11 January 2018
EMA/24903/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: venlafaxine

Procedure no.: PSUSA/00003104/201705

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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
Apclaven XL 150 mg Prolonged release capsule	not available	PL 40378/0190	TORRENT PHARMA LTD.	UK
Apclaven XL 37.5 mg Prolonged release capsule	not available	PL 40378/0188	TORRENT PHARMA LTD.	UK
Apclaven XL 75 mg Prolonged release capsule	not available	PL 40378/0189	TORRENT PHARMA LTD.	UK
Dobupal 37,5 mg comprimidos	ES/H/0154/001	60.666	ALMIRALL, S.A.	ES
Dobupal 50 mg comprimidos	ES/H/0154/002	60.667	ALMIRALL, S.A.	ES
Dobupal 75 mg comprimidos	ES/H/0154/003	60.668	ALMIRALL, S.A.	ES
Dobupal Retard 150 mg cápsulas duras de liberación prolongada	ES/H/0154/005	62.421	ALMIRALL, S.A.	ES
Dobupal Retard 75 mg cápsulas duras de liberación prolongada	ES/H/0154/004	62.420	ALMIRALL, S.A.	ES
Duofaxin	SE/H/0564/004	IS/1/14/052/01	MEDILINK A/S	IS
Duofaxin	SE/H/0564/003	IS/1/14/052/02	MEDILINK A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Duofaxin	SE/H/0564/004	IS/1/14/052/03	MEDILINK A/S	IS
Duofaxin	SE/H/0564/004	IS/1/14/052/01	MEDILINK A/S	IS
Duofaxin	SE/H/0564/003	IS/1/14/052/02	MEDILINK A/S	IS
Duofaxin	SE/H/0564/004	IS/1/14/052/03	MEDILINK A/S	IS
Duofaxin, depottabletter	SE/H/0564/002	39249	MEDILINK A/S	DK
Duofaxin, depottabletter	SE/H/0564/003	39250	MEDILINK A/S	DK
Duofaxin, depottabletter	SE/H/0564/004	39251	MEDILINK A/S	DK
Duofaxin, depottabletter	SE/H/0564/002	39249	MEDILINK A/S	DK
Duofaxin, depottabletter	SE/H/0564/003	39250	MEDILINK A/S	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Duofaxin, depottabletter	SE/H/0564/004	39251	MEDILINK A/S	DK
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/19	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/18	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/21	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/20	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/07	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/08	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/23	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/22	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/09	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/24	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/10	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/11	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/02	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/12	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/04	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/05	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/13	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/03	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/01	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/16	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/14	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/15	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/06	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/17	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/12	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/13	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/10	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/16	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/15	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/14	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/17	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/19	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/02	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/01	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/18	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/20	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/03	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/22	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/07	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/05	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/24	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/21	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/04	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/09	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/23	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/11	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/06	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/25	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/08	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/16	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/02	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/16	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/17	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/01	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/18	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/03	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/19	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/04	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/05	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/21	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/06	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/23	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/08	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/20	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/09	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/07	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/22	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/24	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/11	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/10	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/13	PFIZER EUROPE MA EEIG	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/12	PFIZER EUROPE MA EEIG	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/14	PFIZER EUROPE MA EEIG	RO
Efectin ER 150 mg Hartkapseln, retardiert	SE/H/0936/003	1-23043	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/097	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/087	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/095	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/100	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/101	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/098	PFIZER EUROPE MA EEIG	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
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/107	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/108	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/094	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/103	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/086	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/096	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/106	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/081	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/093	PFIZER EUROPE MA EEIG	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
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/085	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/105	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/082	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/080	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/104	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/077	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/102	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/078	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/076	PFIZER EUROPE MA EEIG	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
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/099	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/089	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/088	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/083	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/092	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/074	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/084	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/090	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/079	PFIZER EUROPE MA EEIG	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
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/073	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/091	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/075	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 150 mg tvrdé tobolky s prodlouženým uvolňováním	SE/H/0936/003	30/687/99-C	PFIZER, SPOL. S R.O.	CZ
Efectin ER 150, 150 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/003	4940	PFIZER EUROPE MA EEIG	PL
Efectin ER 37,5 mg Hartkapseln, retardiert	SE/H/0936/001	1-24637	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/025	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/031	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/021	PFIZER EUROPE MA EEIG	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
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/028	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/026	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/023	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/024	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/015	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/030	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/027	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/018	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/034	PFIZER EUROPE MA EEIG	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
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/019	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/035	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/016	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/020	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/036	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/012	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/022	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/010	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/017	PFIZER EUROPE MA EEIG	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
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/011	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/014	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/032	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/033	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/013	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/029	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/006	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/008	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/007	PFIZER EUROPE MA EEIG	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
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/009	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/001	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/002	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/005	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/004	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/003	PFIZER EUROPE MA EEIG	SI
Efectin ER 37,5, 37,5 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/001	10523	PFIZER EUROPE MA EEIG	PL
Efectin ER 75 mg Hartkapseln, retardiert	SE/H/0936/002	1-23042	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/066	PFIZER EUROPE MA EEIG	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
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/064	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/061	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/065	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/054	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/056	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/057	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/053	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/048	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/063	PFIZER EUROPE MA EEIG	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
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/071	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/043	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/072	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/055	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/068	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/052	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/058	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/051	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/042	PFIZER EUROPE MA EEIG	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
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/069	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/037	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/044	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/049	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/039	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/045	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/050	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/046	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/070	PFIZER EUROPE MA EEIG	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
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/040	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/038	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/067	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/041	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/062	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/060	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/059	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/047	PFIZER EUROPE MA EEIG	SI
EFECTIN ER 75 mg tvrdé tobolky s prodlouženým uvolňováním	SE/H/0936/002	30/686/99-C	PFIZER, SPOL. S R.O.	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
Efectin ER 75, 75 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/002	4939	PFIZER EUROPE MA EEIG	PL
Efexor 150 mg capsule rigide a rilascio prolungato	SE/H/0936/003	028831105	PFIZER ITALIA S.R.L.	IT
Efexor 150 mg capsule rigide a rilascio prolungato	SE/H/0936/003	028831067	PFIZER ITALIA S.R.L.	IT
Efexor 37,5 mg capsule rigide a rilascio prolungato	SE/H/0936/001	028831129	PFIZER ITALIA S.R.L.	IT
Efexor 37,5 mg capsule rigide a rilascio prolungato	SE/H/0936/001	028831117	PFIZER ITALIA S.R.L.	IT
Efexor 37,5 mg capsule rigide a rilascio prolungato	SE/H/0936/001	028831131	PFIZER ITALIA S.R.L.	IT
Efexor 75 mg capsule rigide a rilascio prolungato	SE/H/0936/002	028831093	PFIZER ITALIA S.R.L.	IT
Efexor 75 mg capsule rigide a rilascio prolungato	SE/H/0936/002	028831055	PFIZER ITALIA S.R.L.	IT
Efexor Depot 150 mg depotkapsel, hard	SE/H/0936/003	97-4783	PFIZER AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor Depot 150 mg depotkapsel, hård	SE/H/0936/003	12607	PFIZER OY	FI
Efexor Depot 150 mg depotkapsel, hård	SE/H/0936/003	13321	PFIZER AB	SE
Efexor Depot 150 mg depotkapseli, kova	SE/H/0936/003	12607	PFIZER OY	FI
Efexor Depot 150 mg forðahylki, hart	SE/H/0936/003	970068	PFIZER APS	IS
Efexor Depot 225 mg depotkapsel, hård	SE/H/0936/004	49965	PFIZER AB	SE
Efexor Depot 37,5 mg depotkapsel, hard	SE/H/0936/001	04-3083	PFIZER AS	NO
Efexor Depot 37,5 mg depotkapsel, hård	SE/H/0936/001	20185	PFIZER OY	FI
Efexor Depot 37,5 mg depotkapsel, hård	SE/H/0936/001	21800	PFIZER AB	SE
Efexor Depot 37,5 mg depotkapseli, kova	SE/H/0936/001	20185	PFIZER OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor Depot 37,5 mg forðahylki, hart	SE/H/0936/001	IS/1/06/101/01	PFIZER APS	IS
Efexor Depot 75 mg depotkapsel, hard	SE/H/0936/002	97-4782	PFIZER AS	NO
Efexor Depot 75 mg depotkapsel, hård	SE/H/0936/002	12606	PFIZER OY	FI
Efexor Depot 75 mg depotkapsel, hård	SE/H/0936/002	13320	PFIZER AB	SE
Efexor Depot 75 mg depotkapseli, kova	SE/H/0936/002	12606	PFIZER OY	FI
Efexor Depot 75 mg forðahylki, hart	SE/H/0936/002	970067	PFIZER APS	IS
Efexor Depot, hårde depotkapsler	SE/H/0936/001	37361	PFIZER APS	DK
Efexor Depot, hårde depotkapsler	SE/H/0936/003	18578	PFIZER APS	DK
Efexor Depot, hårde depotkapsler	SE/H/0936/002	18577	PFIZER APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor XL 150 mg prolonged-release capsules, hard	SE/H/0936/003	PA 822/72/3	PFIZER HEALTHCARE IRELAND	IE
Efexor XL 150 mg prolonged-release capsules, hard	SE/H/0936/003	MA505/06202	PFIZER HELLAS, A.E.	MT
Efexor XL 150 mg prolonged-release capsules, hard	SE/H/0936/003	PL 00057/1281	PFIZER LIMITED	UK
Efexor XL 225 mg prolonged-release capsules, hard	SE/H/0936/004	PL 00057/1512	PFIZER LIMITED	UK
Efexor XL 37.5 mg prolonged-release capsules, hard	SE/H/0936/001	PA 822/72/1	PFIZER HEALTHCARE IRELAND	IE
Efexor XL 75 mg prolonged-release capsules, hard	SE/H/0936/002	PA 822/72/2	PFIZER HEALTHCARE IRELAND	IE
Efexor XL 75 mg prolonged-release capsules, hard	SE/H/0936/002	MA505/06201	PFIZER HELLAS, A.E.	MT
Efexor XL 75 mg prolonged-release capsules, hard	SE/H/0936/002	PL 00057/1280	PFIZER LIMITED	UK
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	5417183	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	5770698	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	4273181	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/055	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/071	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/057	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/061	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/052	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/063	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/064	PFIZER EUROPE MA EEIG	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
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/062	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/008	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/059	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/068	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/007	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/070	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/056	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/060	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/067	PFIZER EUROPE MA EEIG	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
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/066	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/065	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/009	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/053	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/069	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/058	PFIZER EUROPE MA EEIG	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/054	PFIZER EUROPE MA EEIG	LT
Efexor XR 150, capsules met verlengde afgifte, hard, 150 mg	SE/H/0936/003	RVG 20863	PFIZER B.V.	NL
Efexor XR 225 mg cápsulas de libertação prolongada	SE/H/0936/004	SE/H/0936/004	LABORATÓRIOS PFIZER, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor XR 37,5 mg cápsulas de libertação prolongada	SE/H/0936/001	4895181	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 37,5 mg cápsulas de libertação prolongada	SE/H/0936/001	4895280	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/001	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/027	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/012	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/015	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/024	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/013	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/031	PFIZER EUROPE MA EEIG	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
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/022	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/029	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/028	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/030	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/003	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/017	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/010	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/018	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/019	PFIZER EUROPE MA EEIG	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
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/021	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/016	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/011	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/002	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/020	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/023	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/026	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/014	PFIZER EUROPE MA EEIG	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/025	PFIZER EUROPE MA EEIG	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
Efexor XR 37,5, capsules met verlengde afgifte, hard, 37,5 mg	SE/H/0936/001	RVG 26661	PFIZER B.V.	NL
Efexor XR 75 mg cápsulas de libertação prolongada	SE/H/0936/002	4499885	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 75 mg cápsulas de libertação prolongada	SE/H/0936/002	4273082	LABORATÓRIOS PFIZER, LDA.	PT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/044	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/038	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/005	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/047	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/006	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/032	PFIZER EUROPE MA EEIG	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
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/051	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/049	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/037	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/042	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/046	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/050	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/033	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/043	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/041	PFIZER EUROPE MA EEIG	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
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/034	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/048	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/045	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/040	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/004	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/039	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/035	PFIZER EUROPE MA EEIG	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/036	PFIZER EUROPE MA EEIG	LT
Efexor XR 75, capsules met verlengde afgifte, hard, 75 mg	SE/H/0936/002	RVG 20862	PFIZER B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Efexor XR, 150 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/003	317700	PFIZER EUROPE MA EEIG	EE
Efexor XR, 37,5 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/001	352801	PFIZER EUROPE MA EEIG	EE
Efexor XR, 75 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/002	317800	PFIZER EUROPE MA EEIG	EE
Efexor® XR 150 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/003	018334	PFIZER HELLAS, A.E.	CY
Efexor® XR 150 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/003	PRD495475	PFIZER HELLAS, A.E.	GR
Efexor® XR 37,5 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/001	20345	PFIZER HELLAS, A.E.	CY
Efexor® XR 37,5 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/001	30731/6-4-2016	PFIZER HELLAS, A.E.	GR
Efexor® XR 75 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/002	018335	PFIZER HELLAS, A.E.	CY
Efexor® XR 75 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/002	30729/6-4-2016	PFIZER HELLAS, A.E.	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
Efexor-Exel 150 mg gélules à libération prolongée	SE/H/0936/003	BE196533	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg gélules à libération prolongée	SE/H/0936/003	BE421994	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg gélules à libération prolongée	SE/H/0936/003	2011030970	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 150 mg harde capsules met verlengde afgifte	SE/H/0936/003	BE421994	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg harde capsules met verlengde afgifte	SE/H/0936/003	BE196533	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg retardierte Hartkapseln	SE/H/0936/003	BE196533	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg retardierte Hartkapseln	SE/H/0936/003	BE421994	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 150 mg retardierte Hartkapseln	SE/H/0936/003	2011030970	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 225 mg gélules à libération prolongée	SE/H/0936/004	BE467333	PFIZER S.A. (BELGIUM)	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
Efexor-Exel 225 mg gélules à libération prolongée	SE/H/0936/004	2015040065	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 225 mg harde capsules met verlengde afgifte	SE/H/0936/004	BE467333	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 225 mg retardierte Hartkapseln	SE/H/0936/004	BE467333	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 225 mg retardierte Hartkapseln	SE/H/0936/004	2015040065	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 37,5 mg gélules à libération prolongée	SE/H/0936/001	BE239337	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 37,5 mg gélules à libération prolongée	SE/H/0936/001	BE422003	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 37,5 mg gélules à libération prolongée	SE/H/0936/001	2011010961	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 37,5 mg harde capsules met verlengde afgifte	SE/H/0936/001	BE239337	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 37,5 mg harde capsules met verlengde afgifte	SE/H/0936/001	BE422003	PFIZER S.A. (BELGIUM)	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
Efexor-Exel 37,5 mg retardierte Hartkapseln	SE/H/0936/001	BE422003	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 37,5 mg retardierte Hartkapseln	SE/H/0936/001	BE239337	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 37,5 mg retardierte Hartkapseln	SE/H/0936/001	2011010961	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 75 mg gélules à libération prolongée	SE/H/0936/002	BE422012	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 75 mg gélules à libération prolongée	SE/H/0936/002	BE196524	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 75 mg gélules à libération prolongée	SE/H/0936/002	2011010962	PFIZER S.A. (BELGIUM)	LU
Efexor-Exel 75 mg harde capsules met verlengde afgifte	SE/H/0936/002	BE196524	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 75 mg harde capsules met verlengde afgifte	SE/H/0936/002	BE422012	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 75 mg retardierte Hartkapseln	SE/H/0936/002	BE422012	PFIZER S.A. (BELGIUM)	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
Efexor-Exel 75 mg retardierte Hartkapseln	SE/H/0936/002	BE196524	PFIZER S.A. (BELGIUM)	BE
Efexor-Exel 75 mg retardierte Hartkapseln	SE/H/0936/002	2011010962	PFIZER S.A. (BELGIUM)	LU
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 562 7 9	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 565 6 9	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 563 3 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 557 3 9	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 556 7 8	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 376 270 4 7	PFIZER HOLDING FRANCE (S.C.A.)	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 555 0 0	PFIZER HOLDING FRANCE (S.C.A.)	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
ELIFY XR 37.5 mg κάψουλες παρατεταμένης αποδέσμευσης, σκληρές	NL/H/0927/001	20208	MEDOCHEMIE LTD.	CY
ЕФЕКТИН ER 150 mg капсули с удължено освобождаване, твърди	SE/H/0936/003	9800389	PFIZER EUROPE MA EEIG	BG
ЕФЕКТИН ER 75 mg капсули с удължено освобождаване, твърди	SE/H/0936/002	9800390	PFIZER EUROPE MA EEIG	BG
Faxiprol 150 mg retard tabletta	SE/564/01-04/DC	OGYI-T-20519/21-30	MEDICO UNO PHARMACEUTICALS SE	HU
Faxiprol 150 mg retard tabletta	SE/564/01-04/DC	OGYI-T-20519/21-30	MEDICO UNO PHARMACEUTICALS SE	HU
Faxiprol 75 mg retard tabletta	SE/564/01-04/DC	OGYI-T-20519/11-20	MEDICO UNO PHARMACEUTICALS SE	HU
Faxiprol 75 mg retard tabletta	SE/564/01-04/DC	OGYI-T-20519/11-20	MEDICO UNO PHARMACEUTICALS SE	HU
Genexin 150 mg comprimidos de libertação prolongada	not available	5099528	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099536	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Genexin 150 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5099544	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 150 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5099551	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 150 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5099569	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 150 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5099510	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 37,5 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5974399	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 37,5 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5974498	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 37,5 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5974597	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 37,5 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5974290	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 75 mg comprimidos de liberta 7̄7 7̄Jo prolongada	not available	5975198	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Genexin 75 mg comprimidos de libertaﾃｧﾃ｣o prolongada	not available	5975297	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 75 mg comprimidos de libertaﾃｧﾃ｣o prolongada	not available	5975396	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Genexin 75 mg comprimidos de libertaﾃｧﾃ｣o prolongada	not available	5975099	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180922	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180930	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180948	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180963	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180955	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180971	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 150 mg comprimidos de libertação prolongada	not available	5181003	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181029	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181011	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181045	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181078	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181110	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181128	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181136	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181144	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 150 mg comprimidos de libertação prolongada	not available	5181151	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181037	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181052	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181060	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181102	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181169	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181177	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181201	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181219	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 225 mg comprimidos de libertação prolongada	not available	5181227	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181235	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181243	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181250	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181268	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181276	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181300	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181318	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181326	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 225 mg comprimidos de libertação prolongada	not available	5181334	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181342	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181359	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181367	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181375	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181409	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181417	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180443	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180435	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180450	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180427	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180476	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180468	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180500	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180518	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180526	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180534	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180542	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180559	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180567	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180575	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180609	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180617	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180625	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180633	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180641	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180658	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 75 mg comprimidos de libertação prolongada	not available	5180674	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180708	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180716	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180724	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180732	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180740	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180757	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180765	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180773	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 75 mg comprimidos de libertação prolongada	not available	5180807	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180815	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180823	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180831	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180666	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180849	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180856	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180864	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180872	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	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
Pracet 75 mg comprimidos de libertação prolongada	not available	5180914	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180906	UCB PHARMA (PRODUTOS FARMACÊUTICOS), LDA. (PACO PT)	PT
Serosmine 150 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/003	68910	ITF HELLAS AE	GR
Serosmine 150 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/003	68910	ITF HELLAS AE	GR
Serosmine 225 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/004	68911	ITF HELLAS AE	GR
Serosmine 225 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/004	68911	ITF HELLAS AE	GR
Serosmine 37,5 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/001	68912	ITF HELLAS AE	GR
Serosmine 37,5 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/001	68912	ITF HELLAS AE	GR
Serosmine 75 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/002	68909	ITF HELLAS AE	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
Serosmine 75 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/002	68909	ITF HELLAS AE	GR
Sunveniz XL 150 mg prolonged-release tablets	NL/H/3948/003	PL31750/0027	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	UK
Sunveniz XL 37.5 mg prolonged-release tablets	NL/H/3948/001	PL31750/0025	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	UK
Sunveniz XL 75 mg prolonged-release tablets	NL/H/3948/002	PL31750/0026	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	UK
Trevilor 150 mg hard prolonged-release capsules	SE/H/0938/003	42827	PFIZER AB	SE
Trevilor 37.5 mg hard prolonged-release capsules	SE/H/0938/001	42825	PFIZER AB	SE
Trevilor 75 mg hard prolonged-release capsules	SE/H/0938/002	42826	PFIZER AB	SE
Trevilor® retard 150 mg Hartkapseln, retardiert	SE/H/0936/003	45300.01.00	PFIZER PHARMA PFE GMBH	DE
Trevilor® retard 37,5 mg Hartkapseln, retardiert	SE/H/0936/001	62104.00.00	PFIZER PHARMA PFE GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Trevilor® retard 75 mg Hartkapseln, retardiert	SE/H/0936/002	45300.00.00	PFIZER PHARMA PFE GMBH	DE
Vandral Retard 150 mg cápsulas duras de liberación prolongada	SE/H/0936/003	62.402	WYETH FARMA, S.A	ES
Vandral Retard 225 mg cápsulas duras de liberación prolongada	SE/H/0936/004	79.960	WYETH FARMA, S.A	ES
Vandral Retard 75 mg cápsulas duras de liberación prolongada	SE/H/0936/002	62.401	WYETH FARMA, S.A	ES
VENLABRAIN retard 150 mg comprimidos de liberación prolongada	SE/H/0582/002	69.751	FERRER INTERNACIONAL, S.A.	ES
VENLABRAIN retard 150 mg comprimidos de liberación prolongada	SE/H/0582/002	69.751	FERRER INTERNACIONAL, S.A.	ES
VENLABRAIN retard 225 mg comprimidos de liberación prolongada	SE/H/0582/003	69.750	FERRER INTERNACIONAL, S.A.	ES
VENLABRAIN retard 225 mg comprimidos de liberación prolongada	SE/H/0582/003	69.750	FERRER INTERNACIONAL, S.A.	ES
VENLABRAIN retard 75 mg comprimidos de liberación prolongada	SE/H/0582/001	69.752	FERRER INTERNACIONAL, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLABRAIN retard 75 mg comprimidos de liberación prolongada	SE/H/0582/001	69.752	FERRER INTERNACIONAL, S.A.	ES
Venlafaxin "Hexal"	NL/H/1307/001	42725	HEXAL A/S	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/001	58739	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/002	58740	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/003	58741	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/001	58739	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/002	58740	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Medical Valley", depottabletter	SE/H/0582/003	58741	MEDICAL VALLEY INVEST AB	DK
Venlafaxin AbZ 150 mg Retardtabletten	DE/H/3592/003	74546.00.00	ABZ-PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin AbZ 225 mg Retardtabletten	DE/H/3592/004	74547.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 37,5 mg Retardtabletten	DE/H/3592/001	74544.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 75 mg Retardtabletten	DE/H/3592/002	74545.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AGP 150 mg depotabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 150 mg depotabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin AGP 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depotabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depotabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depotabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depottabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depottabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depottabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 225 mg depottabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37,5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin AGP 37,5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37,5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37,5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37.5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37.5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 37.5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depotabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depotabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin AGP 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin AGP 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin Heumann 225 mg Hartkapseln, retardiert	DE/H/4271/001	93933.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Venlafaxin Hexal 50 mg tablett	SE/H/0880/003	25910	HEXAL AG	SE
Venlafaxin Medical Valley 150 mg depottabletter	SE/H/0582/002	23690	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 150 mg depottabletter	SE/H/0582/002	23690	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 150 mg forðatöflur	SE/H/0582/002	IS/1/17/021/02	MEDICAL VALLEY INVEST AB	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin Medical Valley 150 mg forðatöflur	SE/H/0582/002	IS/1/17/021/02	MEDICAL VALLEY INVEST AB	IS
Venlafaxin Medical Valley 225 mg depottabletter	SE/H/0582/003	23691	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 225 mg depottabletter	SE/H/0582/003	23691	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 225 mg forðatöflur	SE/H/0582/003	IS/1/17/021/03	MEDICAL VALLEY INVEST AB	IS
Venlafaxin Medical Valley 225 mg forðatöflur	SE/H/0582/003	IS/1/17/021/03	MEDICAL VALLEY INVEST AB	IS
Venlafaxin Medical Valley 37,5 mg depottabletter	SE/H/0582/004	42949	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 37,5 mg depottabletter	SE/H/0582/004	42949	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 75 mg depottabletter	SE/H/0582/001	23689	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 75 mg depottabletter	SE/H/0582/001	23689	MEDICAL VALLEY INVEST AB	SE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin Medical Valley 75 mg forðatöflur	SE/H/0582/001	IS/1/17/021/01	MEDICAL VALLEY INVEST AB	IS
Venlafaxin Medical Valley 75 mg forðatöflur	SE/H/0582/001	IS/1/17/021/01	MEDICAL VALLEY INVEST AB	IS
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 150 MG RETARDTABLETTEN	SE/H/0582/002	66084.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	**MRP/DCP Authorisation number**	**National Authorisation Number**	**MAH of product in the member state**	**Member State where product is authorised**
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 225 MG RETARDTABLETTEN	SE/H/0582/003	66085.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin Winthrop® osmo 37,5 mg Retardtabletten	SE/H/0582/004	79446.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXIN WINTHROP® OSMO 75 MG RETARDTABLETTEN	SE/H/0582/001	66083.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Venlafaxina Aurobindo 37,5 mg cápsulas duras de liberación prolongada	PT/H/0703/001	79.387	LABORATORIOS AUROBINDO S.L.U.	ES
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691015	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691027	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691039	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691041	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691054	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691066	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691078	AUROBINDO PHARMA (ITALIA) S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691080	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691092	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691104	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691116	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691128	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691130	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691142	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Aurobindo 37,5 mg capsule rigide a rilascio prolungato	PT/H/0703/001	041691155	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Venlafaxina Pfizer 150 mg capsule rigide a rilascio prolungato	SE/H/0938/003	028834101	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxina Pfizer 150 mg capsule rigide a rilascio prolungato	SE/H/0938/003	028834063	PFIZER ITALIA S.R.L.	IT
Venlafaxina Pfizer 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834125	PFIZER ITALIA S.R.L.	IT
Venlafaxina Pfizer 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834137	PFIZER ITALIA S.R.L.	IT
Venlafaxina Pfizer 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834113	PFIZER ITALIA S.R.L.	IT
Venlafaxina Pfizer 75 mg capsule rigide a rilascio prolungato	SE/H/0938/002	028834099	PFIZER ITALIA S.R.L.	IT
Venlafaxina Pfizer 75 mg capsule rigide a rilascio prolungato	SE/H/0938/002	028834051	PFIZER ITALIA S.R.L.	IT
Venlafaxina SUN 150 mg comprimidos de liberación prolongada	NL/H/3948/003	76603	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Venlafaxina SUN 37,5 mg comprimidos de liberación prolongada	NL/H/3948/001	76602	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Venlafaxina SUN 75 mg comprimidos de liberación prolongada	NL/H/3948/002	76604	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxina toLife 150 mg comprimidos de libertação prolongada	not available	5073226	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 150 mg comprimidos de libertação prolongada	not available	5073234	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 150 mg comprimidos de libertação prolongada	not available	5073242	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 37,5 mg comprimidos de libertação prolongada	not available	5708995	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 37,5 mg comprimidos de libertação prolongada	not available	5709092	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 75 mg comprimidos de libertação prolongada	not available	5709191	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxina toLife 75 mg comprimidos de libertação prolongada	not available	5709290	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Venlafaxin-CT 150 mg Retardtabletten	DE/H/3593/003	74550.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin-CT 225 mg Retardtabletten	DE/H/3593/004	74551.00.00	ABZ-PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxin-CT 37,5 mg Retardtabletten	DE/H/3593/001	74548.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin-CT 75 mg Retardtabletten	DE/H/3593/002	74549.00.00	ABZ-PHARMA GMBH	DE
Venlafaxine 150 mg Focus, tabletten met verlengde afgifte	not available	RVG 108591	FOCUS CARE PHARMACEUTICALS B.V.	NL
Venlafaxine 225 mg Focus, tabletten met verlengde afgifte	not available	RVG 108592	FOCUS CARE PHARMACEUTICALS B.V.	NL
Venlafaxine 37,5 mg Focus, tabletten met verlengde afgifte	not available	RVG 108589	FOCUS CARE PHARMACEUTICALS B.V.	NL
Venlafaxine 75 mg Focus, tabletten met verlengde afgifte	not available	RVG 108590	FOCUS CARE PHARMACEUTICALS B.V.	NL
Venlafaxine Aurobindo 225 mg tabletten met verlengde afgifte	SE/H/0581/004	RVG 33889	AUROBINDO PHARMA B.V.	NL
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 482 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 483 3 7	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 485 6 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 487 9 5	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 488 5 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 489 1 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 491 6 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 492 2 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 572 358 8 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 494 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 495 1 8	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 496 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 497 4 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 498 0 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 499 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 500 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 501 1 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 572 359 4 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 486 2 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 493 9 6	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 482 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 483 3 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 485 6 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 487 9 5	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 488 5 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 489 1 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 491 6 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 492 2 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 572 358 8 8	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 494 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 495 1 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 496 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 497 4 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 498 0 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 499 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 500 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 501 1 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 572 359 4 9	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 486 2 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 150 mg, comprimé à libération prolongée	SE/H/0581/003	34009 384 493 9 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 502 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 503 4 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 504 0 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 505 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 506 3 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 508 6 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 509 2 7	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 510 0 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 511 7 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 572 360 2 1	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 512 3 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 514 6 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 515 2 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 516 9 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 517 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 518 1 8	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 519 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 520 6 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 521 2 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 572 361 9 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 502 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 503 4 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 504 0 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 505 7 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 506 3 7	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 508 6 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 509 2 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 510 0 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 511 7 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 572 360 2 1	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 512 3 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 514 6 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 515 2 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 516 9 6	MYLAN MEDICAL SAS	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
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 517 5 7	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 518 1 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 519 8 6	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 520 6 8	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 384 521 2 9	MYLAN MEDICAL SAS	FR
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 572 361 9 9	MYLAN MEDICAL SAS	FR
Venlafaxine Liconsa 150 mg depottabletter	SE/H/0581/003	23687	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 150 mg depottabletter	SE/H/0581/003	23687	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 150 mg depottabletter	SE/H/0581/003	23687	LABORATORIOS LICONSA, S.A.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxine Liconsa 150 mg depottabletter	SE/H/0581/003	23687	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 150 mg tabletten met verlengde afgifte	SE/H/0581/003	RVG 33888	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 150 mg tabletten met verlengde afgifte	SE/H/0581/003	RVG 33888	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 150 mg tabletten met verlengde afgifte	SE/H/0581/003	RVG 33888	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 150 mg tabletten met verlengde afgifte	SE/H/0581/003	RVG 33888	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 225 mg depottabletter	SE/H/0581/004	23688	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 225 mg depottabletter	SE/H/0581/004	23688	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 225 mg depottabletter	SE/H/0581/004	23688	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 225 mg depottabletter	SE/H/0581/004	23688	LABORATORIOS LICONSA, S.A.	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxine Liconsa 225 mg tabletten met verlengde afgifte	SE/H/0581/004	RVG 33889	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 225 mg tabletten met verlengde afgifte	SE/H/0581/004	RVG 33889	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 37,5 mg depottabletter	SE/H/0581/001	23685	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 37,5 mg depottabletter	SE/H/0581/001	23685	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 37,5 mg depottabletter	SE/H/0581/001	23685	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 37,5 mg depottabletter	SE/H/0581/001	23685	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 37,5 mg tabletten met verlengde afgifte	SE/H/0581/001	RVG 33886	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 37,5 mg tabletten met verlengde afgifte	SE/H/0581/001	RVG 33886	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 37,5 mg tabletten met verlengde afgifte	SE/H/0581/001	RVG 33886	LABORATORIOS LICONSA, S.A.	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
Venlafaxine Liconsa 37,5 mg tabletten met verlengde afgifte	SE/H/0581/001	RVG 33886	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 75 mg depottabletter	SE/H/0581/002	23686	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 75 mg depottabletter	SE/H/0581/002	23686	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 75 mg depottabletter	SE/H/0581/002	23686	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 75 mg depottabletter	SE/H/0581/002	23686	LABORATORIOS LICONSA, S.A.	SE
Venlafaxine Liconsa 75 mg tabletten met verlengde afgifte	SE/H/0581/002	RVG 33887	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 75 mg tabletten met verlengde afgifte	SE/H/0581/002	RVG 33887	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 75 mg tabletten met verlengde afgifte	SE/H/0581/002	RVG 33887	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 75 mg tabletten met verlengde afgifte	SE/H/0581/002	RVG 33887	LABORATORIOS LICONSA, S.A.	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
Venlafaxine Liconsa, 150 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/003	553307	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 150 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/003	553307	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 150 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/003	553307	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 150 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/003	553307	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 225 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/004	553207	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 225 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/004	553207	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 225 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/004	553207	LABORATORIOS LICONSA, S.A.	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
Venlafaxine Liconsa, 225 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/004	553207	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 37,5 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/001	553407	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 37,5 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/001	553407	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 37,5 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/001	553407	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 37,5 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/001	553407	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 75 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/002	553507	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 75 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/002	553507	LABORATORIOS LICONSA, S.A.	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
Venlafaxine Liconsa, 75 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/002	553507	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa, 75 mg toimeainet prolongeeritult vabastavad tabletid	SE/H/0581/002	553507	LABORATORIOS LICONSA, S.A.	EE
VENLAFAXINE PFIZER L.P. 37,5 mg, gélule à libération prolongée	SE/H/0938/001	34009 346 617 6 1	PFIZER HOLDING FRANCE (S.C.A.)	FR
VENLAFAXINE PFIZER L.P. 37,5 mg, gélule à libération prolongée	SE/H/0938/001	34009 346 618 2 2	PFIZER HOLDING FRANCE (S.C.A.)	FR
VENLAFAXINE PFIZER L.P. 37,5 mg, gélule à libération prolongée	SE/H/0938/001	34009 346 619 9 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34009 346 567 9 8	PFIZER HOLDING FRANCE (S.C.A.)	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34009 346 566 2 0	PFIZER HOLDING FRANCE (S.C.A.)	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34 009 346 568 5 9	PFIZER HOLDING FRANCE (S.C.A.)	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
Venlafaxine SUN 150 mg tabletten met verlengde afgifte	NL/H/3948/003	RVG 105066	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Venlafaxine SUN 37,5 mg tabletten met verlengde afgifte	NL/H/3948/001	RVG 105061	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Venlafaxine SUN 75 mg tabletten met verlengde afgifte	NL/H/3948/002	RVG 105065	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Venlafaxine Teva 150 mg comprimés à libération prolongée	DE/H/3591/003	BE347365	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 150 mg comprimés à libération prolongée	DE/H/3591/003	BE347356	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 150 mg RETARDTABLETTEN	DE/H/3591/003	BE347365	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 150 mg RETARDTABLETTEN	DE/H/3591/003	BE347356	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 150 mg tabletten met verlengde afgifte	DE/H/3591/003	BE347356	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 150 mg tabletten met verlengde afgifte	DE/H/3591/003	BE347365	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxine Teva 225 mg comprimés à libération prolongée	DE/H/3591/004	BE347374	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 225 mg comprimés à libération prolongée	DE/H/3591/004	BE347383	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 225 mg RETARDTABLETTEN	DE/H/3591/004	BE347374	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 225 mg RETARDTABLETTEN	DE/H/3591/004	BE347383	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 225 mg tabletten met verlengde afgifte	DE/H/3591/004	BE347374	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 225 mg tabletten met verlengde afgifte	DE/H/3591/004	BE347383	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 225 mg tabletten met verlengde afgifte	DE/H/3591/001	BE347313	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 37,5 mg comprimés à libération prolongée	DE/H/3591/001	BE347313	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 37,5 mg comprimés à libération prolongée	DE/H/3591/001	BE347322	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAFAXINE TEVA 37,5 mg RETARDTABLETTEN	DE/H/3591/001	BE347313	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 37,5 mg RETARDTABLETTEN	DE/H/3591/001	BE347322	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 37,5 mg tabletten met verlengde afgifte	DE/H/3591/001	BE347322	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 75 mg comprimés à libération prolongée	DE/H/3591/002	BE347331	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 75 mg comprimés à libération prolongée	DE/H/3591/002	BE347347	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 75 mg RETARDTABLETTEN	DE/H/3591/002	BE347331	TEVA PHARMA BELGIUM N.V./S.A	BE
VENLAFAXINE TEVA 75 mg RETARDTABLETTEN	DE/H/3591/002	BE347347	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 75 mg tabletten met verlengde afgifte	DE/H/3591/002	BE347331	TEVA PHARMA BELGIUM N.V./S.A	BE
Venlafaxine Teva 75 mg tabletten met verlengde afgifte	DE/H/3591/002	BE347347	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlafaxine XR Pfizer 150 mg, capsules met verlengde afgifte, hard, 150 mg	SE/H/0938/003	RVG 107029	PFIZER B.V.	NL
Venlafaxine XR Pfizer 37,5 mg, capsules met verlengde afgifte, hard, 37,5 mg	SE/H/0938/001	RVG 107027	PFIZER B.V.	NL
Venlafaxine XR Pfizer 75 mg, capsules met verlengde afgifte, hard, 75 mg	SE/H/0938/002	RVG 107028	PFIZER B.V.	NL
Venlafaxin-neuraxpharm 225 mg retard Retardtabletten	not available	73279.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Venlafaxin-ratiopharm® 150 mg Retardtabletten	DE/H/3591/003	74542.00.00	RATIOPHARM GMBH	DE
Venlafaxin-ratiopharm® 225 mg Retardtabletten	DE/H/3591/004	74543.00.00	RATIOPHARM GMBH	DE
Venlafaxin-ratiopharm® 37,5 mg Retardtabletten	DE/H/3591/001	74540.00.00	RATIOPHARM GMBH	DE
Venlafaxin-ratiopharm® 75 mg Retardtabletten	DE/H/3591/002	74541.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlalic XL 150 mg prolonged-release tablets	SE/H/0581/003	PA1969/001/002	ETHYPHARM UK LTD	IE
Venlalic XL 150 mg prolonged-release tablets	SE/H/0581/003	PA1969/001/002	ETHYPHARM UK LTD	IE
Venlalic XL 150 mg prolonged-release tablets	SE/H/0581/003	PL 42623/0019	ETHYPHARM UK LTD	UK
Venlalic XL 150 mg prolonged-release tablets	SE/H/0581/003	PL 42623/0019	ETHYPHARM UK LTD	UK
Venlalic XL 225 mg prolonged-release tablets	SE/H/0581/004	PA 1969/001/003	ETHYPHARM UK LTD	IE
Venlalic XL 225 mg prolonged-release tablets	SE/H/0581/004	PA 1969/001/003	ETHYPHARM UK LTD	IE
Venlalic XL 225 mg prolonged-release tablets	SE/H/0581/004	PL 42623/0020	ETHYPHARM UK LTD	UK
Venlalic XL 225 mg prolonged-release tablets	SE/H/0581/004	PL 42623/0020	ETHYPHARM UK LTD	UK
Venlalic XL 37.5 mg prolonged-release tablets	SE/H/0581/001	PA1969/001/004	ETHYPHARM UK LTD	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
Venlalic XL 37.5 mg prolonged-release tablets	SE/H/0581/001	PA1969/001/004	ETHYPHARM UK LTD	IE
Venlalic XL 37.5 mg prolonged-release tablets	SE/H/0581/001	PL 42623/0017	ETHYPHARM UK LTD	UK
Venlalic XL 37.5 mg prolonged-release tablets	SE/H/0581/001	PL 42623/0017	ETHYPHARM UK LTD	UK
Venlalic XL 75 mg prolonged-release tablets	SE/H/0581/002	PA1969/001/001	ETHYPHARM UK LTD	IE
Venlalic XL 75 mg prolonged-release tablets	SE/H/0581/002	PA1969/001/001	ETHYPHARM UK LTD	IE
Venlalic XL 75 mg prolonged-release tablets	SE/H/0581/002	PL 42623/0018	ETHYPHARM UK LTD	UK
Venlalic XL 75 mg prolonged-release tablets	SE/H/0581/002	PL 42623/0018	ETHYPHARM UK LTD	UK
Venlaneo XL 150 mg Prolonged Release Capsules	not available	PL 20417/0085	FANNIN (UK) LIMITED	UK
Venlaneo XL 75 mg Prolonged Release Capsules	not available	PL 20417/0087	FANNIN (UK) LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VENLAXIN 150 mg δισκία παρατεταμένης αποδέσμευσης	not available	021936	IASIS PHARMA	CY
VENLAXIN 150 mg δισκία παρατεταμένης αποδέσμευσης	not available	108419/14/18-05-2015	IASIS PHARMA	GR
VENLAXIN 225 mg δισκία παρατεταμένης αποδέσμευσης	not available	021937	IASIS PHARMA	CY
VENLAXIN 225 mg δισκία παρατεταμένης αποδέσμευσης	not available	108419/14/18-05-2015	IASIS PHARMA	GR
VENLAXIN 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	021935	IASIS PHARMA	CY
VENLAXIN 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	108419/14/18-05-2015	IASIS PHARMA	GR
Venlazid 150 mg depottabletter	SE/H/0582/002	16-11408	MEDICAL VALLEY INVEST AB	NO
Venlazid 150 mg depottabletter	SE/H/0582/002	16-11408	MEDICAL VALLEY INVEST AB	NO
Venlazid 225 mg depottabletter	SE/H/0582/003	16-11409	MEDICAL VALLEY INVEST AB	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venlazid 225 mg depottabletter	SE/H/0582/003	16-11409	MEDICAL VALLEY INVEST AB	NO
Venlazid 75 mg depottabletter	SE/H/0582/001	16-11407	MEDICAL VALLEY INVEST AB	NO
Venlazid 75 mg depottabletter	SE/H/0582/001	16-11407	MEDICAL VALLEY INVEST AB	NO
Venxin 150 mg comprimidos de libertação prolongada	not available	5099635	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099643	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099577	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099601	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099619	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099627	DECOMED FARMACÊUTICA, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974795	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974894	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974993	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974696	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975594	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975693	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975792	DECOMED FARMACÊUTICA, S.A.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975495	DECOMED FARMACÊUTICA, S.A.	PT
VENZIP XL 37.5 mg prolonged-release capsules, hard	PT/H/0703/001	MA807/04601	AUROBINDO PHARMA (MALTA) LIMITED	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316218	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316220	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316232	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316244	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316257	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316269	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316271	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316283	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316295	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316307	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316319	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316321	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316333	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316345	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316358	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316360	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316372	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316384	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316396	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316408	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316218	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316220	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316232	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316244	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316257	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316269	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316271	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316283	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316295	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316307	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316319	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316321	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316333	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316345	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316358	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316360	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316372	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316384	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316396	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/02/MR	038316408	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316410	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316422	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316434	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316446	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316459	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316461	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316473	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316485	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316497	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316509	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316511	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316523	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316535	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316547	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316550	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316562	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316574	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316586	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316598	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316600	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316410	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316422	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316434	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316446	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316459	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316461	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316473	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316485	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316497	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316509	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316511	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316523	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316535	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316547	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316550	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316562	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316574	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316586	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316598	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/03/MR	038316600	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316612	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316624	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316636	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316648	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316651	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316663	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316675	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316687	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316699	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316701	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316713	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316725	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316737	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316749	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316752	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316764	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316776	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316788	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316790	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316802	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316612	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316624	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316636	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316648	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316651	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316663	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316675	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316687	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316699	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316701	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316713	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316725	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316737	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316749	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316752	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316764	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316776	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316788	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316790	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/04/MR	038316802	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316016	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316028	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316030	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316042	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316055	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316067	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316079	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316081	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316093	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316105	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316117	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316129	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316131	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316143	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316156	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316168	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316170	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316182	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316194	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316206	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316016	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316028	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316030	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316042	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316055	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316067	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316079	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316081	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316093	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316105	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316117	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316129	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316131	ITALFARMACO S.P.A	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316143	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316156	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316168	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316170	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316182	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316194	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/01/MR	038316206	ITALFARMACO S.P.A	IT
Zarelis Retard 150 mg comprimidos de liberación prolongada	SE/H/0564/003	69.738	ITALFARMACO S.A.	ES
Zarelis Retard 150 mg comprimidos de liberación prolongada	SE/H/0564/003	69.738	ITALFARMACO S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarelis Retard 225 mg comprimidos de liberación prolongada	SE/H/0564/004	69.739	ITALFARMACO S.A.	ES
Zarelis Retard 225 mg comprimidos de liberación prolongada	SE/H/0564/004	69.739	ITALFARMACO S.A.	ES
Zarelis Retard 75 mg comprimidos de liberación prolongada	SE/H/0564/002	69.737	ITALFARMACO S.A.	ES
Zarelis Retard 75 mg comprimidos de liberación prolongada	SE/H/0564/002	69.737	ITALFARMACO S.A.	ES