12 July 2018
EMA/498820/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

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

Procedure no.: PSUSA/00010358/201711

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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
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
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 2 mg+5 mg+0,625 mg tabletes	HU/H/0342/001	14-0025	KRKA, D.D., NOVO MESTO	LV
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

Product 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+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	2014007	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 4 mg/5 mg/1,25 mg таблетки	HU/H/0342/002	2014008	KRKA, D.D., NOVO MESTO	BG
Ко-Амлеса 4 mg/10 mg/1,25 mg таблетки	HU/H/0342/003	2014009	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 mg/2,5 mg таблетки	HU/H/0342/005	2014011	KRKA, D.D., NOVO MESTO	BG
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

Product 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 tablety	HU/H/0342/004	58/0449/13-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-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 2 mg/5 mg/0,625 mg comprimate	HU/H/0342/001	6141/2014/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/03	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/5 mg/1,25 mg comprimate	HU/H/0342/002	6142/2014/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/08	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Co-Amlessa 4 mg/10 mg/1,25 mg comprimate	HU/H/0342/003	6143/2014/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/04	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/5 mg/2,5 mg comprimate	HU/H/0342/004	6144/2014/09	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/01	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/02	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/03	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/04	KRKA, D.D., NOVO MESTO	RO

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/05	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/06	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/07	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/08	KRKA, D.D., NOVO MESTO	RO
Co-Amlessa 8 mg/10 mg/2,5 mg comprimate	HU/H/0342/005	6145/2014/09	KRKA, D.D., NOVO MESTO	RO
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
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

Product 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/14/01901/006	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 mg/1,25 mg tablete	HU/H/0342/003	H/14/01901/021	KRKA, D.D., NOVO MESTO	SI

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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

Product 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/14/01901/038	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
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

Product 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 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 mg/1,25 mg tabletės	HU/H/0342/002	LT/1/13/3462/011	KRKA, D.D., NOVO MESTO	LT
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

Product 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/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
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

Product 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/1/13/3462/035	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 mg/2,5 mg tabletės	HU/H/0342/005	LT/1/13/3462/044	KRKA, D.D., NOVO MESTO	LT
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

Product 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 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
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

Product 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-22577/18	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

Product 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 8 mg/5 mg/2,5 mg tabletta	HU/H/0342/004	OGYI-T-22577/32	KRKA, D.D., NOVO MESTO	HU
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

Product 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 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/01	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/02	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/03	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/04	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/05	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/06	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/07	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/08	KRKA, D.D., NOVO MESTO	HU
Tonanda 2 mg/5 mg/0,625 mg tabletta	not available	OGYI-T-22621/09	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/10	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/11	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/12	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/13	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/14	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
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/15	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/16	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/17	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/5 mg/1,25 mg tabletta	not available	OGYI-T-22621/18	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/19	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/20	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/21	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/22	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/23	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/24	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/25	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/26	KRKA, D.D., NOVO MESTO	HU
Tonanda 4 mg/10 mg/1,25 mg tabletta	not available	OGYI-T-22621/27	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/28	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
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/29	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/30	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/31	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/32	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/33	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/34	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/35	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/5 mg/2,5 mg tabletta	not available	OGYI-T-22621/36	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/37	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/38	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/39	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/40	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/41	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/42	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
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/43	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/44	KRKA, D.D., NOVO MESTO	HU
Tonanda 8 mg/10 mg/2,5 mg tabletta	not available	OGYI-T-22621/45	KRKA, D.D., NOVO MESTO	HU
Co-Dalneva 8 mg/2,5 mg/10 mg tablete	not available	HR-H-180076502	HCS BVBA	HR
Co-Dalneva 8 mg/2,5 mg/5 mg tablete	not available	HR-H-209243724	HCS BVBA	HR
Co-Dalneva 4 mg/1,25 mg/5 mg tablete	not available	HR-H-223376332	HCS BVBA	HR
Co-Dalneva 4 mg/1,25 mg/10 mg tablete	not available	HR-H-622847904	HCS BVBA	HR
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

Product 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 + 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 mg + 5 mg comprimidos	not available	5650866	KRKA, D.D., NOVO MESTO	PT
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
Triplixam, 10 mg/2,5 mg/10 mg õhukese polümeerikattega tabletid	NL/H/2636/005	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/2636/002	835014	LES LABORATOIRES SERVIER (SURESNES)	EE
Triplixam, 10mg/2,5mg/5mg õhukese polümeerikattega tabletid	NL/H/2636/004	835314	LES LABORATOIRES SERVIER (SURESNES)	EE

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
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 mg/10 mg, potahované tablety	NL/H/2636/05/DC	58/103/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
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/2636/004	RVG 112147	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2636/002	RVG 112145	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/005	RVG 112148	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2636/003	RVG 112146	LES LABORATOIRES SERVIER (SURESNES)	NL
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/005	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
Triplixam 10 mg/2,5 mg/5 mg, comprimés pelliculés	NL/H/2636/004	BE448675	SERVIER BENELUX S.A./N.V.	BE
Coverdine 10mg/2.5mg/5mg film-coated tablets]	NL/H/2636/004	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 10mg/2.5mg/10mg film-coated tablets	NL/H/2636/05/DC	PA0568/024/005	LES LABORATOIRES SERVIER (SURESNES)	IE
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/2636/003	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/5 mg, comprimés pelliculés	NL/H/2636/002	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/10 mg, comprimés pelliculés	NL/H/2636/005	2014060178	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg, comprimés pelliculés	NL/H/2636/003	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/5 mg, comprimés pelliculés	NL/H/2636/002	2014060175	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/5 mg, comprimés pelliculés	NL/H/2636/004	2014060177	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2636/005	2014060178	S.A SERVIER BENELUX 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

Product 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, filmomhulde tabletten	NL/H/2636/003	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2636/002	2014060175	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10mg/ 2,5mg/ 10mg filmom obložene tablete	NL/H/2636/005	HR-H-826085198	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5mg/ 1,25mg/ 10mg filmom obložene tablete	NL/H/2636/003	HR-H-175417914	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 10mg/ 2,5mg/ 5mg filmom obložene tablete	NL/H/2636/004	HR-H-919762076	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5mg/ 1,25mg/ 5mg filmom obložene tablete	NL/H/2636/002	HR-H-385858036	SERVIER PHARMA D.O.O - CROATIA	HR
Triplixam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2636/002	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2636/003	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/10 mg Filmtabletten	NL/H/2636/003	BE448666	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/10 mg, comprimés pelliculés	NL/H/2636/005	BE448684	SERVIER BENELUX S.A./N.V.	BE
Triplixam 5 mg/1,25 mg/5 mg Filmtabletten	NL/H/2636/002	BE448657	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/5 mg Filmtabletten	NL/H/2636/004	BE448675	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2636/004	BE448675	SERVIER BENELUX S.A./N.V.	BE
Triplixam 10 mg/2,5 mg/10 mg Filmtabletten	NL/H/2636/005	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
ТРИПЛИКСАМ 10 mg/2,5 mg/10mg филмирани таблетки	NL/H/2636/05/DC	20140079	LES LABORATOIRES SERVIER (SURESNES)	BG
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407243	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407142	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407193	LES LABORATOIRES SERVIER (SURESNES)	IT
Triplixam, 10 mg + 2,5 mg + 10 mg, tabletki powlekane	NL/H/2636/005	21775	LES LABORATOIRES SERVIER (SURESNES)	PL
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407092	LES LABORATOIRES SERVIER (SURESNES)	IT
Triplixam, 5 mg + 1,25 mg + 5 mg, tabletki powlekane	NL/H/2636/002	21772	LES LABORATOIRES SERVIER (SURESNES)	PL
Triplixam, 10 mg + 2,5 mg + 5 mg, tabletki powlekane	NL/H/2636/004	21774	LES LABORATOIRES SERVIER (SURESNES)	PL
Triplixam, 5 mg + 1,25 mg + 10 mg, tabletki powlekane	NL/H/2636/003	21773	LES LABORATOIRES SERVIER (SURESNES)	PL
ТРИПЛИКСАМ 10 mg/2,5 mg/5 mg филмирани таблетки	NL/H/2636/04/DC	20140078	LES LABORATOIRES SERVIER (SURESNES)	BG
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/04	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
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	6484/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
ТРИПЛИКСАМ 5 mg/1,25 mg/5 mg филмирани таблетки	NL/H/2636/02/DC	20140076	LES LABORATOIRES SERVIER (SURESNES)	BG
TRIPLIXAM 10mg/2,5mg/10mg	NL/H/2636/05/DC	58/0026/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
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 tabletes	NL/H/2636/05/DC	14-0060	LES LABORATOIRES SERVIER (SURESNES)	LV
Triplixam 10 mg/2,5 mg/10 mg Filmtablettten	NL/H/2636/005	135513	LES LABORATOIRES SERVIER (SURESNES)	AT
Triplixam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/003	6483/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02	6482/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10mg/2,5mg/5mg	NL/H/2636/04/DC	58/0025/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
TRIPLIXAM 5 mg/1,25 mg/5 mg Filmtabletten	NL/H/2636/02/DC	135510	LES LABORATOIRES SERVIER (SURESNES)	AT
TRIPLIXAM 5mg/1,25mg/10mg	NL/H/2636/03/DC	58/0024/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
Triplixam 10 mg/ 2,5 mg/ 5 mg apvalkotās tabletes	NL/H/2636/04/DC	14-0059	LES LABORATOIRES SERVIER (SURESNES)	LV
TRIPLIXAM 5mg/1,25mg/5mg	NL/H/2636/02/DC	58/0023/14-S	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	SK
TRIPLIXAM 5 mg/1,25 mg/10 mg Filmtabletten	NL/H/2636/03/DC	135511	LES LABORATOIRES SERVIER (SURESNES)	AT
Triplixam 10 mg/2,5 mg/5 mg Filmtablettten	NL/H/2636/004	135512	LES LABORATOIRES SERVIER (SURESNES)	AT
TRIPLIAM 5 mg/1,25 mg/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407155	LES LABORATOIRES SERVIER (SURESNES)	IT
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

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407205	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/5 mg compresse rivestite con film	NL/H/2636/04/DC	042407294	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/10 mg compresse rivestite con film	NL/H/2636/03/DC	042407282	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407104	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407270	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03	6483/2014/01	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
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407066	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02	6482/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
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 rivestite con film	NL/H/2636/02/DC	042407078	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03	6483/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIAM 5 mg/1,25 mg/5 mg compresse rivestite con film	NL/H/2636/02/DC	042407080	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02	6482/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407256	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407306	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/01	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
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	6484/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	6484/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02	6482/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIAM 10 mg/2,5 mg/10 mg compresse rivestite con film	NL/H/2636/05/DC	042407231	LES LABORATOIRES SERVIER (SURESNES)	IT
TRIPLIXAM 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/02	6482/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03	6483/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/005	6484/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	6484/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
TRIPLIXAM 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/03	6483/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2636/004	5616073	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2636/004	5615968	LES LABORATOIRES SERVIER (SURESNES)	PT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Triplixam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2636/004	5615950	LES LABORATOIRES SERVIER (SURESNES)	PT
TRIPLIXAM 10mg/2,5mg/10mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/05/DC	72112/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/007	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10mg/2,5mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/04/DC	72111/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 5mg/1,25mg/10mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/03/DC	72110/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/012	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5mg/1,25mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2636/02/DC	72109/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/022	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/017	LES LABORATOIRES SERVIER	LT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/025	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/021	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/024	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/023	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/008	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/018	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/005	LT/1/14/3522/030	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/006	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/016	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/020	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/027	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/029	LES LABORATOIRES SERVIER	LT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/009	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/028	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/011	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/013	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/004	LT/1/14/3522/019	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/014	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2636/003	LT/1/14/3522/015	LES LABORATOIRES SERVIER	LT
TRIPLIXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2636/002	LT/1/14/3522/010	LES LABORATOIRES SERVIER	LT
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/005	H/14/01566/029	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/005	H/14/01566/028	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/005	H/14/01566/027	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/005	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/2636/005	H/14/01566/026	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/10 mg filmsko obložene tablete	NL/H/2636/005	H/14/01566/030	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg, Filmtabletten	NL/H/2636/002	2014060175	SERVIER BENELUX S.A./N.V.	LU
Triplixam 5 mg/1,25 mg/10 mg, Filmtabletten	NL/H/2636/003	2014060176	SERVIER BENELUX S.A./N.V.	LU
Triplixam 10 mg/2,5 mg/5 mg, Filmtabletten	NL/H/2636/004	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/2636/002	30887	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10 mg/2,5 mg/5 mg kalvopäällysteiset tabletit	NL/H/2636/004	30889	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10 mg/2,5 mg/10 mg kalvopäällysteiset tabletit	NL/H/2636/005	30890	LES LABORATOIRES SERVIER (SURESNES)	FI
Triplixam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2636/005	6485/2014/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2636/002	6482/2014/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2636/004	6484/2014/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Triplixam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2636/003	6483/2014/06	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
Triplixam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2636/005	5616107	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2636/005	5595822	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2636/005	5595830	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2636/003	5616032	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2636/003	5595657	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2636/003	5595665	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2636/002	5616024	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2636/002	5595525	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2636/002	5595517	LES LABORATOIRES SERVIER (SURESNES)	PT
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/023	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/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/019	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/021	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/022	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/024	SERVIER PHARMA D.O.O	SI
Triplixam 10 mg/2,5 mg/5 mg filmsko obložene tablete	NL/H/2636/004	H/14/01566/020	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	H/14/01566/017	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	H/14/01566/018	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	H/14/01566/015	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	H/14/01566/013	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/010	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	H/14/01566/014	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/007	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/009	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/012	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/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/011	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/5 mg filmsko obložene tablete	NL/H/2636/002	H/14/01566/008	SERVIER PHARMA D.O.O	SI
Triplixam 5 mg/1,25 mg/10 mg filmsko obložene tablete	NL/H/2636/003	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/2636/005	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
Triplixam 5mg/1,25mg/10mg επικαλυμμένα με υμένιο δισκία	NL/H/2636/03/DC	022157	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 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
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

Product 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/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 mg/2,5 mg tabletta	not available	OGYI-T-22622/29	KRKA, D.D., NOVO MESTO	HU
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

Product 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 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
Flutensif 1,5 mg / 10 mg δισκία ελεγχόμενης αποδέσμευσης	NL/H/2639/02/DC	51997/13-6-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/06	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
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 10 mg comprimate cu eliberare modificată	NL/H/2639/002	6128/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Flutensif 1,5 mg / 5 mg δισκία ελεγχόμενης αποδέσμευσης	NL/H/2639/01/DC	51996/13-6-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/06	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludexam 1,5 mg / 5 mg comprimate cu eliberare modificată	NL/H/2639/001	6127/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Cardilopin Komb 1,5 mg/5 mg módosított hatóanyagleadású tabletta	NL/H/2639/001/DC	OGYI-T-22562/01	PROTERAPIA HUNGARY	HU
Cardilopin Komb 1,5 mg/10 mg módosított hatóanyagleadású tabletta	NL/H/2639/002/DC	OGYI-T-22562/02	PROTERAPIA HUNGARY	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
FLUTENSIF 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2639/002	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/2639/002	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/2639/001	5594205	LES LABORATOIRES SERVIER (SURESNES)	PT
FLUTENSIF 1,5 mg / 10 mg, tabletten met gereguleerde afgifte	NL/H/2639/002	RVG 112322	LES LABORATOIRES SERVIER	NL
Flutensif, 1,5 mg/10 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2639/002	824413	LES LABORATOIRES SERVIER (SURESNES)	EE
Flutensif, 1,5 mg/5 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2639/001	824513	LES LABORATOIRES SERVIER (SURESNES)	EE

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
FLUTENSIF 1,5 mg / 5 mg, tabletten met gereguleerde afgifte	NL/H/2639/001	RVG 112321	LES LABORATOIRES SERVIER (SURESNES)	NL
FLUTENSIF 1,5 mg/10 mg tablety s riadeným uvoľňovaním	NL/H/2639/002	58/0437/13-S	LES LABORATOIRES SERVIER	SK
FLUTENSIF 1,5 mg/5 mg tablety s riadeným uvoľňovaním	NL/H/2639/001	58/0436/13-S	LES LABORATOIRES SERVIER	SK
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/2639/02/DC	135240	LES LABORATOIRES SERVIER (SURESNES)	AT
Natrixam 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 585 688 9 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Natrixam 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 585 687 5 6	LES LABORATOIRES SERVIER (SURESNES)	FR
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

Product 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 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 275 983 5 7	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	34009 275 982 9 6	LES LABORATOIRES SERVIER (SURESNES)	FR
NATRIXAM 1,5mg/10mg, comprimé à libération modifiée	NL/H/2637/02/DC	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 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 275 977 5 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Natrixam 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 275 976 9 5	LES LABORATOIRES SERVIER (SURESNES)	FR
Natrixam 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 275 975 2 7	LES LABORATOIRES SERVIER (SURESNES)	FR
Natrixam 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	34009 275 974 6 6	LES LABORATOIRES SERVIER (SURESNES)	FR
Natrixam 1,5 mg / 10 mg δισκία ελεγχόμενης αποδέσμευσης	NL/H/2637/02/DC	51979/13-6-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Tertensam 1,5 mg / 10 mg ilgstošās darbības tabletes	NL/H/2637/002	13-0247	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
Fludex Plus 1,5mg/10mg δισκία ελεγχόμενης αποδέσμευσης	NL/H/2637/02/DC	021971	LES LABORATOIRES SERVIER (SURESNES)	CY
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Fludex Plus 1,5mg/5m gδισκία ελεγχόμενηςαποδέσμευσης	NL/H/2637/01/DC	021970	LES LABORATOIRES SERVIER (SURESNES)	CY
Natrixam 1,5 mg / 5 mg δισκία ελεγχόμενης αποδέσμευσης	NL/H/2637/01/DC	51978/13-6-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
Tertensam 1,5 mg / 5 mg ilgstošās darbības tabletes	NL/H/2637/001	13-0246	LES LABORATOIRES SERVIER (SURESNES)	LV
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/05	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/06	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
Natrixam 1,5 mg/10 mg comprimate cu eliberare modificată	NL/H/2637/02/DC	6126/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/01	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/04	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/02	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/03	LES LABORATOIRES SERVIER (SURESNES)	RO
Natrixam 1,5 mg/5 mg comprimate cu eliberare modificată	NL/H/2637/01/DC	6125/2014/06	LES LABORATOIRES SERVIER (SURESNES)	RO
NATRIXAM 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2637/002	5615877	LES LABORATOIRES SERVIER (SURESNES)	PT
NATRIXAM 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2637/002	5594106	LES LABORATOIRES SERVIER (SURESNES)	PT
NATRIXAM 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2637/001	5594148	LES LABORATOIRES SERVIER (SURESNES)	PT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
NATRIXAM 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2637/001	5594163	LES LABORATOIRES SERVIER (SURESNES)	PT
NATRIXAM 1.5 mg / 10 mg comprimidos de libertação modificada	NL/H/2637/002	5594114	LES LABORATOIRES SERVIER (SURESNES)	PT
NATRIXAM 1.5 mg / 5 mg comprimidos de libertação modificada	NL/H/2637/001	5594155	LES LABORATOIRES SERVIER (SURESNES)	PT
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/2637/001	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/2637/001	LT/1/13/3427/001	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/001	LT/1/13/3427/002	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
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/001	LT/1/13/3427/005	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/001	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 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/02/DC	LT/1/13/3427/011	LES LABORATOIRES SERVIER (SURESNES)	LT
NATRIXAM 1,5 mg / 5 mg modifikuoto atpalaidavimo tabletės	NL/H/2637/001	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/2637/002	H/13/01074/011	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/002	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/005	SERVIER PHARMA D.O.O	SI

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/001	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/006	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/002	H/13/01074/012	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/004	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/5 mg tablete s prirejenim sproščanjem	NL/H/2637/001	H/13/01074/003	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/002	H/13/01074/008	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/002	H/13/01074/007	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/002	H/13/01074/010	SERVIER PHARMA D.O.O	SI
Nadexam 1,5 mg/10 mg tablete s prirejenim sproščanjem	NL/H/2637/002	H/13/01074/009	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
NATRIXAM 1,5 mg/5 mg tablete s prilagođenim oslobađanjem	NL/H/2637/001	HR-H-053397374	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg tablete s prilagođenim oslobađanjem	NL/H/2637/002	HR-H-921621992	SERVIER PHARMA D.O.O - CROATIA	HR
NATRIXAM 1,5 mg/10 mg, comprimé à libération modifiée	NL/H/2637/002	2014120414	LES LABORATOIRES SERVIER (SURESNES)	LU
NATRIXAM 1,5 mg/5 mg, comprimé à libération modifiée	NL/H/2637/001	2014120415	LES LABORATOIRES SERVIER (SURESNES)	LU
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297123	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297097	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297085	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297073	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297111	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
Frecanstal 1,5 mg/10 mg compresse a rilascio modificato	NL/H/2637/002	042297109	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297034	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297010	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297046	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297022	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297061	LES LABORATOIRES SERVIER (SURESNES)	IT
Frecanstal 1,5 mg/5 mg compresse a rilascio modificato	NL/H/2637/001	042297059	LES LABORATOIRES SERVIER (SURESNES)	IT
Natrixam 1,5 mg / 10 mg, tablety s řízeným uvolňováním	NL/H/2637/002	58/426/13-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Natrixam 1,5 mg / 5 mg, tablety s řízeným uvolňováním	NL/H/2637/001	58/425/13-C	LES LABORATOIRES SERVIER (SURESNES)	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Натриксам 1,5 mg / 10 mg, таблетки с изменено освобождаване	NL/H/2637/02/DC	20130382	LES LABORATOIRES SERVIER (SURESNES)	BG
Натриксам 1,5 mg / 5 mg, таблетки с изменено освобождаване	NL/H/2637/01/DC	20130381	LES LABORATOIRES SERVIER (SURESNES)	BG
Natrixam 1,5 mg / 10 mg, tabletten met gereguleerde afgifte	NL/H/2637/002	RVG 112320	LES LABORATOIRES SERVIER (SURESNES)	NL
Natrixam 1.5 mg / 10 mg modified-release tablets	NL/H/2637/002	MA066/02102	LES LABORATOIRES SERVIER (SURESNES)	MT
Natrixam, 1,5 mg/5 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2637/001	825313	LES LABORATOIRES SERVIER (SURESNES)	EE
Natrixam 1,5 mg / 5 mg, tabletten met gereguleerde afgifte	NL/H/2637/001	RVG 112319	LES LABORATOIRES SERVIER (SURESNES)	NL
Natrixam 1.5 mg / 5 mg modified-release tablets	NL/H/2637/001	MA066/02101	LES LABORATOIRES SERVIER (SURESNES)	MT
Natrixam, 1,5 mg/10 mg, toimeainet modifitseeritult vabastavad tabletid	NL/H/2637/002	825213	LES LABORATOIRES SERVIER (SURESNES)	EE
NATRIXAM 1,5 mg/10 mg tablety s riadeným uvoľňovaním	NL/H/2637/002	58/0435/13-S	LES LABORATOIRES SERVIER	SK
NATRIXAM 1,5 mg/5 mg tablety s riadeným uvoľňovaním	NL/H/2637/001	58/0434/13-S	LES LABORATOIRES SERVIER	SK

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Tertens-AM, 1,5 mg + 5 mg, tabletki o zmodyfikowanym uwalnianiu	NL/H/2637/001	21557	LES LABORATOIRES SERVIER (SURESNES)	PL
Tertens-AM, 1,5 mg + 10 mg, tabletki o zmodyfikowanym uwalnianiu	NL/H/2637/002	21558	LES LABORATOIRES SERVIER (SURESNES)	PL
NATRIXAM 1,5 mg/5 mg Tabletten mit veränderter Wirkstofffreisetzung	NL/H/2637/01/DC	135237	LES LABORATOIRES SERVIER (SURESNES)	AT
NATRIXAM 1,5 mg/10 mg Tabletten mit veränderter Wirkstofffreisetzung	NL/H/2637/02/DC	135238	LES LABORATOIRES SERVIER (SURESNES)	AT
Arplexam, 10 mg/2,5 mg/5 mg õhukese polümeerikattega tabletid	NL/H/2638/004	834814	LES LABORATOIRES SERVIER (SURESNES)	EE
Arplexam, 5 mg/1,25 mg/5 mg õhukese polümeerikattega tabletid	NL/H/2638/002	835514	LES LABORATOIRES SERVIER (SURESNES)	EE
Arplexam, 5 mg/1,25 mg/10 mg õhukese polümeerikattega tabletid	NL/H/2638/003	835414	LES LABORATOIRES SERVIER (SURESNES)	EE
Arplexam, 10 mg/2,5 mg/10 mg õhukese polümeerikattega tabletid	NL/H/2638/005	835214	LES LABORATOIRES SERVIER (SURESNES)	EE
ARPLEXAM 10 mg/2,5 mg/10 mg, potahované tablety	NL/H/2638/005	58/098/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
ARPLEXAM 5 mg/1,25 mg/5 mg, potahované tablety	NL/H/2638/002	58/095/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Arplexam 10 mg/2,5 mg/5 mg, potahované tablety	NL/H/2638/004	58/097/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Arplexam 5 mg/1,25 mg/10 mg, potahované tablety	NL/H/2638/003	58/096/14-C	LES LABORATOIRES SERVIER (SURESNES)	CZ
Arplexam 10 mg/2,5 mg/10 mg, filmomhulde tabletten	NL/H/2638/005	RVG 112143	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 10 mg/2,5 mg/5 mg, filmomhulde tabletten	NL/H/2638/004	RVG 112142	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 5 mg/1,25 mg/10 mg, filmomhulde tabletten	NL/H/2638/003	RVG 112141	LES LABORATOIRES SERVIER (SURESNES)	NL
Arplexam 5 mg/1,25 mg/5 mg, filmomhulde tabletten	NL/H/2638/002	RVG 112140	LES LABORATOIRES SERVIER (SURESNES)	NL
АРПЛЕКСАМ 5 mg/1,25 mg/10mg филмирани таблетки	NL/H/2638/003	20140068	LES LABORATOIRES SERVIER (SURESNES)	BG
Arplexam 10 mg/ 2,5 mg/ 5 mg apvalkotās tabletes	NL/H/2638/04/DC	14-0054	LES LABORATOIRES SERVIER (SURESNES)	LV
АРПЛЕКСАМ 10 mg/2,5 mg/5 mg филмирани таблетки	NL/H/2638/004	20140069	LES LABORATOIRES SERVIER (SURESNES)	BG
Arplexam 5 mg/ 1,25 mg/ 5 mg apvalkotās tabletes	NL/H/2638/02/DC	14-0052	LES LABORATOIRES SERVIER (SURESNES)	LV
ARPLEXAM 10 mg/2,5 mg/10 mg filmom obalené tablety	NL/H/2638/005	58/0021/14-S	LES LABORATOIRES SERVIER (SURESNES)	SK
АРПЛЕКСАМ 5 mg/1,25 mg/5 mg филмирани таблетки	NL/H/2638/002	20140067	LES LABORATOIRES SERVIER (SURESNES)	BG
Arplexam 5 mg/ 1,25 mg/ 10 mg apvalkotās tabletes	NL/H/2638/03/DC	14-0053	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
ARPLEXAM 10 mg/2,5 mg/5 mg filmom obalené tablety	NL/H/2638/004	58/0020/14-S	LES LABORATOIRES SERVIER (SURESNES)	SK
ARPLEXAM 5mg/1,25mg/5mg filmom obalené tablety	NL/H/2638/002	58/0018/14-S	LES LABORATOIRES SERVIER (SURESNES)	SK
АРПЛЕКСАМ 10 mg/2,5 mg/10 mg филмирани таблетки	NL/H/2638/005	20140070	LES LABORATOIRES SERVIER (SURESNES)	BG
Arplexam 10 mg/ 2,5 mg/ 10 mg apvalkotās tabletes	NL/H/2638/05/DC	14-0055	LES LABORATOIRES SERVIER (SURESNES)	LV
ARPLEXAM 5mg/1,25mg/10mg filmom obalené tablety	NL/H/2638/003	58/0019/14-S	LES LABORATOIRES SERVIER (SURESNES)	SK
Arplexam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2638/005	5616156	LES LABORATOIRES SERVIER	PT
Arplexam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2638/002	5596259	LES LABORATOIRES SERVIER	PT
Covercard Plus 5 mg/1,25 mg/5 mg filmtabletta	NL/H/2638/002/DC	OGYI-T-22626/02	PROTERAPIA HUNGARY	HU
Covercard Plus 10 mg/2,5 mg/5 mg filmtabletta	NL/H/2638/004/DC	OGYI-T-22626/04	PROTERAPIA HUNGARY	HU
Covercard Plus 5 mg/1,25 mg/10 mg filmtabletta	NL/H/2638/003/DC	OGYI-T-22626/03	PROTERAPIA HUNGARY	HU
Covercard Plus 10 mg/2,5 mg/10 mg filmtabletta	NL/H/2638/005/DC	OGYI-T-22626/05	PROTERAPIA HUNGARY	HU
Arplexam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2638/005	5615976	LES LABORATOIRES SERVIER	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
Arplexam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2638/002	5616123	LES LABORATOIRES SERVIER	PT
Arplexam 10mg/2,5mg/10mg comprimidos revestidos por película	NL/H/2638/005	5616008	LES LABORATOIRES SERVIER	PT
Arplexam 5mg/1,25mg/5mg comprimidos revestidos por película	NL/H/2638/002	5596242	LES LABORATOIRES SERVIER	PT
ARPLEXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/002	LT/1/14/3519/007	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/003	LT/1/14/3519/012	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5mg/1,25mg/10mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/03/DC	72115/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10mg/2,5mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/04/DC	72116/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10mg/2,5mg/10mg, επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/05/DC	72117/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/005	LT/1/14/3519/022	LES LABORATOIRES SERVIER	LT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
ARPLEXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/004	LT/1/14/3519/017	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5mg/1,25mg/5mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/2638/02/DC	72114/4-8-2014	SERVIER HELLAS PHARMACEUTICALS LTD	GR
ARPLEXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/005	LT/1/14/3519/025	LES LABORATOIRES SERVIER	LT
ARPLEXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/005	LT/1/14/3519/021	LES LABORATOIRES SERVIER	LT
ARPLEXAM 10 mg / 2,5 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/005	LT/1/14/3519/023	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/002	LT/1/14/3519/006	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/002	LT/1/14/3519/010	LES LABORATOIRES SERVIER	LT
ARPLEXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/004	LT/1/14/3519/016	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/002	LT/1/14/3519/008	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/003	LT/1/14/3519/015	LES LABORATOIRES SERVIER	LT
ARPLEXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/004	LT/1/14/3519/018	LES LABORATOIRES SERVIER	LT
ARPLEXAM 10 mg / 2,5 mg / 5 mg plėvele dengtos tabletės	NL/H/2638/004	LT/1/14/3519/020	LES LABORATOIRES SERVIER	LT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
ARPLEXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/003	LT/1/14/3519/013	LES LABORATOIRES SERVIER	LT
ARPLEXAM 5 mg / 1,25 mg / 10 mg plėvele dengtos tabletės	NL/H/2638/003	LT/1/14/3519/011	LES LABORATOIRES SERVIER	LT
Arplexam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2638/004	5596309	LES LABORATOIRES SERVIER (SURESNES)	PT
Arplexam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2638/004	5616149	LES LABORATOIRES SERVIER (SURESNES)	PT
Arplexam 10mg/2,5mg/5mg comprimidos revestidos por película	NL/H/2638/004	5596317	LES LABORATOIRES SERVIER (SURESNES)	PT
Arplexam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2638/003	5616131	LES LABORATOIRES SERVIER (SURESNES)	PT
Arplexam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2638/003	5596275	LES LABORATOIRES SERVIER (SURESNES)	PT
Arplexam 5mg/1,25mg/10mg comprimidos revestidos por película	NL/H/2638/003	5596267	LES LABORATOIRES SERVIER (SURESNES)	PT
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/04	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 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/10 mg comprimate filmate	NL/H/2638/005	7099/2014/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	7097/2014/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	7098/2014/04	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 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	7098/2014/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	7096/2014/04	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	7097/2014/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	7096/2014/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	7098/2014/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	7097/2014/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	7096/2014/01	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO

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Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	7097/2014/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	7098/2014/06	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/10 mg comprimate filmate	NL/H/2638/003	7097/2014/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	7096/2014/02	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
Norplexam 10 mg/2,5 mg/5 mg comprimate filmate	NL/H/2638/004	7098/2014/03	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO
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Norplexam 5 mg/1,25 mg/5 mg comprimate filmate	NL/H/2638/002	7096/2014/05	ANPHARM PRZEDSIĘBIORSTWO FARMACEUTYCZNE SPÓŁKA AKCYJNA	RO