30 November 2017
EMA/812281/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): moclobemide

Procedure No.: PSUSA/00002079/201704

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Moclobemide 300 mg film-coated tablets	DK/H/0292/002	PL 04416/1420	SANDOZ LTD	UK
Moclobemid HEXAL 300 mg Filmtabletten	DK/H/0292/002	53011.01.00	HEXAL AG	DE
Moclobemid-1 A Pharma 300 mg Filmtabletten	DK/H/0294/002	53015.01.00	1 A PHARMA GMBH	DE
Moclobemid Sandoz 300 mg Filmtabletten	DK/H/0293/002	53013.01.00	HEXAL AG	DE
MOCLAMINE 150 mg, comprimé pelliculé sécable	not available	333 249-3	BIOCODEX	FR
MOCLAMINE 150 mg, comprimé pelliculé sécable	not available	333 250-1	BIOCODEX	FR
Zorix, 150 mg, comprimidos revestidos por película	not available	5393129	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
ZORIX 300 mg comprimidos revestidos por película	not available	5516786	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Zorix, 150 mg, comprimidos revestidos por película	not available	5393111	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
ZORIX 300 mg comprimidos revestidos por película	not available	5516687	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Moclobemide 150 mg film-coated tablet	not available	PL 11311/0493	TILLOMED LABORATORIES LTD	UK
Moclobemide 300 mg film-coated tablet	not available	PL 11311/0494	TILLOMED LABORATORIES LTD	UK
AURORIX® 150 mg - Filmtabletten	not available	1-18878	MEDA PHARMA GMBH	AT
AURORIX® 300 mg - Filmtabletten	not available	1-21026	MEDA PHARMA GMBH	AT
Aurorix 150 mg tabletter, filmdrasjerte	not available	7547	MEDA AS	NO
Aurorix 300 mg tabletter, filmdrasjerte	not available	7983	MEDA AS	NO
Aurorix 150 mg comprimés pelliculés	not available	BE157482	S.A. MEDA PHARMA N.V.	BE
Aurorix 150 mg filmomhulde tabletten	not available	BE157482	S.A. MEDA PHARMA 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
AURORIX 150 mg filmom obložene tablete	not available	HR-H-485452581	MEDICAL INTERTRADE D.O.O.	HR
Aurorix, 150 mg, tabletki powlekane	not available	R/0948	MEDA PHARMA GMBH & CO. KG	PL
Aurorix, 300 mg, tabletki powlekane	not available	R/4198	MEDA PHARMA GMBH & CO. KG	PL
АУРОРИКС 150 mg филмирани таблетки	not available	20010448	MEDA PHARMA GMBH & CO. KG	BG
Aurorix 300 mg filmtabletta	not available	OGYI-T-1809/03	MEDA PHARMA HUNGARY KFT.	HU
Aurorix 150 mg filmtabletta	not available	OGYI-T-1809/02	MEDA PHARMA HUNGARY KFT.	HU
Aurorix 150 mg töflur	not available	880154	MEDA AB	IS
Aurorix 300 mg töflur	not available	930243	MEDA AB	IS
Aurorix 150 mg tablett, filmdragerad	not available	10474	MEDA OY	FI
Aurorix 300 mg tablett, filmdragerad	not available	11544	MEDA OY	FI
Aurorix 150 mg Filmtabletten	not available	BE157482	S.A. MEDA PHARMA N.V.	BE
Aurorix 150 mg Filmtabletten	not available	2008/089875	S.A. MEDA PHARMA N.V.	LU
Manerix 150 mg, comprimidos recubiertos con película	not available	59.169	MEDA PHARMA S.L.	ES
Manerix 300 mg, comprimidos recubiertos con película.	not available	60.824	MEDA PHARMA S.L.	ES
Aurorix® 150; 150 mg Filmtablette	not available	37340.00.00	MEDA PHARMA GMBH & CO. KG	DE
Aurorix 150 mg comprimidos revestidos por película	not available	5831086	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Aurorix 150 mg comprimidos revestidos por película	not available	8757435	MEDA PHARMA – PRODUTOS FARMACÊUTICOS, S.A.	PT
Aurorix® 300; 300 mg Filmtablette	not available	37340.01.00	MEDA PHARMA GMBH & CO. KG	DE
Aurorix 150 mg filmsko obložene tablete	not available	H/93/00241/001	MEDA PHARMA GMBH & CO. KG	SI
Aurorix 300 mg filmsko obložene tablete	not available	H/93/00241/002	MEDA PHARMA GMBH & CO. KG	SI
Aurorix® 150 mg/F.C.Tab	not available	283/23-09-2011	MEDA PHARMACEUTICALS 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
Aurorix 150 mg tabletti, kalvopäällysteinen	not available	10474	MEDA OY	FI
Aurorix 150, tabletten 150 mg	not available	RVG 14904	MEDA PHARMA B.V.	NL
AURORIX 150 mg potahované tablety	not available	30/159/91 – A/C	MEDA PHARMA S.R.O.	CZ
Aurorix 150 mg comprimés pelliculés	not available	2008/089875	S.A. MEDA PHARMA N.V.	LU
Aurorix® 150 mg	not available	30/0159/91-C/S	MEDA PHARMA SPOL. S R.O.	SK
AURORIX 300 mg potahované tablety	not available	30/159/91 – B/C	MEDA PHARMA S.R.O.	CZ
Aurorix® 300 mg/F.C.Tab	not available	87855/12-12-2011	MEDA PHARMACEUTICALS S.A.	GR
Aurorix 150 mg tabletter	not available	11031	MEDA AB	SE
Aurorix 300 mg tabletti, kalvopäällysteinen	not available	11544	MEDA OY	FI
Aurorix 300, tabletten 300 mg	not available	RVG 17493	MEDA PHARMA B.V.	NL
Aurorix® 300 mg	not available	30/0159/91-C/S	MEDA PHARMA SPOL. S R.O.	SK
Aurorix 300 mg tabletter	not available	12051	MEDA AB	SE
Manerix 150 mg Film-coated Tablets	not available	PA 1332/28/1	MEDA HEALTH SALES IRELAND LIMITED	IE
Aurorix, filmovertrukne tabletter	not available	13354	MEDA AS	DK
Aurorix, filmovertrukne tabletter	not available	15758	MEDA AS	DK
MOCLAMINE 150 mg, comprimé pelliculé sécable	not available	333 249-3	BIOCODEX	FR
MOCLAMINE 150 mg, comprimé pelliculé sécable	not available	333 250-1	BIOCODEX	FR