14 September 2023
EMA/419197/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): paroxetine

Procedure No. PSUSA/00002319/202212

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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
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/001	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/002	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/003	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/004	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/005	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/006	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/007	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/008	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/009	MEDOCHEMIE LTD.	LT
ARKETIS 10 mg tabletės	NL/H/0449/001	LT/1/07/0679/010	MEDOCHEMIE LTD.	LT
Arketis 10 mg, tablets	NL/H/0449/001	MA032/06601	MEDOCHEMIE LTD.	MT
Arketis 10 mg, δισκία	NL/H/0449/001	20185	MEDOCHEMIE LTD.	CY
ARKETIS 30 mg tabletes	NL/H/0449/003	LT/1/07/0679/021	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletes	NL/H/0449/003	LT/1/07/0679/022	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletes	NL/H/0449/003	LT/1/07/0679/023	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletes	NL/H/0449/003	LT/1/07/0679/024	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletes	NL/H/0449/003	LT/1/07/0679/025	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletės	NL/H/0449/003	LT/1/07/0679/026	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletės	NL/H/0449/003	LT/1/07/0679/027	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletės	NL/H/0449/003	LT/1/07/0679/028	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletės	NL/H/0449/003	LT/1/07/0679/029	MEDOCHEMIE LTD.	LT
ARKETIS 30 mg tabletės	NL/H/0449/003	LT/1/07/0679/030	MEDOCHEMIE LTD.	LT
Arketis 30 mg, tablets	NL/H/0449/003	MA032/06603	MEDOCHEMIE LTD.	MT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/031	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/032	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/033	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/034	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/035	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/036	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/037	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/038	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/039	MEDOCHEMIE LTD.	LT
ARKETIS 40 mg tabletės	NL/H/0449/004	LT/1/07/0679/040	MEDOCHEMIE LTD.	LT
Arketis 40 mg, tablets	NL/H/0449/004	MA032/06604	MEDOCHEMIE LTD.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Arketis 40 mg, δισκία	NL/H/0449/004	20190	MEDOCHEMIE LTD.	CY
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444013	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444025	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444114	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444037	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444049	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444126	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444052	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444138	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444140	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg	DK/H/0240/001	035444064	AZIENDE CHIMICHE	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			RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444076	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444153	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444088	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444090	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444165	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444177	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444102	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
DAPAROX 20 mg compresse rivestite con film	DK/H/0240/001	035444189	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Daparox 20 mg comprimidos recubiertos con película	ES/H/0406/001	65.259	ANGELINI PHARMA ESPANA 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
DEROXAT 20 mg, comprimé pelliculé sécable	NL/H/0566/001	NL16996	LABORATOIRE GLAXOSMITHKLINE	FR
DEROXAT 20 mg/10 ml, suspension buvable.	NL/H/0566/003	NL22372	LABORATOIRE GLAXOSMITHKLINE	FR
DIVARIUS 20 mg, comprimé pelliculé sécable	not available	3400936011250	CHIESI S.A.S.	FR
DIVARIUS 20 mg, comprimé pelliculé sécable	not available	3400936011311	CHIESI S.A.S.	FR
DIVARIUS 20 mg, comprimé pelliculé sécable	not available	3400956427208	CHIESI S.A.S.	FR
Dropax 10 mg/ml gotas orais, solução	IT/H/0136/001	5928486	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Dropax 10 mg/ml gotas orais, solução	IT/H/0136/001	5928486	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Dropax 10 mg/ml gotas orais, solução	IT/H/0136/001	5928486	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Dropax 10 mg/ml gotas orais, solu｢ｱｱ｣o	IT/H/0136/001	5737333	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Dropax 10 mg/ml gotas orais, solu｢ｱｱ｣o	IT/H/0136/001	5737333	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Dropax 10 mg/ml gotas orais, solu｢ｱｱ｣o	IT/H/0136/001	5737333	ITALFARMACO - PRODUTOS FARMACÊUTICOS, LDA.	PT
Ennos 20 mg-Filmtabletten	DK/H/0240/001	1-24116	G.L. PHARMA GMBH	AT
Euplix 20 mg Filmtabletten	DK/H/0240/001	51352.00.00	SYNTHON BV	DE
Euplix 33,1 mg/ml, orale dråber, opløsning	DK/H/0240/002	38097	SYNTHON BV	DK
Euplix, filmovertrukne tabletter	DK/H/0240/001	31126	SYNTHON BV	DK
Eutimil 2 mg/ml sospensione orale	NL/H/0567/003	027964028	GLAXOSMITHKLINE S.P.A.	IT
EUTIMIL 20 mg compresse rivestite con	NL/H/0567/001	027964030	GLAXOSMITHKLINE S.P.A.	IT

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film				
EUTIMIL 20 mg compresse rivestite con film	NL/H/0567/001	027964042	GLAXOSMITHKLINE S.P.A.	IT
Frosinor 20 mg comprimidos recubiertos con película	NL/H/0567/001	59.466	GLAXOSMITHKLINE, S.A.	ES
Motivan 20 mg comprimidos recubiertos con película	NL/H/0568/001	59.535	GLAXOSMITHKLINE, S.A.	ES
Optipar	DK/H/0237/002	31777	HEXAL A/S	DK
Optipar 40 mg tabletti, kalvopäällysteinen	DK/H/0237/002	16398	HEXAL A/S	FI
Optipar, filmovertrukne tabletter	DK/H/0237/003	35323	HEXAL A/S	DK
Paroxat 10 mg Filmtabletten	DK/H/0233/003	64765.00.00	HEXAL AG	DE
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

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Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/013	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/027	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/028	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/029	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/030	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/031	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/032	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/033	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/034	GLAXOSMITHKLINE TRADING SERVICES	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
			LIMITED	
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/035	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/036	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/037	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/038	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/039	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
PAROXAT 20 mg filmsko obložene tablete	NL/H/0567/001	H/04/01234/052	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 20 mg filmtabletta	NL/H/0567/001	OGYI-T-8263/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Paroxat 20 mg filmtabletta	NL/H/0567/001	OGYI-T-8263/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/041	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/042	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/043	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko	NL/H/0567/002	H/04/01234/048	GLAXOSMITHKLINE	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
obložene tablete			TRADING SERVICES LIMITED	
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/049	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/050	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/051	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg filmsko obložene tablete	NL/H/0567/002	H/04/01234/040	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Paroxat 30 mg Filmtabletten	DK/H/0233/004	64766.00.00	HEXAL AG	DE
Paroxat 40 mg Filmtabletten	DK/H/0233/002	51396.01.00	HEXAL AG	DE
Paroxat Hexal 10 mg - Filmtabletten	DK/H/0233/003	1-26608	HEXAL PHARMA GMBH	AT
Paroxat Hexal 30 mg - Filmtabletten	DK/H/0233/004	1-26609	HEXAL PHARMA GMBH	AT
Paroxat Hexal 40 mg - Filmtabletten	DK/H/0233/002	1-24136	HEXAL PHARMA GMBH	AT
paroxedura 40 mg Tabletten	NL/H/0450/004	66370.00.00	MYLAN GERMANY GMBH	DE
Paroxetin - 1 A Pharma 10 mg Filmtabletten	DK/H/0237/003	64767.00.00	1 A PHARMA GMBH	DE
Paroxetin - 1 A Pharma 30 mg Filmtabletten	DK/H/0237/004	64768.00.00	1 A PHARMA GMBH	DE
Paroxetin - 1 A Pharma 40 mg Filmtabletten	DE/H/6987/002	51402.01.00	1 A PHARMA GMBH	DE
Paroxetin "HEXAL", filmovertrukne tabletter	DK/H/0233/002	31632	HEXAL A/S	DK
Paroxetin "HEXAL", filmovertrukne tabletter	DK/H/0233/003	35317	HEXAL A/S	DK
Paroxetin beta 10 mg Tabletten	DE/H/3070/001	66634.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE

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Paroxetin beta 30 mg Tabletten	DE/H/3070/003	66636.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Paroxetin beta 40 mg Tabletten	DE/H/3070/004	66637.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Paroxetin ratiopharm 40 mg Tabletten	NL/H/0453/001	1-26740	TEVA B.V	AT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos	DK/H/1135/002	DK/H/1135/002	GENERIS 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
revestidos por película				
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurobindo 30 mg comprimidos revestidos por película	DK/H/1135/002	DK/H/1135/002	GENERIS FARMACÊUTICA, S.A.	PT
Paroxetina Aurovitas 30 mg comprimidos recubiertos con película	PT/H/1559/002	81.512	AUROVITAS SPAIN,S.A.U.	ES
paroxetina cinfa 20 mg comprimidos recubiertos con película EFG	not available	66.932	LABORATORIOS CINFA, S.A.	ES
Paroxetina Mabo 30 mg, comprimidos	NL/H/0831/003	68508	MABO-FARMA, S.A	ES
Paroxetina Mabo 40 mg, comprimidos	NL/H/0831/004	68509	MABO-FARMA, S.A	ES
Paroxetina STADA 30 mg comprimidos	not available	76.031	LABORATORIO STADA, S.L.	ES
Paroxetina Stada 40 mg	not available	76.032	LABORATORIO STADA, 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
comprimidos				
Paroxetine 10 mg Film-coated tablets.	not available	PL 30322/0025	ALISSA HEALTHCARE RESEARCH LIMITED	XI
Paroxetine 10 mg Film-Coated Tablets.	not available	PL 17780/0879	ZENTIVA PHARMA UK LIMITED	XI
Paroxetine 10mg film-coated tablets	not available	PL 49565/0093	RUDIPHARM LIMITED	XI
Paroxetine 10mg film-coated tablets	not available	PL 20416/0549	CRESCENT PHARMA LIMITED	XI
Paroxetine 20 mg Film-Coated Tablets	not available	PL 20416/0355	CRESCENT PHARMA LIMITED	XI
Paroxetine 20 mg Film-coated tablets.	not available	PL 30322/0026	ALISSA HEALTHCARE RESEARCH LIMITED	XI
Paroxetine 20 mg Tablets	not available	PL 36687/0342	TORRENT PHARMA (UK) LTD.	XI
Paroxetine 20 mg, Filmomhulde tabletten	DK/H/0240/001	RVG 26382	GENTHON BV	NL
Paroxetine 30 mg Film-coated tablets.	not available	PL 30322/0027	ALISSA HEALTHCARE RESEARCH LIMITED	XI
Paroxetine 30 mg Tablets	not available	PL 36687/0343	TORRENT PHARMA (UK) LTD.	XI
Paroxetine 30mg film-coated tablets	not available	PL 20416/0550	CRESCENT PHARMA LIMITED	XI
Paroxetine 40 mg Film-Coated Tablets	not available	PL 20416/0363	CRESCENT PHARMA LIMITED	XI
Paroxetine 40 mg film-coated tablets	NL/H/3147/004	PL19156/0137	JUBILANT PHARMACEUTICALS NV	XI
Paroxetine 40 mg Film-coated tablets.	not available	PL 30322/0028	ALISSA HEALTHCARE RESEARCH LIMITED	XI
Paroxetine 40 mg Tablets	not available	PL 20117/0105	MORNINGSIDE HEALTHCARE LTD	XI
Paroxetine 40 mg Tablets	not available	PL 51718/0013	CLYDESDALE PHARMA LIMITED	XI
Paroxetine 40 mg, tabletten	NL/H/0449/004	RVG 31774	ICC B.V.	NL
Paroxetine 40 mg, tabletten	NL/H/0831/004	RVG 33148	ICC B.V.	NL
Paroxetine 40 mg,	NL/H/0453/001	RVG 33151	RATIOPHARM GMBH	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
tabletten				
Paroxetine 40 mg, tabletten	NL/H/0450/004	RVG 33137	ICC B.V.	NL
Paroxetine Aurobindo 30 mg film-coated tablets	PT/H/1559/002	MA807/09902	AUROBINDO PHARMA (MALTA) LIMITED	MT
PAROXETINE BIOGARAN 20 mg, comprimé pelliculé sécable	NL/H/0567/001	NL16997	LABORATOIRE GLAXOSMITHKLINE	FR
Paroxetine EG 20 mg comprimés pelliculés	DK/H/0240/001	BE269062	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg comprimés pelliculés	DK/H/0240/001	BE227701	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg comprimés pelliculés	DK/H/0240/001	BE227717	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg comprimés pelliculés	DK/H/0240/001	2003100016	EUROGENERICS N.V./S.A.	LU
Paroxetine EG 20 mg filmomhulde tabletten	DK/H/0240/001	BE227717	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg filmomhulde tabletten	DK/H/0240/001	BE227701	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg filmomhulde tabletten	DK/H/0240/001	BE269062	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg Filmtabletten	DK/H/0240/001	BE227717	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg Filmtabletten	DK/H/0240/001	BE227701	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 20 mg Filmtabletten	DK/H/0240/001	BE269062	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 40 mg comprimés	NL/H/0449/004	BE316197	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 40 mg comprimés	NL/H/0449/004	2010050087	EUROGENERICS N.V./S.A.	LU
Paroxetine EG 40 mg tabletten	NL/H/0449/004	BE316197	EUROGENERICS N.V./S.A.	BE
Paroxetine EG 40 mg Tabletten	NL/H/0449/004	BE316197	EUROGENERICS N.V./S.A.	BE
Paroxetine GSK 2 mg/ml, oral suspension	NL/H/0567/003	RVG 30168	GLAXOSMITHKLINE B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Paroxetine GSK 20 mg tablet, filmomhulde tabletten	NL/H/0568/001	RVG 30169	GLAXOSMITHKLINE B.V.	NL
Paroxetine GSK 20 mg, film coated tablets	NL/H/0567/001	RVG 30166	GLAXOSMITHKLINE B.V.	NL
Paroxetine GSK 30 mg, film coated tablets	NL/H/0567/002	RVG 30167	GLAXOSMITHKLINE B.V.	NL
Paroxetine Jubilant 40 mg filmomhulde tabletten	NL/H/3147/004	RVG 115271	JUBILANT PHARMACEUTICALS NV	NL
Paroxetine Prolepha 40 mg, tabletten	not available	RVG 100091	PROLEPHA RESEARCH B.V.	NL
Paroxetine Sandoz 40 mg filmomhulde tabletten	DK/H/0237/002	BE232285	SANDOZ N.V.	BE
Paroxetine Sandoz 40 mg filmomhulde tabletten	DK/H/0237/002	BE232294	SANDOZ N.V.	BE
Paroxetin-Hormosan 33,1 mg/ml Tropfen zum Einnehmen, Lösung	NL/H/0877/001	67595.00.00	HORMOSAN PHARMA GMBH	DE
Paroxetin-neuraxpharm 10 mg Tabletten	not available	61351.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Paroxetin-neuraxpharm 30 mg Tabletten	not available	61353.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Paroxetin-neuraxpharm 40 mg Tabletten	not available	61354.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Paroxia 40 mg, δισκία	NL/H/0449/004	45309/1-7-2015	MEDOCHEMIE HELLAS SA	GR
SEREUPIN 20 mg compresse rivestite con film	not available	027965033	GLAXOSMITHKLINE S.P.A.	IT
Sereupin 20 mg/10 ml sospensione orale	not available	027965021	GLAXOSMITHKLINE S.P.A.	IT
SEROXAT 10 mg film-coated tablets.	NL/H/0566/04	PA 1077/97/4	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seroxat 10 mg film-coated tablets.	not available	10592/0218	SMITHKLINE BEECHAM LTD	XI
Seroxat 10 mg	NL/H/0566/004	03-1911	GLAXOSMITHKLINE AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmdrasjert tablett				
Seroxat 10 mg filmuhúðaðar töflur.	NL/H/0566/004	IS/1/05/011/01	GLAXOSMITHKLINE PHARMA A/S	IS
Seroxat 10 mg tablet, filmomhulde tabletten	NL/H/0566/004	RVG 29433	GLAXOSMITHKLINE B.V.	NL
Seroxat 2 mg/ml – Suspension zum Einnehmen	NL/H/0566/003	1-22309	GLAXOSMITHKLINE PHARMA GMBH.	AT
Seroxat 2 mg/ml sospensione orale	NL/H/0566/003	027963026	GLAXOSMITHKLINE S.P.A.	IT
Seroxat 2 mg/ml Suspension zum Einnehmen	NL/H/0566/003	44028.00.01	GLAXOSMITHKLINE GMBH & CO. KG	DE
Seroxat 20 mg - Filmtabletten	NL/H/0566/001	1-19931	GLAXOSMITHKLINE PHARMA GMBH.	AT
SEROXAT 20 mg apvalkotas tabletes	NL/H/0566/001	99-1041	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
SEROXAT 20 mg compresse rivestite con film	NL/H/0566/001	027963038	GLAXOSMITHKLINE S.P.A.	IT
SEROXAT 20 mg compresse rivestite con film	NL/H/0566/001	027963040	GLAXOSMITHKLINE S.P.A.	IT
SEROXAT 20 mg comprimate filmate	not available	601/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
SEROXAT 20 mg comprimate filmate	not available	601/2008/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
SEROXAT 20 mg comprimés pelliculés	NL/H/0566/001	BE159774	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 20 mg comprimés pelliculés	NL/H/0566/001	BE285354	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 20 mg comprimés pelliculés	NL/H/0566/001	2011041083	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Seroxat 20 mg comprimidos recubiertos con película	NL/H/0566/01	59.468	GLAXOSMITHKLINE, S.A.	ES
SEROXAT 20 mg	NL/H/0566/001	2313484	GLAXOSMITHKLINE -	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			PRODUTOS FARMACEUTICOS, LDA	
SEROXAT 20 mg comprimidos revestidos por película	NL/H/0566/001	2313385	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
SEROXAT 20 mg comprimidos revestidos por película	NL/H/0566/001	5053780	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
SEROXAT 20 mg film-coated tablets	NL/H/0566/001	PA 1077/97/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
SEROXAT 20 mg film-coated tablets.	NL/H/0566/001	MA192/02501	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Seroxat 20 mg film-coated tablets.	not available	10592/0001	SMITHKLINE BEECHAM LTD	XI
Seroxat 20 mg filmdragerad tablett	NL/H/0566/001	10814	GLAXOSMITHKLINE (IRELAND) LIMITED	FI
Seroxat 20 mg filmdragerad tablett.	NL/H/0566/001	11376	GLAXOSMITHKLINE AB	SE
Seroxat 20 mg filmdrasjert tablett	NL/H/0566/001	00-7840	GLAXOSMITHKLINE AS	NO
Seroxat 20 mg filmom obalené tablety	NL/H/0566/001	30/0859/95-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
SEROXAT 20 mg filmom obložene tablete	not available	HR-H-051250245	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
SEROXAT 20 mg filmomhulde tabletten	NL/H/0566/001	BE159774	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 20 mg filmomhulde tabletten	NL/H/0566/001	BE285354	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/004	GLAXOSMITHKLINE TRADING SERVICES	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
			LIMITED	
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/013	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/014	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/029	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/030	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko	NL/H/0566/001	H/00/01408/031	GLAXOSMITHKLINE	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
obložene tablete			TRADING SERVICES LIMITED	
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/032	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/033	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/034	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/035	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/036	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/037	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/038	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/039	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/040	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/041	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg filmsko obložene tablete	NL/H/0566/001	H/00/01408/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 20 mg Filmtabletten	NL/H/0566/001	44028.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Seroxat 20 mg	NL/H/0566/001	900044	GLAXOSMITHKLINE PHARMA	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
filmuhúðaðar töflur.			A/S	
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/001	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/003	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/004	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/008	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/009	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/011	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/012	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/029	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/031	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plevele dengtos tabletes	NL/H/0566/001	LT/1/06/0597/032	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/002	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/005	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/006	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/007	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/010	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/013	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/030	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/033	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele	NL/H/0566/001	LT/1/06/0597/034	GLAXOSMITHKLINE	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 tabletės			(IRELAND) LIMITED	
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/035	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/036	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/037	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/038	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/039	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/040	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/041	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/042	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg plėvele dengtos tabletės	NL/H/0566/001	LT/1/06/0597/043	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 20 mg potahované tablety	NL/H/0566/001	30/490/96-A/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Seroxat 20 mg tablet, filmomhulde tabletten	NL/H/0566/001	RVG 14668	GLAXOSMITHKLINE B.V.	NL
Seroxat 20 mg tabletti, kalvopäällysteinen	NL/H/0566/001	10814	GLAXOSMITHKLINE (IRELAND) LIMITED	FI
SEROXAT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0566/001	14178	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
SEROXAT 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0566/001	2017401	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
SEROXAT 20 mg, Filmtabletten	NL/H/0566/001	BE159774	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 20 mg, Filmtabletten	NL/H/0566/001	BE285354	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 20 mg, Filmtabletten	NL/H/0566/001	2011041083	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
SEROXAT 20 mg/10 ml	NL/H/0566/003	PA 1077/97/1	GLAXOSMITHKLINE	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
oral suspension.			(IRELAND) LIMITED	
Seroxat 20 mg/10 ml oral suspension.	not available	10592/0092	SMITHKLINE BEECHAM LTD	XI
SEROXAT 30 mg comprimés pelliculés	NL/H/0566/002	BE285363	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 30 mg comprimés pelliculés	NL/H/0566/002	BE285372	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 30 mg comprimés pelliculés	NL/H/0566/002	2011041084	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
SEROXAT 30 mg film coated tablets.	NL/H/0566/002	PA 1077/97/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Seroxat 30 mg film-coated tablets.	not available	10592/0002	SMITHKLINE BEECHAM LTD	XI
Seroxat 30 mg filmom obalené tablety	NL/H/0566/002	30/0339/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
SEROXAT 30 mg filmom obložene tablete	not available	HR-H-284734421	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
SEROXAT 30 mg filmomhulde tabletten	NL/H/0566/002	BE285363	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 30 mg filmomhulde tabletten	NL/H/0566/002	BE285372	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/043	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/044	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/045	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/050	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/051	GLAXOSMITHKLINE TRADING SERVICES LIMITED	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
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/052	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/02	H/00/01408/053	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg filmsko obložene tablete	NL/H/0566/002	H/00/01408/042	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/018	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/019	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/021	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/022	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/044	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/045	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/046	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg plėvele dengtos tabletės	NL/H/0566/002	LT/1/06/0597/047	GLAXOSMITHKLINE (IRELAND) LIMITED	LT
Seroxat 30 mg potahované tablety	NL/H/0566/002	30/490/96-B/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Seroxat 30 mg tablet, filmomhulde tabletten	NL/H/0566/002	RVG 27135	GLAXOSMITHKLINE B.V.	NL
SEROXAT 30 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0566/002	2017402	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
SEROXAT 30 mg, Filmtabletten	NL/H/0566/002	BE285363	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 30 mg, Filmtabletten	NL/H/0566/002	BE285372	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
SEROXAT 30 mg, Filmtabletten	NL/H/0566/002	2011041084	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Seroxat Suspensie 2 mg/ml, suspensie voor oraal gebruik	NL/H/0566/003	RVG 20557	GLAXOSMITHKLINE B.V.	NL
Seroxat Suspensie 2 mg/ml, suspensie voor oraal gebruik	NL/H/0566/003	RVG20557	GLAXOSMITHKLINE B.V.	NL
SEROXAT, 20 mg õhukese polümeerikattega tabletid	NL/H/0566/001	161197	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Seroxat, 20 mg, tabletki powlekane	NL/H/0566/001	R/6405	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Seroxat, filmovertrukne tabletter	NL/H/0566/001	13893	GLAXOSMITHKLINE PHARMA A/S	DK
Seroxat, filmovertrukne tabletter	NL/H/0566/002	33651	GLAXOSMITHKLINE PHARMA A/S	DK
SOLBEN 40mg δισκία	NL/H/0450/004	32477/8-4-2014	GAP S.A.	GR
Syntopar 10 mg, δισκία	NL/H/0831/001	20195	CODAL SYNTO LTD	CY
Syntopar 40 mg, δισκία	NL/H/0831/004	20198	CODAL SYNTO LTD	CY
Парикс 40 mg таблетки	not available	20150151	MEDOCHEMIE LTD.	BG
ПАРОКСАТ 20 mg филмирани таблетки	not available	20030653	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
СЕРОКСАТ 20 mg филмирани таблетки	not available	20000014	GLAXOSMITHKLINE (IRELAND) LIMITED	BG