29 September 2022
EMA/805538/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / atorvastatin

Procedure no.: PSUSA/00000177/202201

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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
Caduet 5 mg/10 mg filmom obložene tablete	not available	HR-H-444337926	PFIZER CROATIA D.O.O.	HR
Caduet 10 mg/10 mg filmom obložene tablete	not available	HR-H-732959683	PFIZER CROATIA D.O.O.	HR
Atorvastatina e amlodipina Gedeon Richter 10 mg/5 mg compresse rivestite con film	HU/H/0322/001	041064015	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 10 mg/5 mg compresse rivestite con film	HU/H/0322/001	041064027	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 10 mg/5 mg compresse rivestite con film	HU/H/0322/001	041064039	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/10 mg compresse rivestite con film	HU/H/0322/004	041064104	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/10 mg compresse rivestite con film	HU/H/0322/004	041064116	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/10 mg compresse rivestite con film	HU/H/0322/004	041064128	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/5 mg compresse rivestite con film	HU/H/0322/003	041064078	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/5 mg compresse rivestite con film	HU/H/0322/003	041064080	GEDEON RICHTER PLC.	IT
Atorvastatina e amlodipina Gedeon Richter 20 mg/5 mg compresse rivestite con film	HU/H/0322/003	041064092	GEDEON RICHTER PLC.	IT
CADUET 10 mg/10 mg comprimate filmate	not available	6945/2006/01	PFIZER EUROPE MA EEIG	RO
CADUET 5 mg/10 mg comprimate filmate	not available	6944/2006/01	PFIZER EUROPE MA EEIG	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
Caduet® 5 mg/10 mg Filmtabletten	FR/H/0280/001	1-26271	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Caduet® 10 mg/10 mg Filmtabletten	FR/H/0280/002	1-26273	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Caduet 10 mg/10 mg apvalkotās tabletes	FR/H/0280/002	06-0021	PFIZER EUROPE MA EEIG	LV
Caduet 10 mg/10 mg filmtabletta	FR/H/0280/002	OGYI-T-10 613/02	PFIZER KFT.	HU
Caduet 5 mg/10 mg apvalkotās tabletes	FR/H/0280/001	06-0020	PFIZER EUROPE MA EEIG	LV
Caduet 10 mg/10 mg comprimidos recubiertos con película.	FR/H/0280/002	67.326	PFIZER, S.L.	ES
Caduet 5 mg/10 mg filmtabletta	FR/H/0280/001	OGYI-T-10 613/01	PFIZER KFT.	HU
Caduet 5 mg/10 mg comprimidos recubiertos con película.	FR/H/0280/001	67.325	PFIZER, S.L.	ES
CADUET 5 mg + 10 mg comprimidos revestidos por película	FR/H/0280/001	5674585	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 10 mg + 10 mg comprimidos revestidos por película	FR/H/0280/002	5675186	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 10 mg + 10 mg comprimidos revestidos por película	FR/H/0280/002	5675087	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 10 mg + 10 mg comprimidos revestidos por película	FR/H/0280/002	5674882	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 5 mg/10 mg potahované tablety	FR/H/0280/001	83/616/05-C	PFIZER, SPOL. S R.O.	CZ
CADUET 5 mg + 10 mg comprimidos revestidos por	FR/H/0280/001	5674387	LABORATÓRIOS PFIZER, 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
película				
CADUET 10 mg/10 mg potahované tablety	FR/H/0280/002	83/617/05-C	PFIZER, SPOL. S R.O.	CZ
CADUET 10 mg + 10 mg comprimidos revestidos por película	FR/H/0280/002	5674981	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 5 mg + 10 mg comprimidos revestidos por película	FR/H/0280/001	5674783	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 10 mg + 10 mg comprimidos revestidos por película	FR/H/0280/002	5675285	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 5 mg + 10 mg comprimidos revestidos por película	FR/H/0280/001	5674486	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 5 mg + 10 mg comprimidos revestidos por película	FR/H/0280/001	5674684	LABORATÓRIOS PFIZER, LDA.	PT
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 300 9 4	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 846 4 9	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 304 4 5	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 303 8 4	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 844 1 0	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 375 754 8 5	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 847 0 0	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 302 1 6	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé	FR/H/0280/001	34009 369 306 7 4	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pelliculé				
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 845 8 8	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 301 5 5	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 843 5 9	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg film-coated tablets.	FR/H/0280/002	MA505/04802	PFIZER HELLAS, A.E.	MT
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 566 848 7 8	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg, comprimé pelliculé	FR/H/0280/001	34009 369 305 0 6	PFIZER HOLDING FRANCE	FR
CADUET 5 mg/10 mg film-coated tablets.	FR/H/0280/001	MA505/04801	PFIZER HELLAS, A.E.	MT
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 838 1 9	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 841 2 0	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 375 755 4 6	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 294 9 4	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 299 0 6	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 295 5 5	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 298 4 5	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 837 5 8	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 842 9 8	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 296 1 6	PFIZER HOLDING FRANCE	FR

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CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 293 2 6	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 840 6 9	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 566 839 8 7	PFIZER HOLDING FRANCE	FR
CADUET 10 mg/10 mg, comprimé pelliculé	FR/H/0280/002	34009 369 297 8 4	PFIZER HOLDING FRANCE	FR
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/06	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/04	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/22	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/09	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/14	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/27	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/13	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/15	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/19	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/03	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/24	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/23	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/12	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/11	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/17	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/25	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/16	PFIZER KFT.	HU

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DICARTIL 10 mg/10 mg filmom obalené tablety	FR/H/0478/002	58/0333/11-S	PFIZER EUROPE MA EEIG	SK
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/18	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/02	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/08	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/26	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/20	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/05	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/01	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/10	PFIZER KFT.	HU
Dicartil 10 mg/10 mg filmtabletta	FR/H/0478/002	OGYI-T-21776/21	PFIZER KFT.	HU
Dicartil 5 mg/10 mg filmtabletta	FR/H/0478/001	OGYI-T-21776/07	PFIZER KFT.	HU
DICARTIL 5 mg/10 mg filmom obalené tablety	FR/H/0478/001	58/0332/11-S	PFIZER EUROPE MA EEIG	SK
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 444 0 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 325 6 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 449 2 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 443 4 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 448 6 8	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé	FR/H/0478/001	34009 219 331 6 1	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pelliculé				
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 328 5 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 329 1 1	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 445 7 8	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 326 2 1	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 332 2 2	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 219 327 9 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 450 0 1	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 5 mg/10 mg, comprimé pelliculé	FR/H/0478/001	34009 581 446 3 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 321 0 2	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 322 7 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 442 8 8	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE	FR/H/0478/002	34009 581 438 0 9	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PFIZER 10 mg/10 mg, comprimé pelliculé				
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 320 4 1	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 316 7 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 323 3 1	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 319 6 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 437 4 8	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 441 1 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 436 8 7	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 440 5 9	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 219 317 3 0	PFIZER HOLDING FRANCE	FR
AMLODIPINE/ATORVASTATINE PFIZER 10 mg/10 mg, comprimé pelliculé	FR/H/0478/002	34009 581 439 7 7	PFIZER HOLDING FRANCE	FR
КАДУЕТ 5 mg/10 mg филмирани таблетки	not available	20060484	PFIZER EUROPE MA EEIG	BG
КАДУЕТ 10 mg/10 mg	not available	20060485	PFIZER EUROPE MA EEIG	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
филмирани таблетки				
TULIP COMBI 5 mg/10 mg potahované tablety	PL/H/0474/001	83/026/17-C	SANDOZ S.R.O.	CZ
TULIP Combi 5 mg /20 mg potahované tablety	PL/H/0474/002	83/027/17-C	SANDOZ S.R.O.	CZ
TULIP Combi 10 mg/20 mg potahované tablety	PL/H/0474/003	83/028/17-C	SANDOZ S.R.O.	CZ
Amlosatin, 5 mg + 10 mg, tabletki powlekane	PL/H/0474/001	24897	SANDOZ GMBH	PL
Amlosatin, 5 mg + 20 mg, tabletki powlekane	PL/H/0474/002	24898	SANDOZ GMBH	PL
Amlosatin, 10 mg + 20 mg, tabletki powlekane	PL/H/0474/003	24899	SANDOZ GMBH	PL
Amlator 20 mg/5 mg filmtabletta	HU/H/0248/003	OGYI-T-21844/03	GEDEON RICHTER PLC.	HU
Amlator 10 mg/10 mg filmtabletta	HU/H/0248/002	OGYI-T-21844/02	GEDEON RICHTER PLC.	HU
Amlator 20 mg/10 mg filmtabletta	HU/H/0248/004	OGYI-T-21844/04	GEDEON RICHTER PLC.	HU
Duplecor 10 mg/5 mg comprimate filmate	HU/H/0248/001	9047/2016/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Duplecor 10 mg/10 mg comprimate filmate	HU/H/0248/002	9048/2016/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Duplecor 20 mg/10 mg comprimate filmate	HU/H/0248/004	9050/2016/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Duplecor 20 mg/5 mg comprimate filmate	HU/H/0248/003	9049/2016/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Amlator, 20 mg + 5 mg, tabletki powlekane	HU/H/0248/003	18575	GEDEON RICHTER PLC.	PL
Amlator 20 mg/5 mg potahované tablety	HU/H/0248/003	58/482/11-C	GEDEON RICHTER PLC.	CZ
Amlator 10 mg/10 mg filmom obalené tablety	HU/H/0248/002	58/0382/11-S	GEDEON RICHTER PLC.	SK
Amlator 20 mg/10 mg filmom obalené tablety	HU/H/0248/004	58/0384/11-S	GEDEON RICHTER PLC.	SK
Amlator, 10 mg + 5 mg, tabletki	HU/H/0248/001	18573	GEDEON RICHTER PLC.	PL

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powlekane				
Amlator, 10 mg + 10 mg, tabletki powlekane	HU/H/0248/002	18574	GEDEON RICHTER PLC.	PL
Amlator 10 mg/5 mg filmtabletta	HU/H/0248/001	OGYI-T-21844/01	GEDEON RICHTER PLC.	HU
Amlator 10 mg/5 mg filmom obalené tablety	HU/H/0248/001	58/0381/11-S	GEDEON RICHTER PLC.	SK
Amlator 10 mg/10 mg potahované tablety	HU/H/0248/002	58/481/11-C	GEDEON RICHTER PLC.	CZ
Amlator 20 mg/5 mg filmom obalené tablety	HU/H/0248/003	58/0383/11-S	GEDEON RICHTER PLC.	SK
Amlator 20 mg/10 mg potahované tablety	HU/H/0248/004	58/483/11-C	GEDEON RICHTER PLC.	CZ
АМЛАТОР 20 mg/5 mg филмирани таблетки	HU/H/0248/003	20110297	GEDEON RICHTER PLC.	BG
АМЛАТОР 10 mg/10 mg филмирани таблетки	HU/H/0248/002	20110296	GEDEON RICHTER PLC.	BG
Duplecor 10 mg/10 mg apvalkotās tabletes	HU/H/0248/002	11-0286	GEDEON RICHTER PLC.	LV
Duplecor,10 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0248/002	740311	GEDEON RICHTER PLC.	EE
Duplecor, 10 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0248/001	740211	GEDEON RICHTER PLC.	EE
Duplecor 10 mg/5 mg apvalkotās tabletes	HU/H/0248/001	11-0287	GEDEON RICHTER PLC.	LV
Duplecor 20 mg/10 mg apvalkotās tabletes	HU/H/0248/004	11-0288	GEDEON RICHTER PLC.	LV
Duplecor, 20 mg/5 mg õhukese polümeerikattega tabletid	HU/H/0248/003	740511	GEDEON RICHTER PLC.	EE
Duplecor 20 mg/5 mg apvalkotās tabletes	HU/H/0248/003	11-0289	GEDEON RICHTER PLC.	LV
Duplecor, 20 mg/10 mg õhukese polümeerikattega tabletid	HU/H/0248/004	740411	GEDEON RICHTER PLC.	EE
АМЛАТОР 10 mg/5 mg филмирани таблетки	HU/H/0248/001	20110295	GEDEON RICHTER PLC.	BG
АМЛАТОР 20 mg/10 mg	HU/H/0248/004	20110298	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
филмирани таблетки				
Amlator 10 mg/5 mg potahované tablety	HU/H/0248/001	58/480/11-C	GEDEON RICHTER PLC.	CZ
Amlator, 20 mg + 10 mg, tabletki powlekane	HU/H/0248/004	18576	GEDEON RICHTER PLC.	PL
Suvarol 5 mg / 10 mg apvalkotās tabletes	PL/H/0399/001	16-0257	WARSZAWSKIE ZAKLADY FARMACEUTYCZNE POLFA S.A.	LV
Suvarol 10 mg / 10 mg apvalkotās tabletes	PL/H/0399/002	16-0258	WARSZAWSKIE ZAKLADY FARMACEUTYCZNE POLFA S.A.	LV
Suvarol 5 mg / 20 mg apvalkotās tabletes	PL/H/0399/003	16-0259	WARSZAWSKIE ZAKLADY FARMACEUTYCZNE POLFA S.A.	LV
Suvarol 10 mg / 20 mg apvalkotās tabletes	PL/H/0399/004	16-0260	WARSZAWSKIE ZAKLADY FARMACEUTYCZNE POLFA S.A.	LV
Amlodipine + Atorvastatin Polpharma, 5 mg + 10 mg, tabletki powlekane	PL/H/0399/001	23797	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Amlodipine + Atorvastatin Polpharma, 10 mg + 10 mg, tabletki powlekane	PL/H/0399/002	23798	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Amlodipine + Atorvastatin Polpharma, 5 mg + 20 mg, tabletki powlekane	PL/H/0399/003	23799	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Amlodipine + Atorvastatin Polpharma, 10 mg + 20 mg, tabletki powlekane	PL/H/0399/004	23800	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Флодил 5 mg/10 mg филмирани таблетки	PL/H/0399/001	20170153	MEDANA PHARMA SPOLKA AKCYJNA	BG
Флодил 10 mg/10 mg филмирани таблетки	PL/H/0399/002	20170154	MEDANA PHARMA SPOLKA AKCYJNA	BG
Флодил 5 mg/20 mg филмирани таблетки	PL/H/0399/003	20170155	MEDANA PHARMA SPOLKA AKCYJNA	BG
Флодил 10 mg/20 mg филмирани таблетки	PL/H/0399/004	20170156	MEDANA PHARMA SPOLKA AKCYJNA	BG
Astucor 10mg/10mg comprimidos recubiertos con	not available	67.593	ALMIRALL, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
película				
Astucor 5mg/10mg comprimidos recubiertos con película	not available	67.594	ALMIRALL, S.A.	ES
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582614	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582606	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582622	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582630	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582648	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582655	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582663	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582838	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582671	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 10 mg comprimidos revestidos por película	PT/H/1713/001	5582705	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por	PT/H/1713/003	5582721	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
película				
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582713	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582739	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582747	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582754	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582762	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582770	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582820	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582804	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 5 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582812	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582853	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582846	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg	PT/H/1713/003	5582861	TETRAFARMA- PRODUTOS	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 revestidos por película			FARMACÊUTICOS, LDA.	
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582879	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582903	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582911	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582929	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582937	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582945	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Trinalion 10 mg + 20 mg comprimidos revestidos por película	PT/H/1713/003	5582952	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Lidorat 10 mg/5 mg Filmtabletten	HU/H/0320/001	1-31754	GEDEON RICHTER PLC.	AT
Lidorat 20mg/5mg Filmtabletten	HU/H/0320/003	1-31755	GEDEON RICHTER PLC.	AT
Lidorat 20 mg/10 mg Filmtabletten	HU/H/0320/004	1-31756	GEDEON RICHTER PLC.	AT
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/01	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/02	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/03	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/04	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/05	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/06	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/07	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
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/08	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/09	GEDEON RICHTER PLC.	HU
Lidorat 10 mg/5 mg filmtabletta	HU/H/0320/001	OGYI-T-22308/10	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/11	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/12	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/13	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/14	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/15	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/16	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/17	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/18	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/19	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/5 mg filmtabletta	HU/H/0320/003	OGYI-T-22308/20	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/21	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/22	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/23	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/24	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/25	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/26	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/27	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/28	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/29	GEDEON RICHTER PLC.	HU
Lidorat 20 mg/10 mg filmtabletta	HU/H/0320/004	OGYI-T-22308/30	GEDEON RICHTER PLC.	HU