01 September 2017
EMA/590630/2017
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: escitalopram

Procedure no.: PSUSA/00001265/201612

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

Product 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 Mylan 5 mg Filmtabletten	DE/H/0594/001	64451.00.00	MYLAN DURA GMBH	DE
Escitalopram Mylan 10 mg Filmtabletten	DE/H/0594/002	64451.01.00	MYLAN DURA GMBH	DE
Escitalopram Mylan 20 mg Filmtabletten	DE/H/0594/004	64451.03.00	MYLAN DURA GMBH	DE
Escitalopram 10 mg film-coated tablets	DE/H/0594/002	PL 04569/0777	GENERICS [UK] LIMITED	UK
Escitalopram 20 mg film-coated tablets	DE/H/0594/004	PL 04569/0778	GENERICS [UK] LIMITED	UK
Escitalopram 5 mg film-coated tablets	DE/H/0594/001	PL 04569/0776	GENERICS [UK] LIMITED	UK
Escitil 15 mg filmtabletta	AT/H/0212/003/DC	OGYI-T-20966/07-09	EGIS PHARMACEUTICALS PLC	HU
Escitalopram Genedec 10 mg comprimidos orodispersíveis	not available	5673736	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Escitalopram Genedec 10 mg comprimidos orodispersíveis	not available	5673744	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Escitalopram Genedec 20 mg comprimidos orodispersíveis	not available	5673777	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Escitalopram Genedec 20 mg comprimidos orodispersíveis	not available	5673801	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
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

Product 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 beta 5 mg Filmtabletten	DE/H/3578/001	68164.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
ESCITALOPRAM SANDOZ 15 mg tabletti, suussa hajoava	PT/H/1134/003	31596	SANDOZ A/S	FI
ESCITALOPRAM SANDOZ 5 mg tabletti, suussa hajoava	PT/H/1134/001	31594	SANDOZ A/S	FI
Escitalopram Sandoz 5 mg – Schmelztabletten	PT/H/1134/001	135734	SANDOZ GMBH	AT
Escitalopram Sandoz 15 mg – Schmelztabletten	PT/H/1134/003	135737	SANDOZ GMBH	AT
Escitalopram Sandoz 20 mg/ml – Tropfen zum Einnehmen, Lösung	NL/H/2512/001	136143	SANDOZ GMBH	AT
Esciprex DisTab 5 mg Orodispersible Tablets	PT/H/1134/001	PA0711/194/005	ROWEX LTD	IE
Esciprex DisTab 15 mg Orodispersible Tablets	PT/H/1134/003	PA0711/194/007	ROWEX LTD	IE
Escitalopram-neuraxpharm 10 mg Filmtabletten	DE/H/1174/002	69884.00.00	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 5 mg Filmtabletten	DE/H/1174/001	69883.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Escitalopram Zentiva 20 mg comprimido revestido por película	UK/H/6146/004	5583265	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Escitalopram Zentiva 10 mg comprimido revestido por	UK/H/6146/002	5583273	SANOFI - 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
película			FARMACEUTICOS LDA	
ESCITALOPRAM ZENTIVA 10 MG COMPRIMIDO REVESTIDO POR PELICULA	UK/H/6146/002	5097225	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
ESCITALOPRAM ZENTIVA 10 MG COMPRIMIDO REVESTIDO POR PELICULA	UK/H/6146/002	5103478	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Escitalopram Zentiva 20 mg comprimido revestido por película	UK/H/6146/004	5097241	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Escitalopram Zentiva 20 mg comprimido revestido por película	UK/H/6146/004	5103510	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
ESCITALOPRAM ZENTIVA 10 MG COMPRIMIDO REVESTIDO POR PELICULA	UK/H/6146/002	5097233	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
ESCITALOPRAM ZENTIVA 10 MG COMPRIMIDO REVESTIDO POR PELICULA	UK/H/6146/002	5103502	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Escitalopram 5 mg film-coated tablets	not available	PL 42930/0019	WILCARE PHARMA LIMITED	UK
Escitalopram 10 mg film-coated tablets	not available	PL 42930/0020	WILCARE PHARMA LIMITED	UK
Escitalopram 20 mg film-coated tablets	not available	PL 42930/0021	WILCARE PHARMA LIMITED	UK
ESCITALOPRAM ZENTIVA LAB 10 mg, comprimé pelliculé sécable	SE/H/0279/002	277 341-0	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 10	SE/H/0279/002	277 342-7	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
mg, comprimé pelliculé sécable				
ESCITALOPRAM ZENTIVA LAB 10 mg, comprimé pelliculé sécable	SE/H/0279/002	277 343-3	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 5 mg, comprimé pelliculé	SE/H/0279/001	277 336-7	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 5 mg, comprimé pelliculé	SE/H/0279/001	277 337-3	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 5 mg, comprimé pelliculé	SE/H/0279/001	277 339-6	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 5 mg, comprimé pelliculé	SE/H/0279/001	277 340-4	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 15 mg, comprimé pelliculé sécable	SE/H/0279/003	277 345-6	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 15 mg, comprimé pelliculé sécable	SE/H/0279/003	277 346-2	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 15 mg, comprimé pelliculé sécable	SE/H/0279/003	277 347-9	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 20 mg, comprimé pelliculé sécable	SE/H/0279/004	277 348-5	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 20 mg, comprimé pelliculé sécable	SE/H/0279/004	277 349-1	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 20 mg, comprimé pelliculé sécable	SE/H/0279/004	277 351-6	H. LUNDBECK A/S	FR
ESCITALOPRAM ZENTIVA LAB 20 mg/ml, solution buvable en gouttes	SE/H/0279/006	277 333-8	H. LUNDBECK A/S	FR
Escitalopram Lundbeck 5 mg tabletti, kalvopäällysteinen	SE/H/0279/001	31841	H. LUNDBECK A/S	FI
Escitalopram Lundbeck 10 mg tabletti, kalvopäällysteinen	SE/H/0279/002	31842	H. LUNDBECK A/S	FI
Escitalopram Lundbeck 15 mg tabletti, kalvopäällysteinen	SE/H/0279/003	31843	H. LUNDBECK A/S	FI
Escitalopram Lundbeck 20 mg tabletti, kalvopäällysteinen	SE/H/0279/004	31844	H. LUNDBECK A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Escitalopram Lundbeck 10 mg Filmtabletten	SE/H/0279/002	90990.00.00	H. LUNDBECK A/S	DE
Escitalopram Lundbeck 20 mg Filmtabletten	SE/H/0279/004	90991.00.00	H. LUNDBECK A/S	DE
Escitalopram Lundbeck 20 mg/ml Tropfen zum Einnehmen, Lösung	SE/H/0279/006	90992.00.00	H. LUNDBECK A/S	DE
Entact 20 mg/ml gocce orali, soluzione	SE/H/0280/006	035768656	H. LUNDBECK A/S	IT
Escitalopram Lundbeck 20 mg/ml orala droppar, lösning	SE/H/0279/006	23324	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 5 mg, filmdragerade tabletter	SE/H/0279/001	17088	H. LUNDBECK A/S	SE
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
Escitalopram Lundbeck 20 mg, filmdragerade tabletter	SE/H/0279/004	17091	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 10 mg munsönderfallande tabletter	SE/H/0279/008	42058	H. LUNDBECK A/S	SE
Escitalopram Lundbeck 20 mg munsönderfallande tabletter	SE/H/0279/009	42059	H. LUNDBECK A/S	SE
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 rivestite con film	SE/H/0280/001	035768035	H. LUNDBECK A/S	IT
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

Product 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 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 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

Product 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	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
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 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

Product 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 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 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

Product 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	035768478	H. LUNDBECK A/S	IT
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 filmtabletta	SE/H/0278/002	OGYI-T-8634/01	H. LUNDBECK A/S	HU
Cipralex MELTZ 10 mg szájban diszpergálódó tabletta	SE/H/0278/008	OGYI-T-8634/05	H. LUNDBECK A/S	HU
Cipralex MELTZ 20 mg szájban diszpergálódó tabletta	SE/H/0278/009	OGYI-T-8634/06	H. LUNDBECK A/S	HU
Cipralex 10 mg filmom obalené tablety	SE/H/278/02	30/0210/07-S	H. LUNDBECK A/S	SK
Lexapro 15 mg film-coated tablets	SE/H/0278/003	PA 805/2/3	H. LUNDBECK A/S	IE
Lexapro 5 mg film-coated tablets	SE/H/0278/001	PA 805/2/1	H. LUNDBECK A/S	IE
Lexapro 10 mg film-coated tablets	SE/H/0278/002	PA 805/2/2	H. LUNDBECK A/S	IE
Lexapro 20 mg film-coated tablets	SE/H/0278/004	PA 805/2/4	H. LUNDBECK A/S	IE
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	H. LUNDBECK A/S	FR
SEROPLEX 5 mg, comprimé pelliculé	SE/H/0278/001	563 706-7	H. LUNDBECK A/S	FR
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

Product 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	563 707-3	H. LUNDBECK A/S	FR
SEROPLEX 15 mg, comprimé pelliculé sécable	SE/H/0278/003	359 939-7	H. LUNDBECK A/S	FR
SEROPLEX 15 mg, comprimé pelliculé sécable	SE/H/0278/003	359 940-5	H. LUNDBECK A/S	FR
SEROPLEX 15 mg, comprimé pelliculé sécable	SE/H/0278/003	570 953-6	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	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/ml, solution buvable en gouttes	SE/H/0278/006	382 045-9	H. LUNDBECK A/S	FR
Cipralex 20 mg/ml Gotas orales en solución	SE/H/0278/006	68811	LUNDBECK ESPANA, ES	ES
CIPRALEX 5 mg, comprimidos recubiertos con película	SE/H/0278/001	65.231	LUNDBECK ESPANA, ES	ES
CIPRALEX 10 mg, comprimidos recubiertos con película	SE/H/0278/002	65.230	LUNDBECK ESPANA, ES	ES
CIPRALEX 15 mg, comprimidos recubiertos con película	SE/H/0278/003	65.234	LUNDBECK ESPANA, ES	ES
CIPRALEX 20 mg, comprimidos recubiertos con película	SE/H/0278/004	65.233	LUNDBECK ESPANA, ES	ES
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

Product 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 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 comprimidos revestidos por película	SE/H/0278/001	4302089	H. LUNDBECK A/S	PT
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 comprimidos revestidos por película	SE/H/0278/002	4305082	H. LUNDBECK A/S	PT
Cipralex 10 mg comprimidos revestidos por película	SE/H/0278/002	5013958	H. LUNDBECK A/S	PT
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

Product 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 comprimidos revestidos por película	SE/H/0278/003	4308185	H. LUNDBECK A/S	PT
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/ml Gotas orais, solução	SE/H/0278/006	5049739	H. LUNDBECK A/S	PT
CIPRALEX 10 mg potahované tablety	SE/H/0278/002	30/276/02-C	H. LUNDBECK A/S	CZ
CIPRALEX Orotab 10 mg tablety dispergovatelné v ústech	SE/H/0278/008	30/486/10-C	H. LUNDBECK A/S	CZ
CIPRALEX Orotab 20 mg tablety dispergovatelné v ústech	SE/H/0278/009	30/487/10-C	H. LUNDBECK A/S	CZ
CIPRALEX 20 mg/ml perorální kapky, roztok	SE/H/0278/006	30/494/07-C	H. LUNDBECK A/S	CZ
Cipralex® 5 mg – Filmtabletten	SE/H/0278/001	1-24549	H. LUNDBECK A/S	AT
Cipralex® 10 mg – Filmtabletten	SE/H/0278/002	1-24550	H. LUNDBECK A/S	AT
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	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 Meltz	SE/H/0278/008	44624	H. LUNDBECK A/S	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Cipralex Meltz	SE/H/0278/009	44625	H. LUNDBECK A/S	DK
Cipralex	SE/H/0278/006	41275	H. LUNDBECK A/S	DK
CIPRALEX 5 mg filmdrasjerte tabletter	SE/H/0278/001	02-712	H. LUNDBECK A/S	NO
CIPRALEX 10 mg filmdrasjerte tabletter	SE/H/0278/002	02-713	H. LUNDBECK A/S	NO
CIPRALEX 15 mg filmdrasjerte tabletter	SE/H/0278/003	02-714	H. LUNDBECK A/S	NO
CIPRALEX 20 mg filmdrasjerte tabletter	SE/H/0278/004	02-715	H. LUNDBECK A/S	NO
Cipralex 20 mg/ml dråper, oppløsning	SE/H/0278/006	07-4892	H. LUNDBECK A/S	NO
CIPRALEX® 5 mg film-coated tablets	SE/H/0278/001	PL 13761/0008	H. LUNDBECK A/S	UK
CIPRALEX® 10 mg film-coated tablets	SE/H/0278/002	PL 13761/0009	H. LUNDBECK A/S	UK
CIPRALEX® 20 mg film-coated tablets	SE/H/0278/004	PL 13761/0011	H. LUNDBECK A/S	UK
CIPRALEX 20 mg/ml oral drops, solution	SE/H/0278/006	PL 13761/0028	H. LUNDBECK A/S	UK
CIPRALEX® 10 mg Filmtabletten	SE/H/0278/002	55880.01.00	H. LUNDBECK A/S	DE
CIPRALEX® 20 mg Filmtabletten	SE/H/0278/004	55880.03.00	H. LUNDBECK A/S	DE
Cipralex® 20 mg/ml Tropfen zum Einnehmen, Lösung	SE/H/0278/006	69241.00.00	H. LUNDBECK A/S	DE
Cipralex 10 mg apvalkotās tabletes	SE/H/0278/002	07 – 0298	H. LUNDBECK A/S	LV
Cipralex 20 mg apvalkotās tabletes	SE/H/0278/004	07-0300	H. LUNDBECK A/S	LV
Sipralexa 5 mg filmomhulde tabletten	SE/H/0278/001	BE238962	H. LUNDBECK A/S	BE
Sipralexa 10 mg filmomhulde tabletten	SE/H/0278/002	BE238971	H. LUNDBECK A/S	BE
Sipralexa 15 mg filmomhulde	SE/H/0278/003	BE238944	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
tabletten				
SIPRALEXA 20 mg, filmomhulde tabletten	SE/H/0278/004	BE238953	H. LUNDBECK A/S	BE
SIPRALEXA 5 mg, comprimés pelliculés.	SE/H/0278/001	BE 238962	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	BE 238944	H. LUNDBECK A/S	BE
SIPRALEXA 20 mg, comprimés pelliculés	SE/H/0278/004	BE 238953	H. LUNDBECK A/S	BE
CIPRALEX 5 mg film-coated tablets	SE/H/0278/001	MA591/00201	H. LUNDBECK A/S	MT
CIPRALEX® 10 mg film-coated tablets	SE/H/0278/002	MA601/00302	H. LUNDBECK A/S	MT
CIPRALEX 15 mg film-coated tablets	SE/H/0278/003	MA601/00303	H. LUNDBECK A/S	MT
CIPRALEX 20 mg film-coated tablets	SE/H/0278/004	MA601/00304	H. LUNDBECK A/S	MT
Sipralexa 5 mg comprimés pelliculés	SE/H/0278/001	BE238962	H. LUNDBECK A/S	LU
Sipralexa 10 mg comprimés pelliculés	SE/H/0278/002	BE238971	H. LUNDBECK A/S	LU
Sipralexa 15 mg comprimés pelliculés	SE/H/0278/003	BE238944	H. LUNDBECK A/S	LU
Sipralexa 20 mg comprimés pelliculés	SE/H/0278/004	BE238953	H. LUNDBECK A/S	LU
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 5 mg filmuhúðaðar töflur	SE/H/0278/001	IS/1/02/019/01	H. LUNDBECK A/S	IS
CIPRALEX 10 mg filmuhúðaðar	SE/H/0278/002	IS/1/02/019/02	H. LUNDBECK A/S	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
töflur				
CIPRALEX 15 mg filmuhúðaðar töflur	SE/H/0278/003	IS/1/02/019/03	H. LUNDBECK A/S	IS
CIPRALEX 20 mg filmuhúðaðar töflur	SE/H/0278/004	IS/1/02/019/04	H. LUNDBECK A/S	IS
CIPRALEX 5 mg, filmdragerade tabletter	SE/H/0278/001	17084	H. LUNDBECK A/S	SE
CIPRALEX 10 mg, filmdragerade tabletter	SE/H/0278/002	17085	H. LUNDBECK A/S	SE
CIPRALEX 15 mg, filmdragerade tabletter	SE/H/0278/003	17086	H. LUNDBECK A/S	SE
CIPRALEX 20 mg, filmdragerade tabletter	SE/H/0278/004	17087	H. LUNDBECK A/S	SE
PRILECT 5 mg, filmdragerade tabletter	SE/H/0280/001	17096	H. LUNDBECK A/S	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 10 mg munsönderfallande tabletter	SE/H/0280/008	42055	H. LUNDBECK A/S	SE
Prilect 20 mg munsönderfallande tabletter	SE/H/0280/009	42056	H. LUNDBECK A/S	SE
Cipralex 10 mg munsönderfallande tabletter	SE/H/0278/008	42055	H. LUNDBECK A/S	SE
Cipralex 20 mg munsönderfallande tabletter	SE/H/0278/009	42056	H. LUNDBECK A/S	SE
CIPRALEX 20 mg/ml orala droppar, lösning	SE/H/0278/006	23323	H. LUNDBECK A/S	SE
PRILECT 20 mg/ml orala droppar, lösning	SE/H/0280/006	23323	H. LUNDBECK A/S	SE
LEXAPRO 20 mg/ml druppels	SE/H/0279/006	RVG 35339	H. LUNDBECK A/S	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
voor oraal gebruik, oplossing				
LEXAPRO 5 mg filmomhulde tabletten	SE/H/0279/001	RVG 30494	H. LUNDBECK A/S	NL
LEXAPRO 10 mg filmomhulde tabletten	SE/H/0279/002	RVG 30495	H. LUNDBECK A/S	NL
LEXAPRO 15 mg filmomhulde tabletten	SE/H/0279/003	RVG 30496	H. LUNDBECK A/S	NL
LEXAPRO 20 mg filmomhulde tabletten	SE/H/0279/004	RVG 30497	H. LUNDBECK A/S	NL
Cipralex 5 mg tabletti, kalvopäällysteinen	SE/H/0278/001	17690	H. LUNDBECK A/S	FI
Cipralex 10 mg tabletti, kalvopäällysteinen	SE/H/0278/002	17691	H. LUNDBECK A/S	FI
Cipralex 15 mg tabletti, kalvopäällysteinen	SE/H/0278/003	17692	H. LUNDBECK A/S	FI
Cipralex 20 mg tabletti, kalvopäällysteinen	SE/H/0278/004	17693	H. LUNDBECK A/S	FI
Cipralex 10 mg tabletti, suussa hajoava	SE/H/0278/008	27218	H. LUNDBECK A/S	FI
Cipralex 20 mg tabletti, suussa hajoava	SE/H/0278/009	27219	H. LUNDBECK A/S	FI
Cipralex 20 mg/ml tipat, liuos	SE/H/0278/006	23395	H. LUNDBECK A/S	FI
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

Product 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 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 dengtos tabletes	SE/H/0278/002	LT/1/07/0832/027	H. LUNDBECK A/S	LT
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 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 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

Product 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 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 film	SE/H/0278/001	035767173/M	H. LUNDBECK A/S	IT
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 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

Product 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 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 compresse rivestite con film	SE/H/0278/002	035767250/M	H. LUNDBECK A/S	IT
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 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

Product 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	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
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 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	035767161/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 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
Cipralex 10 mg/ml gocce orali, soluzione	SE/H/0278/002	035767641/M	H. LUNDBECK A/S	IT
Cipralex MELTZ 10 mg comprimate orodispersabile	SE/H/0278/008	2602/2010/01-03	H. LUNDBECK A/S	RO
Cipralex MELTZ 20 mg comprimate orodispersabile	SE/H/0278/009	2603/2010/01-03	H. LUNDBECK A/S	RO
CIPRALEX 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/001	76959	H. LUNDBECK A/S	GR
CIPRALEX 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/002	76961	H. LUNDBECK A/S	GR
CIPRALEX 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/003	76963	H. LUNDBECK A/S	GR
CIPRALEX 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/004	76964	H. LUNDBECK A/S	GR
ENTACT 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/001	76950/26-11-07	H. LUNDBECK A/S	GR
ENTACT 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/002	76953/26-11-07	H. LUNDBECK A/S	GR
ENTACT 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/003	76954/26-11-07	H. LUNDBECK A/S	GR
ENTACT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0279/004	76955/26-11-07	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
CIPRALEX 10 mg/ml πόσιμες σταγόνες, διάλυμα	SE/H/0278/005	76965	H. LUNDBECK A/S	GR
CIPRALEX 5 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/001	20266	H. LUNDBECK A/S	CY
CIPRALEX 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/002	20267	H. LUNDBECK A/S	CY
CIPRALEX 15 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/003	20268	H. LUNDBECK A/S	CY
CIPRALEX 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0278/004	20269	H. LUNDBECK A/S	CY
Cipralex 20 mg/ml gocce orali, soluzione	SE/H/0278/006	035767654/M	H. LUNDBECK A/S	IT
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
Cipralex 10 mg filmsko obložene tablete	SE/H/0278/002	5363-I-689/07	LUNDBECK PHARMA D.O.O	SI
Lexapro, 5 mg, tabletki powlekane	SE/H/0278/001	14282	H. LUNDBECK A/S	PL
Lexapro, 10 mg, tabletki powlekane	SE/H/0278/002	14283	H. LUNDBECK A/S	PL
ЦИПРАЛЕКС МЕЛЦ 10 mg таблетки, диспергиращи се в устата	SE/H/0278/008	20100096	H. LUNDBECK A/S	BG
ЦИПРАЛЕКС МЕЛЦ 20 mg таблетки, диспергиращи се в устата	SE/H/0278/009	20100097	H. LUNDBECK A/S	BG
ЦИПРАЛЕКС 10 mg филмирани таблетки	SE/H/0278/002	20020663	H. LUNDBECK A/S	BG
CIPRALEX 10 mg comprimate filmate	SE/H/0278/002	9256/2016/01-19	H. LUNDBECK A/S	RO
Zocital 5 mg comprimidos revestidos por película	not available	5122437	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zocital 5 mg comprimidos revestidos por película	not available	5122445	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 10 mg comprimidos revestidos por película	not available	5122460	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 10 mg comprimidos revestidos por película	not available	5122478	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122510	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122528	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122544	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122551	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 5 mg comprimidos revestidos por película	not available	5122429	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 10 mg comprimidos revestidos por película	not available	5122452	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 15 mg comprimidos revestidos por película	not available	5122502	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zocital 20 mg comprimidos revestidos por película	not available	5122536	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Escitalopram-neuraxpharm 10 mg Schmelztabletten	DE/H/5050/002	88039.00.00	NEURAXPHARM ARZNEIMITTEL	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
			GMBH	
Escitalopram-neuraxpharm 20 mg Schmelztabletten	DE/H/5050/004	88041.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Zecidec 5 mg comprimidos revestidos por película	not available	5122577	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 5 mg comprimidos revestidos por película	not available	5122601	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 10 mg comprimidos revestidos por película	not available	5122627	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 10 mg comprimidos revestidos por película	not available	5122635	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 15 mg comprimidos revestidos por película	not available	5122650	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 15 mg comprimidos revestidos por película	not available	5122668	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 20 mg comprimidos revestidos por película	not available	5122700	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 20 mg comprimidos revestidos por película	not available	5122718	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 5 mg comprimidos revestidos por película	not available	5122569	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 10 mg comprimidos revestidos por película	not available	5122619	DECOMED FARMACÊUTICA, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zecidec 15 mg comprimidos revestidos por película	not available	5122643	DECOMED FARMACÊUTICA, S.A.	PT
Zecidec 20 mg comprimidos revestidos por película	not available	5122676	DECOMED FARMACÊUTICA, S.A.	PT
ENLIFT 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	EE/H/0181/002	34924/17-5-13	MEDOCHEMIE HELLAS SA	GR
ENLIFT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	EE/H/0181/004	37662/17-5-13	MEDOCHEMIE HELLAS SA	GR
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
Escitalopram 15 mg Film-Coated Tablets	UK/H/6523/003	PL 25298/0110	BROWN & BURK UK LIMITED	UK
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-ratiopharm® 5 mg Filmtabletten	HU/H/0179/001	2012110001	RATIOPHARM GMBH	LU
Escitalopram-ratiopharm® 15 mg Filmtabletten	HU/H/0179/003	2012100113	RATIOPHARM GMBH	LU