26 January 2023
EMA/115290/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): venlafaxine

Procedure No. PSUSA/00003104/202205

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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
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, depottabletter	SE/H/0564/002	39249	LABORATORIOS LICONSA, S.A.	DK
Duofaxin, depottabletter	SE/H/0564/003	39250	LABORATORIOS LICONSA, S.A.	DK
Duofaxin, depottabletter	SE/H/0564/004	39251	LABORATORIOS LICONSA, S.A.	DK
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/19	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/18	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/21	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/20	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/07	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare	SE/H/0936/003	7977/2015/08	UPJOHN EESV	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
prelungită				
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/23	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/22	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/09	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/24	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/10	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/11	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/02	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/12	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/04	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/05	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/13	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/03	UPJOHN EESV	RO
EFECTIN EP 150 mg	SE/H/0936/003	7977/2015/01	UPJOHN EESV	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
capsule cu eliberare prelungită				
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/16	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/14	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/15	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/06	UPJOHN EESV	RO
EFECTIN EP 150 mg capsule cu eliberare prelungită	SE/H/0936/003	7977/2015/17	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/12	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/13	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/10	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/16	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/15	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/14	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/17	UPJOHN EESV	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	7975/2015/19	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/02	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/01	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/18	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/20	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/03	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/22	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/07	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/05	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/24	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/21	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/04	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare	SE/H/0936/001	7975/2015/09	UPJOHN EESV	RO

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prelungită				
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/23	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/11	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/06	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/25	UPJOHN EESV	RO
EFECTIN EP 37,5 mg capsule cu eliberare prelungită	SE/H/0936/001	7975/2015/08	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/16	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/02	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/15	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/17	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/01	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/18	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/03	UPJOHN EESV	RO
EFECTIN EP 75 mg	SE/H/0936/002	7976/2015/19	UPJOHN EESV	RO

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capsule cu eliberare prelungită				
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/04	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/05	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/21	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/06	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/23	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/08	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/20	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/09	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/07	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/22	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/24	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/11	UPJOHN EESV	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	7976/2015/10	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/13	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/12	UPJOHN EESV	RO
EFECTIN EP 75 mg capsule cu eliberare prelungită	SE/H/0936/002	7976/2015/14	UPJOHN EESV	RO
Efectin ER 150 mg Hartkapseln, retardiert	SE/H/0936/003	1-23043	UPJOHN EESV	AT
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/097	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/087	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/095	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/100	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/101	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/098	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/107	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/108	UPJOHN EESV	SI

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EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/094	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/103	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/086	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/096	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/106	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/081	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/093	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/085	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/105	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/082	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/080	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/104	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim	SE/H/0936/003	H/00/00529/077	UPJOHN EESV	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
sproščanjem				
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/102	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/078	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/076	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/099	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/089	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/088	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/083	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/092	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/074	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/084	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/090	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/079	UPJOHN EESV	SI
EFECTIN ER 150 mg trde	SE/H/0936/003	H/00/00529/073	UPJOHN EESV	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
kapsule s podaljšanim sproščanjem				
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/091	UPJOHN EESV	SI
EFECTIN ER 150 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/003	H/00/00529/075	UPJOHN EESV	SI
EFECTIN ER 150 mg tvrdé tobolky s prodlouženým uvolňováním	SE/H/0936/003	30/687/99-C	UPJOHN EESV	CZ
Efectin ER 150, 150 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/003	4940	UPJOHN EESV	PL
Efectin ER 37,5 mg Hartkapseln, retardiert	SE/H/0936/001	1-24637	UPJOHN EESV	AT
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/025	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/031	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/021	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/028	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/026	UPJOHN EESV	SI
EFECTIN ER 37,5 mg	SE/H/0936/001	H/00/00529/023	UPJOHN EESV	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
trde kapsule s podaljšanim sproščanjem				
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/024	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/015	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/030	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/027	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/018	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/034	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/019	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/035	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim	SE/H/0936/001	H/00/00529/016	UPJOHN EESV	SI

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sproščanjem				
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/020	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/036	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/012	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/022	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/010	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/017	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/011	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/014	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/032	UPJOHN EESV	SI
EFECTIN ER 37,5 mg	SE/H/0936/001	H/00/00529/033	UPJOHN EESV	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
trde kapsule s podaljšanim sproščanjem				
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/013	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/029	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/006	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/008	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/007	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/009	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/001	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/002	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim	SE/H/0936/001	H/00/00529/005	UPJOHN EESV	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
sproščanjem				
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/004	UPJOHN EESV	SI
EFECTIN ER 37,5 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/001	H/00/00529/003	UPJOHN EESV	SI
Efectin ER 37,5, 37,5 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/001	10523	UPJOHN EESV	PL
Efectin ER 75 mg Hartkapseln, retardiert	SE/H/0936/002	1-23042	UPJOHN EESV	AT
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/066	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/064	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/061	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/065	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/054	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/056	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/057	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim	SE/H/0936/002	H/00/00529/053	UPJOHN EESV	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
sproščanjem				
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/048	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/063	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/071	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/043	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/072	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/055	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/068	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/052	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/058	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/051	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/042	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/069	UPJOHN EESV	SI
EFECTIN ER 75 mg trde	SE/H/0936/002	H/00/00529/037	UPJOHN EESV	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
kapsule s podaljšanim sproščanjem				
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/044	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/049	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/039	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/045	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/050	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/046	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/070	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/040	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/038	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/067	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/041	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/062	UPJOHN EESV	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/00/00529/060	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/059	UPJOHN EESV	SI
EFECTIN ER 75 mg trde kapsule s podaljšanim sproščanjem	SE/H/0936/002	H/00/00529/047	UPJOHN EESV	SI
EFECTIN ER 75 mg tvrdé tobolky s prodlouženým uvolňováním	SE/H/0936/002	30/686/99-C	UPJOHN EESV	CZ
Efectin ER 75, 75 mg, kapsułki o przedłużonym uwalnianiu, twarde	SE/H/0936/002	4939	UPJOHN EESV	PL
Efexor 150 mg capsule rigide a rilascio prolungato	SE/H/0936/003	028831105	VIATRIS PHARMA S.R.L.	IT
Efexor 150 mg capsule rigide a rilascio prolungato	SE/H/0936/003	028831067	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831307	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831295	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831283	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831319	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831232	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio	SE/H/0936/004	028831206	VIATRIS PHARMA 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
prolungato				
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831194	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831257	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831182	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831269	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831218	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831271	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831220	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831156	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831170	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831244	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831143	VIATRIS PHARMA S.R.L.	IT
Efexor 225 mg capsule rigide a rilascio prolungato	SE/H/0936/004	028831168	VIATRIS PHARMA S.R.L.	IT
Efexor 37,5 mg capsule	SE/H/0936/001	028831129	VIATRIS PHARMA 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
rigide a rilascio prolungato				
Efexor 37,5 mg capsule rigide a rilascio prolungato	SE/H/0936/001	028831117	VIATRIS PHARMA S.R.L.	IT
Efexor 37,5 mg capsule rigide a rilascio prolungato	SE/H/0936/001	028831131	VIATRIS PHARMA S.R.L.	IT
Efexor 75 mg capsule rigide a rilascio prolungato	SE/H/0936/002	028831093	VIATRIS PHARMA S.R.L.	IT
Efexor 75 mg capsule rigide a rilascio prolungato	SE/H/0936/002	028831055	VIATRIS PHARMA S.R.L.	IT
Efexor Depot 150 mg depotkapsel, hard	SE/H/0936/003	97-4783	UPJOHN EESV	NO
Efexor Depot 150 mg depotkapsel, hård	SE/H/0936/003	12607	UPJOHN EESV	FI
Efexor Depot 150 mg depotkapsel, hård	SE/H/0936/003	13321	UPJOHN EESV	SE
Efexor Depot 150 mg forðahylki, hart.	SE/H/0936/003	970068	UPJOHN EESV	IS
Efexor Depot 225 mg Depotkapsel, hård	SE/H/0936/004	49965	UPJOHN EESV	SE
Efexor Depot 37,5 mg depotkapsel, hard	SE/H/0936/001	04-3083	UPJOHN EESV	NO
Efexor Depot 37,5 mg depotkapsel, hård	SE/H/0936/001	21800	UPJOHN EESV	SE
Efexor Depot 37,5 mg depotkapseli, kova	SE/H/0936/001	20185	UPJOHN EESV	FI
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	UPJOHN EESV	NO
Efexor Depot 75 mg depotkapsel, hård	SE/H/0936/002	12606	UPJOHN EESV	FI
Efexor Depot 75 mg depotkapsel, hård	SE/H/0936/002	13320	UPJOHN EESV	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
Efexor Depot 75 mg forðahylki, hart.	SE/H/0936/002	970067	UPJOHN EESV	IS
Efexor Depot, hårde depotkapsler	SE/H/0936/003	18578	UPJOHN EESV	DK
Efexor Depot, hårde depotkapsler	SE/H/0936/002	18577	UPJOHN EESV	DK
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	PL 50622/0018	UPJOHN UK LIMITED	XI
Efexor XL 225 mg prolonged-release capsules, hard	SE/H/0936/004	PL 50622/0019	UPJOHN UK LIMITED	XI
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	MA1396/01301	UPJOHN HELLAS L.T.D.	MT
Efexor XL 75 mg prolonged-release capsules, hard	SE/H/0936/002	PL 50622/0020	UPJOHN UK LIMITED	XI
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	5417183	UPJOHN EESV	PT
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	5770698	UPJOHN EESV	PT
Efexor XR 150 mg cápsulas de libertação prolongada	SE/H/0936/003	4273181	UPJOHN EESV	PT
Efexor XR 150 mg pailginto atpalaidavimo	SE/H/0936/003	LT/1/2000/1045/055	UPJOHN EESV	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
kietosios kapsulės				
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/071	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/057	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/061	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/052	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/063	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/064	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/062	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/008	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/059	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/068	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/007	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/070	UPJOHN EESV	LT
Efexor XR 150 mg	SE/H/0936/003	LT/1/2000/1045/056	UPJOHN EESV	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
pailginto atpalaidavimo kietosios kapsulės				
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/060	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/067	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/066	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/065	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/009	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/053	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/069	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/058	UPJOHN EESV	LT
Efexor XR 150 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/003	LT/1/2000/1045/054	UPJOHN EESV	LT
Efexor XR 150, capsules met verlengde afgifte, hard, 150 mg	SE/H/0936/003	RVG 20863	VIATRIS NETHERLANDS B.V.	NL
Efexor XR 225 mg cápsulas de libertação prolongada	SE/H/0936/004	SE/H/0936/004	UPJOHN EESV	PT
Efexor XR 37,5 mg cápsulas de libertação prolongada	SE/H/0936/001	4895181	UPJOHN EESV	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	4895280	UPJOHN EESV	PT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/001	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/027	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/012	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/015	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/024	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/013	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/031	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/022	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/029	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/028	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/030	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo	SE/H/0936/001	LT/1/2000/1045/003	UPJOHN EESV	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
kietosios kapsulės				
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/017	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/010	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/018	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/019	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/021	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/016	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/011	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/002	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/020	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/023	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/026	UPJOHN EESV	LT
Efexor XR 37,5 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/001	LT/1/2000/1045/014	UPJOHN EESV	LT
Efexor XR 37,5 mg	SE/H/0936/001	LT/1/2000/1045/025	UPJOHN EESV	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
pailginto atpalaidavimo kietosios kapsulės				
Efexor XR 37,5, capsules met verlengde afgifte, hard, 37,5 mg	SE/H/0936/001	RVG 26661	VIATRIS NETHERLANDS B.V.	NL
Efexor XR 75 mg cápsulas de libertação prolongada	SE/H/0936/002	4499885	UPJOHN EESV	PT
Efexor XR 75 mg cápsulas de libertação prolongada	SE/H/0936/002	4273082	UPJOHN EESV	PT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/044	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/038	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/005	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/047	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/006	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/032	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/051	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/049	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/037	UPJOHN EESV	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/1/2000/1045/042	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/046	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/050	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/033	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/043	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/041	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/034	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/048	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/045	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/040	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/004	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/039	UPJOHN EESV	LT
Efexor XR 75 mg pailginto atpalaidavimo	SE/H/0936/002	LT/1/2000/1045/035	UPJOHN EESV	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
kietosios kapsulės				
Efexor XR 75 mg pailginto atpalaidavimo kietosios kapsulės	SE/H/0936/002	LT/1/2000/1045/036	UPJOHN EESV	LT
Efexor XR 75, capsules met verlengde afgifte, hard, 75 mg	SE/H/0936/002	RVG 20862	VIATRIS NETHERLANDS B.V.	NL
Efexor XR, 150 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/003	317700	UPJOHN EESV	EE
Efexor XR, 37,5 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/001	352801	UPJOHN EESV	EE
Efexor XR, 75 mg toimeainet prolongeeritult vabastavad kõvakapslid	SE/H/0936/002	317800	UPJOHN EESV	EE
Efexor® XR 150 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/003	018334	UPJOHN HELLAS L.T.D.	CY
Efexor® XR 150 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/003	30730/6-4-2016	UPJOHN HELLAS L.T.D.	GR
Efexor® XR 37,5 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/001	20345	UPJOHN HELLAS L.T.D.	CY
Efexor® XR 37,5 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/001	30731/6-4-2016	UPJOHN HELLAS L.T.D.	GR
Efexor® XR 75 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/002	018335	UPJOHN HELLAS L.T.D.	CY
Efexor® XR 75 mg καψάκια παρατεταμένης αποδέσμευσης, σκληρά	SE/H/0936/002	30729/6-4-2016	UPJOHN HELLAS L.T.D.	GR
Efexor-Exel 150 mg	SE/H/0936/003	BE421994	UPJOHN SRL	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
harde capsules met verlengde afgifte				
Efexor-Exel 150 mg harde capsules met verlengde afgifte	SE/H/0936/003	BE196533	UPJOHN SRL	BE
Efexor-Exel 150 mg retardierte Hartkapseln	SE/H/0936/003	2011030970	UPJOHN SRL	LU
Efexor-Exel 225 mg gélules à libération prolongée	SE/H/0936/004	2015040065	UPJOHN SRL	LU
Efexor-Exel 225 mg harde capsules met verlengde afgifte	SE/H/0936/004	BE467333	UPJOHN SRL	BE
Efexor-Exel 37,5 mg gélules à libération prolongée	SE/H/0936/001	2011010961	UPJOHN SRL	LU
Efexor-Exel 37,5 mg harde capsules met verlengde afgifte	SE/H/0936/001	BE239337	UPJOHN SRL	BE
Efexor-Exel 37,5 mg harde capsules met verlengde afgifte	SE/H/0936/001	BE422003	UPJOHN SRL	BE
Efexor-Exel 75 mg harde capsules met verlengde afgifte	SE/H/0936/002	BE196524	UPJOHN SRL	BE
Efexor-Exel 75 mg harde capsules met verlengde afgifte	SE/H/0936/002	BE422012	UPJOHN SRL	BE
Efexor-Exel 75 mg retardierte Hartkapseln	SE/H/0936/002	2010110962	UPJOHN SRL	LU
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 147 6 7	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 585 421 5 2	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération	SE/H/0936/001	34009 273 132 8 8	VIATRIS UP	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
prolongée				
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 129 7 7	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 151 3 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 158 8 7	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 131 1 0	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 160 2 0	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 154 2 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 122 2 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 124 5 8	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 128 0 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 149 9 6	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 125 1 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 276 156 5 8	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg,	SE/H/0936/001	34009 276 144 7 7	VIATRIS UP	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
gélule à libération prolongée				
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 130 5 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 127 4 8	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 585 751 5 0	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 126 8 7	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 273 123 9 7	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 562 7 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 565 6 9	VIATRIS UP	FR
EFFEXOR L.P. 37,5 mg, gélule à libération prolongée	SE/H/0936/001	34009 346 563 3 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 171 4 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 195 0 2	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 187 8 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 168 3 9	VIATRIS UP	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
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 191 5 1	PFIZER PFE FRANCE	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 176 6 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 189 0 1	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 193 8 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 276 178 9 8	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 151 2 1	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 156 4 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 155 8 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 585 755 0 1	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 154 1 1	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 150 6 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 585 753 8 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération	SE/H/0936/002	34009 585 425 0 3	VIATRIS UP	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
prolongée				
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 153 5 0	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 585 424 4 2	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 149 8 8	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 273 152 9 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 557 3 9	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 556 7 8	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 376 270 4 7	VIATRIS UP	FR
EFFEXOR L.P. 75 mg, gélule à libération prolongée	SE/H/0936/002	34009 346 555 0 0	VIATRIS UP	FR
ELIFY XR 37.5 mg κάψουλες παρατεταμένης αποδέσμευσης, σκληρές	NL/H/0927/001	20208	MEDOCHEMIE LTD.	CY
ЕФЕКТИН ER 150 mg твърди капсули с удължено освобождаване	SE/H/0936/003	9800389	UPJOHN EESV	BG
ЕФЕКТИН ER 75 mg твърди капсули с удължено освобождаване	SE/H/0936/002	9800390	UPJOHN EESV	BG
Faxiprol 150 mg retard	SE/564/01-04/DC	OGYI-T-20519/21-30	MEDICO UNO	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletta			PHARMACEUTICALS SE	
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	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099536	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099544	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099551	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099569	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 150 mg comprimidos de libertação prolongada	not available	5099510	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 37,5 mg comprimidos de libertação prolongada	not available	5974399	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 37,5 mg comprimidos de libertação prolongada	not available	5974498	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 37,5 mg comprimidos de libertação prolongada	not available	5974597	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 37,5 mg comprimidos de libertação prolongada	not available	5974290	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 75 mg comprimidos de libertação prolongada	not available	5975198	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 75 mg comprimidos de	not available	5975297	DECOMED FARMACÊUTICA, 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
libertação prolongada				
Genexin 75 mg comprimidos de libertação prolongada	not available	5975396	DECOMED FARMACÊUTICA, LDA.	PT
Genexin 75 mg comprimidos de libertação prolongada	not available	5975099	DECOMED FARMACÊUTICA, LDA.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180930	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180948	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180955	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180963	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180971	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181003	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181011	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181029	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181037	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181045	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg	not available	5181052	LABORATORIOS LICONSA,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de libertação prolongada			S.A.	
Pracet 150 mg comprimidos de libertação prolongada	not available	5181060	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181078	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181102	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181110	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181128	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181136	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181144	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5180922	LABORATORIOS LICONSA, S.A.	PT
Pracet 150 mg comprimidos de libertação prolongada	not available	5181151	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181169	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181177	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181201	LABORATORIOS LICONSA, 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
Pracet 225 mg comprimidos de libertação prolongada	not available	5181219	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181227	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181235	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181243	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181250	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181268	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181276	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181300	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181318	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181326	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181334	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181342	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de	not available	5181359	LABORATORIOS LICONSA, 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
libertação prolongada				
Pracet 225 mg comprimidos de libertação prolongada	not available	5181367	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181375	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181409	LABORATORIOS LICONSA, S.A.	PT
Pracet 225 mg comprimidos de libertação prolongada	not available	5181417	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180633	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180641	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180658	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180427	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180435	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180443	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180450	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180468	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg	not available	5180476	LABORATORIOS LICONSA,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de libertação prolongada			S.A.	
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180500	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180518	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180526	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180534	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180542	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180559	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180567	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180575	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180609	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180617	LABORATORIOS LICONSA, S.A.	PT
Pracet 37,5 mg comprimidos de libertação prolongada	not available	5180625	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180765	LABORATORIOS LICONSA, 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
Pracet 75 mg comprimidos de libertação prolongada	not available	5180773	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180807	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180815	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180823	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180831	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180849	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180856	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180864	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180872	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180906	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180914	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180666	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de	not available	5180674	LABORATORIOS LICONSA, 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
libertação prolongada				
Pracet 75 mg comprimidos de libertação prolongada	not available	5180708	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180716	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180724	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180732	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180740	LABORATORIOS LICONSA, S.A.	PT
Pracet 75 mg comprimidos de libertação prolongada	not available	5180757	LABORATORIOS LICONSA, S.A.	PT
Serosmine 150 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/003	68910/11-09-2013	ITF HELLAS S.A.	GR
Serosmine 225 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/004	68911/11-09-2013	ITF HELLAS S.A.	GR
Serosmine 300 mg δισκίο παρατεταμένης αποδέσμευσης	SE/H/0581/005	62688/16-05-2019	ITF HELLAS S.A.	GR
Serosmine 37,5 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/001	68912/11-09-2013	ITF HELLAS S.A.	GR
Serosmine 75 mg δισκία παρατεταμένης αποδέσμευσης	SE/H/0581/002	68909/11-09-2013	ITF HELLAS S.A.	GR
Sunveniz XL 150 mg prolonged-release tablets	NL/H/3948/003	PL31750/0027	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	XI
Sunveniz XL 37.5 mg	NL/H/3948/001	PL31750/0025	SUN PHARMACEUTICAL	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolonged-release tablets			INDUSTRIES EUROPE B.V.	
Sunveniz XL 75 mg prolonged-release tablets	NL/H/3948/002	PL31750/0026	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	XI
Trevilor, 150 mg depotkapsel, hård	SE/H/0938/003	42827	UPJOHN EESV	SE
Trevilor, 37,5 mg depotkapsel, hård	SE/H/0938/001	42825	UPJOHN EESV	SE
Trevilor, 75 mg depotkapsel, hård	SE/H/0938/002	42826	UPJOHN EESV	SE
Trevilor® retard 150 mg Hartkapseln, retardiert	SE/H/0936/003	45300.01.00	VIATRIS PHARMA GMBH	DE
Trevilor® retard 37,5 mg Hartkapseln, retardiert	SE/H/0936/001	62104.00.00	VIATRIS PHARMA GMBH	DE
Trevilor® retard 75 mg Hartkapseln, retardiert	SE/H/0936/002	45300.00.00	VIATRIS PHARMA GMBH	DE
Vandral Retard 150 mg cápsulas duras de liberación prolongada.	SE/H/0936/003	62.402	VIATRIS HEALTHCARE S.L.	ES
Vandral Retard 225 mg cápsulas duras de liberación prolongada.	SE/H/0936/004	79.960	VIATRIS HEALTHCARE S.L.	ES
Vandral Retard 75 mg cápsulas duras de liberación prolongada.	SE/H/0936/002	62.401	VIATRIS HEALTHCARE S.L.	ES
Velaxin 37,5 mg tablety	HU/H/0102/002	30/0457/05-S	EGIS PHARMACEUTICALS PLC	SK
Velaxin 50 mg tablety	HU/H/0102/003	30/550/05-C	EGIS PHARMACEUTICALS PLC	CZ
Velaxin 75 mg tablety	HU/H/0102/004	30/551/05-C	EGIS PHARMACEUTICALS PLC	CZ
Velaxin 75 mg tablety	HU/H/0102/004	30/0459/05-S	EGIS PHARMACEUTICALS PLC	SK
Venlabrain retard 150 mg comprimidos de liberación prolongada	SE/H/0582/002/DC	69751	EXELTIS HEALTHCARE S.L	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 225 mg comprimidos de liberación prolongada	SE/H/0582/003/DC	69750	EXELTIS HEALTHCARE S.L	ES
Venlabrain retard 300 mg comprimidos de liberación prolongada	SE/H/0582/005/DC	84300	EXELTIS HEALTHCARE S.L	ES
Venlabrain retard 75 mg comprimidos de liberación prolongada	SE/H/0582/001/DC	69752	EXELTIS HEALTHCARE S.L	ES
Venlafaksin Pliva 150 mg tvrde kapsule s produljenim oslobaĎanjem	DE/H/6399/003	HR-H-045409982	PLIVA HRVATSKA D.O.O.	HR
Venlafaksin Pliva 75 mg tvrde kapsule s produljenim oslobaĎanjem	DE/H/6399/002	HR-H-513371317	PLIVA HRVATSKA D.O.O.	HR
Venlafaxin - 1 A Pharma 225 mg Hartkapseln, retardiert	AT/H/1196/004	99251.00.00	1 A PHARMA GMBH	DE
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 300 mg	SE/H/0582/005	60473	MEDICAL VALLEY INVEST AB	DK
Venlafaxin "Sandoz", hårde depotkapsler	AT/H/1196/004	62037	SANDOZ A/S	DK
Venlafaxin ”Hexal”, hårde depotkapsler	BE/H/0360/001	42725	HEXAL A/S	DK
Venlafaxin ”Krka”, hårde depotkapsler	NL/H/0799/001	39041	KRKA SVERIGE AB	DK
Venlafaxin +pharma 150 mg Retardtabletten	SE/H/0564/003	1-27289	+PHARMA ARZNEIMITTEL GMBH	AT
Venlafaxin +pharma 75	SE/H/0564/002	1-27287	+PHARMA ARZNEIMITTEL	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg Retardtabletten			GMBH	
Venlafaxin 1A Farma 150 mg depottabletter	SE/H/0581/003	23687	1A FARMA A/S	SE
Venlafaxin 1A Farma 225 mg depottabletter	SE/H/0581/004	23688	1A FARMA A/S	SE
Venlafaxin 1A Farma 300 mg depottabletter	SE/H/0581/005	57270	1A FARMA A/S	SE
Venlafaxin 1A Farma 37,5 mg depottabletter	SE/H/0581/001	23685	1A FARMA A/S	SE
Venlafaxin 1A Farma 75mg depottabletter	SE/H/0581/002	23686	1A FARMA A/S	SE
Venlafaxin 1A Pharma GmbH 225 mg - Hartkapseln, retardiert	AT/H/1198/004	138100	1A PHARMA GMBH	AT
Venlafaxin AbZ 150 mg Hartkapseln, retardiert	DE/H/6400/003	2204466.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 150 mg Hartkapseln, retardiert	DE/H/6400/003	2204466.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 150 mg Hartkapseln, retardiert	DE/H/6410/003	2204500.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 225 mg Retardtabletten	DE/H/3592/004	74547.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 37,5 mg Hartkapseln, retardiert	DE/H/6400/001	2204464.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 37,5 mg Hartkapseln, retardiert	DE/H/6400/001	2204464.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 37,5 mg Hartkapseln, retardiert	DE/H/6410/001	2204498.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 75 mg Hartkapseln, retardiert	DE/H/6400/002	2204465.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 75 mg Hartkapseln, retardiert	DE/H/6400/002	2204465.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin AbZ 75 mg Hartkapseln, retardiert	DE/H/6410/002	2204499.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin Aristo 225 mg Retardkapseln, Hartkapseln, retardiert	not available	99127.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Venlafaxin Genericon	SE/H/0564/004	1-27291	GENERICON PHARMA	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
225 mg Retardtabletten			GESELLSCHAFT M.B.H.	
Venlafaxin Heumann 225 mg Hartkapseln, retardiert	DE/H/4271/001	93933.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Venlafaxin HEXAL 225 mg Hartkapseln, retardiert	AT/H/1197/004	99258.00.00	HEXAL AG	DE
Venlafaxin Hexal GmbH retard 225 mg – Hartkapseln, retardiert	AT/H/1197/004	138104	HEXAL PHARMA GMBH	AT
Venlafaxin Krka 37,5 mg hard depotkapsel	NL/H/0799/001	06-3972	KRKA SVERIGE AB	NO
Venlafaxin Krka 37,5 mg hård depotkapsel	NL/H/0799/001	23463	KRKA SVERIGE AB	SE
Venlafaxin Krka 37,5 mg kova depotkapseli	NL/H/0799/001	21947	KRKA SVERIGE AB	FI
Venlafaxin Liconsa 150 mg depottabletter	SE/H/0564/003	23683	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin Liconsa 225 mg depottabletter	SE/H/0564/004	23684	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin Liconsa 300 mg depottabletter	SE/H/0564/005	57269	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin Liconsa 37.5 mg depottabletter	SE/H/0564/001	23681	LABORATORIOS LICONSA, S.A.	SE
Venlafaxin Liconsa 75 mg depottabletter	SE/H/0564/002	23682	LABORATORIOS LICONSA, S.A.	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
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 300 mg depottabletter	SE/H/0582/005	57271	MEDICAL VALLEY INVEST AB	SE
Venlafaxin Medical Valley 300 mg forðatafla	SE/H/0582/005/DC	IS/1/19/017/01	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 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 forðatöflur	SE/H/0582/001	IS/1/17/021/01	MEDICAL VALLEY INVEST AB	IS
Venlafaxin Sandoz 225 mg depotkapsel, hard	AT/H/1196/004	18-12644	SANDOZ A/S	NO
Venlafaxin Sandoz GmbH 225 mg – Hartkapseln, retardiert	AT/H/1196/004	138109	SANDOZ GMBH	AT
Venlafaxin Teva 150 mg depotkapsel, hard	DE/H/6399/003	21-14355	TEVA B.V	NO
Venlafaxin Teva 150 mg depotkapseli, kova	DE/H/6399/003	39974	TEVA B.V	FI
Venlafaxin Teva 150 mg hårda depotkapslar	DE/H/6399/003	60250	TEVA B.V	SE
Venlafaxin Teva 150 mg hart forðahylki	DE/H/6399/003	IS/1/22/043/03	TEVA B.V	IS
Venlafaxin Teva 37,5 mg depotkapsel, hard	DE/H/6399/001	21-14353	TEVA B.V	NO
Venlafaxin Teva 37,5 mg depotkapseli, kova	DE/H/6399/001	39972	TEVA B.V	FI
Venlafaxin Teva 37,5 mg hårda depotkapslar	DE/H/6399/001	39972	TEVA B.V	FI
Venlafaxin Teva 37,5 mg hårda depotkapslar	DE/H/6399/001	60248	TEVA B.V	SE
Venlafaxin Teva 37,5 mg hart forðahylki	DE/H/6399/001	IS/1/22/043/01	TEVA B.V	IS
Venlafaxin Teva 75 mg depotkapsel, hard	DE/H/6399/002	21-14354	TEVA B.V	NO
Venlafaxin Teva 75 mg depotkapseli, kova	DE/H/6399/002	39973	TEVA B.V	FI
Venlafaxin Teva 75 mg hårda depotkapslar	DE/H/6399/002	39973	TEVA B.V	FI
Venlafaxin Teva 75 mg hårda depotkapslar	DE/H/6399/003	39974	TEVA B.V	FI
Venlafaxin Teva 75 mg	DE/H/6399/002	60249	TEVA B.V	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
hårda depotkapslar				
Venlafaxin Teva 75 mg hart forðahylki	DE/H/6399/002	IS/1/22/043/02	TEVA B.V	IS
Venlafaxin Winthrop 37,5 mg Hartkapseln, retardiert	DE/H/5043/001	67508.00.00	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 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
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	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
rilascio prolungato				
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 Refta 150 mg cápsulas de libertação prolongada	DE/H/6399/003	5811336	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Venlafaxina Refta 37,5 mg cápsulas de libertação prolongada	DE/H/6399/001	5811302	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Venlafaxina Refta 37,5 mg cápsulas de libertação prolongada	DE/H/6399/001	5811310	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Venlafaxina Refta 75 mg cápsulas de libertação prolongada	DE/H/6399/002	5811328	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Venlafaxina Retard	DE/H/6400/003	87922	LABORATORIOS DAVUR	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
Davurgama 150 mg cápsulas duras de liberación prolongada			S.L.U.,	
Venlafaxina Retard Davurgama 150 mg cápsulas duras de liberación prolongada	DE/H/6400/003	87922	LABORATORIOS DAVUR S.L.U.,	ES
Venlafaxina Retard Davurgama 75 mg cápsulas duras de liberación prolongada	DE/H/6400/002	87923	LABORATORIOS DAVUR S.L.U.,	ES
Venlafaxina Retard Davurgama 75 mg cápsulas duras de liberación prolongada	DE/H/6400/002	87923	LABORATORIOS DAVUR S.L.U.,	ES
Venlafaxina retard Krka 37,5 mg cápsulas duras de liberación prolongada	NL/H/0799/001	69976	KRKA, D.D., NOVO MESTO	ES
Venlafaxina Retard ratio 150 mg cápsulas duras de liberación prolongada	DE/H/6399/003	87919	RATIOPHARM ESPAÑA S.A.,	ES
Venlafaxina Retard ratio 75 mg cápsulas duras de liberación prolongada	DE/H/6399/002	87918	RATIOPHARM ESPAÑA S.A.,	ES
Venlafaxina Retard Tevagen 150 mg cápsulas duras de liberación prolongada	DE/H/6410/003	87921	TEVA B.V	ES
Venlafaxina Retard Tevagen 75 mg cápsulas duras de liberación prolongada	DE/H/6410/002	87920	TEVA B.V	ES
Venlafaxina Salipax 150 mg cápsulas de libertação prolongada	DE/H/6400/003	5811278	MEPHA LDA	PT
Venlafaxina Salipax 150 mg cápsulas de libertação prolongada	DE/H/6400/003	5811278	MEPHA 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
Venlafaxina Salipax 37,5 mg cápsulas de libertação prolongada	DE/H/6400/001	5811229	MEPHA LDA	PT
Venlafaxina Salipax 37,5 mg cápsulas de libertação prolongada	DE/H/6400/001	5811237	MEPHA LDA	PT
Venlafaxina Salipax 37,5 mg cápsulas de libertação prolongada	DE/H/6400/001	5811229	MEPHA LDA	PT
Venlafaxina Salipax 37,5 mg cápsulas de libertação prolongada	DE/H/6400/001	5811237	MEPHA LDA	PT
Venlafaxina Salipax 75 mg cápsulas de libertação prolongada	DE/H/6400/002	5811260	MEPHA LDA	PT
Venlafaxina Salipax 75 mg cápsulas de libertação prolongada	DE/H/6400/002	5811260	MEPHA LDA	PT
Venlafaxina Sandoz 225 mg capsule rigide a rilascio prolungato	NL/H/3993/004	045555051	SANDOZ S.P.A.	IT
Venlafaxina SUN 150 mg comprimidos de liberación prolongada	NL/H/3948/003	76603	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
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	not available	5708995	TOLIFE - PRODUTOS FARMACÊUTICOS, 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
libertação prolongada				
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
Venlafaxina Viatris 150 mg capsule rigide a rilascio prolungato	SE/H/0938/003	028834101	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 150 mg capsule rigide a rilascio prolungato	SE/H/0938/003	028834063	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834125	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834137	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 37,5 mg capsule rigide a rilascio prolungato	SE/H/0938/001	028834113	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 75 mg capsule rigide a rilascio prolungato	SE/H/0938/002	028834099	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Viatris 75 mg capsule rigide a rilascio prolungato	SE/H/0938/002	028834051	VIATRIS PHARMA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881253	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881012	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5	DE/H/5043/001	037881075	ZENTIVA 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
mg capsule rigide a rilascio prolungato				
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881036	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881087	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881291	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881024	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881051	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881048	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881063	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881378	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881341	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zentiva 37,5 mg capsule rigide a rilascio prolungato	DE/H/5043/001	037881303	ZENTIVA ITALIA S.R.L.	IT
Venlafaxina Zidrium 150 mg cápsulas de libertação prolongada	DE/H/6410/003	5810932	TEVA B.V	PT
Venlafaxina Zidrium 37,5 mg cápsulas de libertação prolongada	DE/H/6410/001	5810908	TEVA B.V	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
Venlafaxina Zidrium 37,5 mg cápsulas de libertação prolongada	DE/H/6410/001	5810916	TEVA B.V	PT
Venlafaxina Zidrium 75 mg cápsulas de libertação prolongada	DE/H/6410/002	5810924	TEVA B.V	PT
Venlafaxin-CT 225 mg Retardtabletten	DE/H/3593/004	74551.00.00	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 300 mg Focus, tabletten met verlengde afgifte	not available	RVG 123673	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 ARROW LP 75 mg, gélule à libération prolongée	not available	NL32838	ARROW GENERIQUES	FR
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
VENLAFAXINE BIPHAR	SE/H/0581/003	34009 384 485 6 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
150 mg, comprimé à libération prolongée				
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
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

Product 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 384 500 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
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
VENLAFAXINE BIPHAR 225 mg, comprimé à	SE/H/0581/004	34009 384 510 0 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
libération prolongée				
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
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	SE/H/0581/004	34009 384 501 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
225 mg, comprimé à libération prolongée				
VENLAFAXINE BIPHAR 225 mg, comprimé à libération prolongée	SE/H/0581/004	34009 572 359 4 9	MYLAN MEDICAL SAS	FR
Venlafaxine Liconsa 150 mg tabletten met verlengde afgifte	SE/H/0581/003	RVG 33888	LABORATORIOS LICONSA, S.A.	NL
Venlafaxine Liconsa 300 mg pikendatud vabanemisajaga tablett	SE/H/0581/005	985019	LABORATORIOS LICONSA, S.A.	EE
Venlafaxine Liconsa 300 mg tabletten met verlengde afgifte	SE/H/0581/005	RVG 122326	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 75 mg tabletten met verlengde afgifte	SE/H/0581/002	RVG 33887	LABORATORIOS LICONSA, S.A.	NL
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, 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 PFIZER L.P. 37,5 mg, gélule à	SE/H/0938/001	34009 346 617 6 1	VIATRIS UP	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
libération prolongée				
VENLAFAXINE PFIZER L.P. 37,5 mg, gélule à libération prolongée	SE/H/0938/001	34009 346 618 2 2	VIATRIS UP	FR
VENLAFAXINE PFIZER L.P. 37,5 mg, gélule à libération prolongée	SE/H/0938/001	34009 346 619 9 0	VIATRIS UP	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34009 346 567 9 8	VIATRIS UP	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34009 346 566 2 0	VIATRIS UP	FR
VENLAFAXINE PFIZER L.P. 75 mg, gélule à libération prolongée	SE/H/0938/002	34009 346 568 5 9	VIATRIS UP	FR
Venlafaxine retard Teva 150 mg, harde capsules met verlengde afgifte	DE/H/6399/003	RVG 129032	TEVA B.V	NL
Venlafaxine retard Teva 37,5 mg, harde capsules met verlengde afgifte	DE/H/6399/001	RVG 129029	TEVA B.V	NL
Venlafaxine retard Teva 75 mg, harde capsules met verlengde afgifte	DE/H/6399/002	RVG 129031	TEVA B.V	NL
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	DE/H/3591/003	BE347356	TEVA PHARMA 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
mg comprimés à libération prolongée			N.V./S.A	
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
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 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
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

Product 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 tabletten met verlengde afgifte	DE/H/3591/001	BE347313	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
VENLAFAXINE TEVA L.P. 150 mg, gélule à libération prolongée	DE/H/6399/003	NL53853	TEVA B.V	FR
VENLAFAXINE TEVA L.P. 37,5 mg, gélule à libération prolongée	DE/H/6399/001	NL53851	TEVA B.V	FR
VENLAFAXINE TEVA L.P. 75 mg, gélule à libération prolongée	DE/H/6399/002	NL53852	TEVA B.V	FR
Venlafaxine XL 150 mg prolonged-release tablets	SE/H/0581/003	PL 01883/0340	MACARTHYS LABORATORIES LTD	XI
Venlafaxine XL 225 mg prolonged-release tablets	SE/H/0581/004	PL 01883/0341	MACARTHYS LABORATORIES LTD	XI
Venlafaxine XL 300 mg	SE/H/0581/005	PL 01883/0363	MACARTHYS LABORATORIES	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolonged-release tablet			LTD	
Venlafaxine XL 37.5 mg prolonged-release tablets	SE/H/0581/001	PL 01883/0338	MACARTHYS LABORATORIES LTD	XI
Venlafaxine XL 75 mg prolonged-release tablets	SE/H/0581/002	PL 01883/0339	MACARTHYS LABORATORIES LTD	XI
Venlafaxine XR Viatris 150 mg, capsules met verlengde afgifte, hard, 150 mg	SE/H/0938/003	RVG 107029	VIATRIS NETHERLANDS B.V.	NL
Venlafaxine XR Viatris 37,5 mg, capsules met verlengde afgifte, hard, 37,5 mg	SE/H/0938/001	RVG 107027	VIATRIS NETHERLANDS B.V.	NL
Venlafaxine XR Viatris 75 mg, capsules met verlengde afgifte, hard, 75 mg	SE/H/0938/002	RVG 107028	VIATRIS NETHERLANDS B.V.	NL
Venlafaxin-neuraxpharm 225 mg retard Retardtabletten	not available	73279.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Venlafaxin-neuraxpharm 300 mg Retardtabletten	DE/H/5795/001	2202672.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Venlafaxin-ratiopharm 150 mg Hartkapseln, retardiert	DE/H/6399/003	2204463.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin-ratiopharm 150 mg Hartkapseln, retardiert	DE/H/6399/003	2021090188	RATIOPHARM GMBH	LU
Venlafaxin-ratiopharm 37,5 mg Hartkapseln, retardiert	DE/H/6399/001	2204461.00.00	ABZ-PHARMA GMBH	DE
Venlafaxin-ratiopharm 37,5 mg Hartkapseln, retardiert	DE/H/6399/001	2021090186	RATIOPHARM GMBH	LU
Venlafaxin-ratiopharm 75 mg Hartkapseln,	DE/H/6399/002	2204462.00.00	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
retardiert				
Venlafaxin-ratiopharm 75 mg Hartkapseln, retardiert	DE/H/6399/002	2021090187	RATIOPHARM GMBH	LU
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
Venlatev 150 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/003	TEVA B.V	IE
Venlatev 150 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/003	TEVA B.V	IE
Venlatev 37.5 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/001	TEVA B.V	IE
Venlatev 37.5 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/001	TEVA B.V	IE
Venlatev 75 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/002	TEVA B.V	IE
Venlatev 75 mg hard prolonged-release capsules	DE/H/6400	PA1986/099/002	TEVA B.V	IE
VENLAXIN 150 mg δισκία παρατεταμένης αποδέσμευσης	not available	021936	IASIS PHARMA	CY
Venlaxin 150 mg δισκία παρατεταμένης αποδέσμευσης	not available	84836/20-09-2021	IASIS PHARMA	GR
VENLAXIN 225 mg δισκία παρατεταμένης	not available	021937	IASIS PHARMA	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
αποδέσμευσης				
Venlaxin 225 mg δισκία παρατεταμένης αποδέσμευσης	not available	84836/20-09-2021	IASIS PHARMA	GR
VENLAXIN 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	021935	IASIS PHARMA	CY
Venlaxin 75 mg δισκία παρατεταμένης αποδέσμευσης	not available	84836/20-09-2021	IASIS PHARMA	GR
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
Venlazid 300 mg depottabletter	SE/H/0582/005/DC	17-12032	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, LDA.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099643	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099577	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099601	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099619	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 150 mg comprimidos de libertação prolongada	not available	5099627	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 37,5 mg comprimidos de	not available	5974795	DECOMED FARMACÊUTICA, 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
libertação prolongada				
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974894	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974993	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 37,5 mg comprimidos de libertação prolongada	not available	5974696	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975594	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975693	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975792	DECOMED FARMACÊUTICA, LDA.	PT
Venxin 75 mg comprimidos de libertação prolongada	not available	5975495	DECOMED FARMACÊUTICA, LDA.	PT
VENZIP XL 37.5 mg prolonged-release capsules, hard	PT/H/0703/001	MA807/04601	AUROBINDO PHARMA (MALTA) LIMITED	MT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316218	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316220	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316232	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316244	ITALFARMACO S.P.A	IT
Zarelis 150 mg	SE/H/0581/003	038316257	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
Compressa a rilascio prolungato				
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316269	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316271	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316283	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316295	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316307	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316319	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316321	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316333	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316345	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316358	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316360	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316372	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/0581/003	038316384	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316396	ITALFARMACO S.P.A	IT
Zarelis 150 mg Compressa a rilascio prolungato	SE/H/0581/003	038316408	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316410	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316422	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316434	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316446	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316459	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316461	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316473	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316485	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316497	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio	SE/H/0581/004	038316509	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
prolungato				
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316511	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316523	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316535	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316547	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316550	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316562	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316574	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316586	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316598	ITALFARMACO S.P.A	IT
Zarelis 225 mg Compressa a rilascio prolungato	SE/H/0581/004	038316600	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316814	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316826	ITALFARMACO S.P.A	IT
Zarelis 300 mg	SE/H/0581/005	038316838	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
compressa a rilascio prolungato				
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316840	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316853	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316865	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316877	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316889	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316891	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316903	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316915	ITALFARMACO S.P.A	IT
Zarelis 300 mg compressa a rilascio prolungato	SE/H/0581/005	038316927	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316612	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316624	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316636	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/0581/001	038316648	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316651	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316663	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316675	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316687	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316699	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316701	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316713	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316725	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316737	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316749	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316752	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio	SE/H/0581/001	038316764	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
prolungato				
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316776	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316788	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316790	ITALFARMACO S.P.A	IT
Zarelis 37,5 mg Compressa a rilascio prolungato	SE/H/0581/001	038316802	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316016	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316028	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316030	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316042	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316055	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316067	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316079	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316081	ITALFARMACO S.P.A	IT
Zarelis 75 mg	SE/H/0581/002	038316093	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
Compressa a rilascio prolungato				
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316105	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316117	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316129	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316131	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316143	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316156	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316168	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316170	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316182	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316194	ITALFARMACO S.P.A	IT
Zarelis 75 mg Compressa a rilascio prolungato	SE/H/0581/002	038316206	ITALFARMACO S.P.A	IT
ZARELIS RETARD 150 mg comprimidos de liberación prolongada	SE/H/0564/003	69738	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 300 mg comprimidos de liberación prolongada	SE/H/0564/005	84181	ITALFARMACO S.A.	ES
ZARELIS RETARD 75 mg comprimidos de liberación prolongada	SE/H/0564/002	69737	ITALFARMACO S.A.	ES
Zarelix 150 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/003	5058318	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 150 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/003	5058326	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 225 mg comprimidos de libertação prolongada	SE/H/0581/004	5058334	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 225 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/004	5058334	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 225 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/004	5058342	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 225 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/004	5058342	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 300 mg comprimidos de libertação prolongada	SE/H/0581/005	5768437	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 37.5 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/001	5058250	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 37.5 mg comprimidos de libertaｦｧｦ｣o prolongada	SE/H/0581/001	5143037	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 37.5 mg	SE/H/0581/001	5058268	ITALFARMACO - PRODUTOS	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de libertaﾃｧﾃ｣o prolongada			FARMACÊUTICOS, LDA.	
Zarelix 37.5 mg comprimidos de libertaﾃｧﾃ｣o prolongada	SE/H/0581/001	5143045	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 75 mg comprimidos de libertação prolongada	SE/H/0581/002	5058276	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Zarelix 75 mg comprimidos de libertaﾃｧﾃ｣o prolongada	SE/H/0581/002	5058300	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Лароксин XR 150 mg твърди капсули с удължено освобождаване	DE/H/6399/003	20220205	TEVA B.V	BG
Лароксин XR 37,5 mg твърди капсули с удължено освобождаване	DE/H/6399/001	20220203	TEVA B.V	BG
Лароксин XR 75 mg твърди капсули с удължено освобождаване	DE/H/6399/002	20220204	TEVA B.V	BG