1 September 2017
EMA/583896/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: citalopram

Procedure no.: PSUSA/00000779/201612

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

An agency of the European Union

© European Medicines Agency, 2017. Reproduction is authorised provided the source is acknowledged.

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
Citalostad 20 mg Filmtabletten	not available	1-25317	STADA ARZNEIMITTEL GMBH	AT
Citalopram 20 mg AbZ	NL/H/0468/002	RVG 27643	ABZ-PHARMA GMBH	NL
Citalopram 40 mg AbZ	NL/H/0468/003	RVG 27644	ABZ-PHARMA GMBH	NL
Citalopram AbZ 20 mg Filmtabletten	NL/H/0468/002	58953.01.00	ABZ-PHARMA GMBH	DE
Citalopram AbZ 40 mg Filmtabletten	NL/H/0468/003	58953.02.00	ABZ-PHARMA GMBH	DE
Citalopram Azevedos 10 mg comprimidos revestidos por película	not available	5063805	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 10 mg comprimidos revestidos por película	not available	5063813	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 10 mg comprimidos revestidos por película	not available	5063821	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 20 mg comprimidos revestidos por pelicula	not available	5063847	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 20 mg comprimidos revestidos por pelicula	not available	5063854	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 20 mg comprimidos revestidos por pelicula	not available	5063862	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 40 mg comprimidos revestidos por película	not available	5063904	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 40 mg comprimidos revestidos por película	not available	5063912	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 40 mg comprimidos revestidos por película	not available	5063920	LABORATÓRIOS AZEVEDOS - INDÚSTRIA 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
Citalopram Azevedos 10 mg comprimidos revestidos por película.	not available	5063771	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 20 mg comprimidos revestidos por película.	not available	5063839	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citalopram Azevedos 40 mg comprimidos revestidos por película.	not available	5063870	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Citaloprol 40 mg/ml Πόσιμες σταγόνες διάλυμα	not available	2416/3-3-2011	NOVENDIA PHARMACEUTICALS LTD	GR
LINISAN 10 mg comprimate filmate	not available	7778/2006/01	GEDEON RICHTER ROMÂNIA S.A.	RO
LINISAN 20 mg comprimate filmate	not available	7779/2006/01	GEDEON RICHTER ROMÂNIA S.A.	RO
CITALOPRAM SIGMA-TAU 40 mg/ml gocce orali, soluzione	not available	036723017	SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A.	IT
Citalopram-neuraxpharm 30 mg Filmtabletten	not available	63286.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Citalopram Genedec 10 mg comprimidos revestidos por película	not available	5449491	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 10 mg comprimidos revestidos por película	not available	5449590	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 20 mg comprimidos revestidos por película	not available	5449798	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 20 mg comprimidos revestidos por película	not available	5449897	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
Citalopram Genedec 40 mg comprimidos revestidos por película	not available	5450093	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 40 mg comprimidos revestidos por película	not available	5450192	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 10 mg comprimidos revestidos por película	not available	5449392	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 20 mg comprimidos revestidos por película	not available	5449699	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalopram Genedec 40 mg comprimidos revestidos por película	not available	5449996	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Citalostad 40 mg Filmtabletten	not available	1-25316	STADA ARZNEIMITTEL GMBH	AT
Citalopram 20 mg Holsten	not available	59744.00.00	HOLSTEN PHARMA GMBH	DE
Citalopram-Hormosan 20 mg	not available	54073.01.00	HORMOSAN PHARMA GMBH	DE
Citalopram-Hormosan 40 mg	not available	54073.02.00	HORMOSAN PHARMA GMBH	DE
Citalopram-Hormosan 20 mg/ml Tropfen zum Einnehmen, Lösung	DE/H/4478/001	95290.00.00	HORMOSAN PHARMA GMBH	DE
ROPRAMIN®	not available	38706/12/06-06-2013	IASIS PHARMA	GR
ROPRAMIN	not available	20742	IASIS PHARMA	CY
ROPRAMIN	not available	20743	IASIS PHARMA	CY
ROPRAMIN®	not available	44633/06-06-2013	IASIS PHARMA	GR
ROPRAMIN®	not available	42132/11/23-01-2013	IASIS PHARMA	GR
PREFUCET	not available	13959	RAFARM SA.	GR
PREFUCET	not available	13960	RAFARM SA.	GR
PREFUCET	not available	7784	RAFARM SA.	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
Citalopram Orion 40 mg filmdragerad tablett	FI/H/600/03	20584	ORION CORPORATION	FI
Citalopram Orion 20 mg FILMDRAGERAD TABLETT	FI/H/0600/002	20583	ORION CORPORATION	FI
Citalopram Orion 10 mg filmdragerad tablett	FI/H/600/01	20582	ORION CORPORATION	FI
Citronil 10 mg, tabletki powlekane	FI/H/0600/001	12662	ORION CORPORATION	PL
Citalopram Orion 10 mg potahované tablety	FI/H/0600/001	30/003/07-C	ORION CORPORATION	CZ
Citalopram Orion, filmovertrukne tabletter	FI/H/0600/001	39632	ORION CORPORATION	DK
Citalopram Orion 10 mg tabletti, kalvopäällysteinen	FI/H/600/01	20582	ORION CORPORATION	FI
Citalopram Orion 10 mg filmtabletta	FI/H/0600/001	OGYI-T-20342/01	ORION CORPORATION	HU
Citalopram Orion 10 mg tabletter, filmdrasjerte	FI/H/0600/001	06-4251	ORION CORPORATION	NO
Citalopram Orion 10 mg filmdragerade tabletter	FI/H/0600/001	24108	ORION CORPORATION	SE
Citalopram Orion 10 mg, filmom obalené tablety	FI/H/0600/001	30/0518/06-S	ORION CORPORATION	SK
Citronil 20 mg, tabletki powlekane	FI/H/0600/002	12663	ORION CORPORATION	PL
Citalopram Orion 20 mg potahované tablety	FI/H/0600/002	30/004/07-C	ORION CORPORATION	CZ
Citalopram Orion, filmovertrukne tabletter	FI/H/0600/002	39633	ORION CORPORATION	DK
Citalopram Orion 20 mg tabletti, kalvopäällysteinen	FI/H/0600/002	20583	ORION CORPORATION	FI
Citalopram Orion 20 mg filmtabletta	FI/H/0600/002	OGYI-T-20342/02	ORION CORPORATION	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Orion 20 mg tabletter, filmdrasjerte	FI/H/0600/002	06-4252	ORION CORPORATION	NO
Citalopram Orion 20 mg filmdragerade tabletter	FI/H/0600/002	24109	ORION CORPORATION	SE
Citalopram Orion 20 mg, filmom obalené tablety	FI/H/0600/002	30/0519/06-S	ORION CORPORATION	SK
Citronil 40 mg, tabletki powlekane	FI/H/0600/003	12661	ORION CORPORATION	PL
Citalopram Orion 40 mg potahované tablety	FI/H/0600/003	30/005/07-C	ORION CORPORATION	CZ
Citalopram Orion, filmovertrukne tabletter	FI/H/0600/003	39634	ORION CORPORATION	DK
Citalopram Orion 40 mg tabletti, kalvopäällysteinen	FI/H/0600/003	20584	ORION CORPORATION	FI
Citalopram Orion 40 mg filmtabletta	FI/H/0600/003	OGYI-T-20342/03	ORION CORPORATION	HU
Citalopram Orion 40 mg tabletter, filmdrasjerte	FI/H/0600/003	06-4253	ORION CORPORATION	NO
Citalopram Orion 40 mg filmdragerade tabletter	FI/H/0600/003	24110	ORION CORPORATION	SE
Citalopram Orion 40 mg, filmom obalené tablety	FI/H/0600/003	30/0520/06-S	ORION CORPORATION	SK
Citalopram Generis 10 mg comprimidos revestidos por película	not available	5582143	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 10 mg comprimidos revestidos por película	not available	5582150	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 10 mg comprimidos revestidos por película	not available	5582168	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 20 mg comprimidos revestidos por película	not available	5582176	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
Citalopram Generis 20 mg comprimidos revestidos por película	not available	5582200	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 20 mg comprimidos revestidos por película	not available	5582218	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 40 mg comprimidos revestidos por película	not available	5582473	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 40 mg comprimidos revestidos por película	not available	5582507	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram Generis 40 mg comprimidos revestidos por película	not available	5582515	GENERIS FARMACÊUTICA, S.A.	PT
Citalopram 10 mg Tablets	not available	PL 06464/1971	WAYMADE PLC	UK
Citalopram 20 mg Tablets	not available	PL 06464/1972	WAYMADE PLC	UK
Citalopram 40 mg Tablets	not available	PL 06464/1974	WAYMADE PLC	UK
SEROR 20mg επικαλυμμένα με λεπτό υμένιο δισκία.	not available	20301/17-10-2012	SPECIFAR S.A.	GR
SEROR 40mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	72413/17-10-2012	SPECIFAR S.A.	GR
ZECLICID f.c.tabs 20 mg/tab	not available	43642/12	VERISFIELD (UK) LTD	GR
ZECLICID	not available	43643/12	VERISFIELD (UK) LTD	GR
ZECLICID	not available	43641/12	VERISFIELD (UK) LTD	GR
Citalopram 10mg Film-coated Tablets	not available	PA688/8/1	CHANELLE MEDICAL	IE
Citalopram 20mg Film-coated Tablets	not available	PA688/8/2	CHANELLE MEDICAL	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
Citalopram 40mg Film-coated Tablets	not available	PA688/8/3	CHANELLE MEDICAL	IE
Citalopram Genericon 40 mg Filmtabletten	not available	1-24726	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
SEREGRA 20 mg comprimidos recubiertos con película EFG	not available	69163	ARAFARMA GROUP, S.A	ES
SEREGRA 30 mg comprimidos recubiertos con película EFG	not available	69164	ARAFARMA GROUP, S.A	ES
Citalopram 40mg/ml Oral drops, Solution	not available	PL 00289/1460	TEVA UK LIMITED	UK
Citalopram Hennig® 10 mg Filmtabletten	not available	54076.00.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Citalopram Ranbaxy Italia 40 mg/ml gocce orali, soluzione	not available	036651014	RANBAXY ITALIA S.P.A.	IT
CITALOPRAM EG 20 mg, comprimé pelliculé sécable	not available	NL29726	EG LABO - LABORATOIRES EUROGENERICS	FR
Citalopram TARBIS 20 mg comprimidos recubiertos con película EFG	not available	66.347	TARBIS FARMA, S.L.	ES
Citaxin, 20 mg, tabletki powlekane	not available	16319	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Citalopram "Orifarm", filmovertrukne tabletter	DK/H/1007/001	35254	ORIFARM GENERICS A/S	DK
Citalopram "Orifarm", filmovertrukne tabletter	DK/H/1007/002	35255	ORIFARM GENERICS A/S	DK
Citalopram "Orifarm", filmovertrukne tabletter	DK/H/1007/003	35256	ORIFARM GENERICS A/S	DK
Citalopram "Orifarm", filmovertrukne tabletter	DK/H/1007/004	35257	ORIFARM GENERICS A/S	DK
VAROM ®	not available	41595/12/18-02-2013	LYOFIN LTD, GREECE	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
VAROM ®	not available	41596/12/18-02-2013	LYOFIN LTD, GREECE	GR
CILOPRESS	not available	56706/14-09-2011	SJA PHARM LTD	GR
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	5619960	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	5619978	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	5620000	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	5620018	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL 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
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
CITALOPRAM AUROBINDO ITALIA 40 mg/ml gocce orali, soluzione	not available	036660037	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
CITALOPRAM ALTER 20 mg, comprimé pelliculé sécable	not available	6 284 571 3	LABORATOIRES ALTER	FR
Citalopram Bluefish 10 mg filmomhulde tabletten	NL/H/1601/001	RVG 29755	BLUEFISH PHARMACEUTICALS AB	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
Citalopram Bluefish 20 mg filmomhulde tabletten	NL/H/1601/002	RVG 29756	BLUEFISH PHARMACEUTICALS AB	NL
Citalopram Bluefish 40 mg filmomhulde tabletten	NL/H/1601/003	RVG 29757	BLUEFISH PHARMACEUTICALS AB	NL
Citalopram Bluefish 10 mg Filmtabletten	NL/H/1601/001	1-28965	BLUEFISH PHARMACEUTICALS AB	AT
Citalopram Bluefish 20 mg Filmtabletten	NL/H/1601/002	1-28966	BLUEFISH PHARMACEUTICALS AB	AT
Citalopram Bluefish 40 mg Filmtabletten	NL/H/1601/003	1-28967	BLUEFISH PHARMACEUTICALS AB	AT
CITALOPRAM BLUEFISH 20 mg, comprimé pelliculé sécable	NL/H/1601/002	217 217-1 OU 34009 217 217 1 3	BLUEFISH PHARMACEUTICALS AB	FR
Citalopram Bluefish 10 mg film-coated tablets	NL/H/1601/001	PA 1436/18/1	BLUEFISH PHARMACEUTICALS AB	IE
Citalopram Bluefish 20 mg film-coated tablets	NL/H/1601/002	PA 1436/18/2	BLUEFISH PHARMACEUTICALS AB	IE
Citalopram Bluefish 40 mg film-coated tablets	NL/H/1601/003	PA 1436/18/3	BLUEFISH PHARMACEUTICALS AB	IE
Citalopram Bluefish 10 mg filmdragerade tabletter	NL/H/1601/001	42877	BLUEFISH PHARMACEUTICALS AB	SE
Citalopram Bluefish 20 mg filmdragerade tabletter	NL/H/1601/002	42878	BLUEFISH PHARMACEUTICALS AB	SE
Citalopram Bluefish 40 mg filmdragerade tabletter	NL/H/1601/003	42879	BLUEFISH PHARMACEUTICALS AB	SE
CITALOPRAM BLUEFISH 20 mg, comprimé pelliculé sécable	NL/H/1601/002	34009 217 218 8 1	BLUEFISH PHARMACEUTICALS AB	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
CITALOPRAM BLUEFISH 20 mg, comprimé pelliculé sécable	NL/H/1601/002	34009 217 219 4 2	BLUEFISH PHARMACEUTICALS AB	FR
CITALOPRAM BLUEFISH 20 mg, comprimé pelliculé sécable	NL/H/1601/002	34009 217 220 2 4	BLUEFISH PHARMACEUTICALS AB	FR
CITALOPRAM BLUEFISH 20 mg, comprimé pelliculé sécable	NL/H/1601/002	34009 580 858 6 4	BLUEFISH PHARMACEUTICALS AB	FR
Citalopram Bluefish 10 mg filmuhúðaðar töflur	NL/H/1601/001	IS/1/12/070/01	BLUEFISH PHARMACEUTICALS AB	IS
Citalopram Bluefish 20 mg filmuhúðaðar töflur	NL/H/1601/002	IS/1/12/070/02	BLUEFISH PHARMACEUTICALS AB	IS
Citalopram Bluefish 40 mg filmuhúðaðar töflur	NL/H/1601/003	IS/1/12/070/03	BLUEFISH PHARMACEUTICALS AB	IS
Pram 10 mg apvalkotās tabletes	not available	03-0197	G.L. PHARMA GMBH	LV
Pram 20 mg apvalkotas tabletes	not available	03-0198	G.L. PHARMA GMBH	LV
Pram 40 mg apvalkotas tabletes	not available	03-0199	G.L. PHARMA GMBH	LV
CITALOPRAM DOC Generici 20 mg compresse rivestite con film	not available	036266017	DOC GENERICI S.R.L.	IT
CITALOPRAM DOC Generici 40 mg compresse rivestite con film	not available	036266029	DOC GENERICI S.R.L.	IT
PERCITALE 20 mg compresse rivestite con film	not available	036302026	PIAM FARMACEUTICI S.P.A.	IT
PERCITALE 40 mg compresse rivestite con film	not available	036302038	PIAM FARMACEUTICI S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
PERCITALE 40 mg/ml gocce orali, soluzione	not available	036302014	PIAM FARMACEUTICI S.P.A.	IT
Citalopram Actavis 10 mg filmdragerade tabletter	not available	17718	ACTAVIS NORDIC A/S	SE
Citalopram Actavis 20 mg filmdragerade tabletter	not available	17719	ACTAVIS NORDIC A/S	SE
Citalopram Actavis 40 mg filmdragerade tabletter	not available	17720	ACTAVIS NORDIC A/S	SE
LOPRAXER®, επικαλυμμένα με λεπτό υμένιο δισκία, 20mg/tab	not available	2814/18-07-2013	HELP ABEE	GR
LOPRAXER®, επικαλυμμένα με λεπτό υμένιο δισκία, 30mg/tab	not available	47337/18-07-2013	HELP ABEE	GR
LOPRAXER®, επικαλυμμένα με λεπτό υμένιο δισκία, 40mg/tab	not available	47336/18-07-2013	HELP ABEE	GR
LOPRAXER®, πόσιμες σταγόνες, διάλυμα, 40mg/ml	not available	2812/18-07-2013	HELP ABEE	GR
CITALOPRAM EUROGENERICI 40 mg/ml gocce orali, soluzione	not available	036869016	EG S.P.A.	IT
Zanipram: 40mg/ml πόσιμες σταγόνες, διάλυμα	not available	17537	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
Zanipram: 20mg/Tab επικαλυμμένα με λεπτό υμένιο δισκία	not available	73556/12	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	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
Zanipram: 40mg/Tab επικαλυμμένα με λεπτό υμένιο δισκία	not available	43383	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
Zitolex 10 mg comprimidos revestidos por película	not available	5166202	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 10 mg comprimidos revestidos por película	not available	5166210	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 10 mg comprimidos revestidos por película	not available	5166228	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 10 mg comprimidos revestidos por película	not available	5166236	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 20 mg comprimidos revestidos por película	not available	5166251	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 20 mg comprimidos revestidos por película	not available	5166269	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 20 mg comprimidos revestidos por película	not available	5166277	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 20 mg comprimidos revestidos por película	not available	5166301	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 40 mg comprimidos revestidos por película	not available	5166327	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 40 mg comprimidos revestidos por película	not available	5166335	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
Zitolex 40 mg comprimidos revestidos por película	not available	5166343	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 40 mg comprimidos revestidos por película	not available	5166350	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 10 mg comprimidos revestidos por película	not available	5166178	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 20 mg comprimidos revestidos por película	not available	5166244	DECOMED FARMACÊUTICA, S.A.	PT
Zitolex 40 mg comprimidos revestidos por película	not available	5166319	DECOMED FARMACÊUTICA, S.A.	PT
Citalopram AbZ 10 mg Filmtabletten	DK/H/0752/001	62585.00.00	ABZ-PHARMA GMBH	DE
Loxopram, filmovertrukne tabletter	DK/H/0752/001	34364	RATIOPHARM GMBH	DK
Loxopram, filmovertrukne tabletter	DK/H/0752/002	34365	RATIOPHARM GMBH	DK
Citalopram-ratiopharm 40 mg Filmtabletten	DK/H/0752/003	1-26033	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Loxopram, filmovertrukne tabletter	DK/H/0752/003	34366	RATIOPHARM GMBH	DK
Citalopram-ratiopharm 20 mg filmomhulde tabletten	NL/H/0311/002	BE422021	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 20 mg Filmtabletten	NL/H/0311/002	BE422021	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 20 mg comprimés pelliculés	NL/H/0311/002	BE422021	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 20 mg Filmtabletten	NL/H/0311/002	BE239626	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram-ratiopharm 20 mg comprimés pelliculés	NL/H/0311/002	BE239626	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 20 mg Filmtabletten	NL/H/0311/002	BE239635	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 20 mg comprimés pelliculés	NL/H/0311/002	BE239635	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm 40 mg filmdragerade tabletter	NL/H/0311/003	16863	RATIOPHARM GMBH	FI
Citalopram-ratiopharm 20 mg filmdragerade tabletter	NL/H/0311/002	16862	RATIOPHARM GMBH	FI
Citalopram-ratiopharm 10 mg filmdragerade tabletter	NL/H/0311/001	16861	RATIOPHARM GMBH	FI
Citalopram-ratiopharm® 20 mg Filmtabletten	NL/H/0311/002	53072.01.00	RATIOPHARM GMBH	DE
Citalopram-ratiopharm 20 mg Filmtabletten	NL/H/0311/002	1-24578	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Citalopram-ratiopharm 20 mg tabletti, kalvopäällysteinen	NL/H/0311/002	16862	RATIOPHARM GMBH	FI
Citalopram-ratiopharm 10 mg tabletti, Kalvopäällysteinen	NL/H/0311/001	16861	RATIOPHARM GMBH	FI
Citalopram-ratiopharm 40 mg tabletti, kalvopäällysteinen	NL/H/0311/003	16863	RATIOPHARM GMBH	FI
Citalopram-ratiopharm 20 mg filmomhulde tabletten	NL/H/0311/002	BE239635	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram-ratiopharm® 10 mg Filmtabletten	NL/H/0311/001	53072.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram-ratiopharm® 40 mg Filmtabletten	NL/H/0311/003	53072.02.00	RATIOPHARM GMBH	DE
Citalopram-ratiopharm 10 mg Filmtabletten	NL/H/0311/001	1-24577	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Citalopram-ratiopharm 20 mg filmomhulde tabletten	NL/H/0311/002	BE239626	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram ratiopharm 40 mg, filmomhulde tabletten	NL/H/0311/003	RVG 25152	RATIOPHARM NEDERLAND B.V	NL
Citalopram ratiopharm 20 mg, filmomhulde tabletten	NL/H/0311/002	RVG 25151	RATIOPHARM NEDERLAND B.V	NL
Citalopram ratiopharm 10 mg, filmomhulde tabletten	NL/H/0311/001	RVG 25150	RATIOPHARM NEDERLAND B.V	NL
Citalopram LAREQ 30 mg comprimidos recubiertos con película EFG	not available	66.279	APOTEX ESPAÑA, S.L.	ES
Citalostad 10 mg Filmtabletten	not available	1-25315	STADA ARZNEIMITTEL GMBH	AT
Citalopram Davur 20 mg comprimidos recubiertos con película EFG	not available	66505	LABORATORIOS DAVUR S.L	ES
Citalopram-CT 20 mg Filmtabletten	NL/H/0460/002	58938.01.00	ABZ-PHARMA GMBH	DE
Citalopram 40 ct	NL/H/0460/003	RVG 27620	TEVA PHARMA B.V.	NL
Citalopram-CT 40 mg Filmtabletten	NL/H/0460/003	58938.02.00	ABZ-PHARMA GMBH	DE
Citalopram 20 ct, filmomhulde tabletten	NL/H/0460/002	RVG 27619	TEVA PHARMA B.V.	NL
KAIDOR 40 mg/ml gocce orali, soluzione	not available	036246015	FARTO S.R.L. FARMACO BIOCHIMICO TOSCANO	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
Citalopram Orifarm 10 mg filmdragerade tabletter	not available	19631	ORIFARM GENERICS A/S	SE
Citalopram Orifarm 20 mg filmdragerade tabletter	not available	19632	ORIFARM GENERICS A/S	SE
Citalopram Orifarm 30 mg filmdragerade tabletter	not available	19633	ORIFARM GENERICS A/S	SE
Citalopram Orifarm 40 mg filmdragerade tabletter	not available	19634	ORIFARM GENERICS A/S	SE
LINISAN 10 mg comprimate filmate	not available	7778/2006/01	GEDEON RICHTER ROMÂNIA S.A.	RO
LINISAN 20 mg comprimate filmate	not available	7779/2006/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Pram 10 mg-Filmtabletten	not available	1-24597	G.L. PHARMA GMBH	AT
Pram 20 mg-Filmtabletten	not available	1-24598	G.L. PHARMA GMBH	AT
Pram 40 mg-Filmtabletten	not available	1-24599	G.L. PHARMA GMBH	AT
CITALOPRAM NORMON 20 mg Comprimidos recubiertos con película EFG	not available	66795	LABORATORIOS NORMON, S.A.	ES
CITALOPRAM NORMON 30 mg Comprimidos recubiertos con película EFG	not available	66794	LABORATORIOS NORMON, S.A.	ES
Citalopram Ritisca 10 mg comprimidos revestidos por película	not available	5641840	AUROVITAS UNIPESSOAL, 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
Citalopram Ritisca 10 mg comprimidos revestidos por película	not available	5641857	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 20 mg comprimidos revestidos por película	not available	5641865	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 40 mg comprimidos revestidos por película	not available	5641873	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	5136460	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	5136478	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	5136502	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	5136510	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	5136544	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	5137633	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	5136551	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, 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
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	DE/H/1164/001	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, 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
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	DE/H/1164/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	DE/H/1164/04	AUROVITAS UNIPESSOAL, 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
Citalopram Ritisca 40 mg comprimidos revestidos por película	not available	14/H/0172/003	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 40 mg comprimidos revestidos por película	not available	14/H/0172/003	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 20 mg comprimidos revestidos por película	not available	14/H/0172/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 20 mg comprimidos revestidos por película	not available	14/H/0172/002	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Ritisca 10 mg comprimidos revestidos por película	not available	14/H/0172/001	AUROVITAS UNIPESSOAL, LDA.	PT
CITALOPRAM AUROBINDO 40 mg/ml gocce orali, soluzione	not available	036675039	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
CITALOPRAM AUROBINDO 40 mg/ml gocce orali, soluzione	not available	036675039	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Citalopram Wynn 20 mg comprimidos revestidos por película	not available	5016803	AXONE, LDA.	PT
Citalopram Wynn 40 mg comprimidos revestidos por película	not available	5016811	AXONE, LDA.	PT
Citalopram Wynn 40 mg comprimidos revestidos por película	not available	5053830	AXONE, LDA.	PT
Citalopram Wynn 20 mg comprimidos revestidos por película	not available	5016779	AXONE, 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
CITALOPRAM AUROBINDO ITALIA 20 mg compresse rivestite con film	not available	036660013	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
CITALOPRAM AUROBINDO ITALIA 40 mg compresse rivestite con film	not available	036660025	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
CITALOPRAM RANBAXY 20 mg, comprimé pelliculé sécable	not available	NL 33233	RANBAXY PHARMACIE GENERIQUES	FR
VERUS® Αναβράζον δισκίο 20mg/tab	not available	42754/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
VERUS® Αναβράζον δισκίο 40mg/tab	not available	42745/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
VERUS® Επικαλυμμένα με λεπτό υμένιο δισκία 20mg/tab	not available	42746/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
VERUS® Επικαλυμμένο με λεπτό υμένιο δισκίο 40mg/tab	not available	42748/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
VERUS® Πόσιμες σταγόνες, διάλυμα 40mg/ml	not available	42762/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
VERUS® Πόσιμο διάλυμα σε συσκευασία μιας δόσης 16mg/5ml	not available	42743/10	UNI-PHARMA KLEON TSETIS PHARMACEUTICAL LABORATORIES S.A.	GR
CITALOPRAM EVOLUGEN 20 mg, comprimé pelliculé sécable	not available	34009 491 798 8 6	EVOLUPHARM	FR
CITALOPRAM EVOLUGEN 20 mg, comprimé pelliculé sécable	not available	34009 498 910 8 5	EVOLUPHARM	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
CITALOPRAM EVOLUGEN 20 mg, comprimé pelliculé sécable	not available	34009 577 621 9 3	EVOLUPHARM	FR
CITALOPRAM EVOLUGEN 20 mg, comprimé pelliculé sécable	not available	34009 578 944 6 7	EVOLUPHARM	FR
Citalon® 20 mg Filmtabletten	DE/H/2904/001	55457.00.00	KREWEL MEUSELBACH GMBH	DE
Citalon® 40 mg Filmtabletten	DE/H/2904/002	55457.02.00	KREWEL MEUSELBACH GMBH	DE
Malicon®	not available	34107/12/08-03-2013	NORMA HELLAS S.A.	GR
Malicon®	not available	19169/08-03-2013	NORMA HELLAS S.A.	GR
Citalopram GERMED Pharma 20 mg compresse rivestite con film	not available	036482014	GERMED PHARMA S.R.L.	IT
Citalopram GERMED PHARMA 40 mg compresse rivestite con film	not available	036482026	GERMED PHARMA S.R.L.	IT
Citalopram GERMED PHARMA 40 mg/ml concentrato per soluzione per infusione	not available	036482038	GERMED PHARMA S.R.L.	IT
Citalopram GERMED Pharma 40 mg/ml gocce orali, soluzione	not available	036482040	GERMED PHARMA S.R.L.	IT
Citalopram MABO 10 mg comprimidos recubiertos con película	not available	66.249	MABO-FARMA, S.A.	ES
Citalopram MABO 20 mg comprimidos recubiertos con película EFG	not available	66.251	MABO-FARMA, 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
Citalopram ARISTO 20 mg comprimidos recubiertos con película EFG	not available	66231	ARISTO PHARMA IBERIA, S.L.	ES
Citalopram Aristo 30 mg comprimidos recubiertos con película EFG	not available	66229	ARISTO PHARMA IBERIA, S.L.	ES
Citalopram Farmalider 10 mg comprimidos recubiertos con película	not available	66.346	FARMALIDER, S.A.	ES
PRALOTAM	not available	106197/14/06-04-2015	VENIFAR LTD	GR
PRALOTAM	not available	21927/06-04-2015	VENIFAR LTD	GR
Citalopram Kern Pharma 20 mg comprimidos recubiertos con película EFG	not available	65.977	KERN PHARMA, S.L.	ES
Citalopram Kern Pharma 30 mg comprimidos recubiertos con película EFG	not available	65.993	KERN PHARMA, S.L.	ES
Cital, 20 mg, tabletki powlekane	not available	8955	GLENMARK PHARMACEUTICALS S.R.O.	PL
Citalopram Ranbaxygen 20 mg comprimidos recubiertos con película	UK/H/0672/002	67107	LABORATORIOS RANBAXY S.L.	ES
CITALOPRAM BASICS 10 mg Filmtabletten	UK/H/0672/001	60782.00.00	BASICS GMBH	DE
CITALOPRAM BASICS 20 mg Filmtabletten	UK/H/0672/002	60782.01.00	BASICS GMBH	DE
CITALOPRAM BASICS 40 mg Filmtabletten	UK/H/0672/003	60782.02.00	BASICS GMBH	DE
Citalopram EG 20 mg Filmtabletten	NL/H/0310/002	BE239513	EUROGENERICS N.V./S.A.	BE
Citalopram EG 20 mg Filmtabletten	NL/H/0310/002	BE239522	EUROGENERICS N.V./S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram EG 20 mg filmomhulde tabletten	NL/H/0310/002	BE239513	EUROGENERICS N.V./S.A.	BE
Citalopram EG 20 mg filmomhulde tabletten	NL/H/0310/002	BE239522	EUROGENERICS N.V./S.A.	BE
Citalopram EG 20 mg comprimés pelliculés	NL/H/0310/002	BE239522	EUROGENERICS N.V./S.A.	BE
Citalopram EG 20 mg comprimés pelliculés	NL/H/0310/002	BE239513	EUROGENERICS N.V./S.A.	BE
Citalopram CF 40 mg, filmomhulde tabletten	NL/H/0310/003	RVG 25149	CENTRAFARM B.V.	NL
Citalopram CF 20 mg, filmomhulde tabletten	NL/H/0310/002	RVG 25148	CENTRAFARM B.V.	NL
Citalopram CF 10 mg, filmomhulde tabletten	NL/H/0310/001	RVG 25147	CENTRAFARM B.V.	NL
Citalopram CF 30 mg, filmomhulde tabletten	NL/H/0310/004	RVG 31114	CENTRAFARM B.V.	NL
APO-CITAL 20 mg potahované tablety	CZ/H/0182/001	30/210/07-C	APOTEX EUROPE BV	CZ
Citalopram Apotex 20 mg, filmomhulde tabletten	CZ/H/0182/001	RVG 34746	APOTEX EUROPE B.V.	NL
Citalopram Apotex 40 mg, filmomhulde tabletten	CZ/H/0182/002	RVG 34747	APOTEX EUROPE B.V.	NL
Citalopram Apotex 20 mg comprimidos recubiertos con película EFG	CZ/H/0182/001	70250	APOTEX EUROPE BV	ES
Citalopram Apotex 40 mg comprimidos recubiertos con película EFG	CZ/H/0182/002	70249	APOTEX EUROPE BV	ES
SEROR 20mg επικαλυμμένα με λεπτό υμένιο δισκία.	not available	20301/17-10-2012	SPECIFAR S.A.	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
SEROR 40mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	72413/17-10-2012	SPECIFAR S.A.	GR
CITALOPRAM ACTAVIS 20 mg, comprimé pelliculé sécable	not available	NL 38302	ACTAVIS GROUP PTC EHF.	FR
Citalopram PENSA 20 mg compresse rivestite con film	not available	036392013	PENSA PHARMA S.P.A.	IT
Citalopram Pensa 20 mg compresse rivestite con film	not available	036392025	PENSA PHARMA S.P.A.	IT
Citalopram Pensa Pharma 40 mg/ml gocce orali, soluzione	not available	038199016	PENSA PHARMA S.P.A.	IT
Citalopram PENSA 40 mg compresse rivestite con film	not available	036392037	PENSA PHARMA S.P.A.	IT
CITALOPRAM PENSA 10 mg comprimidos recubiertos con película	not available	68.394	PENSA PHARMA, S.A	ES
CITALOPRAM PENSA 20 mg Comprimidos recubiertos con película EFG	not available	68.395	PENSA PHARMA, S.A	ES
CITALOPRAM PENSA 30 mg Comprimidos recubiertos con película EFG	not available	68.396	PENSA PHARMA, S.A	ES
Citalopram Hennig® 20 mg Filmtabletten	not available	54076.01.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Citacip, filmovertrukne tabletter	DK/H/0750/003	34363	STADA ARZNEIMITTEL AG	DK
Citacip, filmovertrukne tabletter	DK/H/0750/002	34362	STADA ARZNEIMITTEL AG	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
Citacip, filmovertrukne tabletter	DK/H/0750/001	34361	STADA ARZNEIMITTEL AG	DK
Citalopram STADA 10 mg filmtabletta	DK/H/0750/001	OGYI-T-10391/01	STADA ARZNEIMITTEL AG	HU
Citalopram STADA® 20 mg Filmtabletten	DK/H/0750/002	62579.01.00	STADAPHARM GMBH	DE
Citalopram STADA® 40 mg Filmtabletten	DK/H/0750/003	62579.02.00	STADAPHARM GMBH	DE
Citalopram STADA® 10 mg Filmtabletten	DK/H/0750/001	62579.00.00	STADAPHARM GMBH	DE
Citalopram STADA 20 mg filmtabletta	DK/H/0750/002	OGYI-T-10391/02	STADA ARZNEIMITTEL AG	HU
Citalopram STADA 40 mg filmtabletta	DK/H/0750/003	OGYI-T-10391/03	STADA ARZNEIMITTEL AG	HU
CITALOPRAM MOLTENI 40 mg compresse rivestite con film	not available	035976036	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
CITALOPRAM MOLTENI 20 mg compresse rivestite con film	not available	035976012	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
CITALOPRAM MOLTENI 20 mg compresse rivestite con film	not available	035976024	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Citalopram Alternova, filmovertrukne tabletter	not available	36152	ALTERNOVA A/S	DK
ОРОПРАМ 20 MG ФИЛМИРАНИ ТАБЛЕТКИ	not available	20030535	ACTAVIS EAD	BG
Citalopram 40 mg film-coated tablets	not available	PL 20532/0043	AUROBINDO PHARMA LIMITED	UK
Citalopram 10 mg film-coated tablets	not available	PL 20532/0041	AUROBINDO PHARMA LIMITED	UK
Citalopram 20 mg film-coated tablets	not available	PL 20532/0042	AUROBINDO PHARMA 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
CITALOPRAM TEVA SANTE 20 mg, comprimé pelliculé sécable	not available	FR NL27598	TEVA SANTÉ	DE
Citalopram beta 20 mg Filmtabletten	not available	66055.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Citalopram beta 30 mg Filmtabletten	not available	70754.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Citalopram beta 40 mg Filmtabletten	not available	66056.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Citalopram Aristo® 10 mg Filmtabletten	not available	66057.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Citalopram Aristo® 20 mg Filmtabletten	not available	66058.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Citalopram Aristo® 30 mg Filmtabletten	not available	70629.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Citalopram Aristo® 40 mg Filmtabletten	not available	66059.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Acelopam Πόσιμες σταγόνες, διάλυμα 40mg/ml	not available	53774	PHERAKON PC	GR
Citalopram ratiopharm 30 mg comprimidos recubiertos con película EFG	not available	66.267	RATIOPHARM ESPANA SA	ES
Citalopram ratiopharm 10 mg comprimidos recubiertos con película	not available	66264	RATIOPHARM ESPANA SA	ES
Unstress, Oral solution 40mg / ml	not available	41683/12	HEREMCO	GR
Zyloram tablety 20 mg filmom obalené tablety	not available	30/0122/04-S	RANBAXY (UK) LIMITED	SK
Citalopram STADA® 30 mg Filmtabletten	not available	70627.00.00	STADAPHARM GMBH	DE
Citalopram 40mg Tablets	not available	PL 44041/0010	NOUMED LIFE SCIENCES	UK
Citalopram 20mg Tablets	not available	PL 44041/0009	NOUMED LIFE SCIENCES	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
Citalopram 10mg Tablets	not available	PL 44041/0008	NOUMED LIFE SCIENCES	UK
CIPRANED Επικαλυμμένο με λεπτό υμένιο δισκίο, 20 mg/tab	not available	74530/14/06-05-2015	MEDICAL PHARMAQUALITY PHARMACEUTICALS S.A.	GR
CIPRANED Επικαλυμμένο με λεπτό υμένιο δισκίο, 40 mg/tab	not available	30791/06-05-2015	MEDICAL PHARMAQUALITY PHARMACEUTICALS S.A.	GR
VESEMA®	not available	47382/11	PHARMEX S.A.	GR
VESEMA®	not available	47383/11	PHARMEX S.A.	GR
RENEVIL	not available	26687/12	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
RENEVIL	not available	43377/04-06-2013	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
RENEVIL	not available	43377/04-06-2013	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
RETURN 20 mg compresse rivestite con film	not available	036792012	EURO-PHARMA S.R.L.	IT
RETURN 40 mg/ml gocce orali, soluzione	not available	036792036	EURO-PHARMA S.R.L.	IT
SOTOVON 40mg/ML	not available	37619/10	BROS LTD	GR
CITALOPRAM ZYDUS 20 mg, comprimé pelliculé sécable	not available	34009 388 171 6 4	ZYDUS FRANCE	FR
PRAM 20 mg Potahované tablety	not available	30/109/04-C	G.L. PHARMA GMBH	CZ
Oropram 10mg filmuhúðaðar töflur	not available	990322	ACTAVIS HF.	IS
Oropram 20 mg filmuhúðaðar töflur	not available	990323	ACTAVIS HF.	IS
Oropram 40mg filmuhúðaðar töflur	not available	990324	ACTAVIS HF.	IS
Citalonte, filmovertrukne tabletter	DK/H/0751/001	34358	STADA ARZNEIMITTEL AG	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
Citalonte, filmovertrukne tabletter	DK/H/0751/002	34359	STADA ARZNEIMITTEL AG	DK
Citalonte, filmovertrukne tabletter	DK/H/0751/003	34360	STADA ARZNEIMITTEL AG	DK
Citalopram AL 40 mg Filmtabletten	DK/H/0751/003	62582.02.00	ALIUD PHARMA GMBH	DE
Citalopram AL 20 mg Filmtabletten	DK/H/0751/002	62582.01.00	ALIUD PHARMA GMBH	DE
Citalopram AL 10 mg Filmtabletten	DK/H/0751/001	62582.00.00	ALIUD PHARMA GMBH	DE
Citalopram Davur 30 mg comprimidos recubiertos con película EFG	not available	67581	LABORATORIOS DAVUR S.L	ES
Citalopram EG 40 mg comprimés pelliculés	NL/H/0465/003	0019/06010040	EUROGENERICS SA	LU
Citalopram EG 10 mg comprimés pelliculés	NL/H/0465/001	0019/06010038	EUROGENERICS SA	LU
Citalopram EG 20 mg comprimés pelliculés	NL/H/0465/002	0019/07069352	EUROGENERICS SA	LU
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503112	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503175	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503201	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503098	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503213	EG S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503011	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503249	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503163	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503050	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503136	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503187	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503047	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503062	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503124	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503148	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503074	EG S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503086	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503100	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503237	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503035	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503151	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503225	EG S.P.A.	IT
CITALOPRAM EG 40 mg compresse rivestite con film	NL/H/0465/003	036503199	EG S.P.A.	IT
CITALOPRAM EG 20 mg compresse rivestite con film	NL/H/0465/002	036503023	EG S.P.A.	IT
Citalopram "PCD", filmovertrukne tabletter	NL/H/0465/001	35967	PHARMACODANE APS	DK
Citalopram "PCD", filmovertrukne tabletter	NL/H/0465/002	35968	PHARMACODANE APS	DK
CITALOPRAM STADA 20 mg, filmomhulde tabletten	NL/H/0465/002	RVG 27634	STADA ARZNEIMITTEL AG	NL
Citalopram "PCD", filmovertrukne tabletter	NL/H/0465/003	35969	PHARMACODANE APS	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram STADA 40 mg, filmomhulde tabletten	NL/H/0465/003	RVG 27635	STADA ARZNEIMITTEL AG	NL
Ciprotan 10 mg film-coated tablets	NL/H/0465/001	PA0126/131/001	CLONMEL HEALTHCARE LTD.	IE
CITALOPRAM STADA 10 mg, filmomhulde tabletten	NL/H/0465/001	RVG 27633	STADA ARZNEIMITTEL AG	NL
Citalopram STADA 30 mg, filmomhulde tabletten	NL/H/0465/004	RVG 31380	STADA ARZNEIMITTEL AG	NL
Ciprotan 20 mg film-coated tablets	NL/H/0465/002	PA0126/131/002	CLONMEL HEALTHCARE LTD.	IE
Citalopram +pharma 20 mg Filmtabletten	AT/H/0436/002	1-26447	+PHARMA ARZNEIMITTEL GMBH	AT
Citalopram +pharma 40 mg Filmtabletten	AT/H/0436/003	1-26448	+PHARMA ARZNEIMITTEL GMBH	AT
Citalopram Actavis 10 mg Film-coated Tablets	not available	PA 1380/27/1	ACTAVIS GROUP PTC EHF.	IE
Citalopram Actavis 20 mg Film-coated Tablets	not available	PA 1380/27/2	ACTAVIS GROUP PTC EHF.	IE
Citalopram 10mg Tablets	not available	PL 04077/0214	M&A PHARMACHEM LTD	UK
CITALOPRAM 20MG TABLETS	not available	PL 04077/0215	M&A PHARMACHEM LTD	UK
Citalopram 40mg Tablets	not available	PL 04077/0216	M&A PHARMACHEM LTD	UK
Citalopram-ratiopharm® 30 mg Filmtabletten	DE/H/2421/001	64641.00.00	RATIOPHARM GMBH	DE
SINTOPRAM 40 mg/ml gocce orali, soluzione	not available	036327017	BIOMEDICA FOSCAMA GROUP S.P.A.	IT
citalopram cinfa 20 mg comprimidos recubiertos con película EFG	not available	67.804	LABORATORIOS CINFA, S.A.	ES
GALOPRAN	not available	24190/13	GALENUS PHARMACEUTICALS SA	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
GALOPRAN	not available	24191/13	GALENUS PHARMACEUTICALS SA	GR
PRAMITAL	not available	19412	ANFARM HELLAS SA	GR
PRAMITAL	not available	59181/12	ANFARM HELLAS SA	GR
Citalopram-neuraxpharm 10 mg Filmtabletten	DE/H/0776/001	53078.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Citalopram-neuraxpharm 20 mg Filmtabletten	DE/H/0776/002	53078.01.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Citalopram-neuraxpharm 40 mg Filmtabletten	DE/H/0776/003	53078.02.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
PRICITAL	not available	40060/11-9-2008	LAVIPHARM HELLAS AE	GR
CITALOPRAM ALMUS 20 mg, comprimé pelliculé sécable	not available	3400936477131	BIOGARAN	FR
CITALOPRAM ABC 20 mg compresse rivestite con film	not available	036043026	ABC FARMACEUTICI S.P.A.	IT
CITALOPRAM ABC 40 mg compresse rivestite con film	not available	036043038	ABC FARMACEUTICI S.P.A.	IT
CITALOPRAM ABC 40 mg/ml gocce orali, soluzione	not available	036043014	ABC FARMACEUTICI S.P.A.	IT
Citalopram 10mg Tablets	not available	PL 21880/0060	MEDREICH PLC	UK
Citalopram 20mg Tablets	not available	PL 21880/0061	MEDREICH PLC	UK
Citalopram 40mg Tablets	not available	PL 21880/0062	MEDREICH PLC	UK
Citalopram Sandoz 10 mg tabletter, filmdrasjerte	DK/H/0575/001	04-2675	SANDOZ A/S	NO
Citalopram Sandoz 40 mg tabletter, filmdrasjerte	DK/H/0575/004	04-2678	SANDOZ A/S	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
Citalopram Sandoz 20 mg tabletter, filmdrasjerte	DK/H/0575/002	04-2676	SANDOZ A/S	NO
Citalon 20 mg filmom obložene tablete		HR-H-322905718	SANDOZ D.O.O.	HR
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/013	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/014	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/015	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/016	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/017	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/018	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/019	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/020	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/024	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/023	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/021	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/025	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/026	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/027	SANDOZ GMBH	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
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/028	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/029	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/030	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/031	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/032	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/033	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/034	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/035	SANDOZ GMBH	SI
Citalon 40 mg filmsko obložene tablete	NL/H/0308/003	H/04/00394/036	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/001	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/002	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/003	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/004	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/005	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/006	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/007	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/008	SANDOZ GMBH	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
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/009	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/010	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/011	SANDOZ GMBH	SI
Citalon 10 mg filmsko obložene tablete	NL/H/0308/001	H/04/00394/012	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/022	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/013	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/014	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/015	SANDOZ GMBH	SI
Citalon 20 mg filmsko obložene tablete	NL/H/0308/002	H/04/00394/016	SANDOZ GMBH	SI
Citalon 10 mg	NL/H/0308/001	30/018/05-C	SANDOZ GMBH	CZ
Citalon 20 mg	NL/H/0308/002	30/019/05-C	SANDOZ GMBH	CZ
CITALOPRAM HEXAL AG	036662	036662017	HEXAL AG	IT
CITALOPRAM Sandoz GmbH 40 mg/ml gocce orali, soluzione	not available	036039016	SANDOZ GMBH	IT
Citalopram 10 mg Tablets	NL/H/0308/001	PL 04416/0419	SANDOZ LTD	UK
Citalopram 20 mg Tablets	NL/H/0308/002	PL 04416/0420	SANDOZ LTD	UK
Citalopram 40 mg Tablets	NL/H/0308/003	PL 04416/0421	SANDOZ LTD	UK
Citalopram Sandoz 20 mg Filmtabletten	DK/H/0576/002	59370.00.00	HEXAL AG	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
Citalopram - 1 A Pharma® 20 mg Filmtabletten	DK/H/0577/002	59374.01.00	1 A PHARMA GMBH	DE
AUREX 40 40 MG, TABLETKI POWLEKANE	10535	10535	SANDOZ GMBH	PL
Citalopram HEXAL® 10 mg Filmtabletten	DK/H/0575/001	59364.00.00	HEXAL AG	DE
Citalopram HEXAL® 30 mg Filmtabletten	DK/H/0575/003	59364.02.00	HEXAL AG	DE
Citalopram HEXAL 30 mg Filmtabletten	DK/H/0575/003	0715/06030002	HEXAL AG	LU
Citalopram HEXAL 40 mg Filmtabletten	DK/H/0575/004	59364.03.00	HEXAL AG	LU
Citalopram Sandoz 30 mg Filmtabletten	DK/H/0576/003	59370.01.00	HEXAL AG	DE
Citalopram Sandoz 10 mg Filmtabletten	DK/H/0576/001	59369.00.00	HEXAL AG	DE
Citalopram - 1 A Pharma® 30 mg Filmtabletten	DK/H/0577/003	59374.02.00	1 A PHARMA GMBH	DE
Citalopram - 1 A Pharma® 40 mg Filmtabletten	DK/H/0577/004	59374.03.00	1 A PHARMA GMBH	DE
AUREX 20 20 mg, tabletki powlekane	not available	10534	SANDOZ GMBH	PL
Citalopram - 1 A Pharma® 10 mg Filmtabletten	DK/H/0577/001	59374.00.00	1 A PHARMA GMBH	DE
Citalopram Sandoz 20 mg - Filmtabletten	NL/H/0308/002	1-24563	SANDOZ GMBH	AT
Citalopram Sandoz 10 mg - Filmtabletten	NL/H/0308/001	1-24561	SANDOZ GMBH	AT
Citalopram 1A Pharma 20 mg - Filmtabletten	945.527	1-24735	1A PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	not available	363 943-5	SANDOZ	FR
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	not available	363 944-1	SANDOZ	FR
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	NL 27809	363 945-8	SANDOZ	FR
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	not available	363 946-4	SANDOZ	FR
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	not available	565 175-9	SANDOZ	FR
CITALOPRAM SANDOZ 20 mg, comprimé pelliculé sécable	not available	565 176-5	SANDOZ	FR
Citapram 20 mg filmtabletta	NOT APPLICABLE	OGYI-T-8976/01	SANDOZ HUNGÁRIA KFT	HU
Citapram 30 mg filmtabletta	NOT APPLICABLE	OGYI-T-8976/02	SANDOZ HUNGÁRIA KFT	HU
Citapram 40 mg filmtabletta	NOT APPLICABLE	OGYI-T-8976/03	SANDOZ HUNGÁRIA KFT	HU
Citalopram Hexal 30 mg - Filmtabletten	NL/H/0366/002	1-24900	HEXAL PHARMA GMBH	AT
Citalopram Hexal 40 mg - Filmtabletten	NL/H/0366/003	1-24901	HEXAL PHARMA GMBH	AT
Citalopram Hexal 20 mg - Filmtabletten	NL/H/0366/001	1-24899	HEXAL PHARMA GMBH	AT
CITALON 20 mg	NL/H/0308/002	30/0254/04-S	SANDOZ GMBH	SK
Ciral 20 mg	NOT APPLICABLE	417503	SANDOZ PHARMACEUTICALS D.D.	EE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM SANDOZ 20, filmomhulde tabletten 20 mg	NL/H/0308/002	RVG 25142	SANDOZ B.V.	NL
CITALOPRAM SANDOZ 10, filmomhulde tabletten 10 mg	NL/H/0308/001	RVG 25141	SANDOZ B.V.	NL
CITALOPRAM SANDOZ 40, filmomhulde tabletten 40 mg	NL/H/0308/003	RVG 25143	SANDOZ B.V.	NL
Citalopram Sandoz 20 mg, filmomhulde tabletten	NL/H/0366/001	RVG 26734	SANDOZ B.V.	NL
Citalopram Sandoz 30 mg, filmomhulde tabletten	NL/H/0366/002	RVG 26735	SANDOZ B.V.	NL
Citalopram Sandoz 40 mg, filmomhulde tabletten	NL/H/0366/003	RVG 26736	SANDOZ B.V.	NL
Citalopram ”Sandoz”, filmovertrukne tabletter	DK/H/0576/003	34596	SANDOZ A/S	DK
Citalopram ”Sandoz”, filmovertrukne tabletter	DK/H/0576/002	34595	SANDOZ A/S	DK
Citalopram "1A Farma"	DK/H/0577/001	34425	1A FARMA A/S	DK
Citalopram ”Sandoz”, filmovertrukne tabletter	DK/H/0576/004	34597	SANDOZ A/S	DK
Citalopram ”Sandoz”, filmovertrukne tabletter	DK/H/0576/001	34424	SANDOZ A/S	DK
Citalopram "1A Farma", filmovertrukne tabletter	DK/H/0577/003	32868	1A FARMA A/S	DK
Citalopram "1A Farma", filmovertrukne tabletter	DK/H/0577/004	32869	1A FARMA A/S	DK
Citalopram "1A Farma", filmovertrukne tabletter	DK/H/0577/002	32867	1A FARMA 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
Citalopram Sandoz 10 mg filmdragerade tabletter	DK/H/0575/001	21044	SANDOZ A/S	SE
Citalopram Sandoz 20 mg filmdragerade tabletter	DK/H/0575/002	21045	SANDOZ A/S	SE
Citalopram Sandoz 30 mg filmdragerade tabletter	DK/H/0575/003	21046	SANDOZ A/S	SE
Citalopram Sandoz 40 mg filmdragerade tabletter	DK/H/0575/004	21047	SANDOZ A/S	SE
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238198	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238200	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238236	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238251	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238275	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238299	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238301	SANDOZ B.V.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367340/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367353/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367365/M	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367377/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367415/M	SANDOZ S.P.A.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238212	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238174	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238186	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238224	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238287	SANDOZ B.V.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367403/M	SANDOZ S.P.A.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238263	SANDOZ B.V.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367389/M	SANDOZ S.P.A.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/004	038238248	SANDOZ B.V.	IT
CITALOPRAM HEXAL 40 mg, compresse rivestite con film	NL/H/0366/003	036367391/M	SANDOZ S.P.A.	IT
Citalopram Sandoz 20 mg, filmomhulde tabletten	NL/H/0366/001	BE 255936	SANDOZ N.V.	BE
Citalopram Sandoz 20 mg, filmomhulde tabletten	NL/H/0366/001	BE 363054	SANDOZ N.V.	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
Citalopram Sandoz 40 mg, filmomhulde tabletten	NL/H/0366/003	BE363063	SANDOZ N.V.	BE
Citalopram Sandoz 30 mg, filmomhulde tabletten	NL/H/0366/002	BE255911	SANDOZ N.V.	BE
Citalopram Sandoz 40 mg, filmomhulde tabletten	NL/H/0366/003	BE255945	SANDOZ N.V.	BE
Citalopram HEXAL® 20 mg Filmtabletten	DK/H/0575/002	59364.01.00	HEXAL AG	DE
Citalopram HEXAL® 40 mg Filmtabletten	DK/H/0575/004	59364.03.00	HEXAL AG	DE
Citalopram Sandoz 40 mg Filmtabletten	DK/H/0576/004	59370.02.00	HEXAL AG	DE
Ciral 20 mg plėvele dengtos tabletės	not available	LT/1/03/2246/002	SANDOZ PHARMACEUTICALS D.D.	LT
Ciral 20 mg plėvele dengtos tabletės	not available	LT/1/03/2246/001	SANDOZ PHARMACEUTICALS D.D.	LT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238022	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238010	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238059	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238034	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238061	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238109	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238046	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238073	SANDOZ B.V.	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
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238085	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238097	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238123	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238111	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238150	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238135	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238147	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238313	SANDOZ B.V.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367124/M	SANDOZ S.P.A.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238162	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238325	SANDOZ B.V.	IT
CITALOPRAM SANDOZ BV	SE/H/1641/002	038238337	SANDOZ B.V.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367148/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367136/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367163/M	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367175/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367151/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367187/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367225/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367201/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367199/M	SANDOZ S.P.A.	IT
CITALOPRAM HEXAL 20 mg, compresse rivestite con film	NL/H/0366/001	036367213/M	SANDOZ S.P.A.	IT
Citalopram HEXAL 20 mg Filmtabletten	DK/H/0575/002	0715/06030001	HEXAL AG	LU
Citalopram Sandoz 40 mg - Filmtabletten	NL/H/0308/003	1-25730	SANDOZ GMBH	AT
Ciral 40 mg plėvele dengtos tabletės	not available	LT/1/03/2246/003	SANDOZ PHARMACEUTICALS D.D.	LT
Citalopram 20mg Tablets	NL/H/0366/001	PL 04416/0914	SANDOZ LTD	UK
Citalopram 10mg Tablets	DK/H/0575/001	PL 04416/0913	SANDOZ LTD	UK
Citalopram 40mg Tablets	NL/H/0366/003	PL 04416/0978	SANDOZ LTD	UK
CITALOPRAM BEXAL 20 mg comprimidos recubiertos con película EFG	NOT APPLICABLE	66.483	BEXAL FARMACÉUTICA 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
CITALOPRAM BEXAL 30 mg comprimidos recubiertos con película EFG	NOT APPLICABLE	65.994	BEXAL FARMACÉUTICA S.A.	ES
Citalopram Sandoz 30 mg comprimidos recubiertos con película EFG	not available	66.119	SANDOZ FARMACÉUTICA, S.A.	ES
Citalopram Sandoz 20 mg comprimidos recubiertos con película EFG	NOT APPLICABLE	66.120	SANDOZ FARMACÉUTICA, S.A.	ES
CITROL 10mg Film-Coated Tablets	DK/H/0575/001	PA 711/64/1	ROWEX LTD	IE
CITROL 20mg Film-Coated Tablets	DK/H/0575/002	PA 711/64/2	ROWEX LTD	IE
CITROL 30mg Film-Coated Tablets	DK/H/0575/003	PA 711/64/3	ROWEX LTD	IE
APO-CITAL 20 mg potahované tablety	CZ/H/0182/001	30/210/07-C	APOTEX EUROPE BV	CZ
Citalopram Apotex 20 mg, filmomhulde tabletten	CZ/H/0182/001	RVG 34746	APOTEX EUROPE B.V.	NL
Citalopram Apotex 40 mg, filmomhulde tabletten	CZ/H/0182/002	RVG 34747	APOTEX EUROPE B.V.	NL
Citalopram Apotex 20 mg comprimidos recubiertos con película EFG	CZ/H/0182/001	70250	APOTEX EUROPE BV	ES
Citalopram Apotex 40 mg comprimidos recubiertos con película EFG	CZ/H/0182/002	70249	APOTEX EUROPE BV	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
OROPRAM 20MG FILM-COATED TABLETS.	not available	MA155/01401	ACTAVIS HF.	MT
Citalopram Heumann 10 mg Filmtabletten	-	59743.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Citalopram Heumann 20 mg Filmtabletten	-	78443.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Citalopram Heumann 30 mg Filmtabletten	-	70625.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Citalopram Heumann 40 mg Filmtabletten	-	78444.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Citalopram 20 mg Holsten	not available	59744.00.00	HOLSTEN PHARMA GMBH	DE
FELIPRAM 20 mg compresse rivestite con film	not available	036125019	S.F. GROUP SRL	IT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/034	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/031	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/032	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/033	TEVA PHARMA B.V.	LT
Citalopram Teva 20 mg filmomhulde tabletten	NL/H/0692/002	BE292311	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 40 mg filmomhulde tabletten	NL/H/0692/003	BE292327	TEVA PHARMA BELGIUM N.V./S.A	BE
CITALOPRAM TEVA 10 mg FILMTABLETTEN	NL/H/0692/001	BE292302	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 40 mg Filmtabletten	NL/H/0692/003	BE292327	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 10 mg filmomhulde tabletten	NL/H/0692/001	BE292302	TEVA PHARMA BELGIUM N.V./S.A	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM TEVA 20 mg FILMTABLETTEN	NL/H/0692/002	BE292311	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram - Teva 40 mg	NL/H/0692/003	30/0181/06-S	TEVA PHARMACEUTICALS CR, S.R.O.	SK
Citalopram Teva 40 mg filmdrasjerte tabletter	NL/H/0692/003	05-3732	TEVA SWEDEN AB	NO
Citalopram Teva 10 mg filmdrasjerte tabletter	NL/H/0692/001	05-3730	TEVA SWEDEN AB	NO
Citalopram Teva 20 mg filmdrasjerte tabletter	NL/H/0692/002	05-3731	TEVA SWEDEN AB	NO
Citalopram "Teva", filmovertrukne tabletter	NL/H/0692/002	38639	TEVA DENMARK A/S	DK
Citalopram-Teva 40 mg potahované tablet	NL/H/0692/003	30/123/06-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Citalopram "Teva", filmovertrukne tabletter	NL/H/0692/003	38640	TEVA DENMARK A/S	DK
Citalopram Teva 20 mg comprimés pelliculés	NL/H/0692/002	BE292311	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 40 mg comprimés pelliculés	NL/H/0692/003	BE292327	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 40mg Film-coated Tablets	NL/H/0692/003	PA 749/19/3	TEVA PHARMA B.V.	IE
Citalopram 20mg film-coated tablets	NL/H/0692/002	PL 00289/0928	TEVA UK LIMITED	UK
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/026	TEVA PHARMA B.V.	LT
Citalopram 40mg film-coated tablets	NL/H/0692/003	PL 00289/0929	TEVA UK LIMITED	UK
Citalopram-Teva 40 mg, õhukese polümeerikattega tabletid	NL/H/0692/003	512906	TEVA PHARMA B.V.	EE
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/021	TEVA PHARMA B.V.	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
Citalopram Teva 10 mg comprimés pelliculés	NL/H/0692/001	BE292302	TEVA PHARMA BELGIUM N.V./S.A	BE
Citalopram Teva 40 mg filmdragerade tabletter	NL/H/0692/003	23002	TEVA SWEDEN AB	SE
Citalopram - Teva 10 mg	NL/H/0692/001	30/0179/06-S	TEVA PHARMACEUTICALS CR, S.R.O.	SK
Citalopram Teva 20 mg filmdragerade tabletter	NL/H/0692/002	23001	TEVA SWEDEN AB	SE
Citalopram - Teva 20 mg	NL/H/0692/002	30/0180/06-S	TEVA PHARMACEUTICALS CR, S.R.O.	SK
Citalopram-Teva 10 mg potahované tablety	NL/H/0692/001	30/121/06-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Citalopram-Teva 10 mg apvalkotās tabletes	NL/H/0692/001	06-0095	TEVA PHARMA B.V.	LV
Citalopram Teva 20mg Film-coated Tablets	NL/H/0692/002	PA 749/19/2	TEVA PHARMA B.V.	IE
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/015	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg potahované tablety	NL/H/0692/002	30/122/06-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/019	TEVA PHARMA B.V.	LT
Citalopram 10mg film-coated tablets	NL/H/0692/001	PL 00289/0927	TEVA UK LIMITED	UK
Citalopram 40 PCH, filmomhulde tabletten 40 mg	NL/H/0692/003	RVG 31127	PHARMACHEMIE B.V	NL
Citalopram 20 PCH, filmomhulde tabletten 20 mg	NL/H/0692/002	RVG 31126	PHARMACHEMIE B.V	NL
Citalopram-Teva 10 mg, õhukese polümeerikattega tabletid	NL/H/0692/001	513006	TEVA PHARMA B.V.	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram-Teva 20 mg apvalkotās tabletes	NL/H/0692/002	06-0096	TEVA PHARMA B.V.	LV
Citalopram Teva 10 mg filmdragerade tabletter	NL/H/0692/001	23000	TEVA SWEDEN AB	SE
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/023	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/022	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/030	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/025	TEVA PHARMA B.V.	LT
Citalopram Teva 10mg Film-coated Tablets	NL/H/0692/001	PA 749/19/1	TEVA PHARMA B.V.	IE
Citalopram-Teva 20 mg, õhukese polümeerikattega tabletid	NL/H/0692/002	512806	TEVA PHARMA B.V.	EE
Citalopram 10 PCH, filmomhulde tabletten 10 mg	NL/H/0692/001	RVG 31125	PHARMACHEMIE B.V	NL
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/029	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/018	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/020	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/003	LT/1/06/0576/024	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/012	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/017	TEVA B.V	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/013	TEVA PHARMA B.V.	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
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/011	TEVA PHARMA B.V.	LT
Citalopram-Teva 40 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/027	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/014	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/010	TEVA PHARMA B.V.	LT
Citalopram-Teva 20 mg plėvele dengtos tabletės	NL/H/0692/002	LT/1/06/0576/016	TEVA PHARMA B.V.	LT
PRAM 20 mg Potahované tablety	not available	30/109/04-C	G.L. PHARMA GMBH	CZ
PRAM 20 mg Potahované tablety	not available	30/109/04-C	G.L. PHARMA GMBH	CZ
RICAP 40 mg/ml gocce orali, soluzione	not available	036056012	I.B.N. SAVIO S.R.L.	IT
RICAP 20 mg compresse rivestite con film	not available	036056024	I.B.N. SAVIO S.R.L.	IT
RICAP 40 mg compresse rivestite con film	not available	036056036	I.B.N. SAVIO S.R.L.	IT
Oropram 20 mg, tabletki powlekane	DK/H/0762/002	12040	+PHARMA ARZNEIMITTEL GMBH	PL
Citalopram +pharma 20 mg potahované tablety	DK/H/0762/002	30/379/05-C	+PHARMA ARZNEIMITTEL GMBH	CZ
CITALOPRAM ACTAVIS 20 mg apvalkotās tabletes	DK/H/0762/002	05-0462	ACTAVIS NORDIC A/S	LV
Citalopram Actavis 10 mg film-coated tablets	DK/H/0762/001	21105	ACTAVIS NORDIC A/S	FI
Citalopram "Actavis", filmovertrukne tabletter	DK/H/0762/001	32806	ACTAVIS NORDIC A/S	DK
Citalopram Actavis 10 mg plėvele dengtos tabletės	DK/H/0762/001	LT/1/05/0397/001-015	ACTAVIS NORDIC 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
CITALOPRAM ACTAVIS 20 MG TABLETTI, KALVOPÄÄLLYSTEINEN	DK/H/0762/002	21106	ACTAVIS NORDIC A/S	FI
Citalopram "Actavis", filmovertrukne tabletter	DK/H/0762/002	32807	ACTAVIS NORDIC A/S	DK
Citalopram Actavis, 20 mg õhukese polümeerikattega tabletid	DK/H/0762/002	491305	ACTAVIS NORDIC A/S	EE
Citalopram Actavis 20 mg plėvele dengtos tabletės	DK/H/0762/002	LT/1/05/0397/016-030	ACTAVIS NORDIC A/S	LT
Citalopram Actavis 40 mg film-coated tablets	DK/H/0762/003	21107	ACTAVIS NORDIC A/S	FI
Citalopram "Actavis", filmovertrukne tabletter	DK/H/0762/003	32808	ACTAVIS NORDIC A/S	DK
Citalopram Actavis, 40 mg õhukese polümeerikattega tabletid	DK/H/0762/003	491605	ACTAVIS NORDIC A/S	EE
Citalopram Actavis 40 mg plėvele dengtos tabletės	DK/H/0762/003	LT/1/05/0397/031-040	ACTAVIS NORDIC A/S	LT
Citalopram Actavis 40 mg plėvele dengtos tabletės	DK/H/0762/003	LT/1/05/0397/041-045	ACTAVIS NORDIC A/S	LT
Citalopram 20 mg Film-coated tablets	UK/H/0531/002	PL 04569/0480	GENERICS [UK] LIMITED	UK
Ciprager 20 mg Film-coated tablets	UK/H/0531/002	PA0577/047/002	MCDERMOTT LABORATORIES LTD	IE
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046441	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Mylan 20 mg Filmtabletten	UK/H/0531	BE242566	MYLAN DURA GMBH	BE
Citalopram Mylan 20 mg filmomhulde tabletten	UK/H/0531/002	BE242566	MYLAN DURA GMBH	BE
Citalopram Mylan 20 mg filmomhulde tabletten	UK/H/0531/002	BE242557	MYLAN DURA GMBH	BE
Citalopram dura 20 mg Filmtabletten	UK/H/0531/002	55313.01.00	MYLAN DURA GMBH	DE
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	36046249	MYLAN S.P.A.	IT
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	36046237	MYLAN S.P.A.	IT
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	36046225	MYLAN S.P.A.	IT
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	36046213	MYLAN S.P.A.	IT
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	36046201	MYLAN S.P.A.	IT
Citalopram Arcana 10 mg – Filmtabletten	UK/H/0531/001	1-24781	ARCANA ARZNEIMITTEL GMBH	AT
Citalopram dura 10 mg Filmtabletten	UK/H/0531/001	55313.00.00	MYLAN DURA GMBH	DE
Ciprager 10 mg Film-coated tablets	UK/H/0531/001	PA0577/047/001	MCDERMOTT LABORATORIES LTD	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram 10 mg Film-coated tablets	UK/H/0531/001	PL 04569/0479	GENERICS [UK] LIMITED	UK
Citalopram Arcana 20 mg – Filmtabletten	UK/H/0531/002	1-24782	ARCANA ARZNEIMITTEL GMBH	AT
Citalopram Mylan 20 mg comprimés pelliculés	UK/H/0531/002	2003020049	MYLAN BVBA/SPRL	LU
Citalopram Mylan 10 mg filmdragerade tabletter	SE/H/0303/001	17748	MYLAN AB	SE
Citalopram "Mylan", filmovertrukne tabletter	SE/H/0303/001	34311	MYLAN AB	DK
Citalopram Mylan 20 mg filmdragerade tabletter	SE/H/0303/002	17749	MYLAN AB	SE
Citalopram "Mylan", filmovertrukne tabletter	SE/H/0303/002	34312	MYLAN AB	DK
Citalopram Mylan 30 mg filmdragerade tabletter	SE/H/0303/003	18825	MYLAN AB	SE
Citalopram "Mylan", filmovertrukne tabletter	SE/H/0303/003	35533	MYLAN AB	DK
Citalopram Mylan 40 mg filmdragerade tabletter	SE/H/0303/004	18826	MYLAN AB	SE
Citalopram "Mylan", filmovertrukne tabletter	SE/H/0303/004	35534	MYLAN AB	DK
Citalopram Arcana 40 mg – Filmtabletten	UK/H/0531/003	1-24783	ARCANA ARZNEIMITTEL GMBH	AT
Citalopram dura 40 mg Filmtabletten	UK/H/0531/003	55313.02.00	MYLAN DURA GMBH	DE
Ciprager 40 mg Film-coated tablets	UK/H/0531/003	PA0577/047/03	MCDERMOTT LABORATORIES LTD	IE
Citalopram 40 mg Film-coated tablets	UK/H/0531/003	PL 04569/0481	GENERICS [UK] LIMITED	UK
Citagen 10 mg filmtabletta	not available	OGYI-T-9357/01	GENERICS [UK] LIMITED	HU
Citagen 10 mg filmtabletta	not available	OGYI-T-9357/02	GENERICS [UK] LIMITED	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citagen 10 mg filmtabletta	not available	OGYI-T-9357/03	GENERICS [UK] LIMITED	HU
Citagen 10 mg filmtabletta	not available	OGYI-T-9357/04	GENERICS [UK] LIMITED	HU
Citagen 20 mg filmtabletta	not available	OGYI-T-9357/05	GENERICS [UK] LIMITED	HU
Citagen 20 mg filmtabletta	not available	OGYI-T-9357/06	GENERICS [UK] LIMITED	HU
Citagen 20 mg filmtabletta	not available	OGYI-T-9357/07	GENERICS [UK] LIMITED	HU
Citagen 20 mg filmtabletta	not available	OGYI-T-9357/08	GENERICS [UK] LIMITED	HU
Citalopram Mylan Generics Italia 40 mg/ml gocce orali, soluzione	not available	036657017	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046288	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046290	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046302	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046314	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046326	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046338	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046340	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046353	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046365	MYLAN S.P.A.	IT
Citalopram Mylan 40 mg Filmtabletten	UK/H/0531/003	BE265833	MYLAN BVBA/SPRL	BE
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046011	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046023	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046035	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046047	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046050	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046062	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046074	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046086	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046098	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046100	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046112	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046124	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046187	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046199	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046136	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046148	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046151	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046163	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 20 mg compresse rivestite con film	UK/H/0531/002	036046175	MYLAN S.P.A.	IT
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	036046264	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Mylan Generics 20 mg compresse rivestite con film	UK/H/0531/002	036046276	MYLAN S.P.A.	IT
Citalopram Mylan 20 mg comprimés pelliculés	UK/H/0531/002	BE242541	MYLAN BVBA/SPRL	BE
Citalopram Mylan 20 mg comprimés pelliculés	UK/H/0531/002	BE242557	MYLAN BVBA/SPRL	BE
Citalopram Mylan 20 mg comprimés pelliculés	UK/H/0531/002	BE242566	MYLAN BVBA/SPRL	BE
Citalopram MYLAN 20 mg comprimidos recubiertos con película EFG	not available	66095	MYLAN PHARMACEUTICALS S.L.	ES
Citalopram MYLAN 30 mg comprimidos recubiertos con película EFG	not available	67521	MYLAN PHARMACEUTICALS S.L.	ES
Citalox 20 mg filmsko obložene tablete	not available	5363-I-620/10	GENERICS [UK] LIMITED	SI
Citalopram dura 30 mg Filmtabletten	not available	67589.00.00	MYLAN DURA GMBH	DE
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046377	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046389	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046391	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046403	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046415	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046427	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046439	MYLAN S.P.A.	IT
CITALOPRAM MYLAN GENERICS 40 mg compresse rivestite con film	UK/H/0531/003	036046454	MYLAN S.P.A.	IT
Citalopram Mylan 40 mg Filmtabletten	UK/H/0531/003	BE265815	MYLAN BVBA/SPRL	BE
Citalopram Mylan 40 mg Filmtabletten	UK/H/0531/003	BE265824	MYLAN BVBA/SPRL	BE
Citalopram Mylan 20 mg, filmomhulde tabletten	not available	RVG 28672	MYLAN B.V.	NL
Citalopram Mylan 40 mg, filmomhulde tabletten	not available	RVG 28673	MYLAN B.V.	NL
Cilodral 10 mg Δισκία επικαλυμμένα με υμένιο	not available	20434	REMEDICA LTD	CY
Cilodral 20 mg Δισκία επικαλυμμένα με υμένιο	not available	20435	REMEDICA LTD	CY
Cilodral 40 mg Δισκία επικαλυμμένα με υμένιο	not available	20436	REMEDICA LTD	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ADEPRENAL	not available	24597/14/18-09-2015	ADELCO CHROMATOURGIA ATHINON E COLOCOTRONIS BROS SA	GR
Citalopram Alter 30 mg comprimidos recubiertos con película	not available	66.018	LABORATORIOS ALTER, S.A.	ES
Citalopram Alter 20 mg comprimidos recubiertos con película	not available	66.017	LABORATORIOS ALTER, S.A.	ES
Citalopram "STADA" 30mg, filmovertrukne tabletter	not available	38263	STADA ARZNEIMITTEL AG	DK
Citalopram 20 mg film-coated tablets	not available	PL 11311/0416	TILLOMED LABORATORIES LTD	UK
Citalopram 40mg film-coated tablets	not available	PL 11311/0418	TILLOMED LABORATORIES LTD	UK
Citalopram 10mg film-coated Tablets	not available	PL 11311/0415	TILLOMED LABORATORIES LTD	UK
citalopram cinfa 30 mg comprimidos recubiertos con película EFG	not available	67.805	LABORATORIOS CINFA, S.A.	ES
CITALOPRAM ARROW 20 mg, comprimé pelliculé sécable	not available	27291	ARROW GENERIQUES	FR
CITALOPRAM ratiopharm Italia 40 mg/ml gocce orali, soluzione	not available	036038014	RATIOPHARM ITALIA S.R.L.	IT
Citalopram ratiopharm 20 mg compresse rivestite con film	not available	035892013	RATIOPHARM GMBH	IT
Citalopram-Teva 20 mg filmtabletta	not available	OGYI-T-9474/01	TEVA MAGYARORSZÁG ZRT	HU
Citalopram ratiopharm 40 mg compresse rivestite con film	not available	035892052	RATIOPHARM GMBH	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
Citalopram ratiopharm 40 mg compresse rivestite con film	not available	035892064	RATIOPHARM GMBH	IT
Citalopram ratiopharm 20 mg compresse rivestite con film	not available	035892049	RATIOPHARM GMBH	IT
Citalopram ratiopharm 20 mg compresse rivestite con film	not available	035892037	RATIOPHARM GMBH	IT
Citalopram ratiopharm 20 mg compresse rivestite con film	not available	035892025	RATIOPHARM GMBH	IT
CITESINT 20 mg compresse rivestite con film	not available	036484018	GENETIC SPA	IT
CITESINT 40 mg compresse rivestite con film	not available	036484020	GENETIC SPA	IT
CITESINT 40 mg/ml gocce orali, soluzione	not available	036484032	GENETIC SPA	IT
CINAVOL 20 mg compresse rivestite con film	not available	036483016	GENETIC SPA	IT
CINAVOL 40 mg compresse rivestite con film	not available	036483028	GENETIC SPA	IT
CINAVOL 40 mg/ml gocce orali, soluzione	not available	036483030	GENETIC SPA	IT
CINAVOL 40 mg/ml concentrato per soluzione per infusione	not available	036483042	GENETIC SPA	IT
CITALOPRAM ALMUS 20 mg, comprimé pelliculé sécable	not available	3400936477131	BIOGARAN	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
CITALOPRAM AUROBINDO 20 mg compresse rivestite con film	not available	036675015	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Citalopram Hennig® 40 mg Filmtabletten	not available	54076.02.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Pram 10 mg, õhukese polümeerikattega tabletid	not available	421803	G.L. PHARMA GMBH	EE
Pram 20 mg, õhukese polümeerikattega tabletid	not available	422003	G.L. PHARMA GMBH	EE
Pram 40 mg, õhukese polümeerikattega tabletid	not available	421903	G.L. PHARMA GMBH	EE
Citalopram Jubilant 10 mg filmdragerade tabletter	NL/H/2919/001	49278	JUBILANT PHARMACEUTICALS NV	SE
Citalopram Jubilant 20 mg filmdragerade tabletter	NL/H/2919/002	49279	JUBILANT PHARMACEUTICALS NV	SE
Citalopram Jubilant 30 mg filmdragerade tabletter	NL/H/2919/003	49280	JUBILANT PHARMACEUTICALS NV	SE
Citalopram Jubilant 40 mg filmdragerade tabletter	NL/H/2919/004	49281	JUBILANT PHARMACEUTICALS NV	SE
Citalopram "Jubilant", filmovertrukne tabletter	NL/H/2919/001	52337	JUBILANT PHARMACEUTICALS NV	DK
Citalopram "Jubilant", filmovertrukne tabletter	NL/H/2919/002	52338	JUBILANT PHARMACEUTICALS NV	DK
Citalopram "Jubilant", filmovertrukne tabletter	NL/H/2919/003	52339	JUBILANT PHARMACEUTICALS NV	DK
Citalopram "Jubilant", filmovertrukne tabletter	NL/H/2919/004	52340	JUBILANT PHARMACEUTICALS NV	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
CITALVIR 10 mg comprimidos recubiertos con película	not available	66.402	ABABOR PHARMACEUTICALS, S.L.	ES
CITALVIR 20 mg comprimidos recubiertos con película EFG	not available	66.401	ABABOR PHARMACEUTICALS, S.L.	ES
DALSAN 10 mg filmtabletta	HU/H/0109/001/MR	OGYI-T-9985/01	EGIS PHARMACEUTICALS PLC	HU
Dalsan 10 mg apvalkotās tabletes	HU/H/0109/001-003/DC	06-0054	EGIS PHARMACEUTICALS PLC	LV
DALSAN 20 mg filmtabletta	HU/H/0109/002/MR	OGYI-T-9985/02	EGIS PHARMACEUTICALS PLC	HU
Dalsan 20 mg apvalkotās tabletes	HU/H/0109/001-003/MR	06-0055	EGIS PHARMACEUTICALS PLC	LV
DALSAN 40 mg filmtabletta	HU/H/0109/003/MR	OGYI-T-9985/03	EGIS PHARMACEUTICALS PLC	HU
Dalsan 40 mg apvalkotās tabletes	HU/H/0109/001-003/MR	06-0056	EGIS PHARMACEUTICALS PLC	LV
CITALOPRAM 10MG FILM-COATED TABLETS	not available	PL 30306/0618	ACTAVIS GROUP PTC EHF.	UK
CITALOPRAM 20MG FILM-COATED TABLETS	not available	PL 30306/0619	ACTAVIS GROUP PTC EHF.	UK
CITALOPRAM 40MG FILM-COATED TABLETS	not available	PL 30306/0620	ACTAVIS GROUP PTC EHF.	UK
Citalopram 40 mg/ml oral drops, solution	not available	PL 28176/0164	STRIDES ARCOLAB INTERNATIONAL LIMITED	UK
Citalopram ratiopharm 20 mg comprimidos recubiertos con película EFG	not available	64.738	RATIOPHARM ESPAÑA, S.A	ES
citalopram-biomo® 10 mg Filmtabletten	not available	50941.00.00	BIOMO PHARMA GMBH	DE
citalopram-biomo® 20 mg Filmtabletten	not available	50941.01.00	BIOMO PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
citalopram-biomo® 40 mg Filmtabletten	not available	50941.02.00	BIOMO PHARMA GMBH	DE
citalopram-biomo® 30 mg Filmtabletten	not available	62359.00.00	BIOMO PHARMA GMBH	DE
Xadorek 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	65424/15-7-2014	MINERVA PHARMACEUTICAL S.A	GR
Xadorek 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	64572/15-7-2014	MINERVA PHARMACEUTICAL S.A	GR
Citalopram Wynn 10 mg comprimidos revestidos por película	not available	5427265	AXONE, LDA.	PT
Citalopram Wynn 10 mg comprimidos revestidos por película	not available	5427257	AXONE, LDA.	PT
CITALOPRAM ALMUS 20 mg comprimidos recubiertos con película EFG	ES/H/0146/002	70369	ALMUS FARMACEUTICA S.A	ES
CITALOPRAM ALMUS 30 mg comprimidos recubiertos con película EFG	ES/H/0146/003	70370	ALMUS FARMACEUTICA S.A	ES
CITALOPRAM STADA 20 mg comprimidos recubiertos con película EFG	not available	66.035	LABORATORIO STADA, S.L.	ES
Citalopram 10mg Film-coated Tablets	not available	PL 17907/0089	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Citalopram 20mg Film-coated Tablets	not available	PL 17907/0090	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Citalopram 40mg Film-coated Tablets	not available	PL 17907/0091	BRISTOL LABORATORIES LTD (BERKHAMSTED)	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
Citalopram EDIGEN 10 mg comprimidos recubiertos con película	not available	66.278	ARISTO PHARMA IBERIA, S.L.	ES
Citalopram EDIGEN 20 mg comprimidos recubiertos con película EFG	not available	66.277	ARISTO PHARMA IBERIA, S.L.	ES
Citalopram EDIGEN 40 mg comprimidos recubiertos con película EFG	not available	66.280	ARISTO PHARMA IBERIA, S.L.	ES
CITALOPRAM TecniGen 40 mg/ml gocce orali, soluzione	not available	036057014	TECNIGEN S.R.L.	IT
CITALOPRAM MYLAN 20 mg, comprimé pelliculé sécable	not available	NL 27 236	MYLAN S.A.S	FR
Citalopram Q-Pharm 10 mg Filmtabletten	not available	90967.00.00	JUTA PHARMA GMBH	DE
Citalopram Q-Pharm 20 mg Filmtabletten	not available	90968.00.00	JUTA PHARMA GMBH	DE
Citalopram Q-Pharm 40 mg Filmtabletten	not available	90969.00.00	JUTA PHARMA GMBH	DE
FRIMAIND 20 mg compresse rivestite con film	not available	036143016	SO.SE.PHARM S.R.L.	IT
FRIMAIND 40 mg compresse rivestite con film	not available	036143028	SO.SE.PHARM S.R.L.	IT
FRIMAIND 40 mg/ml gocce orali, soluzione	not available	036143055	SO.SE.PHARM S.R.L.	IT
Citalostad 40 mg Filmtabletten	not available	1-25316	STADA ARZNEIMITTEL GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Teva-Rimafar 30 mg comprimidos recubiertos con pelicula EFG	not available	67579	TEVA PHARMA S.L.U	ES
Citabax 10, 10 mg, tabletki powlekane	not available	10915	RANBAXY POLAND SP. ZO.O	PL
Citabax 20, 20 mg, tabletki powlekane	not available	10916	RANBAXY POLAND SP. ZO.O	PL
Citabax 40, 40 mg, tabletki powlekane	not available	10917	RANBAXY POLAND SP. ZO.O	PL
CITALOPRAM DOC 40 mg/ml gocce orali, soluzione	not available	036653018	DOC GENERICI S.R.L.	IT
Citalopram Genericon 10 mg Filmtabletten	NL/H/0312/001	1-24570	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
Citalopram Genericon 20 mg Filmtabletten	NL/H/0312/002	1-24571	GENERICON PHARMA GESELLSCHAFT M.B.H.	AT
CITALOPRAM ACTAVIS 10 MG, FILMOMHULDE TABLETTEN	NL/H/0312/001	RVG 25419	ACTAVIS GROUP PTC EHF.	NL
CITALOPRAM ACTAVIS 20 MG, FILMOMHULDE TABLETTEN	NL/H/0312/002	RVG 25420	ACTAVIS GROUP PTC EHF.	NL
CITALOPRAM ACTAVIS 40 MG, FILMOMHULDE TABLETTEN	NL/H/0312/003	RVG 25421	ACTAVIS GROUP PTC EHF.	NL
Citalopram 10mg Film-Coated Tablets	NL/H/0312/001-003/MR	0142/0543	ACTAVIS UK LTD.	UK
Citalopram 20mg Film-Coated Tablets	NL/H/0312/001-003/MR	0142/0544	ACTAVIS UK LTD.	UK
Citalopram 40mg Film-Coated Tablets	NL/H/0312/001-003/MR	0142/0545	ACTAVIS UK LTD.	UK
Exenadil® 20 mg Επικαλυμμένα με λεπτό υμένιο δισκία	not available	43899/12	TARGET PHARMA SINGLE MEMBER PRIVATE LTD	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
Exenadil® 40 mg Επικαλυμμένα με λεπτό υμένιο δισκία	not available	43900/12	TARGET PHARMA SINGLE MEMBER PRIVATE LTD	GR
Exenadil® 40 mg/ml Πόσιμες σταγόνες, διάλυμα	not available	65428/15	TARGET PHARMA SINGLE MEMBER PRIVATE LTD	GR
Citalopram 10 mg Tablets	not available	PL 36390/0024	CIPLA (EU) LIMITED	UK
Citalopram 20 mg Tablets	not available	PL 36390/0025	CIPLA (EU) LIMITED	UK
Citalopram 40 mg Tablets	not available	PL 36390/0026	CIPLA (EU) LIMITED	UK
PREFUCET	not available	13959	RAFARM SA.	GR
PREFUCET	not available	13960	RAFARM SA.	GR
PREFUCET	not available	7784	RAFARM SA.	GR
Citalopram PENSA 20 mg compresse rivestite con film	not available	036392013	PENSA PHARMA S.P.A.	IT
Citalopram Pensa 20 mg compresse rivestite con film	not available	036392025	PENSA PHARMA S.P.A.	IT
Citalopram Pensa Pharma 40 mg/ml gocce orali, soluzione	not available	038199016	PENSA PHARMA S.P.A.	IT
Citalopram PENSA 40 mg compresse rivestite con film	not available	036392037	PENSA PHARMA S.P.A.	IT
Citalopram AL 30 mg Filmtabletten	not available	67591.00.00	ALIUD PHARMA GMBH	DE
CITALOPRAM ZENTIVA 40 MG/ML GOCCE ORALI, SOLUZIONE	not available	036254011	ZENTIVA ITALIA SRL	IT
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALOPRAM ZENTIVA 20 mg, comprimé pelliculé sécable	not available	367 432-5	SANOFI-AVENTIS FRANCE	FR
CITALOPRAM ZENTIVA 20 mg, comprimé pelliculé sécable	not available	565 603-0	SANOFI-AVENTIS FRANCE	FR
Citalopram 40mg Tablets	not available	PL 17780/0038	WINTHROP PHARMACEUTICALS UK LTD	UK
Citalopram 20mg Tablets	not available	PL 17780/0037	WINTHROP PHARMACEUTICALS UK LTD	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
Citalopram 10mg Tablets	not available	PL 17780/0036	WINTHROP PHARMACEUTICALS UK LTD	UK
CitaLich® 10 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/001	58944.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 10 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/001	58944.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 10 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/001	58944.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 40 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/003	58944.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 40 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/003	58944.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 40 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/003	58944.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 40 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/003	58944.02.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 10 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/001	58944.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 20 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/002	58944.01.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 20 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/002	58944.01.00	WINTHROP ARZNEIMITTEL GMBH	DE
CitaLich® 20 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/002	58944.01.00	WINTHROP ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CitaLich® 20 mg Filmtabletten Wirkstoff: Citalopram	DE/H/1319/002	58944.01.00	WINTHROP ARZNEIMITTEL GMBH	DE
CITALEC 10 Zentiva potahované tablety	CZ/H/0127/001	30/552/05-C	ZENTIVA, A.S.	CZ
CITALEC 10 Zentiva filmom obalené tablety	CZ/H/0127/001	30/0357/06-S	ZENTIVA, A.S.	SK
CITALEC 20, 20 mg õhukese polümeerikattega tabletid	CZ/H/0127/002	403202	ZENTIVA, A.S.	EE
CITALEC 20 Zentiva potahované tablety	CZ/H/0127/002	30/553/05-C	ZENTIVA, A.S.	CZ
CITALEC 20 Zentiva filmom obalené tablety	CZ/H/0127/002	30/0358/06-S	ZENTIVA, A.S.	SK
Citalopram-Zentiva 20 mg filmtabletta	DE/H/1319/002	OGYI-T-10046/01	ZENTIVA HU KFT	HU
Citalopram-Zentiva 40 mg filmtabletta	DE/H/1319/003	OGYI-T-10046/02	ZENTIVA HU KFT	HU
CITALOPRAM ZENTIVA ITALIA 20 mg compresse rivestite con film	not available	036656015	ZENTIVA ITALIA SRL	IT
Citalopram Zentiva 30 mg comprimidos recubiertos con película EFG	not available	67545	ZENTIVA, K.S.	ES
Citalopram Zentiva 20 mg comprimidos recubiertos con película EFG	not available	67544	ZENTIVA, K.S.	ES
Citalopram Zentiva 20 mg comprimidos recubiertos con película EFG	not available	67544	ZENTIVA, K.S.	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
Pram 10 mg plėvele dengtos tabletės	not available	LT/1/03/2944/001	G.L. PHARMA GMBH	LT
Pram 20 mg plevele dengtos tabletes	not available	LT/1/03/2944/002	G.L. PHARMA GMBH	LT
Pram 40 mg plėvele dengtos tabletės	not available	LT/1/03/2944/003	G.L. PHARMA GMBH	LT
TALOSIN 20 mg, Επικαλυμμένα με λεπτό υμένιο δισκία	not available	21091/13/05-06-2014	BENNETT PHARMACEUTICALS SA	GR
TALOSIN 40 mg, Επικαλυμμένα με λεπτό υμένιο δισκία	not available	21092/13/05-06-2014	BENNETT PHARMACEUTICALS SA	GR
CITALOPRAM CRISTERS 20 mg, comprimé pelliculé sécable	not available	34009 366 508 8 6	CRISTERS	FR
CITALOPRAM CRISTERS 20 mg, comprimé pelliculé sécable	not available	34009 366 509 4 7	CRISTERS	FR
CITALOPRAM CRISTERS 20 mg, comprimé pelliculé sécable	not available	34009 565 999 1 2	CRISTERS	FR
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	5619960	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	5619978	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	5620000	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	5620018	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL 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
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 10 mg comprimidos revestidos por película	not available	10/H/0124/001	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 20 mg comprimidos revestidos por película	not available	10/H/0124/002	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL 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
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
Citalopram Aurobindo 40 mg comprimidos revestidos por película	not available	10/H/0124/003	AUROBINDO PHARMA (PORTUGAL), UNIPESSOAL LDA	PT
CITALOPRAM ZYDUS 20 mg, comprimé pelliculé sécable	not available	34009 388 171 6 4	ZYDUS FRANCE	FR
LOPRACIL®, πόσιμες σταγόνες, διάλυμα, 40mg/ml	not available	43228/11/4-5-2012	MEDICUS A.E	GR
LOPRACIL®, επικαλυμμένα με λεπτό υμένιο δισκία, 40mg/tab	not available	43225/11/4-5-2012	MEDICUS A.E	GR
LOPRACIL®, επικαλυμμένα με λεπτό υμένιο δισκία, 20mg/tab	not available	43224/11/4-5-2012	MEDICUS A.E	GR
CITALOPRAM STADA 30 mg comprimidos recubiertos con película EFG	not available	66.034	LABORATORIO STADA, S.L.	ES
Citalopram 10mg Film-Coated Tablets	not available	PL 20416/0301	CRESCENT PHARMA LIMITED	UK
Citalopram 20mg Film-Coated Tablets	not available	PL 20416/0302	CRESCENT PHARMA LIMITED	UK
Citalopram 40mg Film-Coated Tablets	not available	PL 20416/0303	CRESCENT PHARMA 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
CITALOPRAM TecniGen 20 mg compresse rivestite con film	not available	036057040	TECNIGEN S.R.L.	IT
CITALOPRAM TecniGen 40 mg compresse rivestite con film	not available	036057053	TECNIGEN S.R.L.	IT
CITALOPRAM ISOMED 20 mg, comprimé pelliculé sécable	not available	NL27608	TEVA SANTÉ	FR
Citalopram Vitabalans 20 mg kalvopäällysteiset tabletit	SE/H/0818/001	28206	VITABALANS OY	FI
Citalopram Vitabalans 40 mg kalvopäällysteiset tabletit	SE/H/0818/002	28207	VITABALANS OY	FI
Citalopram ”Vitabalans”, filmovertrukne tabletter	SE/H/0818/001	46084	VITABALANS OY	DK
Citalopram ”Vitabalans”, filmovertrukne tabletter	SE/H/0818/002	46085	VITABALANS OY	DK
Citalopram Vitabalans 20 mg fildragerade tabletter	SE/H/0818/001	43373	VITABALANS OY	SE
Citalopram Vitabalans 40 mg fildragerade tabletter	SE/H/0818/002	43374	VITABALANS OY	SE
Citalopram Vitabalans 20 mg filmdrasjerte tabletter	SE/H/818/001-002/DC	09-7212	VITABALANS OY	NO
Citalopram Vitabalans 40 mg filmdrasjerte tabletter	SE/H/818/001-002/DC	09-7213	VITABALANS OY	NO
Citalopram Vitabalans, 20 mg õhukese polümeerikattega tabletid	SE/H/0818/001	732311	VITABALANS OY	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Vitabalans, 40 mg õhukese polümeerikattega tabletid	SE/H/0818/002	732411	VITABALANS OY	EE
Citalopram Vitabalans 20 mg apvalkotas tabletes	SE/H/0818/001	11-0174	VITABALANS OY	LV
Citalopram Vitabalans 40 mg apvalkotās tabletes	SE/H/0818/002	11-0175	VITABALANS OY	LV
Citalopram Vitabalans 20 mg Filmtabletten	SE/H/0818/001	80483.00.00	VITABALANS OY	DE
Citalopram Vitabalans 40 mg Filmtabletten	SE/H/0818/002	80484.00.00	VITABALANS OY	DE
Citalopram Vitabalans 20 mg filmom obalené tablety	SE/H/0818/001	30/0306/11-S	VITABALANS OY	SK
Citalopram Vitabalans 40 mg filmom obalené tablety	SE/H/0818/002	30/0307/11-S	VITABALANS OY	SK
Citalopram Vitabalans 20 mg potahované tablety	SE/H/0818/001	30/111/11-C	VITABALANS OY	CZ
Citalopram Vitabalans 40 mg potahované tablety	SE/H/0818/002	30/112/11-C	VITABALANS OY	CZ
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/001	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/007	VITABALANS OY	LT
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/002	VITABALANS OY	LT
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/003	VITABALANS OY	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
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/004	VITABALANS OY	LT
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/005	VITABALANS OY	LT
Citalopram Vitabalans 20 mg plėvele dengtos tabletės	SE/H/0818/001	LT/1/11/2417/006	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/008	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/009	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/010	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/011	VITABALANS OY	LT
Citalopram Vitabalans 40 mg plėvele dengtos tabletės	SE/H/0818/002	LT/1/11/2417/012	VITABALANS OY	LT
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/01	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/07	VITABALANS OY	HU
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/02	VITABALANS OY	HU
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/03	VITABALANS OY	HU
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/04	VITABALANS OY	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/05	VITABALANS OY	HU
Citalopram Vitabalans 20 mg filmtabletta	SE/H/0818/001	OGYI-T-21674/06	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/08	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/09	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/10	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/11	VITABALANS OY	HU
Citalopram Vitabalans 40 mg filmtabletta	SE/H/0818/002	OGYI-T-21674/12	VITABALANS OY	HU
Citalopram Vitabalans 20 mg filmsko obložene tablete	SE/H/0818/001	H/12/00395/001	VITABALANS OY	SI
Citalopram Vitabalans 40 mg filmsko obložene tablete	SE/H/0818/002	H/12/00395/007	VITABALANS OY	SI
Citalopram Vitabalans 20 mg filmsko obložene tablete	SE/H/0818/001	H/12/00395/002	VITABALANS OY	SI
Citalopram Vitabalans 20 mg filmsko obložene tablete	SE/H/0818/001	H/12/00395/003	VITABALANS OY	SI
Citalopram Vitabalans 40 mg filmsko obložene tablete	SE/H/0818/002	H/12/00395/008	VITABALANS OY	SI
Citalopram Vitabalans 40 mg filmsko obložene tablete	SE/H/0818/002	H/12/00395/009	VITABALANS OY	SI
Citalopram Vitabalans, 20 mg, tabletki powlekane	SE/H/0818/001	20322	VITABALANS OY	PL

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
Citalopram Vitabalans, 40 mg, tabletki powlekane	SE/H/0818/002	20323	VITABALANS OY	PL
Citalopram G.L. 10 mg - Filmtabletten	not available	1-24775	G.L. PHARMA GMBH	AT
Citalopram G.L. 20 mg - Filmtabletten	not available	1-24774	G.L. PHARMA GMBH	AT
Citalopram G.L. 40 mg - Filmtabletten	not available	1-24776	G.L. PHARMA GMBH	AT
Citalopram Qualigen 20 mg comprimidos recubiertos con película	not available	70.893	QUALIGEN, S.L.	ES
Citalopram Qualigen 30 mg comprimidos recubiertos con película	not available	70.894	QUALIGEN, S.L.	ES
Citalopram 10 mg Tablets	not available	PL 20075/0272	ACCORD HEALTHCARE LIMITED	UK
Citalopram 20 mg Tablets	not available	PL 20075/0273	ACCORD HEALTHCARE LIMITED	UK
Citalopram 40 mg Tablets	not available	PL 20075/0274	ACCORD HEALTHCARE LIMITED	UK
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400958178528	BIOGARAN	FR
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400949179718	BIOGARAN	FR
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400930031353	BIOGARAN	FR
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400955011026	BIOGARAN	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
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400930031360	BIOGARAN	FR
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400957762025	BIOGARAN	FR
CITALOPRAM BIOGARAN 20 mg, comprimé pelliculé sécable	not available	3400922033167	BIOGARAN	FR
Citalopram Q-Pharm 10 mg Filmtabletten	not available	90967.00.00	JUTA PHARMA GMBH	DE
Citalopram Q-Pharm 20 mg Filmtabletten	not available	90968.00.00	JUTA PHARMA GMBH	DE
Citalopram Q-Pharm 40 mg Filmtabletten	not available	90969.00.00	JUTA PHARMA GMBH	DE
Citalopram 40mg/ml Oral Drops, solution	UK/H/1238/001	PL 20046/0053	FOCUS PHARMACEUTICALS LIMITED	UK
Citalopram 40mg/ml Oral Drops, solution	UK/H/1238/001	PA 1338/003/001	FOCUS PHARMACEUTICALS LIMITED	IE
Citalopram 10mg Tablets	not available	PL 06453/0056	ATHLONE LABORATORIES LIMITED	UK
Citalopram 20mg Tablets	not available	PL 06453/0057	ATHLONE LABORATORIES LIMITED	UK
Citalopram 40mg Tablets	not available	PL 06453/0058	ATHLONE LABORATORIES LIMITED	UK
Pram 20 mg-Konzentrat zur Herstellung einer Infusionslösung	not available	1-29270	G.L. PHARMA GMBH	AT
Pram 40 mg-Konzentrat zur Herstellung einer Infusionslösung	not available	1-29271	G.L. PHARMA GMBH	AT
Cipramil 10 mg filmuhúðaðar töflur	not available	970141	H. LUNDBECK A/S	IS
Cipramil 20 mg filmuhúðaðar töflur	not available	920052	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
ELOPRAM 20 mg compresse rivestite con film	not available	028681017	LUNDBECK ITALIA SPA, IT	IT
ELOPRAM 20 mg compresse rivestite con film	not available	028681031	LUNDBECK ITALIA SPA, IT	IT
ELOPRAM 40 mg compresse rivestite con film	not available	028681029	LUNDBECK ITALIA SPA, IT	IT
ELOPRAM 40 mg/ml concentrato per soluzione per infusione	not available	028681043	LUNDBECK ITALIA SPA, IT	IT
ELOPRAM 40 mg/ml gocce orali, soluzione	not available	028681056	LUNDBECK ITALIA SPA, IT	IT
Cipramil 40 mg/ml Oral drops, solution.	not available	PA 776/1/4	LUNDBECK LTD., IE	IE
Cipramil, 20 mg, tabletki powlekane	not available	8015	H. LUNDBECK A/S	PL
Cipramil 10 mg film-coated tablets	not available	PA 776/1/1	LUNDBECK LTD., IE	IE
Cipramil 20 mg film-coated tablets	not available	PA 776/1/2	LUNDBECK LTD., IE	IE
PRISDAL 20 mg comprimidos recubiertos con película	not available	60.884	LUNDBECK ESPANA, ES	ES
PRISDAL 30 mg comprimidos recubiertos con película	not available	63.627	LUNDBECK ESPANA, ES	ES
SEROPRAM 20 mg comprimidos recubiertos con película	not available	60.885	LUNDBECK ESPANA, ES	ES
SEROPRAM 30 mg comprimidos recubiertos con película	not available	63.564	LUNDBECK ESPANA, ES	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
SEROPRAM 20 mg potahované tablety	not available	30/122/91-C	H. LUNDBECK A/S	CZ
SEROPRAM 40 mg/ml perorální kapky, roztok	not available	30/209/99-C	H. LUNDBECK A/S	CZ
Seropram 10mg - Filmtabletten	not available	1-19871	LUNDBECK AUSTRIA GMBH	AT
Seropram 20mg - Filmtabletten	not available	1-19326	LUNDBECK AUSTRIA GMBH	AT
Seropram 20 mg – Konzentrat zur Infusionsbereitung	not available	1-19815	LUNDBECK AUSTRIA GMBH	AT
Seropram 40 mg – Konzentrat zur Infusionsbereitung	not available	1-19396	LUNDBECK AUSTRIA GMBH	AT
Cipramil	not available	13248	H. LUNDBECK A/S	DK
Cipramil	not available	13249	H. LUNDBECK A/S	DK
Cipramil	not available	19168	H. LUNDBECK A/S	DK
Cipramil	not available	13250	H. LUNDBECK A/S	DK
SEROPRAM koncentrát pro přípravu infuzního roztoku	not available	30/794/92-C	H. LUNDBECK A/S	CZ
Cipramil 10 mg tabletter, filmdrasjerte	not available	8150	H. LUNDBECK A/S	NO
Cipramil 20 mg tabletter, filmdrasjerte	not available	8151	H. LUNDBECK A/S	NO
Cipramil 30 mg tabletter, filmdrasjerte	not available	97-2985	H. LUNDBECK A/S	NO
Cipramil 40 mg tabletter, filmdrasjerte	not available	8152	H. LUNDBECK A/S	NO
Cipramil® Drops 40 mg/ml, oral drops solution	not available	PL 0458/0071	LUNDBECK LTD. GB	UK
Cipramil® 20 mg, Filmtabletten	not available	43145.01.00	LUNDBECK GMBH, DE	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
Cipramil® 40 mg, Filmtabletten	not available	43145.02.00	LUNDBECK GMBH, DE	DE
Cipramil® Infusionslösungskonzentrat 20 mg, Konzentrat zur Herstellung einer Infusionslösung	not available	43451.00.00	LUNDBECK GMBH, DE	DE
Cipramil 20 mg apvalkotās tablets	not available	01-0421	H. LUNDBECK A/S	LV
Cipramil 20 mg, õhukese polümeerikattega tabletid	not available	227098	H. LUNDBECK A/S	EE
CIPRAMIL 20 mg filmomhulde tabletten	not available	BE151261	LUNDBECK SA, BE	BE
CIPRAMIL 40 mg/ml concentraat voor oplossing voor infusie	not available	BE151164	LUNDBECK SA, BE	BE
CIPRAMIL 20 mg comprimés pelliculés	not available	BE151261	LUNDBECK SA - N.V.	BE
CIPRAMIL 40 mg/ml solution à diluer pour perfusion	not available	BE151164	LUNDBECK SA - N.V.	BE
Citalopram Lundbeck 10 mg filmdragerade tabletter	not available	14271	H. LUNDBECK A/S	SE
Citalopram Lundbeck 20 mg filmdragerade tabletter	not available	14272	H. LUNDBECK A/S	SE
Cipramil 10 mg filmdragerade tabletter	not available	11660	H. LUNDBECK A/S	SE
Cipramil 20 mg filmdragerade tabletter	not available	11661	H. LUNDBECK A/S	SE
CIPRAMIL 20 mg comprimés pelliculés	not available	BE151261	LUNDBECK SA, BE	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
CIPRAMIL 40 mg/ml solution à diluer pour perfusion	not available	BE151164	LUNDBECK SA, BE	LU
Cipramil 20 mg, filmomhulde tabletten 20 mg	not available	RVG 19593	LUNDBECK B.V., NL	NL
Cipramil 40 mg, filmomhulde tabletten 40 mg	not available	RVG 19594	LUNDBECK B.V., NL	NL
Cipramil 40 mg/ml druppels voor oraal gebruik, oplossing	not available	RVG 22687	LUNDBECK B.V., NL	NL
Sepram 10 mg tabletti, kalvopäällysteinen	not available	13378	H. LUNDBECK A/S	FI
Sepram 20 mg tabletti, kalvopäällysteinen	not available	13379	H. LUNDBECK A/S	FI
Sepram 30 mg tabletti, kalvopäällysteinen	not available	15871	H. LUNDBECK A/S	FI
Sepram 40 mg tabletti, kalvopäällysteinen	not available	13380	H. LUNDBECK A/S	FI
SEROPRAM 40 mg/ml gocce orali, soluzione	not available	028759049	LUNDBECK ITALIA SPA, IT	IT
SEROPRAM 20 mg compresse rivestite con film	not available	028759013	LUNDBECK ITALIA SPA, IT	IT
SEROPRAM 20 mg compresse rivestite con film	not available	028759037	LUNDBECK ITALIA SPA, IT	IT
SEROPRAM 40 mg compresse rivestite con film	not available	028759025	LUNDBECK ITALIA SPA, IT	IT
SEROPRAM 40 mg/ml concentrato per soluzione per infusione	not available	028759052	LUNDBECK ITALIA SPA, IT	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
SEROPRAM 20 mg/0,5 ml, solution à diluer pour perfusion	not available	561 558-0	LUNDBECK SAS	FR
SEROPRAM 40 mg/1 ml, solution à diluer pour perfusion	not available	561 559-7	LUNDBECK SAS	FR
SEROPRAM 40 mg/ml, solution buvable	not available	346 537-2	LUNDBECK SAS	FR
Seropram 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45651/23-12-2004	LUNDBECK HELLAS, GR	GR
Seropram 20 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	73634/23-12-2004	LUNDBECK HELLAS, GR	GR
Seropram 40 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	73635/23-12-2004	LUNDBECK HELLAS, GR	GR
Seropram 40 mg/ml Πόσιμες σταγόνες διάλυμα	not available	45649/23-12-2004	LUNDBECK HELLAS, GR	GR
Seropram 40 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	19734/21-5-2001	LUNDBECK HELLAS, GR	GR
SEROPRAM 20 mg, comprimé pelliculé sécable	not available	338 336 -1	LUNDBECK SAS	FR
SEROPRAM 20 mg, comprimé pelliculé sécable	not available	570 510 - 7	LUNDBECK SAS	FR
Seropram	not available	16117	H. LUNDBECK A/S	CY
Cipramil® 10 mg film-coated tablets	not available	PL 0458/0057	LUNDBECK LTD. GB	UK
Cipramil® 20 mg film-coated tablets	not available	PL 0458/0058	LUNDBECK LTD. GB	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
Cipramil® 40 mg film-coated tablets	not available	PL 0458/0059	LUNDBECK LTD. GB	UK
Seropram 40 mg/ml koncentrátum oldatos infúzióhoz	not available	OGYI-T-2100/03	LUNDBECK HUNGARIA KFT.	HU
Seropram 20 mg filmtabletta	not available	OGYI-T-2100/01	LUNDBECK HUNGARIA KFT.	HU
Cipramil 20 mg filmsko obložene tablete	not available	5363-I-537/09	LUNDBECK PHARMA D.O.O	SI
Серопрам 20 mg филмирани таблетки	not available	9600121	H. LUNDBECK A/S	BG
Citalopram esparma 10 mg Filmtabletten	DE/H/1980/001	58947.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Citalopram esparma 20 mg Filmtabletten	DE/H/1980/002	58947.01.00	ARISTO PHARMA GMBH (ART 57)	DE
Citalopram esparma 40 mg Filmtabletten	DE/H/1980/003	58947.02.00	ARISTO PHARMA GMBH (ART 57)	DE
Pram, 20 mg, tabletki powlekane	not available	11052	G.L. PHARMA GMBH	PL
CITALOPRAM ALMUS PHARMA 40 mg/ml gocce orali, soluzione	not available	036971012	ALMUS S.R.L	IT
CITALOPRAM ALMUS 20 mg compresse rivestite con film	not available	036434013	ALMUS S.R.L	IT
Далсан 20 mg филмирани таблетки	not available	20060570	EGIS PHARMACEUTICALS PLC	BG
Далсан 10 mg филмирани таблетки	not available	20060569	EGIS PHARMACEUTICALS PLC	BG
Далсан 40 mg филмирани таблетки	not available	20060571	EGIS PHARMACEUTICALS PLC	BG
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111147	RANBAXY ITALIA S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111186	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111198	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111200	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111174	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111111	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111135	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111123	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111162	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 20 mg compresse rivestite con film	UK/H/806/02/MR	037111150	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111034	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111251	RANBAXY ITALIA S.P.A.	IT
Citalopram10 mg Film-coated Tablets	UK/H/0806/001	PL 14894 / 0163	RANBAXY UK LTD	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
Citalopram 20 mg Film-coated Tablets	UK/H/0806/002	PL 14894/0164	RANBAXY UK LTD	UK
Citalopram 40 mg Film-coated Tablets	UK/H/0806/003	PL 14894/0165	RANBAXY UK LTD	UK
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111046	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111085	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111061	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111109	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111073	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111097	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111022	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111059	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 10 mg compresse rivestite con film	UK/H/0806/001	037111010	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111212	RANBAXY ITALIA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111224	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111236	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111248	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111263	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111287	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111301	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111275	RANBAXY ITALIA S.P.A.	IT
Citalopram Ranbaxy 40 mg compresse rivestite con film	UK/H/806/03/MR	037111299	RANBAXY ITALIA S.P.A.	IT
PRICITAL	not available	40060/11-9-2008	LAVIPHARM HELLAS AE	GR
Citalopram Teva-Rimafar 20 mg comprimidos recubiertos con película EFG	not available	66508	TEVA PHARMA S.L.U	ES
Citalopram Actavis 30 mg comprimidos revestidos por película	DE/H/1164/003	5136536	ACTAVIS GROUP PTC EHF.	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
Citalopram Aurovitas Spain 20 mg comprimidos recubiertos con película EFG	DE/H/1164/002	70440	AUROVITAS SPAIN,S.A.U.	ES
Citalopram Aurovitas Spain 40 mg comprimidos recubiertos con película EFG	DE/H/1164/004	70438	AUROVITAS SPAIN,S.A.U.	ES
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	5136460	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 10 mg comprimidos revestidos por película	DE/H/1164/001	5136478	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	5136502	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 20 mg comprimidos revestidos por película	DE/H/1164/002	5136510	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	5136544	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Aurovitas 40 mg comprimidos revestidos por película	DE/H/1164/004	5136551	AUROVITAS UNIPESSOAL, LDA.	PT
Citalopram Actavis 10 mg Filmtabletten	DE/H/1164/001	1-28112	ACTAVIS GROUP PTC EHF.	AT
Citalopram-Actavis 10 mg Filmtabletten	DE/H/1164/001	69808.00.00	ACTAVIS GROUP PTC EHF.	DE
Citalopram Actavis 20 mg Filmtabletten	DE/H/1164/002	1-28113	ACTAVIS GROUP PTC EHF.	AT
Citalopram-Actavis 20 mg Filmtabletten	DE/H/1164/002	69809.00.00	ACTAVIS GROUP PTC EHF.	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
Citalopram Actavis 40 mg Filmtabletten	DE/H/1164/003	1-28115	ACTAVIS GROUP PTC EHF.	AT
Citalopram-Actavis 40 mg Filmtabletten	DE/H/1164/004	69811.00.00	ACTAVIS GROUP PTC EHF.	DE
Oropram 10 mg filmsko obloiene tablete	DE/H/1164/001	H/09/01191/001	ACTAVIS GROUP PTC EHF.	SI
Oropram 20 mg filmsko obloiene tablete	DE/H/1164/002	H/09/01191/011	ACTAVIS GROUP PTC EHF.	SI
Oropram 40 mg filmsko obloiene tablete	DE/H/1164/003	H/09/01191/022	ACTAVIS GROUP PTC EHF.	SI
SELON	not available	41577	GENEPHARM S.A.	GR
Selon	not available	41578/20-10-2009	GENEPHARM S.A.	GR
Citalopram Farmaprojects 20 mg comprimidos recubiertos con película EFG	NOT APPLICABLE	66266	Farmaprojects S.A.U.	ES