1 September 2022
EMA/724081/2022
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): escitalopram

Procedure No. PSUSA/00001265/202112

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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
Cipralex	SE/H/0278/001	33407	H. LUNDBECK A/S	DK
Cipralex	SE/H/0278/002	33408	H. LUNDBECK A/S	DK
Cipralex	SE/H/0278/003	33409	H. LUNDBECK A/S	DK
Cipralex	SE/H/278/04	33411	H. LUNDBECK A/S	DK
Cipralex	SE/H/0278/006	41275	H. LUNDBECK A/S	DK
Cipralex 10 mg apvalkotās tabletes	SE/H/0278/002	07 – 0298	H. LUNDBECK A/S	LV
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767058/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767060/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767072/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/278/02	035767084/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767514/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767526/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767538/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767235/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767247/M	H. LUNDBECK A/S	IT
Cipralex 10 mg	SE/H/0278/002	035767250/M	H. LUNDBECK A/S	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
compresse rivestite con film				
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767262/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767274/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767286/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767437/M	H. LUNDBECK A/S	IT
Cipralex 10 mg compresse rivestite con film	SE/H/0278/002	035767449/M	H. LUNDBECK A/S	IT
CIPRALEX 10 mg comprimate filmate	SE/H/0278/002	9256/2016/01-19	H. LUNDBECK A/S	RO
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	4304085	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	4304184	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	5026315	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	4304788	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	4304887	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	4304986	H. LUNDBECK A/S	PT
Cipralex 10 mg	SE/H/0278/002	4305082	H. LUNDBECK A/S	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película				
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	5013958	H. LUNDBECK A/S	PT
CIPRALEX 10 mg filmdrasjerte tabletter	SE/H/0278/002	02-713	H. LUNDBECK A/S	NO
Cipralex 10 mg filmom obalené tablety	SE/H/278/02	30/0210/07-S	H. LUNDBECK A/S	SK
Cipralex 10 mg filmom obložene tablete	SE/H/0280/002	UP/I-530-09/13-02/247	LUNDBECK CROATIA D.O.O.	HR
Cipralex 10 mg filmsko obložene tablete	SE/H/0278/002	5363-I-689/07	LUNDBECK PHARMA D.O.O	SI
Cipralex 10 mg filmtabletta	SE/H/0278/002	OGYI-T-8634/01	H. LUNDBECK A/S	HU
CIPRALEX 10 mg filmuhúðaðar töflur	SE/H/0278/002	IS/1/02/019/02	H. LUNDBECK A/S	IS
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/017	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/018	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/019	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/020	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/021	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/022	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/023	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/024	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/025	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/026	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele	SE/H/0278/002	LT/1/07/0832/027	H. LUNDBECK A/S	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
dengtos tabletes				
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/028	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/029	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/030	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/031	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/032	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/066	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/069	H. LUNDBECK A/S	LT
Cipralex 10 mg plevele dengtos tabletes	SE/H/0278/002	LT/1/07/0832/070	H. LUNDBECK A/S	LT
CIPRALEX 10 mg potahované tablety	SE/H/0278/002	30/276/02-C	H. LUNDBECK A/S	CZ
Cipralex 10 mg tabletti, kalvopäällysteinen	SE/H/0278/002	17691	H. LUNDBECK A/S	FI
CIPRALEX 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/002	20267	H. LUNDBECK A/S	CY
CIPRALEX 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/002	76961	H. LUNDBECK A/S	GR
CIPRALEX 10 mg, comprimidos recubiertos con película	SE/H/0278/002	65.230	LUNDBECK ESPANA, S.A.	ES
CIPRALEX 10 mg, filmdragerade tabletter	SE/H/0278/002	17085	H. LUNDBECK A/S	SE
Cipralex 10 mg/ml gocce orali, soluzione	SE/H/0278/002	035767615/M	H. LUNDBECK A/S	IT
Cipralex 10 mg/ml gocce orali, soluzione	SE/H/0278/002	035767627/M	H. LUNDBECK A/S	IT
Cipralex 10 mg/ml gocce orali, soluzione	SE/H/0278/002	035767639/M	H. LUNDBECK A/S	IT

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Cipralex 10 mg/ml gocce orali, soluzione	SE/H/0278/002	035767641/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767096/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767108/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767110/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767122/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767540/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767553/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767565/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767298/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767300/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767312/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767324/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767336/M	H. LUNDBECK A/S	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
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767348/M	H. LUNDBECK A/S	IT
Cipralex 15 mg compresse rivestite con film	SE/H/0278/003	035767452/M	H. LUNDBECK A/S	IT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4307187	H. LUNDBECK A/S	PT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4307286	H. LUNDBECK A/S	PT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4307880	H. LUNDBECK A/S	PT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4307989	H. LUNDBECK A/S	PT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4308086	H. LUNDBECK A/S	PT
Cipralex 15 mg comprimidos revestidos por película	SE/H/0278/003	4308185	H. LUNDBECK A/S	PT
CIPRALEX 15 mg filmdrasjerte tabletter	SE/H/0278/003	02-714	H. LUNDBECK A/S	NO
CIPRALEX 15 mg filmuhúðaðar töflur	SE/H/0278/003	IS/1/02/019/03	H. LUNDBECK A/S	IS
Cipralex 15 mg tabletti, kalvopäällysteinen	SE/H/0278/003	17692	H. LUNDBECK A/S	FI
CIPRALEX 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/003	20268	H. LUNDBECK A/S	CY
CIPRALEX 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/003	76963	H. LUNDBECK A/S	GR
CIPRALEX 15 mg, comprimidos recubiertos	SE/H/0278/003	65.234	LUNDBECK ESPANA, 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
con película				
CIPRALEX 15 mg, filmdragerade tabletter	SE/H/0278/003	17086	H. LUNDBECK A/S	SE
Cipralex 20 mg apvalkotās tabletes	SE/H/0278/004	07-0300	H. LUNDBECK A/S	LV
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767134/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767146/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767159/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767 161/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767577/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767589/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767591/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767351/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767363/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767375/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767387/M	H. LUNDBECK A/S	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
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767399/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767401/M	H. LUNDBECK A/S	IT
Cipralex 20 mg compresse rivestite con film	SE/H/0278/004	035767464/M	H. LUNDBECK A/S	IT
Cipralex 20 mg comprimidos revestidos por película	SE/H/0278/004	4302683	H. LUNDBECK A/S	PT
Cipralex 20 mg comprimidos revestidos por película	SE/H/0278/004	5074711	H. LUNDBECK A/S	PT
Cipralex 20 mg comprimidos revestidos por película	SE/H/0278/004	4303384	H. LUNDBECK A/S	PT
Cipralex 20 mg comprimidos revestidos por película	SE/H/0278/004	4303483	H. LUNDBECK A/S	PT
Cipralex 20 mg comprimidos revestidos por película	SE/H/0278/004	4303582	H. LUNDBECK A/S	PT
CIPRALEX 20 mg film-coated tablets	SE/H/0278/004	MA601/00304	H. LUNDBECK A/S	MT
CIPRALEX 20 mg filmdrasjerte tabletter	SE/H/0278/004	02-715	H. LUNDBECK A/S	NO
CIPRALEX 20 mg filmuhúðaðar töflur	SE/H/0278/004	IS/1/02/019/04	H. LUNDBECK A/S	IS
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/049	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/050	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/051	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/052	H. LUNDBECK A/S	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
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/053	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/054	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/055	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/056	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/057	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/058	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/059	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/060	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/061	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/062	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/063	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/064	H. LUNDBECK A/S	LT
Cipralex 20 mg plevele dengtos tabletes	SE/H/278/04	LT/1/07/0832/072	H. LUNDBECK A/S	LT
Cipralex 20 mg tabletti, kalvopäällysteinen	SE/H/0278/004	17693	H. LUNDBECK A/S	FI
CIPRALEX 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/004	20269	H. LUNDBECK A/S	CY
CIPRALEX 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/004	76964	H. LUNDBECK A/S	GR
CIPRALEX 20 mg, comprimidos recubiertos con película	SE/H/0278/004	65.233	LUNDBECK ESPANA, S.A.	ES
CIPRALEX 20 mg,	SE/H/0278/004	17087	H. LUNDBECK A/S	SE

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filmdragerade tabletter				
Cipralex 20 mg/ml dråper, oppløsning	SE/H/0278/006	07-4892	H. LUNDBECK A/S	NO
Cipralex 20 mg/ml gocce orali, soluzione	SE/H/0278/006	035767654/M	H. LUNDBECK A/S	IT
CIPRALEX 20 mg/ml Gotas orais, solução	SE/H/0278/006	5049739	H. LUNDBECK A/S	PT
Cipralex 20 mg/ml Gotas orales en solución	SE/H/0278/006	68811	LUNDBECK ESPANA, ES	ES
CIPRALEX 20 mg/ml orala droppar, lösning	SE/H/0278/006	23323	H. LUNDBECK A/S	SE
CIPRALEX 20 mg/ml perorální kapky, roztok	SE/H/0278/006	30/494/07-C	H. LUNDBECK A/S	CZ
Cipralex 20 mg/ml tipat, liuos	SE/H/0278/006	23395	H. LUNDBECK A/S	FI
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767019/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767021/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767033/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767045/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767603/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767490/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767502/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con	SE/H/0278/001	035767173/M	H. LUNDBECK A/S	IT

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film				
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767185/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767197/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767209/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767211/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767223/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767413/M	H. LUNDBECK A/S	IT
Cipralex 5 mg compresse rivestite con film	SE/H/0278/001	035767425/M	H. LUNDBECK A/S	IT
Cipralex 5 mg comprimidos revestidos por película	SE/H/0278/001	4301081	H. LUNDBECK A/S	PT
Cipralex 5 mg comprimidos revestidos por película	SE/H/0278/001	4301180	H. LUNDBECK A/S	PT
Cipralex 5 mg comprimidos revestidos por película	SE/H/0278/001	4301784	H. LUNDBECK A/S	PT
Cipralex 5 mg comprimidos revestidos por película	SE/H/0278/001	4301883	H. LUNDBECK A/S	PT
Cipralex 5 mg comprimidos revestidos por película	SE/H/0278/001	4301982	H. LUNDBECK A/S	PT
Cipralex 5 mg	SE/H/0278/001	4302089	H. LUNDBECK A/S	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película				
CIPRALEX 5 mg film-coated tablets	SE/H/0278/001	MA601/00301	H. LUNDBECK A/S	MT
CIPRALEX 5 mg filmdrasjerte tabletter	SE/H/0278/001	02-712	H. LUNDBECK A/S	NO
CIPRALEX 5 mg filmuhúðaðar töflur	SE/H/0278/001	IS/1/02/019/01	H. LUNDBECK A/S	IS
Cipralex 5 mg tabletti, kalvopäällysteinen	SE/H/0278/001	17690	H. LUNDBECK A/S	FI
CIPRALEX 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/001	20266	H. LUNDBECK A/S	CY
CIPRALEX 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/001	76959	H. LUNDBECK A/S	GR
CIPRALEX 5 mg, comprimidos recubiertos con película	SE/H/0278/001	65.231	LUNDBECK ESPANA, S.A.	ES
CIPRALEX 5 mg, filmdragerade tabletter	SE/H/0278/001	17084	H. LUNDBECK A/S	SE
Cipralex, 10 mg õhukese polümeerikattega tabletid	SE/H/0278/002	392502	H. LUNDBECK A/S	EE
Cipralex, 20 mg õhukese polümeerikattega tabletid	SE/H/0278/004	392702	H. LUNDBECK A/S	EE
Cipralex® 10 mg – Filmtabletten	SE/H/0278/002	1-24550	H. LUNDBECK A/S	AT
CIPRALEX® 10 mg film-coated tablets	SE/H/0278/002	MA601/00302	H. LUNDBECK A/S	MT
CIPRALEX® 10 mg Filmtabletten	SE/H/0278/002	55880.01.00	H. LUNDBECK A/S	DE
Cipralex® 15 mg – Filmtabletten	SE/H/0278/003	1-24551	H. LUNDBECK A/S	AT
Cipralex® 20 mg – Filmtabletten	SE/H/0278/004	1-24552	H. LUNDBECK A/S	AT
CIPRALEX® 20 mg	SE/H/0278/004	55880.03.00	H. LUNDBECK A/S	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
Filmtabletten				
Cipralex® 20 mg/ml Tropfen zum Einnehmen, Lösung	SE/H/0278/006	69241.00.00	H. LUNDBECK A/S	DE
Cipralex® 5 mg – Filmtabletten	SE/H/0278/001	1-24549	H. LUNDBECK A/S	AT
ENLIFT 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	EE/H/0181/002	73482/10-06-2019	MEDOCHEMIE HELLAS SA	GR
ENLIFT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	EE/H/0181/004	73483/10-06-2019	MEDOCHEMIE HELLAS SA	GR
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768050	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768062	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768074	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768086	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768528	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768542	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768237	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768249	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768252	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768264	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768276	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768288	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768439	H. LUNDBECK A/S	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
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768441	H. LUNDBECK A/S	IT
Entact 10 mg compresse rivestite con film	SE/H/0280/002	035768530	H. LUNDBECK A/S	IT
ENTACT 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/002	76953/26-11-07	H. LUNDBECK A/S	GR
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768098	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768100	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768112	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768555	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768567	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768579	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768290	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768302	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768314	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768326	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768340	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768338	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768454	H. LUNDBECK A/S	IT
Entact 15 mg compresse rivestite con film	SE/H/0280/003	035768124	H. LUNDBECK A/S	IT
ENTACT 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/003	76954/26-11-07	H. LUNDBECK A/S	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768148	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768151	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768136	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768163	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768581	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768593	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768605	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768353	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768365	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768377	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768389	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768391	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768403	H. LUNDBECK A/S	IT
Entact 20 mg compresse rivestite con film	SE/H/0280/004	035768478	H. LUNDBECK A/S	IT
ENTACT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/004	76955/26-11-07	H. LUNDBECK A/S	GR
Entact 20 mg/ml gocce orali, soluzione	SE/H/0280/006	035768656	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768011	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768023	H. LUNDBECK A/S	IT
Entact 5 mg compresse	SE/H/0280/001	035768035	H. LUNDBECK A/S	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rivestite con film				
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768047	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768492	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768504	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768516	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768175	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768187	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768199	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768201	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768213	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768225	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768415	H. LUNDBECK A/S	IT
Entact 5 mg compresse rivestite con film	SE/H/0280/001	035768427	H. LUNDBECK A/S	IT
ENTACT 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/001	76950/26-11-07	H. LUNDBECK A/S	GR
Escitalopram 10 mg film-coated Tablets	PT/H/2163/001	PL 17780/0293	ZENTIVA PHARMA UK LIMITED	XI
Escitalopram 10 mg film-coated tablets	DE/H/1176/002	PL 36687/0283	TORRENT PHARMA (UK) LTD.	XI
Escitalopram 10 mg film-coated tablets	DE/H/0594/002	PL 04569/0777	GENERICS [UK] LIMITED	XI
Escitalopram 20 mg film-coated Tablets	PT/H/2163/002	PL 17780/0295	ZENTIVA PHARMA UK LIMITED	XI
Escitalopram 20 mg film-coated tablets	DE/H/1176/004	PL 36687/0284	TORRENT PHARMA (UK) LTD.	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
Escitalopram 20 mg film-coated tablets	DE/H/0594/004	PL 04569/0778	GENERICS [UK] LIMITED	XI
Escitalopram 5 mg film-coated Tablets	UK/H/6146/001	PL 17780/0292	ZENTIVA PHARMA UK LIMITED	XI
Escitalopram 5 mg film-coated tablets	DE/H/1176/001	PL 36687/0282	TORRENT PHARMA (UK) LTD.	XI
Escitalopram 5 mg film-coated tablets	DE/H/0594/001	PL 04569/0776	GENERICS [UK] LIMITED	XI
Escitalopram beta 10 mg Filmtabletten	DE/H/3578/002	68165.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Escitalopram beta 15 mg Filmtabletten	DE/H/3578/003	68166.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Escitalopram beta 20 mg Filmtabletten	DE/H/3578/004	68167.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Escitalopram beta 5 mg Filmtabletten	DE/H/3578/001	68164.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Escitalopram Condronac	PT/H/1134/003	5634233	SANDOZ FARMACÊUTICA LDA.	PT
Escitalopram Genedec 10 mg comprimidos orodispersíveis	not available	5673736	DECOMED FARMACÊUTICA, LDA.	PT
Escitalopram Genedec 10 mg comprimidos orodispersíveis	not available	5673744	DECOMED FARMACÊUTICA, LDA.	PT
Escitalopram Genedec 20 mg comprimidos orodispersíveis	not available	5673777	DECOMED FARMACÊUTICA, LDA.	PT
Escitalopram Genedec 20 mg comprimidos orodispersíveis	not available	5673801	DECOMED FARMACÊUTICA, LDA.	PT
Escitalopram Lundbeck 10 mg Filmtabletten	SE/H/0279/002	90990.00.00	H. LUNDBECK A/S	DE
Escitalopram Lundbeck 10 mg, filmdragerade tabletter	SE/H/0279/002	17089	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 15 mg, filmdragerade tabletter	SE/H/0279/003	17090	H. LUNDBECK A/S	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
Escitalopram Lundbeck 20 mg Filmtabletten	SE/H/0279/004	90991.00.00	H. LUNDBECK A/S	DE
Escitalopram Lundbeck 20 mg, filmdragerade tabletter	SE/H/0279/004	17091	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 20 mg/ml orala droppar, lösning	SE/H/0279/006	23324	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 20 mg/ml Tropfen zum Einnehmen, Lösung	SE/H/0279/006	90992.00.00	H. LUNDBECK A/S	DE
Escitalopram Lundbeck 5 mg, filmdragerade tabletter	SE/H/0279/001	17088	H. LUNDBECK A/S	SE
Escitalopram Teva 15 mg filmsko oblozene tablete	HU/H/0179/003	H/09/00573/045	TEVA PHARMA B.V.	SI
Escitalopram Teva 20 mg filmsko oblozene tablete	HU/H/0179/004	H/09/00573/066	TEVA PHARMA B.V.	SI
Escitalopram Zentiva 10 mg comprimido revestido por película	PT/H/2163/001	5103502	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 10 mg comprimido revestido por película	PT/H/2163/001	5103478	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 10 mg comprimido revestido por película	PT/H/2163/001	5097225	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 10 mg comprimido revestido por película	PT/H/2163/001	5583273	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 10 mg comprimido revestido por película	PT/H/2163/001	5097233	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 20 mg comprimido revestido por película	PT/H/2163/002	5097241	ZENTIVA PORTUGAL, 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
Escitalopram Zentiva 20 mg comprimido revestido por película	PT/H/2163/002	5103510	ZENTIVA PORTUGAL, LDA	PT
Escitalopram Zentiva 20 mg comprimido revestido por película	PT/H/2163/002	5583265	ZENTIVA PORTUGAL, LDA	PT
Escitalopram-neuraxpharm 10 mg Filmtabletten	DE/H/1174/002	69884.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-neuraxpharm 10 mg Schmelztabletten	DE/H/5050/002	88039.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-neuraxpharm 15 mg Filmtabletten	DE/H/1174/003	69885.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-neuraxpharm 20 mg Filmtabletten	DE/H/1174/004	69886.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-neuraxpharm 20 mg Schmelztabletten	DE/H/5050/004	88041.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-neuraxpharm 5 mg Filmtabletten	DE/H/1174/001	69883.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram-ratiopharm® 15 mg Filmtabletten	HU/H/0179/003	2012100113	RATIOPHARM GMBH	LU
Escitalopram-ratiopharm® 5 mg Filmtabletten	HU/H/0179/001	2012110001	RATIOPHARM GMBH	LU
Esertia 10 mg comprimidos recubiertos con película	SE/H/0280/002	65.184	ALMIRALL, S.A.	ES
Esertia 15 mg comprimidos recubiertos con película	SE/H/0280/003	65.183	ALMIRALL, S.A.	ES
Esertia 20 mg comprimidos recubiertos	SE/H/0280/004	65.182	ALMIRALL, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
con película				
Esertia 20 mg/ml gotas orales en solución	SE/H/0280/006	68.875	ALMIRALL, S.A.	ES
Lexapro 10 mg film-coated tablets	SE/H/0278/002	PA 805/2/2	H. LUNDBECK A/S	IE
LEXAPRO 10 mg filmomhulde tabletten	SE/H/0279/002	RVG 30495	H. LUNDBECK A/S	NL
Lexapro 15 mg film-coated tablets	SE/H/0278/003	PA 805/2/3	H. LUNDBECK A/S	IE
LEXAPRO 15 mg filmomhulde tabletten	SE/H/0279/003	RVG 30496	H. LUNDBECK A/S	NL
Lexapro 20 mg film-coated tablets	SE/H/0278/004	PA 805/2/4	H. LUNDBECK A/S	IE
LEXAPRO 20 mg filmomhulde tabletten	SE/H/0279/004	RVG 30497	H. LUNDBECK A/S	NL
LEXAPRO 20 mg/ml druppels voor oraal gebruik, oplossing	SE/H/0279/006	RVG 35339	H. LUNDBECK A/S	NL
Lexapro 5 mg film-coated tablets	SE/H/0278/001	PA 805/2/1	H. LUNDBECK A/S	IE
LEXAPRO 5 mg filmomhulde tabletten	SE/H/0279/001	RVG 30494	H. LUNDBECK A/S	NL
Lexapro, 10 mg, tabletki powlekane	SE/H/0278/002	14283	H. LUNDBECK A/S	PL
Premalex 10 mg, filmdragerade tabletter	not available	42255	H. LUNDBECK AB, SE	SE
Premalex 20 mg, filmdragerade tabletter	not available	42256	H. LUNDBECK AB, SE	SE
PRILECT 10 mg, filmdragerade tabletter	SE/H/0280/002	17097	H. LUNDBECK A/S	SE
PRILECT 15 mg, filmdragerade tabletter	SE/H/0280/003	17098	H. LUNDBECK A/S	SE
PRILECT 20 mg, filmdragerade tabletter	SE/H/0280/004	17099	H. LUNDBECK A/S	SE
PRILECT 20 mg/ml orala droppar, lösning	SE/H/0280/006	23323	H. LUNDBECK A/S	SE
PRILECT 5 mg, filmdragerade tabletter	SE/H/0280/001	17096	H. LUNDBECK A/S	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
SEROPLEX 10 mg, comprimé pelliculé sécable	SE/H/0278/002	359 937-4	H. LUNDBECK A/S	FR
SEROPLEX 10 mg, comprimé pelliculé sécable	SE/H/0278/002	359 938-0	H. LUNDBECK A/S	FR
SEROPLEX 10 mg, comprimé pelliculé sécable	SE/H/0278/002	570 951-3	H. LUNDBECK A/S	FR
SEROPLEX 10 mg, comprimé pelliculé sécable	SE/H/0278/002	550 837 1 9	H. LUNDBECK A/S	FR
SEROPLEX 10 mg, comprimé pelliculé sécable	SE/H/0278/002	570 951 3 0	H. LUNDBECK A/S	FR
SEROPLEX 15 mg, comprimé pelliculé sécable	SE/H/0278/003	563 709-6	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	359 941-1	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	359 942-8	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	570 954-2 / 34009 570 954 2 0	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	563 710-4	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	550 837 3 3	H. LUNDBECK A/S	FR
SEROPLEX 20 mg, comprimé pelliculé sécable	SE/H/0278/004	570 954 2 0	H. LUNDBECK A/S	FR
SEROPLEX 20 mg/ml, solution buvable en	SE/H/0278/006	382 045-9	H. LUNDBECK A/S	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
gouttes				
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	364 289-7	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	359 935-1	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	359 936-8	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	570 950-7 / 34009 570 950 7 9	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	563 706-7	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	550 836 9 6	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	570 950 7 9	H. LUNDBECK A/S	FR
Sipralexa 10 mg comprimés pelliculés	SE/H/0278/002	BE238971	H. LUNDBECK A/S	LU
Sipralexa 10 mg filmomhulde tabletten	SE/H/0278/002	BE238971	H. LUNDBECK A/S	BE
SIPRALEXA 10 mg, comprimés pelliculés	SE/H/0278/002	BE 238971	H. LUNDBECK A/S	BE
Sipralexa 15 mg comprimés pelliculés	SE/H/0278/003	BE238944	H. LUNDBECK A/S	LU
Sipralexa 15 mg filmomhulde tabletten	SE/H/0278/003	BE238944	H. LUNDBECK A/S	BE
SIPRALEXA 15 mg, comprimés pelliculés	SE/H/0278/003	BE 238944	H. LUNDBECK A/S	BE
Sipralexa 20 mg comprimés pelliculés	SE/H/0278/004	BE238953	H. LUNDBECK A/S	LU
SIPRALEXA 20 mg, comprimés pelliculés	SE/H/0278/004	BE 238953	H. LUNDBECK A/S	BE
SIPRALEXA 20 mg, filmomhulde tabletten	SE/H/0278/004	BE238953	H. LUNDBECK A/S	BE
Sipralexa 5 mg comprimés pelliculés	SE/H/0278/001	BE238962	H. LUNDBECK A/S	LU
Sipralexa 5 mg filmomhulde tabletten	SE/H/0278/001	BE238962	H. LUNDBECK A/S	BE
SIPRALEXA 5 mg,	SE/H/0278/001	BE 238962	H. LUNDBECK A/S	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
comprimés pelliculés.				
S-Oropram 10 mg film-coated tablets	MT/H/0188/002	MA628/05602	ACTAVIS GROUP PTC EHF.	MT
S-Oropram 15 mg film-coated tablets	MT/H/0188/003	MA628/05603	ACTAVIS GROUP PTC EHF.	MT
S-Oropram 20 mg film-coated tablets	MT/H/0188/004	MA628/05604	ACTAVIS GROUP PTC EHF.	MT
S-Oropram 5 mg film-coated tablets	MT/H/0188/001	MA628/05601	ACTAVIS GROUP PTC EHF.	MT
Zocital 10 mg comprimidos revestidos por película	not available	5122452	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 10 mg comprimidos revestidos por película	not available	5122460	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 10 mg comprimidos revestidos por película	not available	5122478	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122502	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122510	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122528	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122536	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122544	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122551	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 5 mg comprimidos revestidos	not available	5122429	OWLPHARMA - CONSULTING, 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
por película				
Zocital 5 mg comprimidos revestidos por película	not available	5122437	OWLPHARMA - CONSULTING, LDA.	PT
Zocital 5 mg comprimidos revestidos por película	not available	5122445	OWLPHARMA - CONSULTING, LDA.	PT
ЦИПРАЛЕКС 10 mg филмирани таблетки	SE/H/0278/002	20020663	H. LUNDBECK A/S	BG