9 June 2017
EMA/369195/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): amlodipine / perindopril

Procedure No.: PSUSA/00000179/201610

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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
[COVERAM 10 mg/10 mg tabletit]	FR/H/0325/004	23483	LES LABORATOIRES SERVIER	FI
[COVERAM 10 mg/5 mg tabletit]	FR/H/0325/003	23482	LES LABORATOIRES SERVIER	FI
[COVERAM 5 mg/10 mg tabletit]	FR/H/0325/002	23481	LES LABORATOIRES SERVIER	FI
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477511/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477547/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477562/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477461/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477523/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477509/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477550/M	LES LABORATOIRES SERVIER	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
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477535/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477497/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477434/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477459/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477473/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477446/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/10 mg] compresse	FR/H/0325/004	038477485/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477410	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477319	LES LABORATOIRES SERVIER	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
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477422	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477321	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477384	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477372	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477295	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477360	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477396	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477333	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477307	LES LABORATOIRES SERVIER	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
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477408	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477358	LES LABORATOIRES SERVIER	IT
[COVERLAM 10 mg/5 mg] compresse	FR/H/0325/003	038477345	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477156	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477218	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477283	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477269	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477194	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477170	LES LABORATOIRES SERVIER	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
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477206	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477271	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477244	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477232/M	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477170	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477168	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477182	LES LABORATOIRES SERVIER	IT
[COVERLAM 5 mg/10 mg] compresse	FR/H/0325/002	038477220	LES LABORATOIRES SERVIER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483513	IST. FARMACO BIOLOGICO SRL STRODER	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
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483549	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483448	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483451	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483436	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483537	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483552	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483525	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483463	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483501	IST. FARMACO BIOLOGICO SRL STRODER	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
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483487	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483475	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483424	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/10 mg] compresse	FR/H/0326/004	038483499	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483323	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483347	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483297	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483285	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483335	IST. FARMACO BIOLOGICO SRL STRODER	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
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483311	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483412	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483398	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483362	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483350	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483400	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483386	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483374	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 10 mg/5 mg] compresse	FR/H/0326/003	038483309	IST. FARMACO BIOLOGICO SRL STRODER	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
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483234	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483259	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483196	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483184	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483145	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483172	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483210	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483246	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483158	IST. FARMACO BIOLOGICO SRL STRODER	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
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483208	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483222	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483261	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483273	IST. FARMACO BIOLOGICO SRL STRODER	IT
[REAPTAN 5 mg/10 mg] compresse	FR/H/0326/002	038483160	IST. FARMACO BIOLOGICO SRL STRODER	IT
3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/001	LES LABORATOIRES SERVIER (SURESNES)	LT
ACERYCAL 10MG/10MG TABLETS	FR/H/0325/004	PA 568/018/004	LES LABORATOIRES SERVIER (SURESNES)	IE
ACERYCAL 10mg/10mg tablets	FR/H/325/04/DC	PL 05815/0059	LES LABORATOIRES SERVIER (SURESNES)	UK
ACERYCAL 10MG/5MG TABLETS	FR/H/0325/003	PA 568/018/003	LES LABORATOIRES SERVIER (SURESNES)	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
ACERYCAL 10mg/5mg tablets	FR/H/325/03/DC	PL 05815/0058	LES LABORATOIRES SERVIER (SURESNES)	UK
ACERYCAL 5MG/10MG TABLETS	FR/H/0325/002	PA 568/018/002	LES LABORATOIRES SERVIER (SURESNES)	IE
ACERYCAL 5mg/10mg tablets	FR/H/325/02/DC	PL 05815/0057	LES LABORATOIRES SERVIER (SURESNES)	UK
ACERYCAL 5MG/5MG TABLETS	FR/H/0325/001	PA 568/018/001	LES LABORATOIRES SERVIER (SURESNES)	IE
ACERYCAL 5mg/5mg tablets	FR/H/325/01/DC	PL 05815/0056	LES LABORATOIRES SERVIER (SURESNES)	UK
Amlessa 4 mg/10 mg	UK/H/4348/002	58/675/11-C	KRKA, D.D., NOVO MESTO	CZ
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/01	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/02	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/03	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/04	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/05	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/06	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/07	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/08	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/09	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/10	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg comprimate	UK/H/4348/002	4083/2011/11	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/10 mg tabletes	UK/H/4348/002	11-0359	KRKA, D.D., NOVO MESTO	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/007	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/008	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/009	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/010	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/011	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/012	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/007	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/008	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/009	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/010	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/011	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tabletės	not available	LT/1/11/2466/012	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/10 mg tablety	UK/H/4348/002	58/0441/11-S	KRKA, D.D., NOVO MESTO	SK
Amlessa 4 mg/10mg tablete	UK/H/4348/002	H/11/00171/012	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10mg tablete	UK/H/4348/002	H/11/00171/013	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10mg tablete	UK/H/4348/002	H/11/00171/014	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg	UK/H/4348/001/DC	58/674/11-C	KRKA, D.D., NOVO MESTO	CZ
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/01	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/02	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/03	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/04	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/05	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/06	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/07	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/08	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/09	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/10	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/5 mg comprimate	UK/H/4348/001	4082/2011/11	KRKA, D.D., NOVO MESTO	RO
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/004	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/005	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/006	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/007	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/008	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/009	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/010	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/011	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
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/001	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/002	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tablete	UK/H/4348/001	H/11/00171/003	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/5 mg tabletes	UK/H/4348/001	11-0360	KRKA, D.D., NOVO MESTO	LV
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/001	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/002	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/003	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/004	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/005	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/006	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/001	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/002	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/003	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/004	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/005	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tabletės	not available	LT/1/11/2466/006	KRKA, D.D., NOVO MESTO	LT
Amlessa 4 mg/5 mg tablety	UK/H/4348/001	58/0440/11-S	KRKA, D.D., NOVO MESTO	SK
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/015	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
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/016	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/017	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/018	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/019	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/020	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/021	KRKA, D.D., NOVO MESTO	SI
Amlessa 4 mg/10 mg tablete	UK/H/4348/002	H/11/00171/022	KRKA, D.D., NOVO MESTO	SI
Amlessa 4mg/10mg tabletta	UK/H/4348/002	OGYI-T-22145/05	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/10mg tabletta	UK/H/4348/002	OGYI-T-22145/06	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
Amlessa 4mg/10mg tabletta	UK/H/4348/002	OGYI-T-22145/07	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/10mg tabletta	UK/H/4348/002	OGYI-T-22145/08	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/5mg tabletta	UK/H/4348/001/DC	OGYI-T-22145/01	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/5mg tabletta	UK/H/4348/001	OGYI-T-22145/02	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/5mg tabletta	UK/H/4348/001	OGYI-T-22145/03	KRKA, D.D., NOVO MESTO	HU
Amlessa 4mg/5mg tabletta	UK/H/4348/001	OGYI-T-22145/04	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/10 mg	UK/H/4348/004	58/677/11-C	KRKA, D.D., NOVO MESTO	CZ
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/01	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/02	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
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/03	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/04	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/05	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/06	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/07	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/08	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/09	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/10	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/10 mg comprimate	UK/H/4348/004	4085/2011/11	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
Amlessa 8 mg/10 mg tabletes	UK/H/4348/004	11-0361	KRKA, D.D., NOVO MESTO	LV
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/019	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/020	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/021	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/022	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/023	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/024	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/019	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/020	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/021	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/022	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/023	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletės	not available	LT/1/11/2466/024	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/10 mg tabletta	UK/H/4348/004	OGYI-T-22145/13	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/10 mg tabletta	UK/H/4348/004	OGYI-T-22145/14	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/10 mg tabletta	UK/H/4348/004	OGYI-T-22145/15	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/10 mg tabletta	UK/H/4348/004	OGYI-T-22145/16	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/10 mg tablety	UK/H/4348/004	58/0443/11-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
Amlessa 8 mg/5 mg	UK/H/4348/003	58/676/11-C	KRKA, D.D., NOVO MESTO	CZ
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/01	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/02	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/03	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/04	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/05	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/06	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/07	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/08	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
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/09	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/10	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg comprimate	UK/H/4348/003	4084/2011/11	KRKA, D.D., NOVO MESTO	RO
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/026	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/027	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/028	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/029	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/030	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/031	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/032	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/033	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/023	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/024	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tablete	UK/H/4348/003	H/11/00171/025	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/5 mg tabletes	UK/H/4348/003	11-0362	KRKA, D.D., NOVO MESTO	LV
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/013	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/014	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/015	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/016	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/017	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/018	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/013	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/014	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/015	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/016	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/017	KRKA, D.D., NOVO MESTO	LT
Amlessa 8 mg/5 mg tabletės	not available	LT/1/11/2466/018	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 8 mg/5 mg tabletta	UK/H/4348/003	OGYI-T-22145/09	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/5 mg tabletta	UK/H/4348/003	OGYI-T-22145/10	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/5 mg tabletta	UK/H/4348/003	OGYI-T-22145/11	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/5 mg tabletta	UK/H/4348/003	OGYI-T-22145/12	KRKA, D.D., NOVO MESTO	HU
Amlessa 8 mg/5 mg tablety	UK/H/4348/003	58/0442/11-S	KRKA, D.D., NOVO MESTO	SK
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/037	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/038	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/039	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/040	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/041	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/042	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/043	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/044	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/034	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/035	KRKA, D.D., NOVO MESTO	SI
Amlessa 8 mg/10 mg tablete	UK/H/4348/004	H/11/00171/036	KRKA, D.D., NOVO MESTO	SI
Amlessa, 4 mg + 10 mg, tabletki	UK/H/4348/002	18738	KRKA, D.D., NOVO MESTO	PL
Amlessa, 4 mg + 5 mg, tabletki	UK/H/4348/001	18736	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
Amlessa, 8 mg + 10 mg, tabletki	UK/H/4348/004	18739	KRKA, D.D., NOVO MESTO	PL
Amlessa, 8 mg + 5 mg, tabletki	UK/H/4348/003	18737	KRKA, D.D., NOVO MESTO	PL
Amplival 3.5 mg/2.5 mg, comprimé	IT/H/374/01/DC	34009 550 127 1 9	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 3.5 mg/2.5 mg, comprimé	IT/H/374/01/DC	34009 300 357 7 1	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 3.5 mg/2.5 mg, comprimé	IT/H/374/01/DC	34009 550 127 0 2	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 3.5 mg/2.5 mg, comprimé	IT/H/374/01/DC	34009 300 357 8 8	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 3.5 mg/2.5 mg, comprimé	IT/H/374/01/DC	34009 300 357 9 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 1 8	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 0 1	LES LABORATOIRES SERVIER (SURESNES)	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
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 550 127 2 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 2 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 550 127 4 0	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 1 8	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 0 1	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 550 127 2 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 300 358 2 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Amplival 7 mg/5 mg, comprimé	IT/H/0374/002	34009 550 127 4 0	LES LABORATOIRES SERVIER (SURESNES)	FR
Articel-Am 10 mg/10 mg tablete	NL/H/2842/004	HR-H-481082025	PLIVA HRVATSKA 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
Articel-Am 10 mg/5 mg tablete	NL/H/2842/003	HR-H-296952323	PLIVA HRVATSKA D.O.O.	HR
Articel-Am 5 mg/10 mg tablete	NL/H/2842/002	HR-H-549475085	PLIVA HRVATSKA D.O.O.	HR
Articel-Am 5 mg/5 mg tablete	NL/H/2842/001	HR-H-630048648	PLIVA HRVATSKA D.O.O.	HR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 550 127 5 7	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 3 2	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 5 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 4 9	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 550 127 6 4	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 550 127 5 7	LES LABORATOIRES SERVIER (SURESNES)	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
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 3 2	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 5 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 300 358 4 9	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 3.5 mg/2.5 mg, comprimé	IT/H/0375/001	34009 550 127 6 4	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 550 127 7 1	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 550 127 8 8	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 7 0	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 6 3	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 9 4	LES LABORATOIRES SERVIER (SURESNES)	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
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 550 127 7 1	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 550 127 8 8	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 7 0	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 6 3	LES LABORATOIRES SERVIER (SURESNES)	FR
Arvacoram 7 mg/5 mg, comprimé	IT/H/0375/002	34009 300 358 9 4	LES LABORATOIRES SERVIER (SURESNES)	FR
BEATIL 4 mg/10 mg comprimate	HU/H/0311/003	4721/2012/01	GEDEON RICHTER ROMÂNIA S.A.	RO
BEATIL 4 mg/5 mg comprimate	HU/H/0311/001	4719/2012/01	GEDEON RICHTER ROMÂNIA S.A.	RO
BEATIL 8 mg/10 mg comprimate	HU/H/0311/004	4722/2012/01	GEDEON RICHTER ROMÂNIA S.A.	RO
BEATIL 8 mg/5 mg comprimate	HU/H/0311/002	4720/2012/01	GEDEON RICHTER ROMÂNIA 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
Co-Prestarium Initio, 3,5 mg + 2,5 mg, tabletki	IT/H/374/01/DC	22472	LES LABORATOIRES SERVIER (SURESNES)	PL
Co-Prestarium Initio, 7 mg + 5 mg, tabletki	IT/H/374/02/DC	22473	LES LABORATOIRES SERVIER (SURESNES)	PL
Co-Prestarium, 10 mg + 10 mg, tabletki	FR/H/0325/004	14898	LES LABORATOIRES SERVIER	PL
Co-Prestarium, 10 mg + 5 mg, tabletki	FR/H/0325/003	14897	LES LABORATOIRES SERVIER	PL
Co-Prestarium, 5 mg + 10 mg, tabletki	FR/H/0325/002	14899	LES LABORATOIRES SERVIER	PL
Co-Prestarium, 5 mg + 5 mg, tabletki	FR/H/0325/001	14896	LES LABORATOIRES SERVIER	PL
COVERAM 10 mg/ 10 mg comprimidos	FR/H/0325/004	5562947	LES LABORATOIRES SERVIER (SURESNES)	PT
COVERAM 10 mg/ 10 mg comprimidos	FR/H/0325/004	5108907	LES LABORATOIRES SERVIER	PT
COVERAM 10 mg/ 5 mg comprimidos	FR/H/0325/003	5562939	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 10 mg/ 5 mg comprimidos	FR/H/0325/003	5108873	LES LABORATOIRES SERVIER	PT
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 835-0	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 839-6	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 834-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	572 857-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 836-7	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 841-0	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 843-3	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 832-1	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	572 856-8	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	572 855-1	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 840-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 837-3	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 833-8	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10 mg/10 mg, comprimé	FR/H/0325/004	385 842-7	LES LABORATOIRES SERVIER (SURESNES)	FR
Coveram 10 mg/10 mg, comprimés	FR/H/0325/004	BE317886	SERVIER R&D BENELUX N.V./S.A.	BE
Coveram 10 mg/10 mg, comprimés	FR/H/0325/004	0536/09010005	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 10 mg/5 mg, comprimé	FR/H/0325/003	385 829-0	LES LABORATOIRES SERVIER (SURESNES)	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
Coveram 10 mg/5 mg, comprimés	FR/H/0325/003	BE317877	SERVIER R&D BENELUX N.V./S.A.	BE
Coveram 10 mg/5 mg, comprimés	FR/H/0325/003	0536/09010008	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 10mg/10mg tablets	FR/H/0325/004	MA 066/01504	LES LABORATOIRES SERVIER (SURESNES)	MT
COVERAM 10mg/10mg Tabletten	FR/H/0325/004	BE317886	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 10mg/10mg tabletten	FR/H/0325/004	BE317886	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 10mg/10mg tabletten	FR/H/0325/004	0536/09010005	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 10mg/10mg δισκία	FR/H/0325/004	20468	LES LABORATOIRES SERVIER (SURESNES)	CY
COVERAM 10mg/10mg δισκία	FR/H/325/04/DC	3609/14.1.2009	SERVIER HELLAS PHARMACEUTICALS LTD	GR
COVERAM 10mg/10mg-Tabletten	FR/H/0325/004	0536/09010005	SERVIER R&D BENELUX 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
COVERAM 10mg/5mg tablets	FR/H/0325/003	MA 066/01503	LES LABORATOIRES SERVIER (SURESNES)	MT
COVERAM 10mg/5mg tabletten	FR/H/0325/003	BE317877	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 10mg/5mg Tabletten	FR/H/0325/003	BE317877	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 10mg/5mg tabletten	FR/H/0325/003	0536/09010008	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 10mg/5mg δισκία	FR/H/0325/003	20467	LES LABORATOIRES SERVIER (SURESNES)	CY
COVERAM 10mg/5mg δισκία	FR/H/325/03/DC	3608/14.1.2009	SERVIER HELLAS PHARMACEUTICALS LTD	GR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 824-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 830-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 822-6	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 827-8	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 823-2	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	572 854-5	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 826-1	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 828-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	572 853-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 831-5	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 820-3	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	572 852-2	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 10MG/5MG, comprimé	FR/H/0325/003	385 825-5	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 10mg/5mg-Tabletten	FR/H/0325/003	0536/09010008	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5 mg/ 10 mg comprimidos	FR/H/0325/002	5562954	LES LABORATOIRES SERVIER (SURESNES)	PT
COVERAM 5 mg/ 10 mg comprimidos	FR/H/0325/002	5108865	LES LABORATOIRES SERVIER	PT
COVERAM 5 mg/ 5 mg comprimidos	FR/H/0325/001	5562921	LES LABORATOIRES SERVIER (SURESNES)	PT
COVERAM 5 mg/ 5 mg comprimidos	FR/H/0325/001	5108857	LES LABORATOIRES SERVIER	PT
COVERAM 5 mg/ 5 mg comprimidos	FR/H/0325/001	5108840	LES LABORATOIRES SERVIER	PT
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 817-2	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 811-4	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 818-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 813-7	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 819-5	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 807-7	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 810-8	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 812-0	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 808-3	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	572 849-1	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 814-3	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	572 851-6	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	385 816-6	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/10 mg, comprimé	FR/H/0325/002	572 848-5	LES LABORATOIRES SERVIER (SURESNES)	FR
Coveram 5 mg/10 mg, comprimés	FR/H/0325/002	BE317861	SERVIER R&D BENELUX N.V./S.A.	BE
Coveram 5 mg/10 mg, comprimés	FR/H/0325/002	0536/09010006	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5 mg/5 mg tabletit	FR/H/0325/001	23480	LES LABORATOIRES SERVIER	FI
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 799-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 801-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 802-5	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 806-0	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 804-8	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 796-5	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 803-1	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 797-1	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	572 845-6	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 805-4	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	572 846-2	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 798-8	LES LABORATOIRES SERVIER (SURESNES)	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
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	572 847-9	LES LABORATOIRES SERVIER (SURESNES)	FR
COVERAM 5 mg/5 mg, comprimé	FR/H/0325/001	385 800-2	LES LABORATOIRES SERVIER (SURESNES)	FR
Coveram 5 mg/5 mg, comprimés	FR/H/0325/001	BE317852	SERVIER R&D BENELUX N.V./S.A.	BE
Coveram 5 mg/5 mg, comprimés	FR/H/0325/001	0536/09010007	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5mg/10mg tablets	FR/H/0325/002	MA 066/01502	LES LABORATOIRES SERVIER (SURESNES)	MT
COVERAM 5mg/10mg Tabletten	FR/H/0325/002	BE317861	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 5mg/10mg tabletten	FR/H/0325/002	BE317861	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 5mg/10mg tabletten	FR/H/0325/002	0536/09010006	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5mg/10mg δισκία	FR/H/0325/002	20466	LES LABORATOIRES SERVIER (SURESNES)	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COVERAM 5mg/10mg δισκία	FR/H/325/02/DC	3606/14.1.2009	SERVIER HELLAS PHARMACEUTICALS LTD	GR
COVERAM 5mg/10mg-Tabletten	FR/H/0325/002	0536/09010006	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5mg/5mg tablets	FR/H/0325/001	MA 066/01501	LES LABORATOIRES SERVIER (SURESNES)	MT
COVERAM 5mg/5mg Tabletten	FR/H/0325/001	BE317852	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 5mg/5mg tabletten	FR/H/0325/001	BE317852	SERVIER R&D BENELUX N.V./S.A.	BE
COVERAM 5mg/5mg tabletten	FR/H/0325/001	0536/09010007	SERVIER R&D BENELUX N.V./S.A.	LU
COVERAM 5mg/5mg δισκία	FR/H/0325/001	20465	LES LABORATOIRES SERVIER (SURESNES)	CY
COVERAM 5mg/5mg δισκία	FR/H/325/01/DC	3605/14.1.2009	SERVIER HELLAS PHARMACEUTICALS LTD	GR
COVERAM 5mg/5mg-Tabletten	FR/H/0325/001	0536/09010007	SERVIER R&D BENELUX 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
COVERAM arg 10 mg/10 mg tabletten	FR/H/0325/004	RVG 100142	LES LABORATOIRES SERVIER (SURESNES)	NL
COVERAM arg 10 mg/5 mg tabletten	FR/H/0325/003	RVG 100141	LES LABORATOIRES SERVIER (SURESNES)	NL
COVERAM arg 5 mg/10 mg tabletten	FR/H/0325/002	RVG 100140	LES LABORATOIRES SERVIER (SURESNES)	NL
COVERAM arg 5 mg/5 mg tabletten	FR/H/0325/001	RVG 100139	LES LABORATOIRES SERVIER (SURESNES)	NL
Coveram, 10 mg/10 mg tabletid	FR/H/0325/004	584208	LES LABORATOIRES SERVIER (SURESNES)	EE
Coveram, 10 mg/5 mg tabletid	FR/H/0325/003	584408	LES LABORATOIRES SERVIER (SURESNES)	EE
Coveram, 5 mg/10 mg tabletid	FR/H/0325/002	584308	LES LABORATOIRES SERVIER (SURESNES)	EE
Coveram, 5 mg/5 mg tabletid	FR/H/0325/001	584108	LES LABORATOIRES SERVIER (SURESNES)	EE
Covercard 10 mg/10 mg tabletta	FR/H/0326/004/DC	OGYI-T-20572/04	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
Covercard 10 mg/5 mg tabletta	FR/H/0326/003/DC	OGYI-T-20572/03	EGIS PHARMACEUTICALS PLC	HU
Covercard 3,5 mg/2,5 mg tabletta	IT/H/0375/001	OGYI-T-20572/05	EGIS PHARMACEUTICALS PLC	HU
Covercard 3,5 mg/2,5 mg tabletta	IT/H/0375/001/DC	OGYI-T-20572/05	EGIS PHARMACEUTICALS PLC	HU
Covercard 3,5 mg/2,5 mg tabletta	IT/H/0375/001/DC	OGYI-T-20572/05	EGIS PHARMACEUTICALS PLC	HU
Covercard 5 mg/10 mg tabletta	FR/H/0326/002/DC	OGYI-T-20572/02	EGIS PHARMACEUTICALS PLC	HU
Covercard 5 mg/5 mg tabletta	FR/H/0326/001/DC	OGYI-T-20572/01	EGIS PHARMACEUTICALS PLC	HU
Covercard 7 mg/5 mg tabletta	IT/H/0375/002/DC	OGYI-T-20572/06	EGIS PHARMACEUTICALS PLC	HU
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477105/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477030/M	LES LABORATOIRES SERVIER	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
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477028/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477016/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477079/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477081/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477129/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477055/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477117/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477042/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477143/M	LES LABORATOIRES SERVIER	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
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477131/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477093/M	LES LABORATOIRES SERVIER	IT
COVERLAM 5 mg/5 mg compresse	FR/H/0325/001	038477067/M	LES LABORATOIRES SERVIER	IT
Coversical	FR/H/0325/001	41221	LES LABORATOIRES SERVIER (SURESNES)	DK
Coversical	FR/H/0325/002	41220	LES LABORATOIRES SERVIER (SURESNES)	DK
Coversical	FR/H/0325/004	41218	LES LABORATOIRES SERVIER (SURESNES)	DK
Coversical	FR/H/0325/003	41219	LES LABORATOIRES SERVIER (SURESNES)	DK
COVERSICAL 10 mg/10 mg töflur	FR/H/0325/004	IS/1/07/045/04	LES LABORATOIRES SERVIER (SURESNES)	IS
COVERSICAL 10 mg/5 mg töflur	FR/H/0325/003	IS/1/07/045/03	LES LABORATOIRES SERVIER (SURESNES)	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COVERSICAL 5 mg/10 mg töflur	FR/H/0325/002	IS/1/07/045/02	LES LABORATOIRES SERVIER (SURESNES)	IS
COVERSICAL 5 mg/5 mg töflur	FR/H/0325/001	IS/1/07/045/01	LES LABORATOIRES SERVIER (SURESNES)	IS
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/012	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/013	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/014	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/015	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/016	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/017	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/018	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/019	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/020	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/021	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	UK/H/4348/002	LT/1/11/2697/022	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/10 mg tabletes	not available	11-0405	KRKA, D.D., NOVO MESTO	LV
Dalnessa 4 mg/10 mg tabletes	not available	11-0405	KRKA, D.D., NOVO MESTO	LV
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/09	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/10	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/11	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/12	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/13	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/14	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/15	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/16	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/09	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/10	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/11	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/12	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/13	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/14	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/15	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/10 mg tabletta	not available	OGYI-T-21727/16	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletes	not available	11-0406	KRKA, D.D., NOVO MESTO	LV
Dalnessa 4 mg/5 mg tabletes	not available	11-0406	KRKA, D.D., NOVO MESTO	LV
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/001	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/002	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/003	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/004	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/005	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/006	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/007	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/008	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/009	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/010	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletės	UK/H/4348/001	LT/1/11/2697/011	KRKA, D.D., NOVO MESTO	LT
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/01	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
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/02	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/03	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/04	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/05	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/06	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/07	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/08	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/01	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/02	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/03	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/04	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/05	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/06	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/07	KRKA, D.D., NOVO MESTO	HU
Dalnessa 4 mg/5 mg tabletta	not available	OGYI-T-21727/08	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletes	not available	11-0407	KRKA, D.D., NOVO MESTO	LV
Dalnessa 8 mg/10 mg tabletes	not available	11-0407	KRKA, D.D., NOVO MESTO	LV
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/034	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/035	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/036	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/037	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/038	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/039	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/040	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/041	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/042	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/043	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 8 mg/10 mg tabletės	UK/H/4348/004	LT/1/11/2697/044	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/25	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/26	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/27	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/28	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/29	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/30	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/31	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/32	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
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/25	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/26	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/27	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/28	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/29	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/30	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/31	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/10 mg tabletta	not available	OGYI-T-21727/32	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletes	not available	11-0408	KRKA, D.D., NOVO MESTO	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 8 mg/5 mg tabletes	not available	11-0408	KRKA, D.D., NOVO MESTO	LV
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/023	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/024	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/025	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/026	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/027	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/028	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/029	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/030	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/031	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/032	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletės	UK/H/4348/003	LT/1/11/2697/033	KRKA, D.D., NOVO MESTO	LT
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/17	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/18	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/19	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/20	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/21	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/22	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
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/23	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/24	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/17	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/18	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/19	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/20	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/21	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/22	KRKA, D.D., NOVO MESTO	HU
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/23	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
Dalnessa 8 mg/5 mg tabletta	not available	OGYI-T-21727/24	KRKA, D.D., NOVO MESTO	HU
Dalnessa, 4 mg / 10 mg tabletid	not available	802312	KRKA, D.D., NOVO MESTO	EE
Dalnessa, 4 mg / 5 mg tabletid	not available	802212	KRKA, D.D., NOVO MESTO	EE
Dalnessa, 8 mg / 10 mg tabletid	not available	802012	KRKA, D.D., NOVO MESTO	EE
Dalnessa, 8 mg / 5 mg tabletid	not available	802112	KRKA, D.D., NOVO MESTO	EE
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094118	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094120	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094132	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094144	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
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094157	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094169	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094171	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094183	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094195	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094207	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg compresse	UK/H/4348/002	040094423	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/10 mg tablete	not available	UP/I-530-09/10-01/481	KRKA-FARMA D.O.O.	HR
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094017	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
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094029	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094031	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094043	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094056	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094068	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094070	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094082	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094094	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094106	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
Dalneva 4 mg/5 mg compresse	UK/H/4348/001	040094411	KRKA, D.D., NOVO MESTO	IT
Dalneva 4 mg/5 mg tablete	not available	UP/I-530-09/10-01/480	KRKA-FARMA D.O.O.	HR
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094310	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094322	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094334	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094346	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094359	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094361	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094373	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
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094385	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094397	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094409	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg compresse	UK/H/4348/004	040094447	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/10 mg tablete	not available	UP/I-530-09/10-01/483	KRKA-FARMA D.O.O.	HR
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094219	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094221	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094233	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	AIC N. 040094245	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
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094258	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094260	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094272	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094284	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094296	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094308	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg compresse	UK/H/4348/003	040094435	KRKA, D.D., NOVO MESTO	IT
Dalneva 8 mg/5 mg tablete	not available	UP/I-530-09/10-01/482	KRKA-FARMA D.O.O.	HR
Levenor Kombi 4 mg és 10 mg tabletta	not available	OGYI-T-21991/02	ARAMIS PHARMA KFT	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
Levenor Kombi 4 mg és 5 mg tabletta	not available	OGYI-T-21991/01	ARAMIS PHARMA KFT	HU
Levenor Kombi 8 mg és 10 mg tabletta	not available	OGYI-T-21991/04	ARAMIS PHARMA KFT	HU
Levenor Kombi 8 mg és 5 mg tabletta	not available	OGYI-T-21991/03	ARAMIS PHARMA KFT	HU
Mixanval 10 mg/10 mg, comprimés	FR/H/0326/004	BE317843	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 10 mg/10 mg, comprimés	FR/H/0325/004	0605/10080024	EUTHERAPIE BENELUX S.A.	LU
Mixanval 10 mg/5 mg, comprimés	FR/H/0326/003	BE317834	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 10 mg/5 mg, comprimés	FR/H/0325/003	0605/10080025	EUTHERAPIE BENELUX S.A.	LU
MIXANVAL 10mg/10mg tabletten	FR/H/0326/004	BE317843	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 10mg/10mg tabletten	FR/H/0325/004	0605/10080024	EUTHERAPIE BENELUX 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
Mixanval 10mg/10mg-Tabletten	FR/H/0326/004	BE317843	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 10mg/5mg tabletten	FR/H/326/003	BE317834	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 10mg/5mg tabletten	FR/H/0325/003	0605/10080025	EUTHERAPIE BENELUX S.A.	LU
Mixanval 10mg/5mg-Tabletten	FR/H/0326/003	BE317834	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5 mg/10 mg, comprimés	FR/H/0325/002	0605/10080023	EUTHERAPIE BENELUX S.A.	LU
Mixanval 5 mg/5 mg, comprimés	FR/H/0326/001	BE317816	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5 mg/5 mg, comprimés	FR/H/0325/001	0605/10080026	EUTHERAPIE BENELUX S.A.	LU
MIXANVAL 5MG/10MG comprimés	FR/H/326/02/DC	BE317825	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5mg/10mg tabletten	FR/H/326/002	BE317825	EUTHERAPIE BENELUX 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
MIXANVAL 5mg/10mg tabletten	FR/H/0325/002	0605/10080023	EUTHERAPIE BENELUX S.A.	LU
Mixanval 5mg/10mg-Tabletten	FR/H/0326/002	BE317825	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5mg/5mg tabletten	FR/H/0326/001	BE317816	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5mg/5mg tabletten	FR/H/0325/001	0605/10080026	EUTHERAPIE BENELUX S.A.	LU
Mixanval 5mg/5mg-Tabletten	FR/H/0326/001	BE317816	EUTHERAPIE BENELUX S.A.	BE
MIXANVAL 5mg/5mg-Tabletten	FR/H/326/03/DC	0605/10080025	EUTHERAPIE BENELUX S.A.	LU
MIXANVAL 5mg/5mg-Tabletten	FR/H/326/04/DC	0605/10080024	EUTHERAPIE BENELUX S.A.	LU
MIXANVAL 5mg/5mg-Tabletten	FR/H/0326/001	0605/10080023	EUTHERAPIE BENELUX S.A.	LU
MIXANVAL 5mg/5mg-Tabletten	FR/H/326/01/DC	0605/10080026	EUTHERAPIE BENELUX 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
NORPREXANIL 10 mg/10 mg tablete	not available	HR-H-431897129	SERVIER PHARMA D.O.O - CROATIA	HR
NORPREXANIL 10 mg/5 mg tablete	not available	HR-H-713394923	SERVIER PHARMA D.O.O - CROATIA	HR
NORPREXANIL 5 mg/10 mg tablete	not available	HR-H-481735885	SERVIER PHARMA D.O.O - CROATIA	HR
NORPREXANIL 5 mg/5 mg tablete	not available	HR-H-197631019	SERVIER PHARMA D.O.O - CROATIA	HR
Peramlonorm 4 mg/10 mg tabletta	UK/H/4621/002	OGYI-T-22129/06	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/10 mg tabletta	UK/H/4621/002	OGYI-T-22129/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/10 mg tabletta	UK/H/4621/002	OGYI-T-22129/07	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/10 mg tabletta	UK/H/4621/002	OGYI-T-22129/08	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/5 mg tabletta	UK/H/4621/001	OGYI-T-22129/01	TEVA GYÓGYSZERGYÁR ZRT	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
Peramlonorm 4 mg/5 mg tabletta	UK/H/4621/001	OGYI-T-22129/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/5 mg tabletta	UK/H/4621/001	OGYI-T-22129/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 4 mg/5 mg tabletta	UK/H/4621/001	OGYI-T-22129/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/10 mg tabletta	UK/H/4621/004	OGYI-T-22129/13	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/10 mg tabletta	UK/H/4621/004	OGYI-T-22129/15	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/10 mg tabletta	UK/H/4621/004	OGYI-T-22129/16	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/10 mg tabletta	UK/H/4621/004	OGYI-T-22129/14	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/5 mg tabletta	UK/H/4621/003	OGYI-T-22129/10	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/5 mg tabletta	UK/H/4621/003	OGYI-T-22129/09	TEVA GYÓGYSZERGYÁR ZRT	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
Peramlonorm 8 mg/5 mg tabletta	UK/H/4621/003	OGYI-T-22129/11	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramlonorm 8 mg/5 mg tabletta	UK/H/4621/003	OGYI-T-22129/12	TEVA GYÓGYSZERGYÁR ZRT	HU
Peramteva ® 10 mg/10 mg comprimés	NL/H/2842/004	BE451564	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva ® 10 mg/5 mg comprimés	NL/H/2842/003	BE451555	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva ® 5 mg/10 mg comprimés	NL/H/2842/002	BE451546	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva ® 5 mg/5 mg comprimés	NL/H/2842/001	BE451537	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 10 mg/10 mg Tabletten	NL/H/2842/04	BE451564	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 10 mg/10 mg Tabletten	NL/H/2842/004	BE451564	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 10 mg/5 mg Tabletten	NL/H/2842/03	BE451555	TEVA PHARMA BELGIUM 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
Peramteva® 10 mg/5 mg Tabletten	NL/H/2842/03	BE451555	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 5 mg/10 mg Tabletten	NL/H/2842/02	BE451546	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 5 mg/10 mg Tabletten	NL/H/2842/002	BE451546	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 5 mg/5 mg Tabletten	NL/H/2842/01	BE451537	TEVA PHARMA BELGIUM N.V./S.A	BE
Peramteva® 5 mg/5 mg Tabletten	NL/H/2842/001	BE451537	TEVA PHARMA BELGIUM N.V./S.A	BE
Perindopril + Amlodipina Krka 4 mg + 10 mg comprimidos	UK/H/4348/002	5395744	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 4 mg + 10 mg comprimidos	UK/H/4348/002	5395751	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 4 mg + 10 mg comprimidos	UK/H/4348/002	5395769	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 4 mg + 5 mg comprimidos	UK/H/4348/001	5395710	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril + Amlodipina Krka 4 mg + 5 mg comprimidos	UK/H/4348/001	5395728	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 4 mg + 5 mg comprimidos	UK/H/4348/001	5395736	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 10 mg comprimidos	UK/H/4348/004	5395827	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 10 mg comprimidos	UK/H/4348/004	5395835	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 10 mg comprimidos	UK/H/4348/004	5395843	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 5 mg comprimidos	UK/H/4348/003	5395777	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 5 mg comprimidos	UK/H/4348/003	5395801	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Krka 8 mg + 5 mg comprimidos	UK/H/4348/003	5395819	KRKA FARMACÊUTICA, UNIPESSOAL LDA.	PT
Perindopril + Amlodipina Ratiopharm 10 mg + 10 mg Comprimidos	NL/H/2842/004	5615711	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril + Amlodipina Ratiopharm 10 mg + 5 mg Comprimidos	NL/H/2842/03	5615703	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Perindopril + Amlodipina Ratiopharm 5 mg + 10 mg Comprimidos	NL/H/2842/02	5618202	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Perindopril + Amlodipina Ratiopharm 5 mg + 5 mg Comprimidos	NL/H/2842/01	5615653	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Perindopril + Amlodipina Ratiopharm 5 mg + 5 mg Comprimidos	NL/H/2842/01	5622675	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Perindopril + Amlodipina Sandoz 4 mg + 10 mg comprimido	not available	5419254	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 4 mg + 10 mg comprimido	not available	5419262	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 4 mg + 10 mg comprimido	not available	5419270	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 4 mg + 5 mg comprimido	not available	5419221	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 4 mg + 5 mg comprimido	not available	5419239	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril + Amlodipina Sandoz 4 mg + 5 mg comprimido	not available	5419247	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 10 mg comprimido	not available	5419338	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 10 mg comprimido	not available	5419346	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 10 mg comprimido	not available	5419353	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 5 mg comprimido	not available	5419304	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 5 mg comprimido	not available	5419312	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
Perindopril + Amlodipina Sandoz 8 mg + 5 mg comprimido	not available	5419320	NANDOZA - PRODUTOS FARMACÊUTICOS, LDA.	PT
PERINDOPRIL + AMLODIPINA ZENTIVA 4 MG + 10 MG COMPRIMIDOS	CZ/H/0474/002	5657358	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
PERINDOPRIL + AMLODIPINA ZENTIVA 4 MG + 5 MG	CZ/H/0474/001	5657333	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
COMPRIMIDOS				
PERINDOPRIL + AMLODIPINA ZENTIVA 4 MG + 5 MG COMPRIMIDOS	CZ/H/0474/001	5657341	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
PERINDOPRIL + AMLODIPINA ZENTIVA 8 MG + 10 MG COMPRIMIDOS	CZ/H/0474/004	5657408	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
PERINDOPRIL + AMLODIPINA ZENTIVA 8 MG + 5 MG COMPRIMIDOS	CZ/H/0474/003	5657366	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Perindopril arginine 10 mg / Amlodipine 10 mg Servier tablety	FR/H/0326/004	58/210/08-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Perindopril arginine 10 mg / Amlodipine 5 mg Servier tablety	FR/H/0326/003	58/209/08-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Perindopril arginine 5 mg / Amlodipine 10 mg Servier tablety	FR/H/0326/002	58/208/08-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Perindopril arginine 5 mg / Amlodipine 5 mg Servier tablety	FR/H/0326/001	58/207/08-C	LES LABORATOIRES SERVIER (SURESNES)	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
Perindopril arginine/ Amlodipine Servier 10 mg/10 mg tabletten	FR/H/326/04/DC	RVG 100138	LES LABORATOIRES SERVIER (SURESNES)	NL
Perindopril arginine/ Amlodipine Servier 10 mg/5 mg tabletten	FR/H/326/03/DC	RVG 100137	LES LABORATOIRES SERVIER (SURESNES)	NL
Perindopril arginine/ Amlodipine Servier 10mg/10mg tabletes	FR/H/326/04/DC	08-0118	LES LABORATOIRES SERVIER (SURESNES)	LV
Perindopril arginine/ Amlodipine Servier 10mg/5mg tabletes	FR/H/326/03/DC	08-0117	LES LABORATOIRES SERVIER (SURESNES)	LV
Perindopril arginine/ Amlodipine Servier 5 mg/10 mg tabletten	FR/H/326/02/DC	RVG 100136	LES LABORATOIRES SERVIER (SURESNES)	NL
Perindopril arginine/ Amlodipine Servier 5 mg/5 mg tabletten	FR/H/326/01/DC	RVG 100016	LES LABORATOIRES SERVIER (SURESNES)	NL
Perindopril arginine/ Amlodipine Servier 5mg/10mg tabletes	FR/H/326/02/DC	08-0116	LES LABORATOIRES SERVIER (SURESNES)	LV
Perindopril arginine/ Amlodipine Servier 5mg/5mg tabletes	FR/H/326/01/DC	08-0115	LES LABORATOIRES SERVIER (SURESNES)	LV
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594447	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594966	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594508	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594096	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594218	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400957288419	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938593846	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400957288587	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938593907	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594676	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594157	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594386	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400938594737	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/10 mg, comprimé	FR/H/0327/004	3400957288358	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593327	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400957288297	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593556	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938592665	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593037	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593785	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400957288068	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593495	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400957288129	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938592894	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593666	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938592726	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938592955	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 10 mg/5 mg, comprimé	FR/H/0327/003	3400938593617	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938592085	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938592436	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591484	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400957287757	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591316	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938592146	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591545	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938592207	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591774	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400957287986	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938592375	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400957287818	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591835	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/10 mg, comprimé	FR/H/0327/002	3400938591606	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590883	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590425	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400957287467	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938591255	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590135	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400957287528	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938591194	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590654	BIOGARAN	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
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590364	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400957287696	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938591026	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590715	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590944	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE BIOGARAN 5 mg/5 mg, comprimé	FR/H/0327/001	3400938590593	BIOGARAN	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10mg/10mg tablets	FR/H/0326/004	PA 568/19/04	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10mg/5mg tablets	FR/H/0326/003	PA 568/19/03	LES LABORATOIRES SERVIER (SURESNES)	IE
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5mg/10mg tablets	FR/H/0326/002	PA 568/19/02	LES LABORATOIRES SERVIER (SURESNES)	IE
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5mg/5mg tablets	FR/H/0326/001	PA 568/19/01	LES LABORATOIRES SERVIER (SURESNES)	IE
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 890-1	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 895-3	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 894-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	572 872-3	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 893-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 899-9	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 892-4	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 889-3	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 898-2	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 900-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	572 870-0	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 897-6	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	385 891-8	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/10 mg, comprimé	FR/H/0326/004	572 871-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 874-6	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 876-9	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 877-5	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 872-3	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 878-1	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 875-2	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 871-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 879-8	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	572 866-3	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 870-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	572 865-7	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 880-6	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	385 869-2	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 10 mg/5 mg, comprimé	FR/H/0326/003	572 864-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	572 862-8	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 859-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	572 861-1	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 857-4	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 864-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 865-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 863-4	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 862-8	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 858-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 866-3	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 868-6	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 860-5	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	572 863-4	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/10 mg, comprimé	FR/H/0326/002	385 861-1	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 856-8	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	572 859-7	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 845-6	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 855-1	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 846-2	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 848-5	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	572 858-0	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 853-9	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 851-6	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 854-5	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 847-9	LES LABORATOIRES SERVIER (SURESNES)	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
PERINDOPRIL ARGININE/AMLODIPINE SERVIER 5 mg/5 mg, comprimé	FR/H/0326/001	385 852-2	LES LABORATOIRES SERVIER (SURESNES)	FR
PERINDOPRIL e AMLODIPINA DOC Generici 4 mg/10 mg compresse	IT/H/0526/002	044161026	DOC GENERICI S.R.L.	IT
PERINDOPRIL e AMLODIPINA DOC Generici 4 mg/5 mg compresse	IT/H/0526/001	044161014	DOC GENERICI S.R.L.	IT
PERINDOPRIL e AMLODIPINA DOC Generici 8 mg/10 mg compresse	IT/H/0526/004	044161040	DOC GENERICI S.R.L.	IT
PERINDOPRIL e AMLODIPINA DOC Generici 8 mg/5 mg compresse	IT/H/0526/003	044161038	DOC GENERICI S.R.L.	IT
PERINDOPRIL e AMLODIPINA EG 4 mg/10 mg compresse	SE/H/1500/002	043970072	EG S.P.A.	IT
PERINDOPRIL e AMLODIPINA EG 4 mg/10 mg compresse	SE/H/1500/002	043970058	EG S.P.A.	IT
PERINDOPRIL e AMLODIPINA EG 4 mg/10	SE/H/1500/002	043970060	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
mg compresse				
PERINDOPRIL e AMLODIPINA EG 4 mg/5 mg compresse	SE/H/1500/001	043970033	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 4 mg/5 mg compresse	SE/H/1500/001	043970019	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 4 mg/5 mg compresse	SE/H/1500/001	043970021	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 8 mg/10 mg compresse	SE/H/1500/004	043970110	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 8 mg/10 mg compresse	SE/H/1500/004	043970122	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 8 mg/10 mg compresse	SE/H/1500/004	043970134	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 8 mg/5 mg compresse	SE/H/1500/003	043970096	EG SPA	IT
PERINDOPRIL e AMLODIPINA EG 8 mg/5 mg compresse	SE/H/1500/003	043970084	EG SPA	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/004	042569196	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/004	042569172	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/004	042569184	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/004	042569208	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/004	042569160	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/10 mg compresse	NL/H/2842/04	042569259	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569119	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569121	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569133	TEVA 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
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569145	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569158	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 10 mg/5 mg compresse	NL/H/2842/003	042569246	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569107	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569095	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569083	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569069	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569071	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/10 mg compresse	NL/H/2842/002	042569234	TEVA 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
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569018	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569044	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569057	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569020	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569210	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569032	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA TEVA 5 mg/5 mg compresse	NL/H/2842/001	042569222	TEVA ITALIA S.R.L.	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/10 MG COMPRESSE	CZ/H/0474/002	043424047	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/10 MG COMPRESSE	CZ/H/0474/002	043424050	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
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/5 MG COMPRESSE	CZ/H/0474/001	043424035	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/5 MG COMPRESSE	CZ/H/0474/001	043424011	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/5 MG COMPRESSE	CZ/H/0474/001	043424100	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 4 MG/5 MG COMPRESSE	CZ/H/0474/001	043424023	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 8 MG/10 MG COMPRESSE	CZ/H/0474/004	043424098	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 8 MG/10 MG COMPRESSE	CZ/H/0474/004	043424086	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 8 MG/5 MG COMPRESSE	CZ/H/0474/003	043424074	ZENTIVA ITALIA SRL	IT
PERINDOPRIL E AMLODIPINA ZENTIVA 8 MG/5 MG COMPRESSE	CZ/H/0474/003	043424062	ZENTIVA ITALIA SRL	IT
Perindopril tert-butylamine/Amlodipine CF 4/10 mg, tabletten	SE/H/1500/002	RVG 116720	CENTRAFARM B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril tert-butylamine/Amlodipine CF 4/5 mg, tabletten	SE/H/1500/001	RVG 116719	CENTRAFARM B.V.	NL
Perindopril tert-butylamine/Amlodipine CF 8/10 mg, tabletten	SE/H/1500/004	RVG 116722	CENTRAFARM B.V.	NL
Perindopril tert-butylamine/Amlodipine CF 8/5 mg, tabletten	SE/H/1500/003	RVG 116721	CENTRAFARM B.V.	NL
Perindopril tert-butylamine/Amlodipine Stada 4 mg/10 mg tablets	SE/H/1500/002	52343	STADA ARZNEIMITTEL AG	SE
Perindopril tert-butylamine/Amlodipine Stada 4 mg/5 mg tablets	SE/H/1500/001	52342	STADA ARZNEIMITTEL AG	SE
Perindopril tert-butylamine/Amlodipine Stada 8 mg/10 mg tablets	SE/H/1500/004	52345	STADA ARZNEIMITTEL AG	SE
Perindopril tert-butylamine/Amlodipine Stada 8 mg/5 mg tablets	SE/H/1500/003	52344	STADA ARZNEIMITTEL AG	SE
Perindopril tosilaat/Amlodipine Teva 10 mg/10 mg, tabletten	NL/H/2842/004	RVG 113219	TEVA NEDERLAND B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril tosilaat/Amlodipine Teva 10 mg/5 mg, tabletten	NL/H/2842/003	RVG 113218	TEVA NEDERLAND B.V.	NL
Perindopril tosilaat/Amlodipine Teva 5 mg/10 mg, tabletten	NL/H/2842/002	RVG 113217	TEVA NEDERLAND B.V.	NL
Perindopril tosilaat/Amlodipine Teva 5 mg/5 mg, tabletten	NL/H/2842/001	RVG 113216	TEVA NEDERLAND B.V.	NL
Perindopril tosilat/Amlodipină Teva 10 mg/5 mg comprimate	NL/H/2842/003	6799/2014/02	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/10mg comprimate	NL/H/2842/04	6800/2014/01	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/10mg comprimate	NL/H/2842/04	6800/2014/02	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/10mg comprimate	NL/H/2842/04	6800/2014/05	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/10mg comprimate	NL/H/2842/04	6800/2014/03	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/10mg comprimate	NL/H/2842/04	6800/2014/04	TEVA PHARMACEUTICALS 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
Perindopril tosilat/Amlodipină Teva 10mg/5mg comprimate	NL/H/2842/03	6799/2014/01	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/5mg comprimate	NL/H/2842/03	6799/2014/03	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/5mg comprimate	NL/H/2842/03	6799/2014/04	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 10mg/5mg comprimate	NL/H/2842/03	6799/2014/05	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/10 mg comprimate	NL-H-2842-01-04	6798/2014/01	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/10 mg comprimate	NL-H-2842-01-04	6798/2014/05	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/10 mg comprimate	NL-H-2842-01-04	6798/2014/04	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/10 mg comprimate	NL-H-2842-01-04	6798/2014/03	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/10 mg comprimate	NL-H-2842-01-04	6798/2014/02	TEVA PHARMACEUTICALS 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
Perindopril tosilat/Amlodipină Teva 5 mg/5 mg comprimate	NL-H-2842-01-04	6797/2014/01	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/5 mg comprimate	NL-H-2842-01-04	6797/2014/03	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/5 mg comprimate	NL-H-2842-01-04	6797/2014/02	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/5 mg comprimate	NL-H-2842-01-04	6797/2014/04	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilat/Amlodipină Teva 5 mg/5 mg comprimate	NL-H-2842-01-04	6797/2014/05	TEVA PHARMACEUTICALS S.R.L	RO
Perindopril tosilate/ Amlodipine Teva 10 mg/10 mg Tablets	NL/H/2842/004	PA 749/197/004	TEVA PHARMA B.V.	IE
Perindopril tosilate/ Amlodipine Teva 10 mg/5 mg Tablets	NL/H/2842/003	PA 749/197/003	TEVA PHARMA B.V.	IE
Perindopril tosilate/ Amlodipine Teva 5 mg/10 mg Tablets	NL/H/2842/002	PA 749/197/002	TEVA PHARMA B.V.	IE
Perindopril tosilate/ Amlodipine Teva 5 mg/5 mg Tablets	NL/H/2842/001	PA0749/197/001	TEVA PHARMA B.V.	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
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/015	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/016	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/017	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/018	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/019	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/020	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/021	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/022	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/012	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/013	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/10 mg tablete	UK/H/4621/002	H/12/01244/014	KRKA, D.D., NOVO MESTO	SI
Perindopril/Amlodipin Krka 4 mg/10 mg tablety	UK/H/4621/002	58/0078/12-S	KRKA, D.D., NOVO MESTO	SK
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/004	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/005	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/006	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/007	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/008	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/009	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
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/010	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/011	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/001	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/002	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 4 mg/5 mg tablete	UK/H/4621/001	H/12/01244/003	KRKA, D.D., NOVO MESTO	SI
Perindopril/Amlodipin Krka 4 mg/5 mg tablety	UK/H/4621/001	58/0077/12-S	KRKA, D.D., NOVO MESTO	SK
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/037	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/038	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/039	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
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/040	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/041	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/042	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/043	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004/DC	H/12/01244/044	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004	H/12/01244/034	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004	H/12/01244/035	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/10 mg tablete	UK/H/4621/004	H/12/01244/036	KRKA, D.D., NOVO MESTO	SI
Perindopril/Amlodipin Krka 8 mg/10 mg tablety	UK/H/4621/004	58/0080/12-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
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/026	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/027	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/028	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/029	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/030	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/031	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/032	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/033	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/023	KRKA, D.D., NOVO MESTO	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/024	KRKA, D.D., NOVO MESTO	SI
Perindopril/amlodipin Krka 8 mg/5 mg tablete	UK/H/4621/003	H/12/01244/025	KRKA, D.D., NOVO MESTO	SI
Perindopril/Amlodipin Krka 8 mg/5 mg tablety	UK/H/4621/003	58/0079/12-S	KRKA, D.D., NOVO MESTO	SK
Perindopril/Amlodipin ratiopharm 10 mg/10 mg tabletit	NL/H/2842/004	31292	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 10 mg/10 mg tabletit	NL-H-2842-01-04	31292	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 10 mg/5 mg tabletit	NL/H/2842/003	31291	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 10 mg/5 mg tabletit	NL-H-2842-01-04	31291	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 5 mg/10 mg tabletit	NL/H/2842/002	31290	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 5 mg/10 mg tabletit	NL/H/2842/002	31290	RATIOPHARM GMBH	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipin ratiopharm 5 mg/5 mg tabletit	NL/H/2842/001	31289	RATIOPHARM GMBH	FI
Perindopril/Amlodipin ratiopharm 5 mg/5 mg tabletit	NL-H-2842-01-04	31289	RATIOPHARM GMBH	FI
Perindopril/amlodipin STADA 4 mg/10 mg tablete	SE/H/1500/002	H/16/02206/004	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 4 mg/10 mg tablete	SE/H/1500/002	H/16/02206/006	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 4 mg/10 mg tablete	SE/H/1500/002	H/16/02206/005	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 4 mg/5 mg tablete	SE/H/1500/001	H/16/02206/001	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 4 mg/5 mg tablete	SE/H/1500/001	H/16/02206/002	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 4 mg/5 mg tablete	SE/H/1500/001	H/16/02206/003	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 8 mg/10 mg tablete	SE/H/1500/004	H/16/02206/009	STADA ARZNEIMITTEL AG	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
Perindopril/amlodipin STADA 8 mg/10 mg tablete	SE/H/1500/004	H/16/02206/010	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 8 mg/10 mg tablete	SE/H/1500/004	H/16/02206/011	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 8 mg/5 mg tablete	SE/H/1500/003	H/16/02206/008	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin STADA 8 mg/5 mg tablete	SE/H/1500/003	H/16/02206/007	STADA ARZNEIMITTEL AG	SI
Perindopril/amlodipin Teva 10 mg/10 mg tablety	NL/H/2842/04	58/224/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Perindopril/amlodipin Teva 10 mg/5 mg tablety	NL/H/2842/03	58/223/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Perindopril/amlodipin Teva 5 mg/10 mg tablety	NL/H/2842/02	58/222/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Perindopril/amlodipin Teva 5 mg/5 mg tablety	NL/H/2842/01	58/221/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Perindopril/Amlodipine 4 mg/10 mg tablets	UK/H/4620/002	PL01656/0134	KRKA, D.D., NOVO MESTO	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine 4 mg/10 mg tablets	UK/H/4348/002	PL 01656/0115	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 4 mg/10 mg tablets	UK/H/4621/002	PL01656/0138	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 4 mg/5 mg tablets	UK/H/4620/001	PL01656/0133	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 4 mg/5 mg tablets	UK/H/4348/001	PL01656/0114	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 4 mg/5 mg tablets	UK/H/4621/001	PL01656/0137	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 8 mg/10 mg tablets	UK/H/4620/004	PL01656/0136	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 8 mg/10 mg tablets	UK/H/4348/004	PL01656/0117	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 8 mg/10 mg tablets	UK/H/4621/004	PL01656/0140	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 8 mg/5 mg tablets	UK/H/4620/003	PL01656/0135	KRKA, D.D., NOVO MESTO	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine 8 mg/5 mg tablets	UK/H/4348/003	PL 01656/0116	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine 8 mg/5 mg tablets	UK/H/4621/003	PL01656/0139	KRKA, D.D., NOVO MESTO	UK
Perindopril/Amlodipine EG 4 mg/10 mg comprimés	SE/H/1500/002	BE499217	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 4 mg/10 mg Tabletten	SE/H/1500/002	BE499217	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 4 mg/10 mg tabletten	SE/H/1500/002	BE499217	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 4 mg/5 mg comprimés	SE/H/1500/001	BE499191	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 4 mg/5 mg Tabletten	SE/H/1500/001	BE499191	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 4 mg/5 mg tabletten	SE/H/1500/001	BE499191	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 8 mg/10 mg comprimés	SE/H/1500/004	BE499235	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
Perindopril/Amlodipine EG 8 mg/10 mg Tabletten	SE/H/1500/004	BE499235	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 8 mg/10 mg tabletten	SE/H/1500/004	BE499235	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 8 mg/5 mg comprimés	SE/H/1500/003	BE499226	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 8 mg/5 mg Tabletten	SE/H/1500/003	BE499226	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine EG 8 mg/5 mg tabletten	SE/H/1500/003	BE499226	EUROGENERICS N.V./S.A.	BE
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/012	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/013	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/014	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/015	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/016	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/017	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/018	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/019	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/020	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/021	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/10 mg tabletės	UK/H/4621/002	LT/1/11/2707/022	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletes	UK/H/4621/001	LT/1/11/2707/006	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/001	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/002	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/003	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/004	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/005	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/007	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/008	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/009	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/010	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 4 mg/5 mg tabletės	UK/H/4621/001	LT/1/11/2707/011	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/034	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/035	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/036	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/037	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/038	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/039	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/040	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/041	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/042	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/043	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/10 mg tabletės	UK/H/4621/004	LT/1/11/2707/044	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/023	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/024	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/025	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/026	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/027	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/028	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/029	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/030	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/031	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/032	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka 8 mg/5 mg tabletės	UK/H/4621/003	LT/1/11/2707/033	KRKA, D.D., NOVO MESTO	LT
Perindopril/Amlodipine Krka, 4 mg + 10 mg, tabletki	UK/H/4621/002	18773	KRKA, D.D., NOVO MESTO	PL
Perindopril/Amlodipine Krka, 4 mg + 5 mg, tabletki	UK/H/4621/001	18772	KRKA, D.D., NOVO MESTO	PL
Perindopril/Amlodipine Krka, 8 mg + 10 mg, tabletki	UK/H/4621/004	18775	KRKA, D.D., NOVO MESTO	PL
Perindopril/Amlodipine Krka, 8 mg + 5 mg, tabletki	UK/H/4621/003	18774	KRKA, D.D., NOVO MESTO	PL
Perindopril/Amlodipine STADA 4 mg/10 mg tablety	SE/H/1500/002	58/299/16-C	STADA ARZNEIMITTEL AG	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
Perindopril/Amlodipine STADA 4 mg/5 mg tablety	SE/H/1500/001	58/298/16-C	STADA ARZNEIMITTEL AG	CZ
Perindopril/Amlodipine STADA 8 mg/10 mg tablety	SE/H/1500/004	58/301/16-C	STADA ARZNEIMITTEL AG	CZ
Perindopril/Amlodipine STADA 8 mg/5 mg tablety	SE/H/1500/003	58/300/16-C	STADA ARZNEIMITTEL AG	CZ
Perindopril/Amlodipine Teva 10 mg/10 mg tabletes	NL/H/2842/004	14-0087	TEVA PHARMA B.V.	LV
Perindopril/Amlodipine Teva 10 mg/5 mg tabletes	NL/H/2842/003	14-0086	TEVA PHARMA B.V.	LV
Perindopril/Amlodipine Teva 5 mg/10 mg tabletes	NL/H/2842/002	14-0085	TEVA PHARMA B.V.	LV
Perindopril/Amlodipine Teva 5 mg/5 mg tabletes	NL/H/2842/01	14-0084	TEVA PHARMA B.V.	LV
Perindopril/Amlodipine Teva, 10 mg/10 mg tabletid	NL/H/2842/04	838414	TEVA PHARMA B.V.	EE
Perindopril/Amlodipine Teva, 10 mg/5 mg tabletid	NL/H/2842/03	838714	TEVA PHARMA B.V.	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Perindopril/Amlodipine Teva, 5 mg/10 mg tabletid	NL/H/2842/02	838614	TEVA PHARMA B.V.	EE
Perindopril/Amlodipine Teva, 5 mg/5 mg tabletid	NL/H/2842/01	838814	TEVA PHARMA B.V.	EE
Perindopril+Amlodipine/Teva (10+10) mg Δισκία	NL/H/2842/04	65761/09-07-2014	TEVA PHARMA B.V.	GR
Perindopril+Amlodipine/Teva (10+5) mg Δισκία	NL/H/2842/03	65760/09-07-2014	TEVA PHARMA B.V.	GR
Perindopril+Amlodipine/Teva (5+10) mg Δισκία	NL/H/2842/02	65759/09-07-2014	TEVA PHARMA B.V.	GR
Perindopril+Amlodipine/Teva (5+5) mg Δισκία	NL/H/2842/01	65758/09-07-2014	TEVA PHARMA B.V.	GR
Perindopril-tozilát/Amlodipin-Teva 10 mg/10 mg tabletta	not available	OGYI-T-22808/23	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/10 mg tabletta	not available	OGYI-T-22808/24	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/10 mg tabletta	not available	OGYI-T-22808/25	TEVA GYÓGYSZERGYÁR ZRT	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
Perindopril-tozilát/Amlodipin-Teva 10 mg/10 mg tabletta	not available	OGYI-T-22808/20	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/10 mg tabletta	not available	OGYI-T-22808/22	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/14	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/15	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/16	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/19	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/18	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 10 mg/5 mg tabletta	not available	OGYI-T-22808/17	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/12	TEVA GYÓGYSZERGYÁR ZRT	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
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/13	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/11	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/08	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/09	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/10 mg tabletta	not available	OGYI-T-22808/10	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/02	TEVA GYÓGYSZERGYÁR ZRT	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
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/07	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/06	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva 5 mg/5 mg tabletta	not available	OGYI-T-22808/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Perindopril-tozilát/Amlodipin-Teva10 mg/10 mg tabletta	not available	OGYI-T-22808/21	TEVA GYÓGYSZERGYÁR ZRT	HU
Perodilam 10 mg/10 mg tablete	NL/H/2842/04	H/15/01979/016	TEVA PHARMA B.V.	SI
Perodilam 10 mg/5 mg tablete	NL/H/2842/03	H/15/01979/011	TEVA PHARMA B.V.	SI
Perodilam 5 mg/5 mg tablete	NL-H-2842-01-04	H/15/01979/001	TEVA PHARMA B.V.	SI
Perodilam 5 mg/5 mg tablete	NL/H/2842/001	H/15/01979/006	TEVA PHARMA B.V.	SI
Prestalia 3,5 mg /2,5 mg comprimidos	IT/H/0375/001	5646450	LES LABORATOIRES SERVIER (SURESNES)	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
Prestalia 3,5 mg /2,5 mg comprimidos	IT/H/0375/001	5646468	LES LABORATOIRES SERVIER (SURESNES)	PT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148016	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148030	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148093	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148028	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148105	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg compresse	IT/H/0375/001	043148168	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 3,5 mg/2,5 mg comprimidos	IT/H/375/01/DC	Nº 79869	LES LABORATOIRES SERVIER (SURESNES)	ES
Prestalia 3,5 mg/2,5 mg tabletes	IT/H/375/01/DC	15-0092	LES LABORATOIRES SERVIER (SURESNES)	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
Prestalia 3,5 mg/2,5 mg tabletit	IT/H/0375/001	31869	LES LABORATOIRES SERVIER (SURESNES)	FI
Prestalia 3,5 mg/2,5 mg Tabletten	IT/H/375/01/DC	136182	LES LABORATOIRES SERVIER (SURESNES)	AT
Prestalia 3,5 mg/2,5 mg δισκία	IT/H/375/01/DC	31132/07-04-2016	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Prestalia 3,5 mg/2,5 mg, tabletid	IT/H/0375/001	876715	LES LABORATOIRES SERVIER (SURESNES)	EE
Prestalia 3,5 mg/2,5 mg, tabletten	IT/H/0375/001	RVG 114413	LES LABORATOIRES SERVIER (SURESNES)	NL
Prestalia 3,5 mg/2,5 mg, tablety	IT/H/0374/001	58/257/15-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Prestalia 7 mg / 5 mg comprimidos	IT/H/0375/002	5646476	LES LABORATOIRES SERVIER (SURESNES)	PT
Prestalia 7 mg / 5 mg comprimidos	IT/H/0375/002	5646500	LES LABORATOIRES SERVIER (SURESNES)	PT
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148067	LES LABORATOIRES SERVIER (SURESNES)	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
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148079	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148081	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148093	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148105	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 7 mg/5 mg compresse	IT/H/0375/002	043148168	LES LABORATOIRES SERVIER (SURESNES)	IT
Prestalia 7 mg/5 mg comprimidos	IT/H/375/02/DC	Nº 79870	LES LABORATOIRES SERVIER (SURESNES)	ES
Prestalia 7 mg/5 mg tabletes	IT/H/375/02/DC	15-0093	LES LABORATOIRES SERVIER (SURESNES)	LV
Prestalia 7 mg/5 mg tabletit	IT/H/0375/002	31870	LES LABORATOIRES SERVIER (SURESNES)	FI
Prestalia 7 mg/5 mg Tabletten	IT/H/375/02/DC	136183	LES LABORATOIRES SERVIER (SURESNES)	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
Prestalia 7 mg/5 mg δισκία	IT/H/375/02/DC	31133/07-04-2016	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Prestalia 7 mg/5 mg, tabletid	IT/H/0375/002	876815	LES LABORATOIRES SERVIER (SURESNES)	EE
Prestalia 7 mg/5 mg, tabletten	IT/H/0375/002	RVG 114414	LES LABORATOIRES SERVIER (SURESNES)	NL
Prestalia 7 mg/5 mg, tablety	IT/H/0374/002	58/258/15-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
PRESTANCE (10MG/5MG)	FR/H/325/03/DC	58/0222/08-S	LES LABORATOIRES SERVIER	SK
PRESTANCE (5MG/10MG)	FR/H/325/02/DC	58/0221/08-S	LES LABORATOIRES SERVIER	SK
PRESTANCE (5MG/5MG)	FR/H/325/01/DC	58/0220/08-S	LES LABORATOIRES SERVIER	SK
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/05	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/02	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/08	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/06	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/10	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/13	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/03	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/09	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/07	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/12	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/04	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/11	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/01	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg comprimate	FR/H/0325/004	742/2008/14	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/047	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/051	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/050	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/055	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/048	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/052	SERVIER PHARMA D.O.O	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
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/049	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/045	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/046	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/053	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/056	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/054	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/044	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablete	FR/H/0325/004	H/08/01281/043	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/10 mg tablety	FR/H/0325/004	58/206/08-C	LES LABORATOIRES SERVIER	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
PRESTANCE 10 mg/10 mg tablety	FR/H/0325/004	58/0223/08-S	LES LABORATOIRES SERVIER (SURESNES)	SK
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/08	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/06	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/05	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/09	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/02	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/14	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/10	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/12	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/13	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/04	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/11	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/01	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/07	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTANCE 10 mg/5 mg comprimate perindopril arginină/amlodipină	FR/H/0325/003	741/2008/03	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/033	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/035	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/040	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/031	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/038	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/032	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/036	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/041	SERVIER PHARMA D.O.O	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
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/034	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/042	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/037	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/039	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/030	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablete	FR/H/0325/003	H/08/01281/029	SERVIER PHARMA D.O.O	SI
PRESTANCE 10 mg/5 mg tablety	FR/H/0325/003	58/205/08-C	LES LABORATOIRES SERVIER	CZ
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/05	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/07	LES LABORATOIRES SERVIER	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
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/09	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/14	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/02	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/10	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/11	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/01	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/06	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/13	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/08	LES LABORATOIRES SERVIER	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
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/03	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/04	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg comprimate	FR/H/0325/002	740/2008/12	LES LABORATOIRES SERVIER	RO
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/028	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/026	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/018	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/019	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/022	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/017	SERVIER PHARMA D.O.O	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
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/020	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/027	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/023	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/025	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/021	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/024	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/015	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablete	FR/H/0325/002	H/08/01281/016	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/10 mg tablety	FR/H/0325/002	58/204/08-C	LES LABORATOIRES SERVIER	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
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/09	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/08	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/07	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/01	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/10	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/325/01/DC	739/2009/03	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/13	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/05	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/12	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/11	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/02	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/06	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/14	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg comprimate	FR/H/0325/001	739/2008/04	LES LABORATOIRES SERVIER (SURESNES)	RO
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/012	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/004	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/008	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/005	SERVIER PHARMA D.O.O	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
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/007	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/003	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/006	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/011	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/009	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/325/01/DC	22/09/2008	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/010	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/013	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablete	FR/H/0325/001	H/08/01281/002	SERVIER PHARMA D.O.O	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
PRESTANCE 5 mg/5 mg tablete	FR/H/325/01/DC	H/08/01281/001	SERVIER PHARMA D.O.O	SI
PRESTANCE 5 mg/5 mg tablety	FR/H/0325/001	58/203/08-C	LES LABORATOIRES SERVIER	CZ
Presteram 10 mg/10 mg tabletes	FR/H/0325/004	08-0114	LES LABORATOIRES SERVIER (SURESNES)	LV
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/053	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/051	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/045	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/052	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/048	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/050	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/056	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/046	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/044	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/054	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/055	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/047	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/043	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/10 mg tabletės	FR/H/0325/004	LT/1/08/1187/049	LES LABORATOIRES SERVIER (SURESNES)	LT
Presteram 10 mg/5 mg tabletes	FR/H/0325/003	08-0113	LES LABORATOIRES SERVIER (SURESNES)	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
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/031	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/041	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/040	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/032	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/034	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/033	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/035	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/037	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/042	LES LABORATOIRES SERVIER	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
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/036	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/029	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/030	LES LABORATOIRES SERVIER (SURESNES)	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/039	LES LABORATOIRES SERVIER	LT
PRESTERAM 10 mg/5 mg tabletės	FR/H/0325/003	LT/1/08/1187/038	LES LABORATOIRES SERVIER	LT
Presteram 5 mg/10 mg tabletes	FR/H/0325/002	08-0112	LES LABORATOIRES SERVIER (SURESNES)	LV
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/024	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/027	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/022	LES LABORATOIRES SERVIER	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
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/020	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/021	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/026	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/018	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/028	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002/	LT/1/08/1187/023	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/016	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/017	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/025	LES LABORATOIRES SERVIER	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
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/019	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/10 mg tabletės	FR/H/0325/002	LT/1/08/1187/015	LES LABORATOIRES SERVIER	LT
Presteram 5 mg/5 mg tabletes	FR/H/0325/001	08-0111	LES LABORATOIRES SERVIER (SURESNES)	LV
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/012	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/006	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/005	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/004	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/013	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/008	LES LABORATOIRES SERVIER	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
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/009	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/007	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/014	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/010	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/001	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/002	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/011	LES LABORATOIRES SERVIER	LT
PRESTERAM 5 mg/5 mg tabletės	FR/H/0325/001	LT/1/08/1187/003	LES LABORATOIRES SERVIER	LT
Prestozek 4 mg / 10 mg tablety	PL/H/0378/002	58/450/16-C	ADAMED	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Prestozek 4 mg / 10 mg tablety	PL/H/0378/002	58/450/16-C	ADAMED	CZ
Prestozek 4 mg / 5 mg tablety	PL/H/0378/001	58/449/16-C	ADAMED	CZ
Prestozek 4 mg / 5 mg tablety	PL/H/0378/001	58/449/16-C	ADAMED	CZ
Prestozek 4 mg/10 mg tabletes	PL/H/0378/002	16-0163	ADAMED	LV
Prestozek 4 mg/10 mg tabletes	PL/H/0378/002	16-0163	ADAMED	LV
Prestozek 4 mg/10 mg tablety	PL/H/0378/002	58/0260/16-S	ADAMED	SK
Prestozek 4 mg/10 mg tablety	PL/H/0378/002	58/0260/16-S	ADAMED	SK
Prestozek 4 mg/5 mg tabletes	PL/H/0378/001	16-0162	ADAMED	LV
Prestozek 4 mg/5 mg tabletes	PL/H/0378/001	16-0162	ADAMED	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
Prestozek 4 mg/5 mg tablety	PL/H/0378/001	58/0259/16-S	ADAMED	SK
Prestozek 4 mg/5 mg tablety	PL/H/0378/001	58/0259/16-S	ADAMED	SK
Prestozek 8 mg / 10 mg tablety	PL/H/0378/004	58/452/16-C	ADAMED	CZ
Prestozek 8 mg / 10 mg tablety	PL/H/0378/004	58/452/16-C	ADAMED	CZ
Prestozek 8 mg / 5 mg tablety	PL/H/0378/003	58/451/16-C	ADAMED	CZ
Prestozek 8 mg / 5 mg tablety	PL/H/0378/003	58/451/16-C	ADAMED	CZ
Prestozek 8 mg/10 mg tablety	PL/H/0378/004	58/0262/16-S	ADAMED	SK
Prestozek 8 mg/10 mg tablety	PL/H/0378/004	58/0262/16-S	ADAMED	SK
Prestozek 8 mg/10mg tabletes	PL/H/0378/004	16-0165	ADAMED	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
Prestozek 8 mg/10mg tabletes	PL/H/0378/004	16-0165	ADAMED	LV
Prestozek 8 mg/5 mg tablety	PL/H/0378/003	58/0261/16-S	ADAMED	SK
Prestozek 8 mg/5 mg tablety	PL/H/0378/003	58/0261/16-S	ADAMED	SK
Prestozek 8 mg/5mg tabletes	PL/H/0378/003	16-0164	ADAMED	LV
Prestozek 8 mg/5mg tabletes	PL/H/0378/003	16-0164	ADAMED	LV
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/08	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/09	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/10	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/11	ADAMED	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
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/12	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/13	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/14	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/08	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/09	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/10	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/11	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/12	ADAMED	HU
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/13	ADAMED	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
Prestozek Combi 4 mg/10 mg tabletta	PL/H/0378/002	OGYI-T-23034/14	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/01	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/02	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/05	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/03	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/04	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/06	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/07	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/01	ADAMED	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
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/02	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/05	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/03	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/04	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/06	ADAMED	HU
Prestozek Combi 4 mg/5 mg tabletta	PL/H/0378/001	OGYI-T-23034/07	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/22	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/25	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/24	ADAMED	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
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/26	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/23	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/28	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/27	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/22	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/25	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/24	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/26	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/23	ADAMED	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
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/28	ADAMED	HU
Prestozek Combi 8 mg/10 mg tabletta	PL/H/0378/004	OGYI-T-23034/27	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/15	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/17	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/16	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/21	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/18	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/20	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/19	ADAMED	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
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/001-004/DC	OGYI-T-23034/15	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/17	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/16	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/21	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/18	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/20	ADAMED	HU
Prestozek Combi 8 mg/5 mg tabletta	PL/H/0378/003	OGYI-T-23034/19	ADAMED	HU
Prestozek Combi, 4 mg + 10 mg, tabletki	PL/H/0378/001	23157	ADAMED	PL
Prestozek Combi, 4 mg + 10 mg, tabletki	PL/H/0378/001	23157	ADAMED	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
Prestozek Combi, 4 mg + 5 mg, tabletki	PL/H/0378/001	23156	ADAMED	PL
Prestozek Combi, 4 mg + 5 mg, tabletki	PL/H/0378/001	23156	ADAMED	PL
Prestozek Combi, 8 mg + 10 mg, tabletki	PL/H/0378/001	23159	ADAMED	PL
Prestozek Combi, 8 mg + 10 mg, tabletki	PL/H/0378/001	23159	ADAMED	PL
Prestozek Combi, 8 mg + 5 mg, tabletki	PL/H/0378/001	23158	ADAMED	PL
Prestozek Combi, 8 mg + 5 mg, tabletki	PL/H/0378/001	23158	ADAMED	PL
PRIAMLO 4 MG/ 10 MG TABLETES	CZ/H/0474/002	15-0205	ZENTIVA, K.S.	LV
PRIAMLO 4 MG/ 10 MG TABLETES	CZ/H/0474/002	15-0205	ZENTIVA, K.S.	LV
PRIAMLO 4 MG/ 5 MG TABLETES	CZ/H/0474/001	15-0204	ZENTIVA, K.S.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PRIAMLO 4 MG/ 5 MG TABLETES	CZ/H/0474/001	15-0204	ZENTIVA, K.S.	LV
PRIAMLO 4 MG/ 5 MG TABLETES	CZ/H/0474/001	15-0204	ZENTIVA, K.S.	LV
PRIAMLO 4 MG/ 5 MG TABLETES	CZ/H/0474/001	15-0204	ZENTIVA, K.S.	LV
PRIAMLO 4 MG/10 MG TABLETID	CZ/H/0474/002	876915	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/10 MG TABLETID	CZ/H/0474/002	876915	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/10 MG TABLETY	CZ/H/0474/002	58/461/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/10 MG TABLETY	CZ/H/0474/002	58/461/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/10 MG TABLETY	CZ/H/0474/002	58/0288/15-S	ZENTIVA, K.S.	SK
PRIAMLO 4 MG/10 MG TABLETY	CZ/H/0474/002	58/0288/15-S	ZENTIVA, K.S.	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
PRIAMLO 4 MG/5 MG TABLETID	CZ/H/0474/001	877015	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/5 MG TABLETID	CZ/H/0474/001	877015	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/5 MG TABLETID	CZ/H/0474/001	877015	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/5 MG TABLETID	CZ/H/0474/001	877015	ZENTIVA, K.S.	EE
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/460/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/460/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/460/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/460/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/0286/15-S	ZENTIVA, K.S.	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
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/0286/15-S	ZENTIVA, K.S.	SK
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/0286/15-S	ZENTIVA, K.S.	SK
PRIAMLO 4 MG/5 MG TABLETY	CZ/H/0474/001	58/0286/15-S	ZENTIVA, K.S.	SK
PRIAMLO 8 MG/ 10 MG TABLETES	CZ/H/0474/004	15-0207	ZENTIVA, K.S.	LV
PRIAMLO 8 MG/ 10 MG TABLETES	CZ/H/0474/004	15-0207	ZENTIVA, K.S.	LV
PRIAMLO 8 MG/ 5 MG TABLETES	CZ/H/0474/003	15-0206	ZENTIVA, K.S.	LV
PRIAMLO 8 MG/ 5 MG TABLETES	CZ/H/0474/003	15-0206	ZENTIVA, K.S.	LV
PRIAMLO 8 MG/10 MG TABLETID	CZ/H/0474/004	877115	ZENTIVA, K.S.	EE
PRIAMLO 8 MG/10 MG TABLETID	CZ/H/0474/004	877115	ZENTIVA, K.S.	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PRIAMLO 8 MG/10 MG TABLETY	CZ/H/0474/004	58/463/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 8 MG/10 MG TABLETY	CZ/H/0474/004	58/463/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 8 MG/10 MG TABLETY	CZ/H/0474/004	58/0290/15-S	ZENTIVA, K.S.	SK
PRIAMLO 8 MG/10 MG TABLETY	CZ/H/0474/004	58/0290/15-S	ZENTIVA, K.S.	SK
PRIAMLO 8 MG/5 MG TABLETID	CZ/H/0474/003	877215	ZENTIVA, K.S.	EE
PRIAMLO 8 MG/5 MG TABLETID	CZ/H/0474/003	877215	ZENTIVA, K.S.	EE
PRIAMLO 8 MG/5 MG TABLETY	CZ/H/0474/003	58/462/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 8 MG/5 MG TABLETY	CZ/H/0474/003	58/462/15-C	ZENTIVA, K.S.	CZ
PRIAMLO 8 MG/5 MG TABLETY	CZ/H/0474/003	58/0289/15-S	ZENTIVA, K.S.	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
PRIAMLO 8 MG/5 MG TABLETY	CZ/H/0474/003	58/0289/15-S	ZENTIVA, K.S.	SK
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483057	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483095	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483018	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483032	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483071	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483119	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483069	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483121	IST. FARMACO BIOLOGICO SRL STRODER	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
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483020	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483044	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483107	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483083	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483133	IST. FARMACO BIOLOGICO SRL STRODER	IT
REAPTAN 5 mg/5 mg compresse	FR/H/0326/001	038483564	IST. FARMACO BIOLOGICO SRL STRODER	IT
Tonarssa 4 mg/10 mg tablety	not available	58/439/11-C	KRKA, D.D., NOVO MESTO	CZ
Tonarssa 4 mg/5 mg tablety	not available	58/438/11-C	KRKA, D.D., NOVO MESTO	CZ
Tonarssa 8 mg/10 mg tablety	not available	58/441/11-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
Tonarssa 8 mg/5 mg tablety	not available	58/440/11-C	KRKA, D.D., NOVO MESTO	CZ
Viacoram	IT/H/0374/001	53247	LES LABORATOIRES SERVIER (SURESNES)	DK
Viacoram	IT/H/0374/002	53248	LES LABORATOIRES SERVIER (SURESNES)	DK
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/003	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/005	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/004	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/011	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/002	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 3,5 mg / 2,5 mg tabletės	IT/H/0374/001	LT/1/15/3727/013	LES LABORATOIRES SERVIER (SURESNES)	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
Viacoram 3,5 mg /2,5 mg comprimidos	IT/H/0374/001	5646419	LES LABORATOIRES SERVIER (SURESNES)	PT
Viacoram 3,5 mg /2,5 mg comprimidos	IT/H/0374/001	5646427	LES LABORATOIRES SERVIER (SURESNES)	PT
Viacoram 3,5 mg /2,5 mg comprimidos	IT/H/0374/001	5679253	LES LABORATOIRES SERVIER (SURESNES)	PT
VIACORAM 3,5 mg/2,5 mg	IT/H/374/01/DC	58/0163/15-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147014	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147026	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147053	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147040	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147038	LES LABORATOIRES SERVIER (SURESNES)	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
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147166	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg compresse	IT/H/0374/001	043147180	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 3,5 mg/2,5 mg comprimate	IT/H/374/01/DC	7997/2015/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 3,5 mg/2,5 mg comprimate	IT/H/374/01/DC	7997/2015/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 3,5 mg/2,5 mg comprimate	IT/H/374/01/DC	7997/2015/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 3,5 mg/2,5 mg comprimate	IT/H/374/01/DC	7997/2015/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 3,5 mg/2,5 mg comprimate	IT/H/374/01/DC	7997/2015/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 3,5 mg/2,5 mg comprimidos	IT/H/374/01/DC	79871	LES LABORATOIRES SERVIER (SURESNES)	ES
Viacoram 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	HR-H-686029633	SERVIER PHARMA D.O.O - CROATIA	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
VIACORAM 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	H/15/02053/001	SERVIER PHARMA D.O.O	SI
VIACORAM 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	H/15/02053/002	SERVIER PHARMA D.O.O	SI
VIACORAM 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	H/15/02053/003	SERVIER PHARMA D.O.O	SI
VIACORAM 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	H/15/02053/005	SERVIER PHARMA D.O.O	SI
VIACORAM 3,5 mg/2,5 mg tablete	IT/H/374/01/DC	H/15/02053/004	SERVIER PHARMA D.O.O	SI
Viacoram 3,5 mg/2,5 mg tablete	IT/H/0374/001	H/15/02053/012	SERVIER PHARMA D.O.O	SI
Viacoram 3,5 mg/2,5 mg tablete	IT/H/0374/001	H/15/02053/011	SERVIER PHARMA D.O.O	SI
Viacoram 3,5 mg/2,5 mg tabletes	IT/H/374/01/DC	15-0094	LES LABORATOIRES SERVIER (SURESNES)	LV
Viacoram 3,5 mg/2,5 mg tabletit	IT/H/374/01/DC	31864	LES LABORATOIRES SERVIER (SURESNES)	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Viacoram 3,5 mg/2,5 mg Tabletten	IT/H/374/01/DC	136180	LES LABORATOIRES SERVIER (SURESNES)	AT
Viacoram 3,5 mg/2,5 mg Tabletten	IT/H/0374/001	90826.00.00	LES LABORATOIRES SERVIER (SURESNES)	DE
Viacoram 3,5 mg/2,5 mg δισκία	IT/H/374/01/DC	32863/13-4-2016	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Viacoram 3,5 mg/2,5 mg, tabletid	IT/H/0374/001	877715	LES LABORATOIRES SERVIER (SURESNES)	EE
Viacoram 3,5 mg/2,5 mg, tabletten	IT/H/374/01/DC	RVG 114410	LES LABORATOIRES SERVIER (SURESNES)	NL
VIACORAM 3,5mg/2,5mg, comprimés	IT/H/374/01/DC	BE474560	SERVIER BENELUX S.A./N.V.	BE
VIACORAM 3,5mg/2,5mg, comprimés	IT/H/374/01/DC	2016060226	SERVIER BENELUX S.A./N.V.	LU
VIACORAM 3,5mg/2,5mg, tabletten	IT/H/374/01/DC	BE474560	SERVIER BENELUX S.A./N.V.	BE
VIACORAM 3,5mg/2,5mg, Tabletten	IT/H/374/01/DC	BE474560	SERVIER BENELUX S.A./N.V.	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
VIACORAM 3,5mg/2,5mg, tabletten	IT/H/374/01/DC	2016060226	SERVIER BENELUX S.A./N.V.	LU
VIACORAM 3,5mg/2,5mg, Tabletten	IT/H/374/01/DC	2016060226	SERVIER BENELUX S.A./N.V.	LU
Viacoram 3.5 mg/2.5 mg tablets	IT/H/0374/001	PA 0568/027/001	LES LABORATOIRES SERVIER (SURESNES)	IE
Viacoram 3.5 mg/2.5 mg tablets	IT/H/374/01/DC	MA066/02301	LES LABORATOIRES SERVIER (SURESNES)	MT
Viacoram 7 mg / 5 mg comprimidos	IT/H/0374/002	5646435	LES LABORATOIRES SERVIER (SURESNES)	PT
Viacoram 7 mg / 5 mg comprimidos	IT/H/0374/002	5679261	LES LABORATOIRES SERVIER (SURESNES)	PT
Viacoram 7 mg / 5 mg comprimidos	IT/H/0374/002	5646443	LES LABORATOIRES SERVIER (SURESNES)	PT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/006	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/008	LES LABORATOIRES SERVIER (SURESNES)	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
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/007	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/010	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/014	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/012	LES LABORATOIRES SERVIER (SURESNES)	LT
Viacoram 7 mg / 5 mg tabletės	IT/H/0374/002	LT/1/15/3727/009	LES LABORATOIRES SERVIER (SURESNES)	LT
VIACORAM 7 mg/5 mg	IT/H/374/02/DC	58/0164/15-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147065	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147077	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147089	LES LABORATOIRES SERVIER (SURESNES)	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
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147091	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147178	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147103	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg compresse	IT/H/0374/002	043147192	LES LABORATOIRES SERVIER (SURESNES)	IT
Viacoram 7 mg/5 mg comprimate	IT/H/374/02/DC	7998/2015/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 7 mg/5 mg comprimate	IT/H/374/02/DC	7998/2015/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 7 mg/5 mg comprimate	IT/H/374/02/DC	7998/2015/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 7 mg/5 mg comprimate	IT/H/374/02/DC	7998/2015/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacoram 7 mg/5 mg comprimate	IT/H/374/02/DC	7998/2015/05	LES LABORATOIRES SERVIER (SURESNES)	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
Viacoram 7 mg/5 mg comprimidos	IT/H/374/02/DC	79868	LES LABORATOIRES SERVIER (SURESNES)	ES
Viacoram 7 mg/5 mg tablete	IT/H/374/02/DC	HR-H-050504425	SERVIER PHARMA D.O.O - CROATIA	HR
VIACORAM 7 mg/5 mg tablete	IT/H/374/02/DC	H/15/02053/006	SERVIER PHARMA D.O.O	SI
VIACORAM 7 mg/5 mg tablete	IT/H/374/02/DC	H/15/02053/007	SERVIER PHARMA D.O.O	SI
VIACORAM 7 mg/5 mg tablete	IT/H/374/02/DC	H/15/02053/010	SERVIER PHARMA D.O.O	SI
VIACORAM 7 mg/5 mg tablete	IT/H/374/02/DC	H/15/02053/009	SERVIER PHARMA D.O.O	SI
VIACORAM 7 mg/5 mg tablete	IT/H/374/02/DC	H/15/02053/008	SERVIER PHARMA D.O.O	SI
Viacoram 7 mg/5 mg tablete	IT/H/0374/002	H/15/02053/013	SERVIER PHARMA D.O.O	SI
Viacoram 7 mg/5 mg tablete	IT/H/0374/002	H/15/02053/014	SERVIER PHARMA D.O.O	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
Viacoram 7 mg/5 mg tabletes	IT/H/374/02/DC	15-0095	LES LABORATOIRES SERVIER (SURESNES)	LV
Viacoram 7 mg/5 mg tabletit	IT/H/0374/002	31865	LES LABORATOIRES SERVIER (SURESNES)	FI
Viacoram 7 mg/5 mg tablets	IT/H/0374/002	PA 0568/027/002	LES LABORATOIRES SERVIER (SURESNES)	IE
Viacoram 7 mg/5 mg tablets	IT/H/374/02/DC	MA066/02302	LES LABORATOIRES SERVIER (SURESNES)	MT
Viacoram 7 mg/5 mg Tabletten	IT/H/374/02/DC	136181	LES LABORATOIRES SERVIER (SURESNES)	AT
Viacoram 7 mg/5 mg Tabletten	IT/H/0374/002	90827.00.00	LES LABORATOIRES SERVIER (SURESNES)	DE
Viacoram 7 mg/5 mg δισκία	IT/H/374/02/DC	32864/13-4-2016	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Viacoram 7 mg/5 mg, tabletid	IT/H/0374/002	877815	LES LABORATOIRES SERVIER (SURESNES)	EE
Viacoram 7 mg/5 mg, tabletten	IT/H/374/02/DC	RVG 114411	LES LABORATOIRES SERVIER (SURESNES)	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VIACORAM 7mg/5mg, comprimés	IT/H/374/02/DC	BE474577	SERVIER BENELUX S.A./N.V.	BE
VIACORAM 7mg/5mg, comprimés	IT/H/374/02/DC	2016060227	SERVIER BENELUX S.A./N.V.	LU
VIACORAM 7mg/5mg, Tabletten	IT/H/374/02/DC	BE474577	SERVIER BENELUX S.A./N.V.	BE
VIACORAM 7mg/5mg, tabletten	IT/H/374/02/DC	BE474577	SERVIER BENELUX S.A./N.V.	BE
VIACORAM 7mg/5mg, tabletten	IT/H/374/02/DC	2016060227	SERVIER BENELUX S.A./N.V.	LU
VIACORAM 7mg/5mg, Tabletten	IT/H/374/02/DC	2016060227	SERVIER BENELUX S.A./N.V.	LU
Vialibram 3,5 mg/2,5 mg, tablety	IT/H/0375/001	58/259/15-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Vialibram 7 mg/5 mg, tablety	IT/H/0375/002	58/260/15-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Vidonorm 4 mg/10 mg tabletes	HU/H/0311/003	12-0295	GEDEON RICHTER PLC.	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
VIDONORM 4 mg/10 mg tabletės	HU/H/0311/003	LT/1/12/3029/005	GEDEON RICHTER PLC.	LT
VIDONORM 4 mg/10 mg tabletės	HU/H/0311/003	LT/1/12/3029/006	GEDEON RICHTER PLC.	LT
Vidonorm 4 mg/10 mg tabletta	HU/H/0311/003	OGYI-T-22147/02	GEDEON RICHTER PLC.	HU
Vidonorm 4 mg/10 mg tablety	HU/H/0311/003	58/293/12-C	GEDEON RICHTER PLC.	CZ
Vidonorm 4 mg/10 mg tablety	HU/H/0311/003	58/0169/12-S	GEDEON RICHTER PLC.	SK
Vidonorm 4 mg/5 mg tabletes	HU/H/0311/001	12-0294	GEDEON RICHTER PLC.	LV
VIDONORM 4 mg/5 mg tabletės	HU/H/0311/001	LT/1/12/3029/001	GEDEON RICHTER PLC.	LT
VIDONORM 4 mg/5 mg tabletės	HU/H/0311/001	LT/1/12/3029/002	GEDEON RICHTER PLC.	LT
Vidonorm 4 mg/5 mg tabletta	HU/H/0311/001	OGYI-T-22147/01	GEDEON RICHTER 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
Vidonorm 4 mg/5 mg tablety	HU/H/0311/001	58/291/12-C	GEDEON RICHTER PLC.	CZ
Vidonorm 4 mg/5 mg tablety	HU/H/0311/001	58/0167/12-S	GEDEON RICHTER PLC.	SK
Vidonorm 8 mg/10 mg tabletes	HU/H/0311/004	12-0297	GEDEON RICHTER PLC.	LV
Vidonorm 8 mg/10 mg tabletés	HU/H/0311/004	LT/1/12/3029/007	GEDEON RICHTER PLC.	LT
Vidonorm 8 mg/10 mg tabletés	HU/H/0311/004	LT/1/12/3029/008	GEDEON RICHTER PLC.	LT
Vidonorm 8 mg/10 mg tabletta	HU/H/0311/004	OGYI-T-22147/04	GEDEON RICHTER PLC.	HU
Vidonorm 8 mg/10 mg tablety	HU/H/0311/004	58/294/12-C	GEDEON RICHTER PLC.	CZ
Vidonorm 8 mg/10 mg tablety	HU/H/0311/004	58/0170/12-S	GEDEON RICHTER PLC.	SK
Vidonorm 8 mg/5 mg tabletes	HU/H/0311/002	12-0296	GEDEON RICHTER PLC.	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
VIDONORM 8 mg/5 mg tabletės	HU/H/0311/002	LT/1/12/3029/003	GEDEON RICHTER PLC.	LT
VIDONORM 8 mg/5 mg tabletės	HU/H/0311/002	LT/1/12/3029/004	GEDEON RICHTER PLC.	LT
Vidonorm 8 mg/5 mg tabletta	HU/H/0311/002	OGYI-T-22147/03	GEDEON RICHTER PLC.	HU
Vidonorm 8 mg/5 mg tablety	HU/H/0311/002	58/292/12-C	GEDEON RICHTER PLC.	CZ
Vidonorm 8 mg/5 mg tablety	HU/H/0311/002	58/0168/12-S	GEDEON RICHTER PLC.	SK
Vilpin Combi, 10 mg + 10 mg, tabletki	NL/H/2842/004	21863	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Vilpin Combi, 10 mg + 5 mg, tabletki	NL/H/2842/003	21862	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Vilpin Combi, 5 mg + 10 mg, tabletki	NL/H/2842/002	21861	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Vilpin Combi, 5 mg + 5 mg, tabletki	NL/H/2842/001	21860	TEVA PHARMACEUTICALS POLSKA 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
Амлеса 4 mg/10 mg таблетки	not available	20120557	KRKA, D.D., NOVO MESTO	BG
Амлеса 4 mg/10 mg таблетки	not available	20120557	KRKA, D.D., NOVO MESTO	BG
Амлеса 4 mg/5 mg таблетки	not available	20120556	KRKA, D.D., NOVO MESTO	BG
Амлеса 4 mg/5 mg таблетки	not available	20120556	KRKA, D.D., NOVO MESTO	BG
Амлеса 8 mg/10 mg таблетки	not available	20120559	KRKA, D.D., NOVO MESTO	BG
Амлеса 8 mg/10 mg таблетки	not available	20120559	KRKA, D.D., NOVO MESTO	BG
Амлеса 8 mg/5 mg таблетки	not available	20120558	KRKA, D.D., NOVO MESTO	BG
Амлеса 8 mg/5 mg таблетки	not available	20120558	KRKA, D.D., NOVO MESTO	BG
Видонорм 4 mg/10 mg таблетки	HU/H/0311/003	20120238	GEDEON RICHTER PLC.	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
Видонорм 4 mg/5 mg таблетки	HU/H/0311/001	20120236	GEDEON RICHTER PLC.	BG
Видонорм 8 mg/10 mg таблетки	HU/H/0311/004	20120239	GEDEON RICHTER PLC.	BG
Видонорм 8 mg/5 mg таблетки	HU/H/0311/002	20120237	GEDEON RICHTER PLC.	BG
ЗАПРИНЕЛ А 10 mg/10 mg таблетки	NL-H-2842-01-04	20140127	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
ЗАПРИНЕЛ А 10 mg/5 mg таблетки	NL/H/2842/003	20140140	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
ЗАПРИНЕЛ А 5 mg/10 mg таблетки	NL-H-2842-01-04	20140136	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
ЗАПРИНЕЛ А 5 mg/5 mg таблетки	NL-H-2842-01-04	20140135	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Престариум Ко старт 3,5 mg/2,5 mg таблетки	IT/H/374/01/DC	20150182	LES LABORATOIRES SERVIER (SURESNES)	BG
Престариум Ко старт 7 mg/5 mg таблетки	IT/H/374/02/DC	20150183	LES LABORATOIRES SERVIER (SURESNES)	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
ПРЕСТАРИУМ-КО 10mg/10mg таблетки	FR/H/325/04/DC	20080116	LES LABORATOIRES SERVIER	BG
ПРЕСТАРИУМ-КО 10mg/5mg таблетки	FR/H/325/03/DC	20080115	LES LABORATOIRES SERVIER	BG
ПРЕСТАРИУМ-КО 5mg/10mg таблетки	FR/H/325/02/DC	20080114	LES LABORATOIRES SERVIER	BG
ПРЕСТАРИУМ-КО 5mg/5mg таблетки	FR/H/325/01/DC	20080113	LES LABORATOIRES SERVIER	BG
Престозек Комби 4 mg / 10 mg таблетки	PL/H/0378/002	20160254	ADAMED	BG
Престозек Комби 4 mg / 10 mg таблетки	PL/H/0378/002	20160254	ADAMED	BG
Престозек Комби 4 mg / 5 mg таблетки	PL/H/0378/001	20160253	ADAMED	BG
Престозек Комби 4 mg / 5 mg таблетки	PL/H/0378/001	20160253	ADAMED	BG
Престозек Комби 8 mg / 10 mg таблетки	PL/H/0378/004	20160256	ADAMED	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
Престозек Комби 8 mg / 10 mg таблетки	PL/H/0378/004	20160256	ADAMED	BG
Престозек Комби 8 mg / 5 mg таблетки	PL/H/0378/003	20160255	ADAMED	BG
Престозек Комби 8 mg / 5 mg таблетки	PL/H/0378/003	20160255	ADAMED	BG
ПРИАМЛО 4 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/001	20150280	ZENTIVA, K.S.	BG
ПРИАМЛО 4 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/001	20150280	ZENTIVA, K.S.	BG
ПРИАМЛО 4 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/001	20150280	ZENTIVA, K.S.	BG
ПРИАМЛО 4 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/001	20150280	ZENTIVA, K.S.	BG
ПРИАМЛО 8 MG/10 MG ТАБЛЕТКИ	CZ/H/0474/004	20150282	ZENTIVA, K.S.	BG
ПРИАМЛО 8 MG/10 MG ТАБЛЕТКИ	CZ/H/0474/004	20150282	ZENTIVA, K.S.	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
ПРИАМЛО 8 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/003	20150281	ZENTIVA, K.S.	BG
ПРИАМЛО 8 MG/5 MG ТАБЛЕТКИ	CZ/H/0474/003	20150281	ZENTIVA, K.S.	BG