ S.R.L.	RO
Prylar 5 mg/10 mg capsule ramipril/amlodipina	SI/H/0231/004	13824/2021/01	S.C. SANDOZ S.R.L.	RO
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Tamayra 5 mg/5 mg hard capsules	BG/H/0106/001	20160224	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Tamayra 10 mg/5 mg hard capsules	BG/H/0106/002	20160225	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Tamayra 5 mg/5 mg tvrde kapsuly	BG/H/0106/001/DC	58/0399/16-S	SWYSSI AG	SK
Tamayra 10 mg/5 mg tvrde kapsuly	BG/H/0106/002/DC	58/0400/16-S	SWYSSI AG	SK
Coreyra 5 mg/5 mg Hartkapseln	BG/H/0106/001/DC	137604	SWYSSI AG	AT
Coreyra 10 mg/5 mg Hartkapseln	BG/H/0106/002/DC	137603	SWYSSI AG	AT
Tamayra 5 mg/5 mg καψάκιο σκληρό	BG/H/0106/001/DC	312480101	SWYSSI AG	GR
Tamayra 10 mg/5 mg καψάκιο σκληρό	BG/H/0106/002/DC	312480201	SWYSSI AG	GR
Coreyra 5 mg/5 mg capsule	BG/H/0106/001	9323/2016/01	SWYSSI AG	RO
Coreyra 10 mg/5 mg capsule	BG/H/0106/002	9324/2016/01	SWYSSI AG	RO
Tamayra 5 mg/5 mg cápsulas duras	BG/H/0106/001	5720701	SWYSSI AG	PT
Tamayra 10 mg/5 mg cápsulas duras	BG/H/0106/002	5720669	SWYSSI AG	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tamayra 5 mg/5 mg tvrdé tobolky	BG/H/0106/001/R/001	58/416/16-C	SWYSSI AG	CZ
Tamayra 10 mg/5 mg tvrde tobolky	BG/H/0106/002/R/002/001	58/417/16-C	SWYSSI AG	CZ
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
SARASVATI 5 mg/5 mg/12,5 mg capsule rigide	not available	045873027	NEOPHARMED GENTILI SPA	IT
SARASVATI 5 mg/5 mg/25 mg capsule rigide	not available	045873039	NEOPHARMED GENTILI SPA	IT
SARASVATI 10 mg/5 mg/25 mg capsule rigide	not available	045873041	NEOPHARMED GENTILI SPA	IT
SARASVATI 10 mg/10 mg/25 mg capsule rigide	not available	045873066	NEOPHARMED GENTILI SPA	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
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/0303/001-005/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
Egiramlon 2,5 mg/2,5 mg Tvrdé kapsuly	HU/H/0303/001-005/DC	58/0743/11-S	EGIS PHARMACEUTICALS PLC	SK
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg tvrdé kapsuly	HU/H/0303/002	58/0744/11-S	EGIS PHARMACEUTICALS PLC	SK
Егирамлон 10 mg/5 mg твърди капсули	HU/H/0303/001-005/DC	20120048	EGIS PHARMACEUTICALS PLC	BG
Egiramlon 10 mg/5 mg kemény kapszula	HU/H/0303/001-005/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/0303/001-005/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/5 mg capsule	HU/H/0303/001-005/DC	9583/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/5 mg tvrdé kapsuly	HU/H/0303/004	58/0745/11-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 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/0303/001-005/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/0303/001-005/DC	LT/1/12/2892/011-015	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 5 mg/10 mg capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/10 mg tvrdé kapsuly	HU/H/0303/003	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/0303/001-005/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/0303/001-005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg tvrdé kapsuly	HU/H/0303/005	58/0747/11-S	EGIS PHARMACEUTICALS PLC	SK
Egiramlon 10 mg/5 mg tvrdé tobolky	HU/H/0303/003	58/840/11-C	EGIS PHARMACEUTICALS PLC	CZ
Egiramlon 5 mg/10 mg capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/10 mg capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/10 mg capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	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 capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/10 mg capsule	HU/H/0303/001-005/DC	9582/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/13-15;19	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 10 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/13-15;19	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 10 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/13-15;19	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg capsule	HU/H/0303/001-005/DC	9581/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/10 mg capsule	HU/H/0303/001-005/DC	9584/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 5 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/04-06;16	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 5 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/04-06;16	EGIS PHARMACEUTICALS PLC	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
Egiramlon 5 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/04-06;16	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 5 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/07-09;17	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 5 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/07-09;17	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 5 mg/10 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/07-09;17	EGIS PHARMACEUTICALS PLC	HU
Ramlon 10 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/011-015	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/011-015	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/021-025	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/10 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/011-015	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Ramlon 5 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/006-010	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/5 mg capsule	HU/H/0303/001-005/DC	9583/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/5 mg capsule	HU/H/0303/001-005/DC	9583/2017/01-02-03-04-05-06	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 10 mg/5 mg capsule	HU/H/0303/001-005/DC	9583/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Egiramlon 10 mg/5 mg capsule	HU/H/0303/001-005/DC	9583/2017/01-02-03-04-05-06	EGIS PHARMACEUTICALS PLC	RO
Ramlon 10 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Ramlon 10 mg/5 mg kietosios kapsulės	HU/H/0303/001-005/DC	LT/1/12/2892/016-020	EGIS PHARMACEUTICALS PLC	LT
Egiramlon 10 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/10-12;18	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 10 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/10-12;18	EGIS PHARMACEUTICALS PLC	HU
Egiramlon 10 mg/5 mg kemény kapszula	HU/H/0303/001-005/DC	OGYI-T-22016/10-12;18	EGIS PHARMACEUTICALS PLC	HU
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 463 4 9	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 461 1 0	ZENTIVA FRANCE	FR

Product 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 / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 462 8 8	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 464 0 0	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 469 2 9	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 468 6 8	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 583 071 7 1	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 465 7 8	ZENTIVA FRANCE	FR
RAMIPRIL / HYDROCHLOROTHIAZIDE ZENTIVA 5 mg/12.5 mg, comprimé	not available	34009 224 466 3 9	ZENTIVA FRANCE	FR
PREMINOR 5 mg/10 mg, gélule	DK/H/2750/003	34009 301 545 5 7	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 5 mg/10 mg, gélule	DK/H/2750/003	34009 301 760 6 1	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 5 mg/10 mg, gélule	DK/H/2750/003	34009 301 760 7 8	LABORATOIRES LEURQUIN MEDIOLANUM	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril HCT-Zentiva 5 mg/25 mg tabletta	DE/H/2629/002	OGYI-T-10606/01	ZENTIVA, K.S.	HU
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742020	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742018	ZENTIVA ITALIA SRL	IT
RamiLich comp 2,5 mg/12,5 mg Tabletten	DE/H/2629/001	62981.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
RamiLich comp 5 mg/25 mg Tabletten	DE/H/2629/002	62982.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742119	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742057	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742121	ZENTIVA ITALIA SRL	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 IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742095	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742196	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742133	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742210	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742160	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742069	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742044	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742222	ZENTIVA ITALIA SRL	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 IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742083	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742158	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742107	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742172	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742208	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742246	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742032	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742071	ZENTIVA ITALIA SRL	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 IDROCLOROTIAZIDE ZENTIVA 2,5 mg/ 12,5 mg compresse	DE/H/2629/001	037742145	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742234	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742184	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742261	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742273	ZENTIVA ITALIA SRL	IT
RAMIPRIL E IDROCLOROTIAZIDE ZENTIVA 5 mg / 25 mg compresse	DE/H/2629/002	037742259	ZENTIVA ITALIA SRL	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
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
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

Product 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 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
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
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/4683/001	5715107	KRKA, D.D., NOVO MESTO	PT
Ramipril/amlodipine Krka 5 mg/5 mg hard capsules	DE/H/4683/004	PA1347/072/001	KRKA, D.D., NOVO MESTO	IE
Ramipril/amlodipine Krka 5 mg/10 mg hard capsules	DE/H/4683/003	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/4683/001	PA1347/072/004	KRKA, D.D., NOVO MESTO	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
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/4683/003	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/4683/001	H/17/02378/001	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/002	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/003	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/004	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/005	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/006	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/007	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/008	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/009	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/5 mg trde kapsule	DE/H/4683/001	H/17/02378/010	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/011	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/012	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/013	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/014	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/015	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/4683/002	H/17/02378/016	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/017	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/018	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/019	KRKA, D.D., NOVO MESTO	SI
Rameam 5 mg/10 mg trde kapsule	DE/H/4683/002	H/17/02378/020	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/021	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/022	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/023	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/024	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/025	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/026	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/027	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/028	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/029	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/5 mg trde kapsule	DE/H/4683/003	H/17/02378/030	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/031	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/032	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/4683/004	H/17/02378/033	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/034	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/035	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/036	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/037	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/038	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	H/17/02378/039	KRKA, D.D., NOVO MESTO	SI
Rameam 10 mg/10 mg trde kapsule	DE/H/4683/004	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/4683/001	10071/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/5 mg capsule	DE/H/4683/001	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

Product 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/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/4683/002	10072/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 5 mg/10 mg capsule	DE/H/4683/002	10072/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/06	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/5 mg capsule	DE/H/4683/003	10073/2017/10	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/01	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/02	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/03	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/04	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/05	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/06	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/07	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/08	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	10074/2017/09	KRKA, D.D., NOVO MESTO	RO
Ramipril/amlodipină Krka 10 mg/10 mg capsule	DE/H/4683/004	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

Product 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 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
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

Product 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	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
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/4683/002	045268214	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	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/4683/002	045268238	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268240	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268253	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268265	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268277	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268289	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268291	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/5 mg capsule rigide	DE/H/4683/002	045268303	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268315	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268327	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268339	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/4683/001	045268341	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268354	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268366	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268378	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268380	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268392	KRKA, D.D., NOVO MESTO	IT
Ramipril e Amlodipina Krka 10 mg/10 mg capsule rigide	DE/H/4683/001	045268404	KRKA, D.D., NOVO MESTO	IT
Rameam 10 mg/10 mg Hartkapseln	DE/H/4683/001	97123.00.00	TAD PHARMA GMBH	DE
Rameam 10 mg/5 mg Hartkapseln	DE/H/4683/002	97124.00.00	TAD PHARMA GMBH	DE
Rameam 5 mg/10 mg Hartkapseln	DE/H/4683/003	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/4683/001	24021	KRKA, D.D., NOVO MESTO	PL
Ramladio, 10 mg + 5 mg, kapsułki, twarde	DE/H/4683/002	24022	KRKA, D.D., NOVO MESTO	PL
Ramladio, 5 mg + 10 mg, kapsułki, twarde	DE/H/4683/003	24023	KRKA, D.D., NOVO MESTO	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramladio, 5 mg + 5 mg, kapsułki, twarde	DE/H/4683/004	24024	KRKA, D.D., NOVO MESTO	PL
Rimal 5 mg / 5 mg cietās kapsulas	LV/H/0188/001	16-0104	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 5 mg / 10 mg cietās kapsulas	LV/H/0188/002	16-0105	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 10 mg / 5 mg cietās kapsulas	LV/H/0188/003	16-0106	MEDANA PHARMA SPOLKA AKCYJNA	LV
Rimal 10 mg / 10 mg cietās kapsulas	LV/H/0188/004	16-0107	MEDANA PHARMA SPOLKA AKCYJNA	LV
Полпрам 5 mg/5 mg капсули, твърди	LV/H/0188/001	20160220	POLPHARMA	BG
Полпрам 10 mg/5 mg капсули, твърди	LV/H/0188/003	20160222	POLPHARMA	BG
Полпрам 10 mg/10 mg капсули, твърди	LV/H/0188/004	20160223	POLPHARMA	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
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
AMORA 2,5 mg/5 mg tvrde kapsule	not available	HR-H-481003043	BELUPO D.D.	HR
AMORA 2,5 mg/5 mg tvrde kapsule	not available	HR-H-481003043	BELUPO D.D.	HR
AMORA 5 mg/5 mg tvrde kapsule	not available	HR-H-582136867	BELUPO D.D.	HR
AMORA 5 mg/5 mg tvrde kapsule	not available	HR-H-582136867	BELUPO D.D.	HR
AMORA 5 mg/10 mg tvrde kapsule	not available	HR-H-729149816	BELUPO D.D.	HR
AMORA 5 mg/10 mg tvrde kapsule	not available	HR-H-729149816	BELUPO D.D.	HR
AMORA 10 mg/5 mg tvrde kapsule	not available	HR-H-100700683	BELUPO D.D.	HR
AMORA 10 mg/5 mg tvrde kapsule	not available	HR-H-100700683	BELUPO D.D.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
AMORA 10 mg/10 mg tvrde kapsule	not available	HR-H-774707043	BELUPO D.D.	HR
AMORA 10 mg/10 mg tvrde kapsule	not available	HR-H-774707043	BELUPO D.D.	HR
Ramipril-ratiopharm® comp. 5 mg/12,5 mg Tabletten	not available	84993.00.00	RATIOPHARM GMBH	DE
Рамира 5 mg/5 mg твърди капсули	not available	20180197	STADA ARZNEIMITTEL AG	BG
Рамира 10 mg/5 mg твърди капсули	not available	20180198	STADA ARZNEIMITTEL AG	BG
Рамира 10 mg/10 mg твърди капсули	not available	20180199	STADA ARZNEIMITTEL AG	BG
RAMANDIUR 5 mg/5 mg/12,5 mg capsule rigide	not available	046736029	ERREKAPPA EUROTERAPICI SPA	IT
RAMANDIUR 5 mg/5 mg/25 mg capsule rigide	not available	046736031	ERREKAPPA EUROTERAPICI SPA	IT
RAMANDIUR 10 mg/5 mg/25 mg capsule rigide	not available	046736056	ERREKAPPA EUROTERAPICI SPA	IT
RAMANDIUR 10 mg/10 mg/25 mg capsule rigide	not available	046736068	ERREKAPPA EUROTERAPICI SPA	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramizek Combi, 2,5 mg + 5 mg, kapsułki, twarde	SE/H/1336/001	21904	ADAMED PHARMA S.A.	PL
Ramizek Combi, 5 mg + 5 mg, kapsułki, twarde	SE/H/1336/002	21905	ADAMED PHARMA S.A.	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
Ramizek Combi, 5 mg + 10 mg, kapsułki, twarde	SE/H/1336/004	21907	ADAMED PHARMA S.A.	PL
Ramizek Combi, 10 mg + 5 mg, kapsułki, twarde	SE/H/1336/005	21906	ADAMED PHARMA S.A.	PL
Ramizek Combi, 10 mg + 10 mg, kapsułki, twarde	SE/H/1336/005	21908	ADAMED PHARMA S.A.	PL
Ramipril/Amlodipine Adamed, 2,5 mg/5 mg, hårda kapslar	SE/H/1336/001	49203	ADAMED PHARMA S.A.	SE
Ramipril/Amlodipine Adamed, 5 mg/5 mg, hårda kapslar	SE/H/1336/002	49204	ADAMED PHARMA S.A.	SE
Ramipril/Amlodipine Adamed, 5 mg/10 mg, hårda kapslar	SE/H/1336/004	49206	ADAMED PHARMA S.A.	SE
Ramipril/Amlodipine Adamed, 10 mg/5 mg, hårda kapslar	SE/H/1336/003	49205	ADAMED PHARMA S.A.	SE
Ramipril/Amlodipine Adamed, 10 mg/10 mg, hårda kapslar	SE/H/1336/005	49207	ADAMED PHARMA S.A.	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

Product 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 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 PHARMA S.A.	CZ
RAMIZEK 10 mg/5 mg tvrdé tobolky	SE/H/1336/003	58/379/14-C	ADAMED PHARMA S.A.	CZ
Ramizek 5 mg/10 mg tvrdé tobolky	SE/H/1336/004	58/380/14-C	ADAMED PHARMA S.A.	CZ
Ramizek 10 mg/10 mg tvrdé tobolky	SE/H/1336/005	58/381/14-C	ADAMED PHARMA S.A.	CZ
RAMIZEK 2,5 mg/5 mg tvrdé kapsuly	SE/H/1336/001	58/0026/15-S	ADAMED PHARMA S.A.	SK
RAMIZEK 5 mg/5 mg tvrdé kapsuly	SE/H/1336/002	58/0027/15-S	ADAMED PHARMA S.A.	SK
RAMIZEK 5 mg/10 mg tvrdé kapsuly	SE/H/1336/004	58/0028/15-S	ADAMED PHARMA S.A.	SK
RAMIZEK 10 mg/5 mg tvrdé kapsuly	SE/H/1336/003	58/0029/15-S	ADAMED PHARMA S.A.	SK
RAMIZEK 10 mg/10 mg tvrdé kapsuly	SE/H/1336/005	58/0030/15-S	ADAMED PHARMA S.A.	SK
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

Product 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 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
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

Product 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	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
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
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

Product 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/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 PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/02	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/03	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/04	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/05	ADAMED PHARMA S.A.	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	7199/2014/06	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/07	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/08	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/09	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 2.5 mg/5 mg capsule	SE/H/1336/001	7199/2014/10	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/01	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/02	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/03	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/04	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/05	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/06	ADAMED PHARMA S.A.	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	7200/2014/07	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/08	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/09	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/5 mg capsule	SE/H/1336/002	7200/2014/10	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/01	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/02	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/03	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/04	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/05	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/06	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/07	ADAMED PHARMA S.A.	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	7201/2014/08	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/09	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 5 mg/10 mg capsule	SE/H/1336/003	7201/2014/10	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/01	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/02	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/03	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/04	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/05	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/06	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/07	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/001-005/DC	7202/2014/08	ADAMED PHARMA S.A.	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/1336/004	7202/2014/09	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/5 mg capsule	SE/H/1336/004	7202/2014/10	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/01	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/02	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/03	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/04	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/05	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/06	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/07	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/08	ADAMED PHARMA S.A.	RO
Ramipril/Amlodipina Adamed 10 mg/10 mg capsule	SE/H/1336/005	7203/2014/09	ADAMED PHARMA S.A.	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	7203/2014/10	ADAMED PHARMA S.A.	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
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 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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tonotec 5 mg/5 mg Hartkapseln	DE/H/3919/002	88599.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec 5 mg/10 mg Hartkapseln	DE/H/3919/003	88600.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec 10 mg/5 mg Hartkapseln	DE/H/3919/004	88601.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec 10 mg/10 mg Hartkapseln	DE/H/3919/005	88602.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipin KLINGE PHARMA 10 mg/10 mg Hartkapseln	DK/H/2751/005	2200210.00.00	ADAMED PHARMA S.A.	DE
Ramipril/Amlodipin KLINGE PHARMA 10 mg/5 mg Hartkapseln	DK/H/2751/004	2200209.00.00	ADAMED PHARMA S.A.	DE
Ramipril/Amlodipin KLINGE PHARMA 5 mg/10 mg Hartkapseln	DK/H/2751/003	2200208.00.00	ADAMED PHARMA S.A.	DE
Ramipril/Amlodipin KLINGE PHARMA 5 mg/5 mg Hartkapseln	DK/H/2751/002	2200207.00.00	ADAMED PHARMA S.A.	DE
Ramipril/Amlodipin KLINGE PHARMA 2,5 mg/5 mg Hartkapseln	DK/H/2751/001	2200206.00.00	ADAMED PHARMA S.A.	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
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 Duo, 5 mg + 5 mg, kapsułki twarde	AT/H/0402/002	20264	SANDOZ GMBH	PL
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

Product 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 HEXAL plus Amlodipin 5 mg/5 mg Hartkapseln	AT/H/0402/002	84285.00.00	HEXAL AG	DE
SAMBETAN 2,5 mg + 5 mg capsule rigide	not available	044544017	CIPROS S.R.L.	IT
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 300 073 5 8	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 300 073 4 1	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 630 9 2	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 627 8 1	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 631 5 3	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 633 8 2	SANOFI-AVENTIS FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 629 0 3	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 567 144 3 8	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 628 4 2	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 626 1 3	SANOFI-AVENTIS FRANCE	FR
COTRIATEC 5 mg/12,5 mg, comprimé	FR/H/0574/001	34009 369 632 1 4	SANOFI-AVENTIS FRANCE	FR
Ramipril/Amlodipin STADA 10 mg/10 mg Hartkapseln	DE/H/6271/004	138767	STADA ARZNEIMITTEL GMBH	AT
Ramipril/Amlodipin STADA 5 mg/10 mg Hartkapseln	DE/H/6271/002	138765	STADA ARZNEIMITTEL GMBH	AT
Ramipril/Amlodipin STADA 10 mg/5 mg Hartkapseln	DE/H/6271/003	138766	STADA ARZNEIMITTEL GMBH	AT
Ramipril/Amlodipin STADA 5 mg/5 mg Hartkapseln	DE/H/6271/001	138764	STADA ARZNEIMITTEL GMBH	AT
Ramipril/Amlodipin AL 10 mg/5 mg Hartkapseln	DE/H/6271/003	99884.00.00	ALIUD PHARMA GMBH	DE
Ramipril/Amlodipin AL 5 mg/5 mg Hartkapseln	DE/H/6271/001	99882.00.00	ALIUD PHARMA GMBH	DE
Ramipril/Amlodipin AL 5 mg/10 mg Hartkapseln	DE/H/6271/002	99883.00.00	ALIUD PHARMA GMBH	DE
Ramipril/Amlodipin AL 10 mg/10 mg Hartkapseln	DE/H/6271/004	99885.00.00	ALIUD PHARMA GMBH	DE
Ramipril/Amlodipine EG 5 mg/5 mg Hartkapseln	DE/H/6271/001	BE540640	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/5 mg harde capsules	DE/H/6271/003	BE540666	EUROGENERICS N.V./S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipine EG 5 mg/5 mg gélules	DE/H/6271/001	BE540640	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/10 mg gélules	DE/H/6271/004	BE540675	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 5 mg/10 mg gélules	DE/H/6271/002	BE540657	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/10 mg harde capsules	DE/H/6271/004	BE540675	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/5 mg gélules	DE/H/6271/003	BE540666	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/10 mg Hartkapseln	DE/H/6271/004	BE540675	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 5 mg/10 mg harde capsules	DE/H/6271/002	BE540657	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 5 mg/10 mg Hartkapseln	DE/H/6271/002	BE540657	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 10 mg/5 mg Hartkapseln	DE/H/6271/003	BE540666	EUROGENERICS N.V./S.A.	BE
Ramipril/Amlodipine EG 5 mg/10 mg harde capsules	DE/H/6271/001	BE540640	EUROGENERICS N.V./S.A.	BE
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721369	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721078	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721066	EG 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
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721104	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721332	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721015	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721054	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721080	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721344	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721041	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721357	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721039	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721167	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721128	EG 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
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721320	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721155	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721181	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721092	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721116	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721217	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 5 mg capsule rigide	DE/H/6271/001	046721027	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721229	EG S.P.A.	IT
Ramipril e Amlodipina EG 5 mg + 10 mg capsule rigide	DE/H/6271/002	046721130	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721268	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721142	EG 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
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721256	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721179	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721282	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721193	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 5 mg capsule rigide	DE/H/6271/003	046721205	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721318	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721306	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721231	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721270	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721243	EG S.P.A.	IT
Ramipril e Amlodipina EG 10 mg + 10 mg capsule rigide	DE/H/6271/004	046721294	EG S.P.A.	IT
Ramipril/Amlodipine EG 5 mg/10 mg gélules	DE/H/6271/002	2019110256	EUROGENERICS N.V./S.A.	LU

Product 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 EG 10 mg/5 mg gélules	DE/H/6271/003	2019110257	EUROGENERICS N.V./S.A.	LU
Ramipril/Amlodipine EG 10 mg/10 mg gélules	DE/H/6271/004	2019110258	EUROGENERICS N.V./S.A.	LU
Ramipril/Amlodipine EG 5 mg/5 mg gélules	DE/H/6271/001	2019110255	EUROGENERICS N.V./S.A.	LU
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril/Amlodipine Apotex 5 mg/5 mg harde capsules	SE/H/1396/001	BE514595	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 5 mg/10 mg harde capsules	SE/H/1396/003	BE514604	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 10 mg/5 mg harde capsules	SE/H/1396/002	BE514613	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 10 mg/10 mg, harde capsule	SE/H/1396/004	BE514622	APOTEX EUROPE B.V.	BE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramomark 5 mg/5 mg tvrdé tobolky	SE/H/1337/001	58/281/14-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Ramipril/Amlodipine Glenmark, 10 mg/10 mg, hårda kapslar	SE/H/1337/004	49211	GLENMARK ARZNEIMITTEL GMBH	SE
Ramipril/Amlodipine Glenmark, 5 mg/5 mg, hårda kapslar	SE/H/1337/001	49208	GLENMARK ARZNEIMITTEL GMBH	SE
Ramipril/Amlodipine Glenmark, 10 mg/5 mg, hårda kapslar	SE/H/1337/002	49209	GLENMARK ARZNEIMITTEL GMBH	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 Glenmark, 5 mg/10 mg, hårda kapslar	SE/H/1337/003	49210	GLENMARK ARZNEIMITTEL GMBH	SE
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 5 mg/5 mg tvrdé kapsuly	SE/H/1337/001	58/0204/14-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
PREMINOR 10 mg/10 mg, gélule	DK/H/2750/005	34009 301 545 2 6	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 10 mg/10 mg, gélule	DK/H/2750/005	34009 301 761 3 9	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 10 mg/10 mg, gélule	DK/H/2750/005	34009 301 761 4 6	LABORATOIRES LEURQUIN MEDIOLANUM	FR
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 Duo, 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
Ramipril/Amlodipin/HCT 1A Pharma 10 mg/5 mg/25 mg – Hartkapseln	PL/H/0490/005	138680	1A PHARMA GMBH	AT
Ramipril/Amlodipin/HCT 1A Pharma 5 mg/5 mg/12,5 mg – Hartkapseln	PL/H/0490/002	138678	1A PHARMA GMBH	AT
Ramipril/Amlodipin/HCT 1A Pharma 5 mg/5 mg/25 mg – Hartkapseln	PL/H/0490/003	138675	1A PHARMA GMBH	AT
Ramipril/Amlodipin/HCT 1A Pharma 10 mg/10 mg/25 mg – Hartkapseln	PL/H/0490/006	138679	1A PHARMA GMBH	AT
Prylar H 10 mg/5 mg/25 mg tvrde kapsule	PL/H/0490/005	HR-H-395953294	SANDOZ D.O.O.	HR
Prylar H 5 mg/5 mg/12.5 mg tvrde kapsule	PL/H/0490/002	HR-H-115715303	SANDOZ D.O.O.	HR
Prylar H 5 mg/5 mg/25 mg tvrde kapsule	PL/H/0490/003	HR-H-378651202	SANDOZ D.O.O.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Prylar H 10 mg/10 mg/25 mg tvrde kapsule	PL/H/0490/006	HR-H-394687607	SANDOZ D.O.O.	HR
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/001	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/002	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/003	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/004	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/005	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/006	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/10 mg/25 mg kietosios kapsulės	PL/H/0490/006	LT/1/19/4357/007	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/001	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/003	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/004	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/005	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg /25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/006	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg /25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/007	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/001	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/002	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/003	SANDOZ PHARMACEUTICALS D.D.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/004	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/005	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/006	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/12,5 mg kietosios kapsulės	PL/H/0490/002	LT/1/19/4354/007	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/001	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/002	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/003	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/004	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/005	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/006	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 5 mg/5 mg/25 mg kietosios kapsulės	PL/H/0490/003	LT/1/19/4355/007	SANDOZ PHARMACEUTICALS D.D.	LT
Prylar 10 mg/5 mg /25 mg kietosios kapsulės	PL/H/0490/005	LT/1/19/4356/002	SANDOZ PHARMACEUTICALS D.D.	LT
Ramdacordia HCT 10 mg/10 mg/25 mg cietās kapsulas	PL/H/0490/006	19-0051	SANDOZ PHARMACEUTICALS D.D.	LV
Ramdacordia HCT 5 mg/5 mg/12,5 mg cietās kapsulas	PL/H/0490/002	19-0048	SANDOZ PHARMACEUTICALS D.D.	LV
Ramdacordia HCT 5 mg/5 mg/25 mg cietās kapsulas	PL/H/0490/003	19-0049	SANDOZ PHARMACEUTICALS D.D.	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
Ramdacordia HCT 10 mg/5 mg/25 mg cietās kapsulas	PL/H/0490/005	19-0050	SANDOZ PHARMACEUTICALS D.D.	LV
Sumilar HCT, 5 mg + 5 mg + 25 mg, kapsułki, twarde	PL/H/0490/003	25403	SANDOZ GMBH	PL
Sumilar HCT, 5 mg + 5 mg + 12,5 mg, kapsułki, twarde	PL/H/0490/002	25402	SANDOZ GMBH	PL
Sumilar HCT, 10 mg + 5 mg + 25 mg, kapsułki, twarde	PL/H/0490/005	25404	SANDOZ GMBH	PL
Sumilar HCT, 10 mg + 10 mg + 25 mg, kapsułki, twarde	PL/H/0490/006	25405	SANDOZ GMBH	PL
Appunto 10 mg/10 mg/25 mg Hartkapseln	PL/H/0490/006	2200508.00.00	HEXAL AG	DE
Appunto 10 mg/5 mg/25 mg Hartkapseln	PL/H/0490/005	2200507.00.00	HEXAL AG	DE
Appunto 5 mg/5 mg/12,5 mg Hartkapseln	PL/H/0490/002	2200504.00.00	HEXAL AG	DE
Appunto 5 mg/5 mg/25 mg Hartkapseln	PL/H/0490/003	2200505.00.00	HEXAL AG	DE
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	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
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	958/22-7-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+5) mg, καψάκια, σκληρά	EL/H/0179/002	6234/22-07-2015	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	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
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	32088/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	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**
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	36547/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	39917/22-07-15	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	GR
Diaspil, (2,5+5) mg, καψάκια, σκληρά	EL/H/0179/001	023131	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	CY
Diaspil, (5+5 mg, καψάκια, σκληρά	EL/H/0179/002	023132	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	CY
Diaspil, (5+10) mg, καψάκια, σκληρά	EL/H/0179/004	023134	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	CY
Diaspil, (10+5) mg, καψάκια, σκληρά	EL/H/0179/003	023133	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	CY
Diaspil, (10+10) mg, καψάκια, σκληρά	EL/H/0179/005	023135	ANGELINI PHARMA HELLAS S.A. HELLAS S.A.	CY
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Mepranil 2,5 mg + 5 mg cápsula	DK/H/2750/001/DC	5765474	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Mepranil 2,5 mg + 5 mg cápsula	DK/H/2750/001	5755350	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Mepranil 5 mg + 5 mg cápsula	DK/H/2750/002	5755368	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Mepranil 5 mg + 10 mg cápsula	DK/H/2750/003	5755913	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Mepranil 10 mg + 5 mg cápsula	DK/H/2750/004	5755905	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Mepranil 10 mg + 10 mg cápsula	DK/H/2750/005	5755921	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Ramipril/HCT Actavis 5 mg/12,5 mg Tabletten	not available	1-29014	ACTAVIS GROUP PTC EHF.	AT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
TRITAZIDE 5 MG/25 MG TABLETE	not available	HR-H-378024673-01	SANOFI-AVENTIS CROATIA D.O.O.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tritazide 2,5 mg/12,5 mg tablete	not available	HR-H-290180500-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606011	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606023	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606035	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606047	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606050	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606062	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606074	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606086	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606098	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606100	ZENTIVA, K.S.	IT
Bisar 2,5 mg/5 mg capsule rigide	SE/H/1340/001	045606516	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606112	ZENTIVA, K.S.	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
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606124	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606136	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606528	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606148	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606151	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606163	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606175	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606187	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606199	ZENTIVA, K.S.	IT
Bisar 5 mg/5 mg capsule rigide	SE/H/1340/002	045606201	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606415	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606427	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606439	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606555	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606441	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606454	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606466	ZENTIVA, K.S.	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
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606478	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606480	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606492	ZENTIVA, K.S.	IT
Bisar 10 mg/10 mg capsule rigide	SE/H/1340/005	045606504	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606314	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606326	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606338	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606542	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606340	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606353	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606365	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606377	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606389	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606391	ZENTIVA, K.S.	IT
Bisar 10 mg/5 mg capsule rigide	SE/H/1340/003	045606403	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606213	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606225	ZENTIVA, K.S.	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
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606237	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606530	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606249	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606252	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606264	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606276	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606288	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606290	ZENTIVA, K.S.	IT
Bisar 5 mg/10 mg capsule rigide	SE/H/1340/004	045606302	ZENTIVA, K.S.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
RamiLich comp 5 mg/12,5 mg Tabletten	DE/H/6479/001	92543.00.00	ZENTIVA PHARMA GMBH	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Triatec Composto 2,5 mg +12,5 mg comprimidos	PT/H/2457/001	5907084	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Triatec Composto 2,5 mg +12,5 mg comprimidos	PT/H/2457/001	2311181	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Triatec Composto Forte 5 mg + 25 mg comprimidos	PT/H/2457/002	5907183	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Triatec Composto Forte 5 mg + 25 mg comprimidos	PT/H/2457/002	2311389	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Triatec Composto 2,5 mg +12,5 mg comprimidos	PT/H/2457/001	2311082	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Triatec Composto Forte 5 mg + 25 mg comprimidos	PT/H/2457/002	2311280	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
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
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/01	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/02	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/03	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/04	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/5 mg kemény kapszula	not available	OGYI-T-23118/05	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/06	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/07	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/08	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/09	KRKA, D.D., NOVO MESTO	HU
Rawuro 5 mg/10 mg kemény kapszula	not available	OGYI-T-23118/10	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/11	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/12	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/13	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/14	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/5 mg kemény kapszula	not available	OGYI-T-23118/15	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/16	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/17	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/18	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/19	KRKA, D.D., NOVO MESTO	HU
Rawuro 10 mg/10 mg kemény kapszula	not available	OGYI-T-23118/20	KRKA, D.D., NOVO MESTO	HU
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
PREMINOR 5 mg/5 mg, gélule	DK/H/2750/002	34009 301 545 3 3	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 5 mg/5 mg, gélule	DK/H/2750/002	34009 301 761 1 5	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 5 mg/5 mg, gélule	DK/H/2750/002	34009 301 761 2 2	LABORATOIRES LEURQUIN MEDIOLANUM	FR
Ramipril/Amlodipin Hexal 2,5 mg/2,5 mg - Hartkapseln	AT/H/0402/001	1-31215	HEXAL PHARMA GMBH	AT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril-comp PUREN 5 mg/12,5 mg Tabletten	DE/H/1597/001	72682.00.00	PUREN PHARMA GMBH & CO. KG	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril comp. AbZ 5 mg/12,5 mg Tabletten	not available	84995.00.00	ABZ-PHARMA GMBH	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
PREMINOR 10 mg/5 mg, gélule	DK/H/2750/004	34009 301 545 4 0	LABORATOIRES LEURQUIN MEDIOLANUM	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PREMINOR 10 mg/5 mg, gélule	DK/H/2750/004	34009 301 760 8 5	LABORATOIRES LEURQUIN MEDIOLANUM	FR
PREMINOR 10 mg/5 mg, gélule	DK/H/2750/004	34009 301 760 9 2	LABORATOIRES LEURQUIN MEDIOLANUM	FR
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
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

Product 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 5 mg/10 mg capsule rigide	AT/H/0469/003	042384053	PFIZER ITALIA S.R.L.	IT
Icomb 10 mg/10 mg capsule rigide	AT/H/0469/005	042384091	PFIZER ITALIA S.R.L.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril Aristo plus Amlodipin 5 mg/5 mg Hartkapseln	SE/H/1340/002	2200100.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 5 mg/10 mg Hartkapseln	DE/H/7023/003	2200102.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 10 mg/5 mg Hartkapseln	SE/H/1340/004	2200101.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 10 mg/10 mg Hartkapseln	SE/H/1340/005	2200103.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Ramipril Aristo plus Amlodipin 2,5 mg/5 mg Hartkapseln	SE/H/1340/001	2200099.00.00	ARISTO PHARMA GMBH (ART 57)	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
Amlodipino/ramipril Aristo 10 mg/5 mg cápsulas duras	SE/H/1340/004	82.689	ARISTO PHARMA GMBH (ART 57)	ES
Amlodipino/ramipril Aristo 10 mg/10 mg cápsulas duras	SE/H/1340/005	82.690	ARISTO PHARMA GMBH (ART 57)	ES
Amlodipino/ramipril Aristo 5 mg/10 mg cápsulas duras	SE/H/1340/003	82.688	ARISTO PHARMA GMBH (ART 57)	ES
Amlodipino/ramipril Aristo 5 mg/2,5 mg cápsulas duras	SE/H/1340/001	82.686	ARISTO PHARMA GMBH (ART 57)	ES
Amlodipino/ramipril Aristo 5 mg/5 mg cápsulas duras	SE/H/1340/002	82.687	ARISTO PHARMA GMBH (ART 57)	ES
Pinmirol 5 mg/5 mg tvrde kapsule	not available	HR-H-700051702	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Pinmirol 5 mg/10 mg tvrde kapsule	not available	HR-H-694862571	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Pinmirol 10 mg/5 mg tvrde kapsule	not available	HR-H-156765293	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Pinmirol 10 mg/10 mg tvrde kapsule	not available	HR-H-015536210	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
RAMIPRIL HCT DOC Generici 2,5 mg/12,5 mg, tabletten	NL/H/0721/001	RVG 30670	DOC GENERICI S.R.L.	NL
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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/1468/003	51824	ACTAVIS GROUP PTC EHF.	SE
Ramipril/Amlodipine Actavis 5 mg/10 mg, capsule, hard	SE/H/1468/004	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
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 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/06	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/5 mg capsule	SE/H/1468/002	13848/2021/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/5 mg capsule	SE/H/1468/002	13848/2021/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 5 mg/10 mg capsule	SE/H/1468/003	13849/2021/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/04	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/5 mg capsule	SE/H/1468/004	13850/2021/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/5 mg capsule	SE/H/1468/004	13850/2021/10	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/01	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/02	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/03	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/04	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/05	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/06	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/07	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/08	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/09	ACTAVIS GROUP PTC EHF.	RO
Vivace Plus 10 mg/10 mg capsule	SE/H/1468/005	13851/2021/10	ACTAVIS GROUP PTC EHF.	RO
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	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
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Pinmirol Plus 5 mg/5 mg/12,5 mg tvrde kapsule	PL/H/0487/002	HR-H-258726339	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Pinmirol Plus 5 mg/5 mg/25 mg tvrde kapsule	PL/H/0487/003	HR-H-892087317	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Pinmirol Plus 10 mg/5 mg/25 mg tvrde kapsule	PL/H/0487/005	HR-H-988558187	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Pinmirol Plus 10 mg/10 mg/25 mg tvrde kapsule	PL/H/0487/006	HR-H-356194718	GENERICON PHARMA GESELLSCHAFT M.B.H.	HR
Ramipril/Amlodipin/HCT Genericon 5 mg/5 mg/12,5 mg Hartkapseln	PL/H/0487/002	138772	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin/HCT Genericon 5 mg/5 mg/25 mg Hartkapseln	PL/H/0487/003	138770	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin/HCT Genericon 10 mg/5 mg/25 mg Hartkapseln	PL/H/0487/005	138774	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Ramipril/Amlodipin/HCT Genericon 10 mg/10 mg/25 mg Hartkapseln	PL/H/0487/006	138773	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
RAMLOID, capsule rigide da 5 mg/5 mg/12,5 mg	PL/H/0487/002	048190019	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/12,5 mg	PL/H/0487/002	048190021	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/12,5 mg	PL/H/0487/002	048190033	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/12,5 mg	PL/H/0487/002	048190045	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/12,5 mg	PL/H/0487/002	048190058	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/25 mg	PL/H/0487/003	048190072	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/25 mg	PL/H/0487/003	048190084	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/25 mg	PL/H/0487/003	048190096	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 5 mg/5 mg/25 mg	PL/H/0487/003	048190108	ADAMED 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
RAMLOID, capsule rigide da 5 mg/5 mg/25 mg	PL/H/0487/003	048190110	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/5 mg/25 mg	PL/H/0487/005	048190122	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/5 mg/25 mg	PL/H/0487/005	048190134	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/5 mg/25 mg	PL/H/0487/005	048190146	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/5 mg/25 mg	PL/H/0487/005	048190159	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/5 mg/25 mg	PL/H/0487/005	048190161	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/10 mg/25 mg	PL/H/0487/006	048190173	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/10 mg/25 mg	PL/H/0487/006	048190185	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/10 mg/25 mg	PL/H/0487/006	048190197	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/10 mg/25 mg	PL/H/0487/006	048190209	ADAMED S.R.L.	IT
RAMLOID, capsule rigide da 10 mg/10 mg/25 mg	PL/H/0487/006	048190211	ADAMED S.R.L.	IT
Paniltri, 5 mg/5 mg/12,5 mg cápsulas	PL/H/0487/002	5773122	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 5 mg/5 mg/12,5 mg cápsulas	PL/H/0487/002	5773130	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 5 mg/5 mg/12,5 mg cápsulas	PL/H/0487/002	5773148	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 10 mg/5 mg/25 mg cápsulas	PL/H/0487/005	5773866	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Paniltri, 10 mg/5 mg/25 mg cΤ psulas	PL/H/0487/005	5773874	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 10 mg/5 mg/25 mg cΤ psulas	PL/H/0487/005	5773908	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 10 mg/10 mg/25 mg cΤ psulas	PL/H/0487/006	5773940	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
Paniltri, 10 mg/10 mg/25 mg cΤ psulas	PL/H/0487/006	5773957	LABORATÓRIO MEDINFAR - PRODUTOS FARMACÊUTICOS, S.A.	PT
RAMI-AMLO PLUS, 5 mg/5 mg/25 mg, καψάκιο, σκληρό	PL/H/0487/003	103899/03-09-2019	IASIS PHARMA	GR
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
RAMIPRIL/AMLODIPINA/IDROCLOROTIAZIDE DOC 5 mg/5 mg/12,5 mg capsule rigide	not available	046501021	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/AMLODIPINA/IDROCLOROTIAZIDE DOC 5 mg/5 mg/25 mg capsule rigide	not available	046501033	DOC GENERICI S.R.L.	IT
RAMIPRIL/AMLODIPINA/IDROCLOROTIAZIDE DOC 10 mg/5 mg/25 mg capsule rigide	not available	046501058	DOC GENERICI S.R.L.	IT
RAMIPRIL/AMLODIPINA/IDROCLOROTIAZIDE DOC 10 mg/10 mg/25 mg capsule rigide	not available	046501060	DOC GENERICI S.R.L.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Piramil Combi 5 mg/10 mg tvrdé tobolky	AT/H/0402/004	58/363/12-C	SANDOZ S.R.O.	CZ
Sumilar Duo, 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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/amlodipin Pliva 10 mg/5 mg tvrde kapsule	DE/H/5152/003	HR-H-620376275	PLIVA HRVATSKA D.O.O.	HR
Ramipril/amlodipin Pliva 5 mg/10 mg tvrde kapsule	DE/H/5152/004	HR-H-750926919	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
Ramipril/Amlodipin AbZ 5 mg/10 mg Hartkapseln	DE/H/5152/003	99417.00.00	ABZ-PHARMA GMBH	DE
Ramipril/Amlodipin AbZ 5 mg/5 mg Hartkapseln	DE/H/5152/002	99415.00.00	ABZ-PHARMA GMBH	DE
Ramipril/Amlodipin AbZ 2,5 mg/5 mg Hartkapseln	DE/H/5152/001	99414.00.00	ABZ-PHARMA GMBH	DE
Ramipril/Amlodipin AbZ 10 mg/5 mg Hartkapseln	DE/H/5152/004	99418.00.00	ABZ-PHARMA GMBH	DE
Ramipril/Amlodipin AbZ 10 mg/10 mg Hartkapseln	DE/H/5152/005	99419.00.00	ABZ-PHARMA GMBH	DE
Ramipril/amlodipin Pliva 5 mg/5 mg tvrde kapsule	DE/H/5152/002	HR-H-023212604	PLIVA HRVATSKA D.O.O.	HR
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570114	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570138	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570126	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570064	TEVA B.V	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 Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570090	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570241	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570037	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570088	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570052	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570239	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570013	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570049	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570215	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570254	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570153	TEVA B.V	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 Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570266	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/10 mg capsule rigide	DE/H/5152/004	045570140	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570102	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570025	TEVA B.V	IT
Ramipril e Amlodipina Teva 5 mg/5 mg capsule rigide	DE/H/5152/002	045570227	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/5 mg capsule rigide	DE/H/5152/003	045570076	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570189	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570165	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570203	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570280	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570191	TEVA B.V	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 Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570177	TEVA B.V	IT
Ramipril e Amlodipina Teva 10 mg/10 mg capsule rigide	DE/H/5152/005	045570278	TEVA B.V	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Piramil Combi 10 mg/5 mg tvrdé tobolky	AT/H/0402/003	58/362/12-C	SANDOZ S.R.O.	CZ
Sumilar Duo, 5 mg + 10 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
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril/Amlodipine Apotex 5 mg/5 mg harde capsules	SE/H/1396/001	BE514595	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 5 mg/10 mg harde capsules	SE/H/1396/003	BE514604	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 10 mg/5 mg harde capsules	SE/H/1396/002	BE514613	APOTEX EUROPE B.V.	BE
Ramipril/Amlodipine Apotex 10 mg/10 mg, harde capsule	SE/H/1396/004	BE514622	APOTEX EUROPE B.V.	BE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
RAMIPRIL E AMLODIPINA DOC Generici 2,5 mg + 5 mg capsule rigide	not available	044802015	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 2,5 mg + 5 mg capsule rigide	not available	044802027	DOC GENERICI S.R.L.	IT
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
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
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

Product 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
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
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

Product 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 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 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, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	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, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 mg, kapsułki, twarde	CZ/H/0674/004	22125	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 5 mg + 10 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
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
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
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

Product 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 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, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Amrap, 10 mg + 5 mg, kapsułki, twarde	CZ/H/0674/003	22124	SANOFI-AVENTIS SP Z.O.O.	PL
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 5 mg/5 mg твърди капсули	CZ/H/0674/002	20140233	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Тритейс комбо 10 mg/5 mg твърди капсули	CZ/H/0674/003	20140227	SANOFI BULGARIA EOOD	BG
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405428	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405479	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405378	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405277	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405455	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405505	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405289	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405416	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405467	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405442	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405430	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405493	SANOFI S.P.A	IT
TRIAMLO 10 mg/10 mg, capsule rigide	CZ/H/0674/005	043405481	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405253	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405291	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405303	SANOFI 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
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405238	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405214	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405265	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405240	SANOFI S.P.A	IT
TRIAMLO 5 mg/10 mg, capsule rigide	CZ/H/0674/004	043405226	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405366	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405315	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405327	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405380	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405392	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405404	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405354	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405341	SANOFI S.P.A	IT
TRIAMLO 10 mg/5 mg, capsule rigide	CZ/H/0674/003	043405339	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405051	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405101	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405075	SANOFI 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
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405149	SANOFI S.R.L.	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405036	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405087	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405012	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405048	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405024	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405099	SANOFI S.P.A	IT
TRIAMLO 2,5 mg/5 mg, capsule rigide	CZ/H/0674/001	043405063	SANOFI S.P.A	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405137	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405190	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405125	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405176	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405164	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405113	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405188	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405202	SANOFI S.R.L.	IT
TRIAMLO 5 mg/5 mg, capsule rigide	CZ/H/0674/002	043405152	SANOFI 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
Parvati 5 mg + 5 mg capsule rigide	not available	043322027	NEOPHARMED GENTILI SPA	IT
Parvati 2,5 mg + 5 mg capsule rigide	not available	043322015	NEOPHARMED GENTILI SPA	IT
Parvati 5 mg + 10 mg capsule rigide	not available	043322041	NEOPHARMED GENTILI SPA	IT
Parvati 10 mg + 5 mg capsule rigide	not available	043322039	NEOPHARMED GENTILI SPA	IT
Parvati 10 mg + 10 mg capsule rigide	not available	043322054	NEOPHARMED GENTILI SPA	IT
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Tonotec HCT 5 mg/5 mg/12,5 mg Hartkapseln	DE/H/6378/001	2200498.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec HCT 5 mg/5 mg/ 25 mg Hartkapseln	DE/H/6378/002	2200499.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec HCT 10 mg/5 mg/25 mg Hartkapseln	DE/H/6378/003	2200501.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Tonotec HCT 10 mg/10 mg/25 mg Hartkapseln	DE/H/6378/004	2200502.00.00	APONTIS PHARMA GMBH & CO. KG	DE
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT
Ramipril/Amlodipín/Hydrochlorotiazid Adamed 5 mg/5 mg/12,5 mg tvrdé kapsuly	PL/H/0487/002	58/0196/19-S	ADAMED PHARMA S.A.	SK
Ramipril/Amlodipín/Hydrochlorotiazid Adamed 5 mg/5 mg/25 mg tvrdé kapsuly	PL/H/0487/003	58/0197/19-S	ADAMED PHARMA S.A.	SK
Ramipril/Amlodipín/Hydrochlorotiazid Adamed 10 mg/5 mg/25 mg tvrdé kapsuly	PL/H/0487/005	58/0198/19-S	ADAMED PHARMA S.A.	SK
Ramipril/Amlodipín/Hydrochlorotiazid Adamed 10 mg/10 mg/25 mg tvrdé kapsuly	PL/H/0487/006	58/0199/19-S	ADAMED PHARMA S.A.	SK
Ramipril + Amlodipine + Hydrochlorothiazide Adamed, 10 mg + 10 mg + 25 mg, kapsułki, twarde	PL/H/0487/006	25568	ADAMED PHARMA S.A.	PL
Ramipril + Amlodipine + Hydrochlorothiazide Adamed, 5 mg + 5 mg + 25 mg, kapsułki, twarde	PL/H/0487/003	25565	ADAMED PHARMA S.A.	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 + Amlodipine + Hydrochlorothiazide Adamed, 5 mg + 5 mg + 12,5 mg, kapsułki, twarde	PL/H/0487/002	25564	ADAMED PHARMA S.A.	PL
Ramipril + Amlodipine + Hydrochlorothiazide Adamed, 10 mg + 5 mg + 25 mg, kapsułki, twarde	PL/H/0487/005	25567	ADAMED PHARMA S.A.	PL
Ramipril + Amlodipine + Hydrochlorothiazide Adamed, 2,5 mg + 5 mg + 12,5 mg, kapsułki, twarde	PL/H/0487/001	25563	ADAMED PHARMA S.A.	PL
Ramipril + Amlodipine + Hydrochlorothiazide Adamed, 5 mg + 10 mg + 25 mg, kapsułki, twarde	PL/H/0487/004	25566	ADAMED PHARMA S.A.	PL
Рамиприл/Амлодипин/Хидрохлоротиазид Адамед 5 mg/5 mg/25 mg твърди капсули	PL/H/0487/003	20190236	ADAMED PHARMA S.A.	BG
Рамиприл/Амлодипин/Хидрохлоротиазид Адамед 10 mg/10 mg/25 mg твърди капсули	PL/H/0487/006	20190238	ADAMED PHARMA S.A.	BG
Рамиприл/Амлодипин/Хидрохлоротиазид Адамед 10 mg/5 mg/25 mg твърди капсули	PL/H/0487/005	20190237	ADAMED PHARMA S.A.	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Рамиприл/Амлодипин/Хидрохлоротиазид Адамед 5 mg/5 mg/12,5 mg твърди капсули	PL/H/0487/002	20190235	ADAMED PHARMA S.A.	BG
Ramipril + Amlodipin + Hydrochlorotiazid Adamed 5 mg + 5 mg + 12,5 mg Hartkapseln	PL/H/0487/002	2200492.00.00	ADAMED PHARMA S.A.	DE
Ramipril + Amlodipin + Hydrochlorotiazid Adamed 5 mg + 5 mg + 25 mg Hartkapseln	PL/H/0487/003	2200493.00.00	ADAMED PHARMA S.A.	DE
Ramipril + Amlodipin + Hydrochlorotiazid Adamed 10 mg + 5 mg + 25 mg Hartkapseln	PL/H/0487/005	2200495.00.00	ADAMED PHARMA S.A.	DE
Ramipril + Amlodipin + Hydrochlorotiazid Adamed 10 mg + 10 mg + 25 mg Hartkapseln	PL/H/0487/006	2200496.00.00	ADAMED PHARMA S.A.	DE
Ramipril/Amlodipine/Hydrochlorothiazide Adamed 10 mg/10 mg/25 mg tvrdé tobolky	PL/H/0487/006	58/196/17-C	ADAMED PHARMA S.A.	CZ
Ramipril/Amlodipine/Hydrochlorothiazide Adamed 10 mg/5 mg/25 mg tvrdé tobolky	PL/H/0487/005	58/195/17-C	ADAMED PHARMA S.A.	CZ
Ramipril/Amlodipine/Hydrochlorothiazide Adamed 5 mg/5 mg/12,5 mg tvrdé tobolky	PL/H/0487/002	58/192/17-C	ADAMED PHARMA S.A.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ramipril/Amlodipine/Hydrochlorothiazide Adamed 5 mg/5 mg/25 mg tvrdé tobolky	PL/H/0487/003	58/193/17-C	ADAMED PHARMA S.A.	CZ
RAMI-AMLO PLUS, 5 mg/5 mg/12,5 mg, καψάκιο, σκληρό	PL/H/0487/002	103898/03-09-2019	IASIS PHARMA	GR
RAMI-AMLO PLUS, 10 mg/5 mg/25 mg, καψάκιο, σκληρό	PL/H/0487/005	103900/03-09-2019	IASIS PHARMA	GR
RAMI-AMLO PLUS, 10 mg/10 mg/25 mg, καψάκιο, σκληρό	PL/H/0487/006	103901/03-09-2019	IASIS PHARMA	GR
Vesdil® 5 plus 5 mg/25 mg Tabletten	DE/H/2813/002	26623.00.00	ASTRAZENECA GMBH	DE
Vesdil® 2,5 plus 2,5 mg/12,5 mg Tabletten	DE/H/2813/001	26623.01.00	ASTRAZENECA GMBH	DE
UNIPRILDIUR 2,5 mg + 12,5 mg compresse	DE/H/2813/001	028532012	ASTRAZENECA S.P.A.	IT
UNIPRILDIUR 5 mg + 25 mg compresse	DE/H/2813/002	028532024	ASTRAZENECA S.P.A.	IT