14 April 2023
EMA/327585/2023
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / indapamide, amlodipine / indapamide / perindopril

Procedure no.: PSUSA/00010358/202211

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact Telephone +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PERAMIND 4 mg/1,25 mg/5 mg compresse	IT/H/0877/001	049310016	DOC GENERICI S.R.L.	IT
PERAMIND 4 mg/1,25 mg/10 mg compresse	IT/H/0877/002	049310028	DOC GENERICI S.R.L.	IT
PERAMIND 8 mg/2,5 mg/5 mg compresse	IT/H/0877/003	049310030	DOC GENERICI S.R.L.	IT
PERAMIND 8 mg/2,5 mg/10 mg compresse	IT/H/0877/004	049310042	DOC GENERICI S.R.L.	IT
Co-Amlessa 2 mg + 0,625 mg + 5 mg comprimidos	not available	5650676	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 2 mg + 0,625 mg + 5 mg comprimidos	not available	5650700	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 2 mg + 0,625 mg + 5 mg comprimidos	not available	5650718	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 5 mg comprimidos	not available	5650726	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 5 mg comprimidos	not available	5650734	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 5 mg comprimidos	not available	5650742	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 10 mg comprimidos	not available	5650833	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 10 mg comprimidos	not available	5650841	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 4 mg + 1,25 mg + 10 mg comprimidos	not available	5650858	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 8 mg + 2,5	not available	5650866	KRKA, D.D., NOVO MESTO	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg + 5 mg comprimidos				
Co-Amlessa 8 mg + 2,5 mg + 5 mg comprimidos	not available	5650874	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 8 mg + 2,5 mg + 5 mg comprimidos	not available	5650908	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 8 mg + 2,5 mg + 10 mg comprimidos	not available	5650957	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 8 mg + 2,5 mg + 10 mg comprimidos	not available	5650965	KRKA, D.D., NOVO MESTO	PT
Co-Amlessa 8 mg + 2,5 mg + 10 mg comprimidos	not available	5650973	KRKA, D.D., NOVO MESTO	PT
Co-Dalneva 8 mg/2,5 mg/10 mg tablete	not available	HR-H-180076502	KRKA-FARMA D.O.O.	HR
Co-Dalneva 8 mg/2,5 mg/5 mg tablete	not available	HR-H-209243724	KRKA-FARMA D.O.O.	HR
Co-Dalneva 4 mg/1,25 mg/5 mg tablete	not available	HR-H-223376332	KRKA-FARMA D.O.O.	HR
Co-Dalneva 4 mg/1,25 mg/10 mg tablete	not available	HR-H-622847904	KRKA-FARMA D.O.O.	HR
Viacorlix 7 mg/5 mg/2,5 mg comprimate filmate	NL/H/3968/001	14832/2022/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacorlix 7 mg/5 mg/2,5 mg comprimate filmate	NL/H/3968/001	14832/2022/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacorlix 7 mg/5 mg/2,5 mg comprimate filmate	NL/H/3968/001	14832/2022/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacorlix 7 mg/5 mg/2,5 mg comprimate filmate	NL/H/3968/001	14832/2022/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Viacorlix 7 mg/5 mg/2,5 mg comprimate filmate	NL/H/3968/001	14832/2022/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRICORLIX 7 mg/5	NL/H/3968/001/DC	34009 301 452 0 3	LES LABORATOIRES	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
mg/2,5 mg, comprimé pelliculé			SERVIER (SURESNES)	
TRICORLIX 7 mg/5 mg/2,5 mg, comprimé pelliculé	NL/H/3968/001/DC	34009 550 543 8 2	LES LABORATOIRES SERVIER (SURESNES)	FR
TRICORLIX 7 mg/5 mg/2,5 mg, comprimé pelliculé	NL/H/3968/001/DC	34009 550 543 9 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRICORLIX 7 mg/5 mg/2,5 mg, comprimé pelliculé	NL/H/3968/001/DC	34009 301 451 9 7	LES LABORATOIRES SERVIER (SURESNES)	FR
TRICORLIX 7 mg/5 mg/2,5 mg, comprimé pelliculé	NL/H/3968/001/DC	34009 550 544 0 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Tricorlix 7 mg/5 mg/2.5 mg apvalkotās tabletes	NL/H/3968/001/DC	18-0074	LES LABORATOIRES SERVIER (SURESNES)	LV
VACORIND 7,5 mg/5 mg/2,5 mg Filmtabletten	NL/H/3968/001/DC	99565.00.00	LES LABORATOIRES SERVIER (SURESNES)	DE
Viacorlix 7 mg/5 mg/2,5 mg comprimidos recubiertos con película	NL/H/3968/001/DC	83.231	LES LABORATOIRES SERVIER (SURESNES)	ES
Tricorlix 7 mg/5 mg/2,5 mg filmomhulde tabletten	NL/H/3968/001/DC	RVG 120767	LES LABORATOIRES SERVIER (SURESNES)	NL
Prindal 4 mg/1,25 mg/10 mg tabletes	CZ/H/0998/002	21-0128	ZENTIVA, K.S.	LV
Prindal 4 mg/1,25 mg/5 mg tabletes	CZ/H/0998/001	21-0127	ZENTIVA, K.S.	LV
Prindal 8 mg/2,5 mg/5 mg tabletes	CZ/H/0998/003	21-0129	ZENTIVA, K.S.	LV
Prindal 8 mg/2,5 mg/10 mg tabletes	CZ/H/0998/004	21-0130	ZENTIVA, K.S.	LV
Perindopril/Indapamid/A	CZ/H/0998/002	58/044/20-C	ZENTIVA, K.S.	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
mlodipin Zentiva 4 mg/1,25 mg/10 mg tablety				
Perindopril/Indapamid/Amlodipin Zentiva 4 mg/1,25 mg/5 mg tablety	CZ/H/0998/001	58/043/20-C	ZENTIVA, K.S.	CZ
Perindopril/Indapamid/Amlodipin Zentiva 8 mg/2,5 mg/5 mg tablety	CZ/H/0998/003	58/045/20-C	ZENTIVA, K.S.	CZ
Perindopril/Indapamid/Amlodipin Zentiva 8 mg/2,5 mg/10 mg tablety	CZ/H/0998/004	58/046/20-C	ZENTIVA, K.S.	CZ
Prindal 4 mg/1,25 mg/10 mg tabletid	CZ/H/0998/002	1040721	ZENTIVA, K.S.	EE
Prindal 4 mg/1,25 mg/5 mg tablettid	CZ/H/0998/001	1040621	ZENTIVA, K.S.	EE
Prindal 8 mg/2,5 mg/5 mg tabletid	CZ/H/0998/003	1040821	ZENTIVA, K.S.	EE
Prindal 8 mg/2,5 mg/10 mg tabletid	CZ/H/0998/004	1040921	ZENTIVA, K.S.	EE
Prindal 4 mg/1,25 mg/5 mg comprimate	CZ/H/0998/001	14129/2021/01	ZENTIVA, K.S.	RO
Prindal 4 mg/1,25 mg/5 mg comprimate	CZ/H/0998/001	14129/2021/02	ZENTIVA, K.S.	RO
Prindal 4 mg/1,25 mg/5 mg comprimate	CZ/H/0998/001	14129/2021/03	ZENTIVA, K.S.	RO
Prindal 4 mg/1,25 mg/10 mg comprimate	CZ/H/0998/002	14130/2021/01	ZENTIVA, K.S.	RO
Prindal 4 mg/1,25 mg/10 mg comprimate	CZ/H/0998/002	14130/2021/02	ZENTIVA, K.S.	RO
Prindal 4 mg/1,25 mg/10	CZ/H/0998/002	14130/2021/03	ZENTIVA, K.S.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg comprimate				
Prindal 8 mg/2,5 mg/5 mg comprimate	CZ/H/0998/003	14131/2021/01	ZENTIVA, K.S.	RO
Prindal 8 mg/2,5 mg/5 mg comprimate	CZ/H/0998/003	14131/2021/02	ZENTIVA, K.S.	RO
Prindal 8 mg/2,5 mg/5 mg comprimate	CZ/H/0998/003	14131/2021/03	ZENTIVA, K.S.	RO
Prindal 8 mg/2,5 mg/10 mg comprimate	CZ/H/0998/004	14132/2021/01	ZENTIVA, K.S.	RO
Prindal 8 mg/2,5 mg/10 mg comprimate	CZ/H/0998/004	14132/2021/02	ZENTIVA, K.S.	RO
Prindal 8 mg/2,5 mg/10 mg comprimate	CZ/H/0998/004	14132/2021/03	ZENTIVA, K.S.	RO
Prindal 4 mg + 1,25 mg + 10 mg comprimidos	CZ/H/0998/002	5823851	ZENTIVA PORTUGAL, LDA	PT
Prindal 4 mg + 1,25 mg + 10 mg comprimidos	CZ/H/0998/002	5823869	ZENTIVA PORTUGAL, LDA	PT
Prindal 8 mg + 2,5 mg + 5 mg comprimidos	CZ/H/0998/003	5823877	ZENTIVA PORTUGAL, LDA	PT
Prindal 8 mg + 2,5 mg + 5 mg comprimidos	CZ/H/0998/003	5823901	ZENTIVA PORTUGAL, LDA	PT
Prindal 8 mg + 2,5 mg + 10 mg comprimidos	CZ/H/0998/004	5823919	ZENTIVA PORTUGAL, LDA	PT
Prindal 8 mg + 2,5 mg + 10 mg comprimidos	CZ/H/0998/004	5823927	ZENTIVA PORTUGAL, LDA	PT
Prindal 4 mg + 1,25 mg + 5 mg comprimidos	CZ/H/0998/001	5823836	ZENTIVA PORTUGAL, LDA	PT
Prindal 4 mg + 1,25 mg + 5 mg comprimidos	CZ/H/0998/001	5823844	ZENTIVA PORTUGAL, LDA	PT
NATRIXAM 1,5mg/5mg, comprimé à libération modifiée	NL/H/2637/01/DC	34009 585 686 9 5	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/5mg,	NL/H/2637/01/DC	34009 585 687 5 6	LES LABORATOIRES	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
comprimé à libération modifiée			SERVIER (SURESNES)	
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 275 985 8 6	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 275 984 1 8	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 275 983 5 7	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5 mg/10 mg, comprimé à libération modifiée	NL/H/2637/002	34009 275 982 9 6	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5 mg/10 mg, comprimé à libération modifiée	NL/H/2637/002	34009 585 691 2 8	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 585 690 6 7	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/5mg, comprimé à libération modifiée	NL/H/2637/01/DC	34009 275 977 5 6	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/5mg, comprimé à libération modifiée	NL/H/2637/01/DC	34009 275 976 9 5	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/5mg, comprimé à libération modifiée	NL/H/2637/01/DC	34009 275 975 2 7	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/5mg, comprimé à libération modifiée	NL/H/2637/01/DC	34009 275 974 6 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Tertensam 1,5 mg / 10	NL/H/2637/02/DC	13-0247	LES LABORATOIRES	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
mg ilgstošās darbības tabletes			SERVIER (SURESNES)	
Tertensam 1,5 mg / 5 mg ilgstošās darbības tabletes	NL/H/2637/01/DC	13-0246	LES LABORATOIRES SERVIER (SURESNES)	LV
NATRIXAM 1,5 mg / 10 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/007	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 10 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/008	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/004	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 10 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/012	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/001	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/002	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/005	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/003	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 10 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/009	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 10	NL/H/2637/02/DC	LT/1/13/3427/011	LES LABORATOIRES	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
mg modifikuoto atpalaidavimo tabletės			SERVIER (SURESNES)	
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/01/DC	LT/1/13/3427/006	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 10 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/010	LES LABORATOIRES SERVIER (SURESNES)	LT
NADEXAM 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/02/DC	H/13/01074/011	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/002	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/005	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/001	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/006	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/02/DC	H/13/01074/012	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/004	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/01/DC	H/13/01074/003	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/10 mg	NL/H/2637/02/DC	H/13/01074/008	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
tablete s prirejenim sproščanjem				
NADEXAM 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/02/DC	H/13/01074/007	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/02/DC	H/13/01074/010	SERVIER PHARMA D.O.O	SI
NADEXAM 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/02/DC	H/13/01074/009	SERVIER PHARMA D.O.O	SI
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg, comprimé à libération modifiée	NL/H/2637/02/DC	2014120415	LES LABORATOIRES SERVIER (SURESNES)	LU
NATRIXAM 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/01/DC	2014120414	LES LABORATOIRES SERVIER (SURESNES)	LU
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2637/002	13491/2020/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2637/002	13491/2020/04	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
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2637/002	13491/2020/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2637/002	13491/2020/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2637/001	13490/2020/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2637/002	13491/2020/02	LES LABORATOIRES SERVIER (SURESNES)	RO
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-03	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-04	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-02	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
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-06	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-05	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/01/DC	HR-H-053397374-01	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-03	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-01	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-02	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-05	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-06	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/02/DC	HR-H-921621992-04	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
Натриксам 1,5 mg / 10 mg таблетки с изменено освобождаване	NL/H/2637/002	20130382	LES LABORATOIRES SERVIER (SURESNES)	BG
Натриксам 1,5 mg / 5 mg таблетки с изменено освобождаване	NL/H/2637/001	20130381	LES LABORATOIRES SERVIER (SURESNES)	BG
NATRIXAM 1,5 mg / 10 mg, tabletten met gereguleerde afgifte	NL/H/2637/02/DC	RVG 112320	LES LABORATOIRES SERVIER (SURESNES)	NL
Natrixam 1.5 mg / 10 mg modified-release tablets	NL/H/2637/02/DC	MA066/02102	LES LABORATOIRES SERVIER (SURESNES)	MT
Natrixam, 1,5 mg/5 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2637/01/DC	825313	LES LABORATOIRES SERVIER (SURESNES)	EE
NATRIXAM 1,5 mg / 5 mg, tabletten met gereguleerde afgifte	NL/H/2637/01/DC	RVG 112319	LES LABORATOIRES SERVIER (SURESNES)	NL
Natrixam 1.5 mg / 5 mg modified-release tablets	NL/H/2637/01/DC	MA066/02101	LES LABORATOIRES SERVIER (SURESNES)	MT
Natrixam, 1,5 mg/10 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2637/02/DC	825213	LES LABORATOIRES SERVIER (SURESNES)	EE
Natrixam 1,5 mg/10 mg tablety s riadeným uvoľňovaním	NL/H/2637/02/DC	58/0435/13-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Natrixam 1,5 mg/5 mg tablety s riadeným uvoľňovaním	NL/H/2637/01/DC	58/0434/13-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Tertens-AM, 1,5 mg + 5 mg, tabletki o	NL/H/2637/01/DC	21557	LES LABORATOIRES SERVIER (SURESNES)	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
zmodyfikowanym uwalnianiu				
Tertens-AM, 1,5 mg + 10 mg, tabletki o zmodyfikowanym uwalnianiu	NL/H/2637/02/DC	21558	LES LABORATOIRES SERVIER (SURESNES)	PL
Arplexam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2638/05/DC	RVG 112143	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2638/04/DC	RVG 112142	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2638/03/DC	RVG 112141	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2638/02/DC	RVG 112140	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 10 mg/ 2,5 mg/ 5 mg apvalkotās tabletes	NL/H/2638/04/DC	14-0054	LES LABORATOIRES SERVIER (SURESNES)	LV
Arplexam 5 mg/ 1,25 mg/ 10 mg apvalkotās tabletes	NL/H/2638/03/DC	14-0053	LES LABORATOIRES SERVIER (SURESNES)	LV
Arplexam 10 mg/ 2,5 mg/ 10 mg apvalkotās tabletes	NL/H/2638/05/DC	14-0055	LES LABORATOIRES SERVIER (SURESNES)	LV
Covercard Plus 5 mg/1,25 mg/5 mg filmtabletta	NL/H/2638/001-005/DC	OGYI-T-22626/02	EGIS PHARMACEUTICALS PLC	HU
Covercard Plus 10 mg/2,5 mg/5 mg filmtabletta	NL/H/2638/001-005/DC	OGYI-T-22626/04	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 Plus 5 mg/1,25 mg/10 mg filmtabletta	NL/H/2638/001-005/DC	OGYI-T-22626/03	EGIS PHARMACEUTICALS PLC	HU
Covercard Plus 10 mg/2,5 mg/10 mg filmtabletta	NL/H/2638/001-005/DC	OGYI-T-22626/05	EGIS PHARMACEUTICALS PLC	HU
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/007	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/012	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/022	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/017	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/025	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/021	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/023	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/006	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/010	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
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/016	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/008	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/015	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/018	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/004	LT/1/14/3519/020	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/013	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/011	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/030	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/038	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/024	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2638/05/DC	LT/1/14/3519/037	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
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/029	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/036	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/035	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2638/04/DC	LT/1/14/3519/019	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/028	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/033	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/034	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/031	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2638/03/DC	LT/1/14/3519/014	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/009	LES LABORATOIRES SERVIER (SURESNES)	LT
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/032	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
ARPLEXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2638/02/DC	LT/1/14/3519/027	LES LABORATOIRES SERVIER (SURESNES)	LT
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	11977/2019/07	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	11976/2019/07	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/05/DC	11977/2019/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/05/DC	11977/2019/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	11977/2019/08	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	11977/2019/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/05/DC	11977/2019/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/10 mg comprimate	NL/H/2638/05/DC	11977/2019/04	ANPHARM PRZEDSIĘBIORSTWO	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
filmate			FARMACEUTYCZNE SPÓŁKA AKCYJNA	
NORPLEXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/05/DC	11977/2019/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/04/DC	11976/2019/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	11976/2019/08	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/04/DC	11976/2019/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/04/DC	11976/2019/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/04/DC	11976/2019/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	11976/2019/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/04/DC	11976/2019/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA	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
			AKCYJNA	
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	11975/2019/07	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	11974/2019/07	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/03/DC	11975/2019/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/03/DC	11975/2019/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	11975/2019/08	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/03/DC	11975/2019/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/03/DC	11975/2019/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/03/DC	11975/2019/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	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
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	11975/2019/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/02/DC	11974/2019/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	11974/2019/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/02/DC	11974/2019/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	11974/2019/08	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/02/DC	11974/2019/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/02/DC	11974/2019/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
NORPLEXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/02/DC	11974/2019/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
ARPLEXAM	NL/H/2638/02/DC	73478/07-06-2019	SERVIER HELLAS	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
5mg/1,25mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία			PHARMACEUTICALS LTD	
ARPLEXAM 5mg/1,25mg/10mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/03/DC	73479/07-06-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10mg/2,5mg/10mg, επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/05/DC	73481/07-06-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10mg/2,5mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/04/DC	73480/07-06-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Arplexam 5 mg/ 1,25 mg/ 5 mg apvalkotās tabletes	NL/H/2638/02/DC	14-0052	LES LABORATOIRES SERVIER (SURESNES)	LV
Lopridam 4 mg/1,25 mg/10 mg tablety	CZ/H/0699/002	58/0095/18-S	ZENTIVA, K.S.	SK
Lopridam 8 mg/2,5 mg/10 mg tablety	CZ/H/0699/004	58/0097/18-S	ZENTIVA, K.S.	SK
Lopridam 4 mg/1,25 mg/5 mg tablety	CZ/H/0699/001	58/0094/18-S	ZENTIVA, K.S.	SK
Lopridam 8 mg/2,5 mg/5 mg tablety	CZ/H/0699/003	58/0096/18-S	ZENTIVA, K.S.	SK
Приндал 4 mg / 1,25 mg / 5 mg таблетки	CZ/H/0699/001	20180099	ZENTIVA, K.S.	BG
Приндал 8 mg / 2,5 mg / 5 mg таблетки	CZ/H/0699/003	20180100	ZENTIVA, K.S.	BG
Lopridam 8 mg/2,5 mg/10 mg tablety	CZ/H/0699/004	58/1016/16-C	ZENTIVA, K.S.	CZ
Lopridam 8 mg/2,5 mg/5	CZ/H/0699/003	58/1015/16-C	ZENTIVA, K.S.	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
mg tablety				
Lopridam 4 mg/1,25 mg/10 mg tablety	CZ/H/0699/002	58/1014/16-C	ZENTIVA, K.S.	CZ
Lopridam 4 mg/1,25 mg/5 mg tablety	CZ/H/0699/001	58/1013/16-C	ZENTIVA, K.S.	CZ
Lopridam 4 mg/ 1,25 mg/ 5 mg compresse	CZ/H/0699/001	045549019	ZENTIVA ITALIA S.R.L.	IT
Lopridam 4 mg/ 1,25 mg/ 5 mg compresse	CZ/H/0699/001	045549021	ZENTIVA ITALIA S.R.L.	IT
Lopridam 4 mg/ 1,25 mg/ 5 mg compresse	CZ/H/0699/001	045549033	ZENTIVA ITALIA S.R.L.	IT
Lopridam 4 mg/ 1,25 mg/ 10 mg compresse	CZ/H/0699/002	045549045	ZENTIVA ITALIA S.R.L.	IT
Lopridam 4 mg/ 1,25 mg/ 10 mg compresse	CZ/H/0699/002	045549058	ZENTIVA ITALIA S.R.L.	IT
Lopridam 4 mg/ 1,25 mg/ 10 mg compresse	CZ/H/0699/002	045549060	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 5 mg compresse	CZ/H/0699/003	045549072	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 5 mg compresse	CZ/H/0699/003	045549084	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 5 mg compresse	CZ/H/0699/003	045549096	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 10 mg compresse	CZ/H/0699/004	045549108	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 10 mg compresse	CZ/H/0699/004	045549110	ZENTIVA ITALIA S.R.L.	IT
Lopridam 8 mg/ 2,5 mg/ 10 mg compresse	CZ/H/0699/004	045549122	ZENTIVA ITALIA S.R.L.	IT
Tonanda 2 mg/5 mg/0,625 mg tablety	HU/H/0342/001	58/346/14-C	KRKA, D.D., NOVO MESTO	CZ
Tonanda 4 mg/5 mg/1,25 mg tablety	HU/H/0342/002	58/347/14-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
Tonanda 4 mg/10 mg/1,25 mg tablety	HU/H/0342/003	58/348/14-C	KRKA, D.D., NOVO MESTO	CZ
Tonanda 8 mg/5 mg/2,5 mg tablety	HU/H/0342/004	58/349/14-C	KRKA, D.D., NOVO MESTO	CZ
Tonanda 8 mg/10 mg/2,5 mg tablety	HU/H/0342/005	58/350/14-C	KRKA, D.D., NOVO MESTO	CZ
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/001	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/010	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/019	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/028	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/037	KRKA, D.D., NOVO MESTO	SI
Co-Amlessa 4 mg+5 mg+1,25 mg tabletes	HU/H/0342/002	14-0026	KRKA, D.D., NOVO MESTO	LV
Co-Amlessa 4 mg+10 mg+1,25 mg tabletes	HU/H/0342/003	14-0027	KRKA, D.D., NOVO MESTO	LV
Co-Amlessa 8 mg+5 mg+2,5 mg tabletes	HU/H/0342/004	14-0028	KRKA, D.D., NOVO MESTO	LV
Co-Amlessa 8 mg+10 mg+2,5 mg tabletes	HU/H/0342/005	14-0029	KRKA, D.D., NOVO MESTO	LV
Ко-Амлеса 2 mg/5 mg/0,625 mg таблетки	HU/H/0342/001	20140007	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 4 mg/5 mg/1,25 mg таблетки	HU/H/0342/002	20140008	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 4 mg/10 mg/1,25 mg таблетки	HU/H/0342/003	20140009	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 8 mg/5 mg/2,5 mg таблетки	HU/H/0342/004	20140010	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 8 mg/10	HU/H/0342/005	20140011	KRKA, D.D., NOVO MESTO	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
mg/2,5 mg таблетки				
Co-Amlessa, 2 mg + 5 mg + 0,625 mg, tabletki	HU/H/0342/001	21639	KRKA, D.D., NOVO MESTO	PL
Co-Amlessa, 4 mg + 5 mg + 1,25 mg, tabletki	HU/H/0342/002	21640	KRKA, D.D., NOVO MESTO	PL
Co-Amlessa, 4 mg + 10 mg + 1,25 mg, tabletki	HU/H/0342/003	21641	KRKA, D.D., NOVO MESTO	PL
Co-Amlessa, 8 mg + 5 mg + 2,5 mg, tabletki	HU/H/0342/004	21642	KRKA, D.D., NOVO MESTO	PL
Co-Amlessa, 8 mg + 10 mg + 2,5 mg, tabletki	HU/H/0342/005	21643	KRKA, D.D., NOVO MESTO	PL
Co-Amlessa 2 mg/5 mg/0,625 mg tablety	HU/H/0342/001	58/0445/13-S	KRKA, D.D., NOVO MESTO	SK
Co-Amlessa 4 mg/5 mg/1,25 mg tablety	HU/H/0342/002	58/0446/13-S	KRKA, D.D., NOVO MESTO	SK
Co-Amlessa 4 mg/10 mg/1,25 mg tablety	HU/H/0342/003	58/0447/13-S	KRKA, D.D., NOVO MESTO	SK
Co-Amlessa 8 mg/5 mg/2,5 mg tablety	HU/H/0342/004	58/0448/13-S	KRKA, D.D., NOVO MESTO	SK
Co-Amlessa 8 mg/10 mg/2,5 mg tablety	HU/H/0342/005	58/0449/13-S	KRKA, D.D., NOVO MESTO	SK
Co Dalnessa 4 mg/10 mg/1,25 mg tabletid	HU/H/0342/003	832813	KRKA, D.D., NOVO MESTO	EE
Co Dalnessa 2 mg/5 mg/0,625 mg tabletid	HU/H/0342/001	833013	KRKA, D.D., NOVO MESTO	EE
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletid	HU/H/0342/002	833113	KRKA, D.D., NOVO MESTO	EE
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletid	HU/H/0342/005	833213	KRKA, D.D., NOVO MESTO	EE
Co Dalnessa 8 mg/5 mg/2,5 mg tabletid	HU/H/0342/004	833313	KRKA, D.D., NOVO MESTO	EE
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/002	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
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/003	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/004	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/005	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/006	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/007	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/008	KRKA, D.D., NOVO MESTO	SI
Amlewel 2 mg/5 mg/0,625 mg tablete	HU/H/0342/001	H/14/01901/009	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/011	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/012	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/013	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/014	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/015	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/016	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/017	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/5 mg/1,25 mg tablete	HU/H/0342/002	H/14/01901/018	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/020	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10	HU/H/0342/003	H/14/01901/021	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
mg/1,25 mg tablete				
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/022	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/023	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/024	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/025	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/026	KRKA, D.D., NOVO MESTO	SI
Amlewel 4 mg/10 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/027	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/029	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/030	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/031	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/032	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/033	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/034	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/035	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/5 mg/2,5 mg tablete	HU/H/0342/004	H/14/01901/036	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/038	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/039	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
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/040	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/041	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/042	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/043	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/044	KRKA, D.D., NOVO MESTO	SI
Amlewel 8 mg/10 mg/2,5 mg tablete	HU/H/0342/005	H/14/01901/045	KRKA, D.D., NOVO MESTO	SI
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/001	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/002	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/003	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/004	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/005	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/006	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/007	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/008	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 2 mg/5 mg/0,625 mg tabletės	HU/H/0342/001	LT/1/13/3462/009	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/010	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5	HU/H/0342/002	LT/1/13/3462/011	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
mg/1,25 mg tabletės				
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/012	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/013	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/014	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/015	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/016	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/017	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/5 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/018	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/019	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/020	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/021	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/022	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/023	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/024	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/025	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/026	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 4 mg/10 mg/1,25 mg tabletės	HU/H/0342/003	LT/1/13/3462/027	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
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/028	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/029	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/030	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/031	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/032	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/033	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/034	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/035	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/5 mg/2,5 mg tabletės	HU/H/0342/004	LT/1/13/3462/036	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/037	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/038	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/039	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/040	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/041	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/042	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/043	KRKA, D.D., NOVO MESTO	LT
Co-Amlessa 8 mg/10	HU/H/0342/005	LT/1/13/3462/044	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
mg/2,5 mg tabletės				
Co-Amlessa 8 mg/10 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/045	KRKA, D.D., NOVO MESTO	LT
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/01	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/02	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/03	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/04	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/05	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/06	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/07	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/08	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 2 mg/5 mg/0,625 mg tabletta	HU/H/0342/001	OGYI-T-22577/09	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/10	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/11	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/12	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/13	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/14	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/15	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
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/16	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/17	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/5 mg/1,25 mg tabletta	HU/H/0342/002	OGYI-T-22577/18	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/19	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/20	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/21	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/22	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/23	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/24	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/25	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/26	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 4 mg/10 mg/1,25 mg tabletta	HU/H/0342/003	OGYI-T-22577/27	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/28	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/29	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/30	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/31	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5	HU/H/0342/004	OGYI-T-22577/32	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
mg/2,5 mg tabletta				
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/33	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/34	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/35	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/36	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/37	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/38	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/39	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/40	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/41	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/42	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/43	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/44	KRKA, D.D., NOVO MESTO	HU
Co-Dalnessa 8 mg/10 mg/2,5 mg tabletta	HU/H/0342/005	OGYI-T-22577/45	KRKA, D.D., NOVO MESTO	HU
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/02	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
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	11463/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/06	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	11464/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	11465/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5	HU/H/0342/004	11466/2019/05	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/2,5 mg comprimate				
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	11466/2019/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	11467/2019/09	KRKA, D.D., NOVO MESTO	RO
Cardilopin Komb 1,5 mg/5 mg módosított hatóanyagleadású tabletta	NL/H/2639/001-002/DC	OGYI-T-22562/01	EGIS PHARMACEUTICALS PLC	HU
Cardilopin Komb 1,5 mg/10 mg módosított	NL/H/2639/001-002/DC	OGYI-T-22562/02	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
hatóanyagleadású tabletta				
FLUTENSIF 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2639/02/DC	5615943	LES LABORATOIRES SERVIER (SURESNES)	PT
Flutensif 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2639/001	5594171	LES LABORATOIRES SERVIER (SURESNES)	PT
Flutensif 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2639/002	5594239	LES LABORATOIRES SERVIER (SURESNES)	PT
FLUTENSIF 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2639/02/DC	5594221	LES LABORATOIRES SERVIER (SURESNES)	PT
Flutensif 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2639/001	5594213	LES LABORATOIRES SERVIER (SURESNES)	PT
FLUTENSIF 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2639/01/DC	5594205	LES LABORATOIRES SERVIER (SURESNES)	PT
FLUTENSIF 1,5 mg / 10 mg, tabletten met gereguleerde afgifte	NL/H/2639/02/DC	RVG 112322	LES LABORATOIRES SERVIER (SURESNES)	NL
Flutensif, 1,5 mg/10 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2639/02/DC	824413	LES LABORATOIRES SERVIER (SURESNES)	EE
Flutensif, 1,5 mg/5 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2639/01/DC	824513	LES LABORATOIRES SERVIER (SURESNES)	EE
FLUTENSIF 1,5 mg / 5 mg, tabletten met	NL/H/2639/01/DC	RVG 112321	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
gereguleerde afgifte				
FLUDEXAM 1,5 mg/5 mg Tabletten mit veränderter Wirkstofffreisetzung	NL/H/2639/01/DC	135239	LES LABORATOIRES SERVIER (SURESNES)	AT
Fludexam 1,5 mg/10 mg Tabletten mit veränderter Wirkstofffreisetzung	NL/H/2639/002	135240	LES LABORATOIRES SERVIER (SURESNES)	AT
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/01	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/02	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/03	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/04	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/05	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/06	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/07	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/08	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22622/09	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/10	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/11	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/12	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
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/13	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/14	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/15	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/16	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/17	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22622/18	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/19	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/20	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/21	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/22	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/23	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/24	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/25	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/26	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22622/27	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/28	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5	not available	OGYI-T-22622/29	KRKA, D.D., NOVO MESTO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/2,5 mg tabletta				
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/30	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/31	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/32	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/33	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/34	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/35	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22622/36	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/37	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/38	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/39	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/40	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/41	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/42	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/43	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/44	KRKA, D.D., NOVO MESTO	HU
Dalnecombi 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22622/45	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
Triplixam 10 mg/2,5 mg/10 mg õhukese polümeerikattega tabletid	NL/H/2636/05/DC	835714	LES LABORATOIRES SERVIER (SURESNES)	EE
Triplixam, 5 mg/1,25 mg/10 mg õhukese polümeerikattega tabletid	NL/H/2636/003	834914	LES LABORATOIRES SERVIER (SURESNES)	EE
Triplixam 5 mg/1,25 mg/5 mg õhukese polümeerikattega tabletid	NL/H/2636/02/DC	835014	LES LABORATOIRES SERVIER (SURESNES)	EE
Triplixam 10mg/2,5mg/5mg õhukese polümeerikattega tabletid	NL/H/2636/04/DC	835314	LES LABORATOIRES SERVIER (SURESNES)	EE
TRIPLIXAM 10mg/2.5mg/5mg film-coated tablets	NL/H/2636/04/DC	MA066/02004	LES LABORATOIRES SERVIER (SURESNES)	MT
TRIPLIXAM 10mg/2.5mg/10mg film-coated tablets	NL/H/2636/05/DC	MA066/02005	LES LABORATOIRES SERVIER (SURESNES)	MT
TRIPLIXAM 5mg/1.25mg/10mg film-coated tablets	NL/H/2636/03/DC	MA066/02003	LES LABORATOIRES SERVIER (SURESNES)	MT
TRIPLIXAM 5mg/1.25mg/5mg film-coated tablets	NL/H/2636/02/DC	MA066/02002	LES LABORATOIRES SERVIER (SURESNES)	MT
TRIPLIXAM 10 mg/2,5 mg/5 mg potahované tablety	NL/H/2636/04/DC	58/102/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
TRIPLIXAM 10 mg/2,5	NL/H/2636/05/DC	58/103/14-C	LES LABORATOIRES	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
mg/10 mg potahované tablety			SERVIER (SURESNES)	
TRIPLIXAM 5 mg/1,25 mg/10 mg potahované tablety	NL/H/2636/03/DC	58/101/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
TRIPLIXAM 5 mg/1,25 mg/5 mg potahované tablety	NL/H/2636/02/DC	58/100/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Triplixam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2636/04/DC	RVG 112147	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2636/02/DC	RVG 112145	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/05/DC	RVG 112148	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2636/03/DC	RVG 112146	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/05/DC	BE448684	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/5 mg, comprimés pelliculés	NL/H/2636/04/DC	BE448675	SERVIER BENELUX S.A./N.V.	BE
COVERDINE 10mg/2.5mg/5mg film-coated tablets	NL/H/2636/04/DC	PA0568/024/004	LES LABORATOIRES SERVIER (SURESNES)	IE
COVERDINE 5mg/1.25mg/10mg film-coated tablets	NL/H/2636/03/DC	PA0568/024/003	LES LABORATOIRES SERVIER (SURESNES)	IE
COVERDINE	NL/H/2636/05/DC	PA0568/024/005	LES LABORATOIRES	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
10mg/2.5mg/10mg film-coated tablets			SERVIER (SURESNES)	
COVERDINE 5mg/1.25mg/5mg film-coated tablets	NL/H/2636/02/DC	PA0568/024/002	LES LABORATOIRES SERVIER (SURESNES)	IE
Triplixam 5 mg/1,25 mg/10 mg, comprimés pelliculés	NL/H/2636/03/DC	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/5 mg, comprimés pelliculés	NL/H/2636/02/DC	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/10 mg, comprimés pelliculés	NL/H/2636/05/DC	2014060178	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg, comprimés pelliculés	NL/H/2636/03/DC	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/5 mg, comprimés pelliculés	NL/H/2636/02/DC	2014060175	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/5 mg, comprimés pelliculés	NL/H/2636/04/DC	2014060177	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/005	2014060178	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2636/004	2014060177	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2636/003	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25	NL/H/2636/002	2014060175	SERVIER BENELUX S.A./N.V.	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
mg/5 mg, filmomhulde tabletten				
Triplixam 10 mg/2,5 mg/10 mg filmom obložene tablete	NL/H/2636/05/DC	HR-H-826085198	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5 mg/1,25 mg/10 mg filmom obložene tablete	NL/H/2636/03/DC	HR-H-175417914	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 10 mg/2,5 mg/5 mg filmom obložene tablete	NL/H/2636/04/DC	HR-H-919762076	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5 mg/1,25 mg/5 mg filmom obložene tablete	NL/H/2636/02/DC	HR-H-385858036	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2636/02/DC	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2636/03/DC	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/10 mg Filmtabletten	NL/H/2636/03/DC	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/10 mg, comprimés pelliculés	NL/H/2636/05/DC	BE448684	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/5 mg Filmtabletten	NL/H/2636/02/DC	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/5 mg Filmtabletten	NL/H/2636/04/DC	BE448675	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2636/04/DC	BE448675	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5	NL/H/2636/05/DC	BE448684	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
mg/10 mg Filmtabletten				
ТРИПЛИКСАМ 10 mg/2,5 mg/10mg филмирани таблетки	NL/H/2636/005	20140079	LES LABORATOIRES SERVIER (SURESNES)	BG
Triplixam, 10 mg + 2,5 mg + 10 mg, tabletki powlekane	NL/H/2636/05/DC	21775	LES LABORATOIRES SERVIER (SURESNES)	PL
Triplixam, 5 mg + 1,25 mg + 5 mg, tabletki powlekane	NL/H/2636/02/DC	21772	LES LABORATOIRES SERVIER (SURESNES)	PL
Triplixam, 10 mg + 2,5 mg + 5 mg, tabletki powlekane	NL/H/2636/04/DC	21774	LES LABORATOIRES SERVIER (SURESNES)	PL
Triplixam, 5 mg + 1,25 mg + 10 mg, tabletki powlekane	NL/H/2636/03/DC	21773	LES LABORATOIRES SERVIER (SURESNES)	PL
ТРИПЛИКСАМ 10 mg/2,5 mg/5 mg филмирани таблетки	NL/H/2636/004	20140078	LES LABORATOIRES SERVIER (SURESNES)	BG
ТРИПЛИКСАМ 5 mg/1,25 mg/5 mg филмирани таблетки	NL/H/2636/02/DC	20140076	LES LABORATOIRES SERVIER (SURESNES)	BG
ТРИПЛИКСАМ 5 mg/1,25 mg/10 mg филмирани таблетки	NL/H/2636/03/DC	20140077	LES LABORATOIRES SERVIER (SURESNES)	BG
Triplixam 5 mg/ 1,25 mg/ 10 mg apvalkotās tabletes	NL/H/2636/03/DC	14-0058	LES LABORATOIRES SERVIER (SURESNES)	LV
Triplixam 5 mg/ 1,25 mg/ 5 mg apvalkotās tabletes	NL/H/2636/02/DC	14-0057	LES LABORATOIRES SERVIER (SURESNES)	LV
Triplixam 10 mg/ 2,5 mg/ 10 mg apvalkotās	NL/H/2636/05/DC	14-0060	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
tabletes				
Triplixam 10 mg/ 2,5 mg/ 5 mg apvalkotās tabletes	NL/H/2636/04/DC	14-0059	LES LABORATOIRES SERVIER (SURESNES)	LV
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407128	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407130	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407116	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407179	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407181	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407167	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407229	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407066	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407217	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse	NL/H/2636/02/DC	042407078	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
rivestite con film				
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407080	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407231	LES LABORATOIRES SERVIER (SURESNES)	IT
Triplixam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2636/04/DC	5615968	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2636/04/DC	5615950	LES LABORATOIRES SERVIER (SURESNES)	PT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/007	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/012	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/022	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/017	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/025	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/021	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
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/024	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/023	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/008	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/018	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/030	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/006	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/016	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/020	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/027	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/029	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/009	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
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/028	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/011	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/013	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/019	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/014	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/015	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/010	LES LABORATOIRES SERVIER (SURESNES)	LT
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/029	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/028	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/027	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/025	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
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/026	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/030	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg, Filmtabletten	NL/H/2636/02/DC	2014060175	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg, Filmtabletten	NL/H/2636/03/DC	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/5 mg, Filmtabletten	NL/H/2636/04/DC	2014060177	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg kalvopäällysteiset tabletit	NL/H/2636/003	30888	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 5 mg/1,25 mg/5 mg kalvopäällysteiset tabletit	NL/H/2636/02/DC	30887	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10 mg/2,5 mg/5 mg kalvopäällysteiset tabletit	NL/H/2636/04/DC	30889	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10 mg/2,5 mg/10 mg kalvopäällysteiset tabletit	NL/H/2636/05/DC	30890	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2636/05/DC	5595822	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2636/05/DC	5595830	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam	NL/H/2636/03/DC	5595657	LES LABORATOIRES	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
5mg/1,25mg/10mg comprimidos revestidos por película			SERVIER (SURESNES)	
Triplixam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2636/03/DC	5595665	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2636/02/DC	5595525	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2636/02/DC	5595517	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/023	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/019	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/021	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/022	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/024	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/020	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
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/017	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/018	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/015	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/013	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/010	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/014	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/007	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/009	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/012	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/011	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/008	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
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/016	SERVIER PHARMA D.O.O	SI
TRIPLIXAM 10 mg/2,5 mg/ 10 mg, comprimé pelliculé	NL/H/2636/05/DC	34009 278 164 5 1	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2.5 mg/5 mg, comprimé pelliculé	NL/H/2636/04/DC	34009 278 162 2 2	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2.5 mg/5 mg, comprimé pelliculé	NL/H/2636/04/DC	34009 278 161 6 1	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2,5 mg/ 10 mg, comprimé pelliculé	NL/H/2636/05/DC	34009 278 163 9 0	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/10 mg, comprimé pelliculé	NL/H/2636/03/DC	34009 278 158 5 0	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/10 mg, comprimé pelliculé	NL/H/2636/03/DC	34009 278 159 1 1	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/5 mg, comprimé pelliculé	NL/H/2636/02/DC	34009 278 157 9 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/5 mg, comprimé pelliculé	NL/H/2636/02/DC	34009 278 156 2 1	LES LABORATOIRES SERVIER (SURESNES)	FR
Triplixam 10 mg/2,5 mg/10 mg, Filmtabletten	NL/H/2636/05/DC	2014060178	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5mg/1,25mg/5mg επικαλυμμένα με υμένιο δισκία	NL/H/2636/02/DC	022156	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
Triplixam 5mg/1,25mg/10mg επικαλυμμένα με υμένιο δισκία	NL/H/2636/03/DC	022157	LES LABORATOIRES SERVIER (SURESNES)	CY
Triplixam 10mg/2,5mg/5mg επικαλυμμένα με υμένιο δισκία	NL/H/2636/04/DC	022158	LES LABORATOIRES SERVIER (SURESNES)	CY
Triplixam 10mg/2,5mg/10mg επικαλυμμένα με υμένιο δισκία	NL/H/2636/05/DC	022159	LES LABORATOIRES SERVIER (SURESNES)	CY
TRIPLIXAM 10 mg/2,5 mg/10 mg filmom obalené tablety	NL/H/2636/005	58/0026/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
TRIPLIXAM 10 mg/2,5 mg/5 mg filmom obalené tablety	NL/H/2636/004	58/0025/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
TRIPLIXAM 5 mg/1,25 mg/10 mg filmom obalené tablety	NL/H/2636/003	58/0024/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
TRIPLIXAM 5 mg/1,25 mg/5 mg filmom obalené tablety	NL/H/2636/002	58/0023/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/038	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele	NL/H/2636/04/DC	LT/1/14/3522/035	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
dengtos tabletės				
TRIPLIXAM 10 mg/2.5 mg/5 mg plėvele dengtos tabletės	NL/H/2636/04/DC	LT/1/14/3522/036	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 10 mg/2.5 mg/10 mg plėvele dengtos tabletės	NL/H/2636/05/DC	LT/1/14/3522/037	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/032	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/034	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/5 mg plėvele dengtos tabletės	NL/H/2636/02/DC	LT/1/14/3522/031	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIXAM 5 mg/1.25 mg/10 mg plėvele dengtos tabletės	NL/H/2636/03/DC	LT/1/14/3522/033	LES LABORATOIRES SERVIER (SURESNES)	LT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407371	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407383	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407357	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407369	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/10 mg compresse	NL/H/2636/03/DC	042407332	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
rivestite con film				
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407344	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407320	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407318	LES LABORATOIRES SERVIER (SURESNES)	IT
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/031	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/02/DC	H/14/01566/032	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/033	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/035	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/05/DC	H/14/01566/037	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/04/DC	H/14/01566/036	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/03/DC	H/14/01566/034	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko	NL/H/2636/05/DC	H/14/01566/038	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
obložene tablete				
TRIPLIXAM 10 mg/2,5 mg/10 mg, comprimé pelliculé	NL/H/2636/005	34009 301 637 7 1	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2,5 mg/10 mg, comprimé pelliculé	NL/H/2636/005	34009 550 601 0 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2.5 mg/5 mg, comprimé pelliculé	NL/H/2636/04/DC	34009 550 600 9 3	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2,5 mg/5 mg, comprimé pelliculé	NL/H/2636/004	34009 301 637 6 4	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1,25 mg/5 mg, comprimé pelliculé	NL/H/2636/002	34009 550 600 7 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1,25 mg/5 mg, comprimé pelliculé	NL/H/2636/002	34009 301 637 4 0	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1,25 mg/10 mg, comprimé pelliculé	NL/H/2636/003	34009 278 159 1 1	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1,25 mg/10 mg, comprimé pelliculé	NL/H/2636/003	34009 550 600 8 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Triplixam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	11961/2019/07	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	11960/2019/07	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate	NL/H/2636/05/DC	11961/2019/06	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
filmate				
Triplixam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	11961/2019/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/05/DC	11961/2019/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	11961/2019/08	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/05/DC	11961/2019/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/05/DC	11961/2019/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/05/DC	11961/2019/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/003	11959/2019/07	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	11960/2019/08	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/04/DC	11960/2019/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	11960/2019/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate	NL/H/2636/04/DC	11960/2019/03	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
filmate				
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/04/DC	11960/2019/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/002	11958/2019/08	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/04/DC	11960/2019/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/04/DC	11960/2019/05	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03/DC	11959/2019/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/003	11959/2019/08	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03/DC	11959/2019/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/003	11959/2019/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03/DC	11959/2019/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03/DC	11959/2019/04	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate	NL/H/2636/03/DC	11959/2019/02	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
filmate				
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02/DC	11958/2019/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02/DC	11958/2019/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02/DC	11958/2019/04	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02/DC	11958/2019/05	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02/DC	11958/2019/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/002	11958/2019/07	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/002	11958/2019/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10mg/2,5mg/10mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/05/DC	100188/16-08-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 10mg/2,5mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/04/DC	100187/16-08-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 5mg/1,25mg/10mg επικαλυμμένα με λεπτό	NL/H/2636/03/DC	100186/16-08-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
υμένιο δισκία				
TRIPLIXAM 5mg/1,25mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/02/DC	100185/16-08-2019	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 10 mg/2.5 mg/5 mg, comprimé pelliculé	NL/H/2636/04/DC	34009 301 637 6 4	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/10 mg, comprimé pelliculé	NL/H/2636/03/DC	34009 301 637 5 7	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/5 mg, comprimé pelliculé	NL/H/2636/02/DC	34009 550 600 7 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/10 mg, comprimé pelliculé	NL/H/2636/03/DC	34009 550 600 8 6	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 5 mg/1.25 mg/5 mg, comprimé pelliculé	NL/H/2636/02/DC	34009 301 637 4 0	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2,5 mg/ 10 mg, comprimé pelliculé	NL/H/2636/05/DC	34009 550 601 0 9	LES LABORATOIRES SERVIER (SURESNES)	FR
TRIPLIXAM 10 mg/2,5 mg/ 10 mg, comprimé pelliculé	NL/H/2636/05/DC	34009 301 637 7 1	LES LABORATOIRES SERVIER (SURESNES)	FR