



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

1 September 2017
EMA/641616/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: topiramate

Procedure no.: PSUSA/00002996/201701



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
Acomil 100 mg comprimidos recubiertos con película EFG	not available	69.661	ADAMED LABORATORIOS, S.L.U.	ES
Acomil 200 mg comprimidos recubiertos con película EFG	not available	69.662	ADAMED LABORATORIOS, S.L.U.	ES
Acomil 25 mg comprimidos recubiertos con película EFG	not available	69.659	ADAMED LABORATORIOS, S.L.U.	ES
Acomil 50 mg comprimidos recubiertos con película EFG	not available	69.660	ADAMED LABORATORIOS, S.L.U.	ES
ECURAM 100 mg compresse rivestite con film	AT/H/0612/003	039937115	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 100 mg compresse rivestite con film	AT/H/0612/003	039937127	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 100 mg compresse rivestite con film	AT/H/0612/003	039937139	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 100 mg compresse rivestite con film	AT/H/0612/003	039937141	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 100 mg compresse rivestite con film	AT/H/0612/003	039937154	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 200 mg compresse rivestite con film	AT/H/0612/004	039937166	BLUEFISH PHARMACEUTICALS AB	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
ECURAM 200 mg compresse rivestite con film	AT/H/0612/004	039937178	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 200 mg compresse rivestite con film	AT/H/0612/004	039937180	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 200 mg compresse rivestite con film	AT/H/0612/004	039937192	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 200 mg compresse rivestite con film	AT/H/0612/004	039937204	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937014	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937216	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937026	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937038	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937040	BLUEFISH PHARMACEUTICALS AB	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
ECURAM 25 mg compresse rivestite con film	AT/H/0612/001	039937053	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 50 mg compresse rivestite con film	AT/H/0612/002	039937065	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 50 mg compresse rivestite con film	AT/H/0612/002	039937077	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 50 mg compresse rivestite con film	AT/H/0612/002	039937089	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 50 mg compresse rivestite con film	AT/H/0612/002	039937091	BLUEFISH PHARMACEUTICALS AB	IT
ECURAM 50 mg compresse rivestite con film	AT/H/0612/002	039937103	BLUEFISH PHARMACEUTICALS AB	IT
Epiramat 100 mg filmom obložene tablete	not available	HR-H-575505055	PLIVA HRVATSKA D.O.O.	HR
Epiramat 200 mg filmom obložene tablete	not available	HR-H-907959213	PLIVA HRVATSKA D.O.O.	HR
Epiramat 25 mg filmom obložene tablete	not available	HR-H-363242016	PLIVA HRVATSKA D.O.O.	HR

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
Epiramat 50 mg filmom obložene tablete	not available	HR-H-856634477	PLIVA HRVATSKA D.O.O.	HR
Epiramat, 100 mg, tabletki powlekane	not available	12581	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Epiramat, 200 mg, tabletki powlekane	not available	12583	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Epiramat, 25 mg, tabletki powlekane	not available	12582	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
Epiramat, 50 mg, tabletki powlekane	not available	12580	TEVA PHARMACEUTICALS POLSKA SP. Z O.O.	PL
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	342 647-8	JANSSEN-CILAG	FR
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	342 645-5	JANSSEN-CILAG	FR
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	356 345-9	JANSSEN-CILAG	FR
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	342 648-4	JANSSEN-CILAG	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
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	342 646-1	JANSSEN-CILAG	FR
EPITOMAX 100 mg, comprimé pelliculé	SE/H/0110/003	356 344-2	JANSSEN-CILAG	FR
EPITOMAX 15 mg, gélule	SE/H/0110/007	348 109-8	JANSSEN-CILAG	FR
EPITOMAX 15 mg, gélule	SE/H/0110/007	348 305-1	JANSSEN-CILAG	FR
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	341 148-8	JANSSEN-CILAG	FR
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	341 149-4	JANSSEN-CILAG	FR
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	341 227-5	JANSSEN-CILAG	FR
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	356 347-1	JANSSEN-CILAG	FR
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	341 226-9	JANSSEN-CILAG	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
EPITOMAX 200 mg, comprimé pelliculé	SE/H/0110/004	356 346-5	JANSSEN-CILAG	FR
EPITOMAX 25 mg, gélule	SE/H/0110/008	348 110-6	JANSSEN-CILAG	FR
EPITOMAX 25 mg, gélule	SE/H/0110/008	348 306-8	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	341 147-1	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	356 343-6	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	341 146-5	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	341 224-6	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	341 225-2	JANSSEN-CILAG	FR
EPITOMAX 50 mg, comprimé pelliculé	SE/H/0110/002	356 341-3	JANSSEN-CILAG	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
EPITOMAX 50 mg, gélule	SE/H/0110/009	348 111-2	JANSSEN-CILAG	FR
EPITOMAX 50 mg, gélule	SE/H/0110/009	348 307-4	JANSSEN-CILAG	FR
Epitoram, 100 mg, tabletki powlekane	NL/H/1231/003	14665	APOTEX EUROPE B.V.	PL
Epitoram, 200 mg, tabletki powlekane	NL/H/1231/004	14664	APOTEX EUROPE B.V.	PL
Epitoram, 25 mg, tabletki powlekane	NL/H/1231/001	14667	APOTEX EUROPE B.V.	PL
Epitoram, 50 mg, tabletki powlekane	NL/H/1231/002	14666	APOTEX EUROPE B.V.	PL
Erravia 100 mg filmom obalené tablety	DK/H/0930/003	21/0512/08-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK
Erravia 200 mg filmom obalené tablety	DK/H/0930/004	21/0513/08-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK
Erravia 25 mg filmom obalené tablety	DK/H/0930/001	21/0510/08-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK

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
Erravia 50 mg filmom obalené tablety	DK/H/0930/002	21/0511/08-S	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	SK
Etopro 100 mg filmtabletta	DK/H/0930/003	OGYI-T- 20648/05	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 100 mg filmtabletta	DK/H/0930/003	OGYI-T- 20648/06	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 200 mg filmtabletta	DK/H/0930/004	OGYI-T- 20648/08	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 200 mg filmtabletta	DK/H/0930/004	OGYI-T- 20648/07	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 25 mg filmtabletta	DK/H/0930/001	OGYI-T- 20648/01	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 25 mg filmtabletta	DK/H/0930/001	OGYI-T- 20648/02	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 50 mg filmtabletta	DK/H/0930/002	OGYI-T- 20648/03	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	HU
Etopro 50 mg filmtabletta	DK/H/0930/002	OGYI-T- 20648/04	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	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
Etopro, 100 mg, tabletki powlekane	DK/H/0930/003	12895	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Etopro, 200 mg, tabletki powlekane	DK/H/0930/004	12896	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Etopro, 25 mg, tabletki powlekane	DK/H/0930/001	12893	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
Etopro, 50 mg, tabletki powlekane	DK/H/0930/002	12894	PHARMASWISS ČESKÁ REPUBLIKA S.R.O.	PL
FAGODOL 25 mg comprimidos recubiertos con película EFG	not available	69001	ARAFARMA GROUP, S.A	ES
FAGODOL 25 mg comprimidos recubiertos con película EFG	not available	69001	ARAFARMA GROUP, S.A	ES
FAGODOL100 mg comprimidos recubiertos con película EFG	not available	69003	ARAFARMA GROUP, S.A	ES
FAGODOL100 mg comprimidos recubiertos con película EFG	not available	69003	ARAFARMA GROUP, S.A	ES
FAGODOL200 mg comprimidos recubiertos con película EFG	not available	69004	ARAFARMA GROUP, 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
FAGODOL200 mg comprimidos recubiertos con película EFG	not available	69004	ARAFARMA GROUP, S.A	ES
FAGODOL50 mg comprimidos recubiertos con película EFG	not available	69002	ARAFARMA GROUP, S.A	ES
FAGODOL50 mg comprimidos recubiertos con película EFG	not available	69002	ARAFARMA GROUP, S.A	ES
JADIX® 100mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/0199/003	89042/ 28-12-2010	PHARMAZAC SA	GR
JADIX® 200mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/0199/004	89043/ 28-12-2010	PHARMAZAC SA	GR
JADIX® 25mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/0199/001	50711/ 29-12-2010	PHARMAZAC SA	GR
JADIX® 50mg Επικαλυμμένα με λεπτό υμένιο δισκία	PT/H/0199/002	89039/ 28-12-2010	PHARMAZAC SA	GR
LUSITRAX 100 mg film-coated tablets	PT/H/0198/003	6591/2014/01-03	ROMASTRU TRADING SRL	RO
LUSITRAX 200 mg film-coated tablets	PT/H/0198/004	6592/2014/01-03	ROMASTRU TRADING SRL	RO

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
LUSITRAX 25 mg film-coated tablets	PT/H/0198/001	6589/2014/01-03	ROMASTRU TRADING SRL	RO
LUSITRAX 50 mg film-coated tablets	PT/H/0198/002	6590/2014/01-03	ROMASTRU TRADING SRL	RO
Maritop, filmovertrokne tabletter	DK/H/1354/001	41827	G.L. PHARMA GMBH	DK
Maritop, filmovertrokne tabletter	DK/H/1354/002	41828	G.L. PHARMA GMBH	DK
Maritop, filmovertrokne tabletter	DK/H/1354/003	41829	G.L. PHARMA GMBH	DK
Maritop, filmovertrokne tabletter	DK/H/1354/004	41830	G.L. PHARMA GMBH	DK
Oritop, 100 mg, tabletki powlekane	FI/H/0800/003	15634	ORION CORPORATION	PL
Oritop, 25 mg, tabletki powlekane	FI/H/0800/001	15632	ORION CORPORATION	PL
Oritop, 50 mg, tabletki powlekane	FI/H/0800/002	15633	ORION CORPORATION	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
Piraleps 100 mg Film-coated tablets	not available	ML 21350	DELORBIS PHARMACEUTICALS LTD	CY
Piraleps 200 mg Film-coated tablets	not available	ML 21351	DELORBIS PHARMACEUTICALS LTD	CY
Piraleps 25 mg Film-coated tablets	not available	ML 21348	DELORBIS PHARMACEUTICALS LTD	CY
Piraleps 50 mg Film-coated tablets	not available	ML 21349	DELORBIS PHARMACEUTICALS LTD	CY
Pirantal 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/1355/003	59454_26-07-2016	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
Pirantal 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/1355/004	59455_26-07-2016	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
Pirantal 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/1355/001	59452_26-07-2016	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
Pirantal 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/1355/002	59453_26-07-2016	PHARMANEL COMMERCIAL PHARMACEUTICAL S.A.	GR
Pirepil 100 mg comprimidos revestidos por película	not available	5138821	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
Pirepil 100 mg comprimidos revestidos por película	not available	5138839	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 100 mg comprimidos revestidos por película	not available	5138847	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 100 mg comprimidos revestidos por película	not available	5138813	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 200 mg comprimidos revestidos por película	not available	5138862	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 200 mg comprimidos revestidos por película	not available	5138870	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 200 mg comprimidos revestidos por película	not available	5138904	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 200 mg comprimidos revestidos por película	not available	5138854	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 25 mg comprimidos revestidos por película	not available	5138722	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 25 mg comprimidos revestidos por película	not available	5138730	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
Pirepil 25 mg comprimidos revestidos por película	not available	5138748	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 25 mg comprimidos revestidos por película	not available	5138714	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 50 mg comprimidos revestidos por película	not available	5138763	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 50 mg comprimidos revestidos por película	not available	5138771	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 50 mg comprimidos revestidos por película	not available	5138805	DECOMED FARMACÊUTICA, S.A.	PT
Pirepil 50 mg comprimidos revestidos por película	not available	5138755	DECOMED FARMACÊUTICA, S.A.	PT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707205	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707229	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707395	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
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707559	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707585	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707217	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707193	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707231	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707546	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707561	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707573	SANDOZ S.P.A.	IT
Ramas 100 mg compresse rivestite con film	FI/H/0599/003	037707534	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
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707623	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707282	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707294	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707306	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707320	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707407	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707597	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707611	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707318	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
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707609	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707635	SANDOZ S.P.A.	IT
Ramas 200 mg compresse rivestite con film	FI/H/0599/004	037707647	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707054	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707371	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707421	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707433	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707027	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707039	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
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707460	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707041	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707015	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707419	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707445	SANDOZ S.P.A.	IT
Ramas 25 mg compresse rivestite con film	FI/H/0599/001	037707458	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707104	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707130	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707508	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
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707116	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707142	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707472	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707510	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707522	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707128	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707496	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707484	SANDOZ S.P.A.	IT
Ramas 50 mg compresse rivestite con film	FI/H/0599/002	037707383	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
SINCRONIL 100 mg compresse rivestite con film	DK/H/1355/003	039469085	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 100 mg compresse rivestite con film	DK/H/1355/003	039469097	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 100 mg compresse rivestite con film	DK/H/1355/001-004/MR	039469073	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 200 mg compresse rivestite con film	DK/H/1355/004	039469111	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 200 mg compresse rivestite con film	DK/H/1355/004	039469123	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 200 mg compresse rivestite con film	DK/H/1355/001-004/MR	039469109	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 25 mg compresse rivestite con film	DK/H/1355/001	039469022	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 25 mg compresse rivestite con film	DK/H/1355/001	039469034	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 25 mg compresse rivestite con film	DK/H/1355/001-004/MR	039469010	FIDIA 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
SINCRONIL 50 mg compresse rivestite con film	DK/H/1355/002	039469059	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 50 mg compresse rivestite con film	DK/H/1355/002	039469061	FIDIA FARMACEUTICI S.P.A	IT
SINCRONIL 50 mg compresse rivestite con film	DK/H/1355/001-004/MR	039469046	FIDIA FARMACEUTICI S.P.A	IT
Talopam 100 mg film-coated tablets	PT/H/0231/003	MA628/06803	ACTAVIS GROUP PTC EHF.	MT
Talopam 100 mg filmom obalené tablety	PT/H/0231/003	21/0757/09-S	ACTAVIS GROUP PTC EHF.	SK
Talopam 200 mg film-coated tablets	PT/H/0231/004	MA628/06804	ACTAVIS GROUP PTC EHF.	MT
Talopam 25 mg film-coated tablets	PT/H/0231/001	MA628/06801	ACTAVIS GROUP PTC EHF.	MT
Talopam 25 mg filmom obalené tablety	PT/H/0231/001	21/0755/09-S	ACTAVIS GROUP PTC EHF.	SK
Talopam 50 mg film-coated tablets	PT/H/0231/002	MA628/06802	ACTAVIS GROUP PTC EHF.	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Talopam 50 mg filmom obalené tablety	PT/H/0231/002	21/0756/09-S	ACTAVIS GROUP PTC EHF.	SK
TIRAMAT 100 mg filmom obložene tablete	not available	UP/I-530-09/12-02/194	BELUPO D.D.	HR
TIRAMAT 25 mg filmom obložene tablete	not available	UP/I-530-09/12-02/192	BELUPO D.D.	HR
TIRAMAT 50 mg filmom obložene tablete	not available	UP/I-530-09/12-02/193	BELUPO D.D.	HR
Topaben 100 mg filmovertrukne tabletter	DK/H/1355/003	41833	PHARMATHEN S.A.	DK
Topaben 100 mg filmovertrukne tabletter	DK/H/1355/003	41833	PHARMATHEN S.A.	DK
Topaben 200 mg filmovertrukne tabletter	DK/H/1355/004	41834	PHARMATHEN S.A.	DK
Topaben 200 mg filmovertrukne tabletter	DK/H/1355/004	41834	PHARMATHEN S.A.	DK
Topaben 25 mg filmovertrukne tabletter	DK/H/1355/001	41831	PHARMATHEN S.A.	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
Topaben 25 mg filmovertrukne tabletter	DK/H/1355/001	41831	PHARMATHEN S.A.	DK
Topaben 50 mg filmovertrukne tabletter	DK/H/1355/002	41832	PHARMATHEN S.A.	DK
Topaben 50 mg filmovertrukne tabletter	DK/H/1355/002	41832	PHARMATHEN S.A.	DK
Topamac 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/003	36170/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Topamac 15 mg καψάκια, σκληρά	SE/H/0110/007	36172/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Topamac 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/004	36179/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Topamac 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/001	36176/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Topamac 25 mg καψάκια, σκληρά	SE/H/0110/008	36173/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
Topamac 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/002	36178/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	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
Topamac 50 mg καψάκια, σκληρά	SE/H/0110/009	36175/26.05.2015	JANSSEN-CILAG PHARMACEUTICAL S.A.C.I.	GR
TOPAMAX	SE/H/0110/007	018/00305	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/009	018/00307	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/003	018/00303	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/008	018/00306	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/001	018/00301	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/004	018/00304	JANSSEN-CILAG INTERNATIONAL NV	MT
TOPAMAX	SE/H/0110/002	018/00302	JANSSEN-CILAG INTERNATIONAL NV	MT
Topamax 100	SE/H/0110/003	21/0121/11-S	JOHNSON & JOHNSON, S.R.O	SK

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
Topamax 100 mg apvalkotās tabletes	SE/H/0110/003	00-0122	UAB JOHNSON & JOHNSON	LV
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023552	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023285	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023261	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023741	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023549	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023828	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023273	JANSSEN-CILAG SPA	IT
Topamax 100 mg comprese rivestite con film	SE/H/0110/003	032023501	JANSSEN-CILAG SPA	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
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023537	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023246	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023499	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023525	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023780	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023513	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023259	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023234	JANSSEN-CILAG SPA	IT
Topamax 100 mg compresse rivestite con film	SE/H/0110/003	032023032	JANSSEN-CILAG SPA	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
Topamax 100 mg comprimate filmate	SE/H/0110/003	4506/2012/01-17	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	RO
Topamax 100 mg comprimés pelliculés	SE/H/0110/003	1997120543	JANSSEN-CILAG NV	LU
Topamax 100 mg comprimés pelliculés	SE/H/0110/003	0162/97/12/0543	JANSSEN-CILAG NV	LU
Topamax 100 mg comprimidos recubiertos con película	SE/H/0110/003	61.875	JANSSEN-CILAG S.A.	ES
Topamax 100 mg comprimidos revestidos por película	SE/H/0110/003	3481298	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 100 mg comprimidos revestidos por película	SE/H/0110/003	5320197	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 100 mg comprimidos revestidos por película	SE/H/0110/003	3481199	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 100 mg film-coated tablets	SE/H/0110/003	PA 0748/012/003	JANSSEN-CILAG LIMITED	IE
Topamax 100 mg filmom obložene tablete	not available	HR-H-839682871	JOHNSON & JOHNSON S.E. D.O.O.	HR

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
Topamax 100 mg filmomhulde tabletten	SE/H/0110/003	BE184371	JANSSEN-CILAG NV	BE
Topamax 100 mg filmomhulde tabletten	SE/H/0110/003	BE398596	JANSSEN-CILAG NV	BE
Topamax 100 mg filmtabletta	SE/H/0110/003	OGYI-T-6268/08	JANSSEN-CILAG KFT.	HU
Topamax 100 mg filmtabletta	SE/H/0110/003	OGYI-T-6268/07	JANSSEN-CILAG KFT.	HU
Topamax 100 mg Filmtabletten	SE/H/0110/003	1-22142	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 100 mg Filmtabletten	SE/H/0110/003	35027.02.00	JANSSEN-CILAG GMBH	DE
Topamax 100 mg plėvele dengtos tabletės	SE/H/0110/003	LT/1/98/0093/003	UAB JOHNSON & JOHNSON	LT
Topamax 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/003	18275	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax 100 mg potahované tablety	SE/H/0110/003	21/512/97-C	JANSSEN-CILAG S.R.O	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
Topamax 100 mg, filmomhulde tabletten	SE/H/0110/003	RVG 24167	JANSSEN-CILAG BV	NL
Topamax 15 mg cápsulas	SE/H/0110/007	3221595	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 15 mg cápsulas	SE/H/0110/007	3221496	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 15 mg capsule rigide	SE/H/0110/007	032023640	JANSSEN-CILAG SPA	IT
Topamax 15 mg capsule rigide	SE/H/0110/007	032023638	JANSSEN-CILAG SPA	IT
Topamax 15 mg capsule rigide	SE/H/0110/007	032023653	JANSSEN-CILAG SPA	IT
Topamax 15 mg capsule rigide	SE/H/0110/007	032023083	JANSSEN-CILAG SPA	IT
Topamax 15 mg gélules	SE/H/0110/007	0162/00/01/0043	JANSSEN-CILAG NV	LU
Topamax 15 mg Granulat in Kapseln	SE/H/0110/007	1-23233	JANSSEN-CILAG 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
Topamax 15 mg harde capsules	SE/H/0110/007	BE207426	JANSSEN-CILAG NV	BE
Topamax 15 mg καψάκια, σκληρά	SE/H/0110/007	19558	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax 15 mg, tvrdé tobolky	SE/H/0110/007	21/653/99-C	JANSSEN-CILAG S.R.O	CZ
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023311	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023335	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023754	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023626	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023588	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023576	JANSSEN-CILAG SPA	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
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023297	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023830	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023347	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023614	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023792	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023590	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023309	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023323	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023602	JANSSEN-CILAG SPA	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
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023564	JANSSEN-CILAG SPA	IT
Topamax 200 mg compresse rivestite con film	SE/H/0110/004	032023044	JANSSEN-CILAG SPA	IT
Topamax 200 mg comprimés pelliculés	SE/H/0110/004	0162/97/12/0544	JANSSEN-CILAG NV	LU
Topamax 200 mg comprimés pelliculés	SE/H/0110/004	1997120544	JANSSEN-CILAG NV	LU
Topamax 200 mg comprimidos recubiertos con película	SE/H/0110/004	61.876	JANSSEN-CILAG S.A.	ES
Topamax 200 mg comprimidos revestidos por película	SE/H/0110/004	3481496	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 200 mg comprimidos revestidos por película	SE/H/0110/004	3481397	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 200 mg film-coated tablets	SE/H/0110/004	PA 0748/012/004	JANSSEN-CILAG LIMITED	IE
Topamax 200 mg filmom obložene tablete	not available	HR-H-668959661	JOHNSON & JOHNSON S.E. D.O.O.	HR

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
Topamax 200 mg filmomhulde tabletten	SE/H/0110/004	BE184362	JANSSEN-CILAG NV	BE
Topamax 200 mg filmomhulde tabletten	SE/H/0110/004	BE398605	JANSSEN-CILAG NV	BE
Topamax 200 mg filmsko obložene tablete	SE/H/0110/004	H/97/01540/056	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	SI
Topamax 200 mg filmtabletta	SE/H/0110/004	OGYI-T-6268/09	JANSSEN-CILAG KFT.	HU
Topamax 200 mg filmtabletta	SE/H/0110/004	OGYI-T-6268/10	JANSSEN-CILAG KFT.	HU
Topamax 200 mg Filmtabletten	SE/H/0110/004	1-22143	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 200 mg Filmtabletten	SE/H/0110/004	35027.03.00	JANSSEN-CILAG GMBH	DE
Topamax 200 mg plėvele dengtos tabletės	SE/H/0110/004	LT/1/98/0093/004	UAB JOHNSON & JOHNSON	LT
Topamax 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/004	18278	JANSSEN-CILAG INTERNATIONAL NV	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
Topamax 200 mg, filmomhulde tabletten	SE/H/0110/004	RVG 24168	JANSSEN-CILAG BV	NL
Topamax 25	SE/H/0110/001	21/0108/98-S	JOHNSON & JOHNSON, S.R.O	SK
Topamax 25 mg	SE/H/0110/001	21/510/97-C	JANSSEN-CILAG S.R.O	CZ
Topamax 25 mg	SE/H/0110/008	21/654/99-C	JANSSEN-CILAG S.R.O	CZ
Topamax 25 mg apvalkotās tabletes	SE/H/0110/001	00-0120	UAB JOHNSON & JOHNSON	LV
Topamax 25 mg cápsulas	SE/H/0110/008	3221694	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 25 mg capsule rigide	SE/H/0110/008	032023677	JANSSEN-CILAG SPA	IT
Topamax 25 mg capsule rigide	SE/H/0110/008	032023689	JANSSEN-CILAG SPA	IT
Topamax 25 mg capsule rigide	SE/H/0110/008	032023665	JANSSEN-CILAG SPA	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
Topamax 25 mg capsule rigide	SE/H/0110/008	032023095	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023362	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023400	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023133	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023412	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023158	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023727	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023804	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023121	JANSSEN-CILAG SPA	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
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023160	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023386	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023350	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023374	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023766	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023119	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023398	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023145	JANSSEN-CILAG SPA	IT
Topamax 25 mg compresse rivestite con film	SE/H/0110/001	032023071	JANSSEN-CILAG SPA	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
Topamax 25 mg comprimate filmate	SE/H/0110/001	4504/2012/01-17	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	RO
Topamax 25 mg comprimés pelliculés	SE/H/0110/001	0162/97/12/0591	JANSSEN-CILAG NV	LU
Topamax 25 mg comprimés pelliculés	SE/H/0110/001	1997120591	JANSSEN-CILAG NV	LU
Topamax 25 mg comprimidos recubiertos con película	SE/H/0110/001	61.873	JANSSEN-CILAG S.A.	ES
Topamax 25 mg comprimidos revestidos por película	SE/H/0110/001	3480894	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 25 mg comprimidos revestidos por película	SE/H/0110/001	3480795	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 25 mg film-coated tablets	SE/H/0110/001	PA 0748/012/001	JANSSEN-CILAG LIMITED	IE
Topamax 25 mg filmom obložene tablete	not available	HR-H-239773821	JOHNSON & JOHNSON S.E. D.O.O.	HR
Topamax 25 mg filmomhulde tabletten	SE/H/0110/001	BE398571	JANSSEN-CILAG NV	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
Topamax 25 mg filmomhulde tabletten	SE/H/0110/001	BE184405	JANSSEN-CILAG NV	BE
Topamax 25 mg filmtabletta	SE/H/0110/001	OGYI-T-6268/03	JANSSEN-CILAG KFT.	HU
Topamax 25 mg filmtabletta	SE/H/0110/001	OGYI-T-6268/04	JANSSEN-CILAG KFT.	HU
Topamax 25 mg Filmtabletten	SE/H/0110/001	1-22139	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 25 mg Filmtabletten	SE/H/0110/001	35027.00.00	JANSSEN-CILAG GMBH	DE
Topamax 25 mg gélules	SE/H/0110/008	0162/00/01/0044	JANSSEN-CILAG NV	LU
Topamax 25 mg Granulat in Kapseln	SE/H/0110/008	1-23234	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 25 mg harde capsules	SE/H/0110/008	BE207435	JANSSEN-CILAG NV	BE
Topamax 25 mg Hartkapseln	SE/H/0110/008	35033.00.00	JANSSEN-CILAG 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
Topamax 25 mg plèvele dengtos tabletės	SE/H/0110/001	LT/1/98/0093/001	UAB JOHNSON & JOHNSON	LT
Topamax 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/001	18277	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax 25 mg καψάκια σκληρά	SE/H/0110/008	19576	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax 25 mg, filmomhulde tabletten	SE/H/0110/001	RVG 24165	JANSSEN-CILAG BV	NL
Topamax 50	SE/H/0110/002	21/0120/11-S	JOHNSON & JOHNSON, S.R.O	SK
Topamax 50 mg	SE/H/0110/002	21/511/97-C	JANSSEN-CILAG S.R.O	CZ
Topamax 50 mg apvalkotās tabletes	SE/H/0110/002	00-0121	UAB JOHNSON & JOHNSON	LV
Topamax 50 mg cápsulas	SE/H/0110/009	3221793	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 50 mg capsule rigide	SE/H/0110/009	032023703	JANSSEN-CILAG SPA	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
Topamax 50 mg capsule rigide	SE/H/0110/009	032023691	JANSSEN-CILAG SPA	IT
Topamax 50 mg capsule rigide	SE/H/0110/009	032023715	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023222	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023196	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023816	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023463	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023739	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023487	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023448	JANSSEN-CILAG SPA	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
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023451	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023475	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023208	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023778	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023210	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023184	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023172	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023424	JANSSEN-CILAG SPA	IT
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023436	JANSSEN-CILAG SPA	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
Topamax 50 mg compresse rivestite con film	SE/H/0110/002	032023020	JANSSEN-CILAG SPA	IT
Topamax 50 mg comprimate filmate	SE/H/0110/002	4505/2012/01-17	JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O.	RO
Topamax 50 mg comprimés pelliculés	SE/H/0110/002	1997120542	JANSSEN-CILAG NV	LU
Topamax 50 mg comprimés pelliculés	SE/H/0110/002	0162/97/12/0542	JANSSEN-CILAG NV	LU
Topamax 50 mg comprimidos recubiertos con película	SE/H/0110/002	61.874	JANSSEN-CILAG S.A.	ES
Topamax 50 mg comprimidos revestidos por película	SE/H/0110/002	3481090	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 50 mg comprimidos revestidos por película	SE/H/0110/002	3480993	JANSSEN-CILAG FARMACÊUTICA, LDA.	PT
Topamax 50 mg film-coated tablets	SE/H/0110/002	PA 0748/012/002	JANSSEN-CILAG LIMITED	IE
Topamax 50 mg filmom obložene tablete	not available	HR-H-610277393	JOHNSON & JOHNSON S.E. D.O.O.	HR

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
Topamax 50 mg filmomhulde tabletten	SE/H/0110/002	BE184387	JANSSEN-CILAG NV	BE
Topamax 50 mg filmomhulde tabletten	SE/H/0110/002	BE398587	JANSSEN-CILAG NV	BE
Topamax 50 mg filmtabletta	SE/H/0110/002	OGYI-T-6268/06	JANSSEN-CILAG KFT.	HU
Topamax 50 mg filmtabletta	SE/H/0110/002	OGYI-T-6268/05	JANSSEN-CILAG KFT.	HU
Topamax 50 mg Filmtabletten	SE/H/0110/002	1-22141	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 50 mg Filmtabletten	SE/H/0110/002	35027.01.00	JANSSEN-CILAG GMBH	DE
Topamax 50 mg gélules	SE/H/0110/009	0162/00/01/0045	JANSSEN-CILAG NV	LU
Topamax 50 mg Granulat in Kapseln	SE/H/0110/009	1-23235	JANSSEN-CILAG PHARMA GMBH	AT
Topamax 50 mg harde capsules	SE/H/0110/009	BE207444	JANSSEN-CILAG NV	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
Topamax 50 mg Hartkapseln	SE/H/0110/009	35033.01.00	JANSSEN-CILAG GMBH	DE
Topamax 50 mg plévele dengtos tabletės	SE/H/0110/002	LT/1/98/0093/002	UAB JOHNSON & JOHNSON	LT
Topamax 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	SE/H/0110/002	18276	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax 50 mg καψάκια σκληρά	SE/H/0110/009	19577	JANSSEN-CILAG INTERNATIONAL NV	CY
Topamax dispersable 15 mg cápsulas duras	SE/H/0110/007	63.957	JANSSEN-CILAG S.A.	ES
Topamax dispersable 25 mg cápsulas duras	SE/H/0110/008	63.959	JANSSEN-CILAG S.A.	ES
Topamax dispersable 50 mg cápsulas duras	SE/H/0110/009	63.960	JANSSEN-CILAG S.A.	ES
Topamax Sprinkle 15 mg hard capsules	SE/H/0110/007	PA 0748/012/007	JANSSEN-CILAG LIMITED	IE
Topamax Sprinkle 15 mg, harde capsules	SE/H/0110/007	RVG 25836	JANSSEN-CILAG BV	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
Topamax Sprinkle 25 mg, harde capsules	SE/H/0110/008	RVG 25837	JANSSEN-CILAG BV	NL
Topamax Sprinkle 50 mg, harde capsules	SE/H/0110/009	RVG 25838	JANSSEN-CILAG BV	NL
TOPAMAX, 100 mg õhukese polümeerikattega tabletid	SE/H/0110/003	314400	UAB JOHNSON & JOHNSON	EE
Topamax, 100 mg, tabletki powlekane	SE/H/0110/003	7590	JANSSEN-CILAG INTERNATIONAL NV	PL
Topamax, 15 mg, kapsułki twarde	SE/H/0110/007	8615	JANSSEN-CILAG INTERNATIONAL NV	PL
TOPAMAX, 200 mg õhukese polümeerikattega tabletid	SE/H/0110/004	314100	UAB JOHNSON & JOHNSON	EE
Topamax, 200 mg, tabletki powlekane	SE/H/0110/004	7591	JANSSEN-CILAG INTERNATIONAL NV	PL
TOPAMAX, 25 mg õhukese polümeerikattega tabletid	SE/H/0110/001	314500	UAB JOHNSON & JOHNSON	EE
Topamax, 25 mg, kapsułki twarde	SE/H/0110/008	8616	JANSSEN-CILAG INTERNATIONAL NV	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
Topamax, 25 mg, tabletki powlekane	SE/H/0110/001	7588	JANSSEN-CILAG INTERNATIONAL NV	PL
TOPAMAX, 50 mg õhukese polümeerikatttega tabletid	SE/H/0110/002	314300	UAB JOHNSON & JOHNSON	EE
Topamax, 50 mg, tabletki powlekane	SE/H/0110/002	7589	JANSSEN-CILAG INTERNATIONAL NV	PL
Topamax® 100 mg film-coated tablets	SE/H/0110/003	PL 00242/0303	JANSSEN-CILAG LIMITED	UK
Topamax® 200 mg film-coated tablets	SE/H/0110/004	PL 00242/0304	JANSSEN-CILAG LIMITED	UK
Topamax® 25 mg film-coated tablets	SE/H/0110/001	PL 00242/0301	JANSSEN-CILAG LIMITED	UK
Topamax® 50 mg film-coated tablets	SE/H/0110/002	PL 00242/0302	JANSSEN-CILAG LIMITED	UK
Topamax® Sprinkle 15 mg hard capsules	SE/H/0110/007	PL 00242/0348	JANSSEN-CILAG LIMITED	UK
Topamax® Sprinkle 25 mg hard capsules	SE/H/0110/008	PA 0748/012/008	JANSSEN-CILAG LIMITED	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
Topamax® Sprinkle 25 mg hard capsules	SE/H/0110/008	PL 00242/0349	JANSSEN-CILAG LIMITED	UK
Topamax® Sprinkle 50 mg hard capsules	SE/H/0110/009	PA 0748/012/009	JANSSEN-CILAG LIMITED	IE
Topamax® Sprinkle 50 mg hard capsules	SE/H/0110/009	PL 00242/0350	JANSSEN-CILAG LIMITED	UK
Topepsil 100 mg filmtabletta	NL/H/0863/003/DC	OGYI-T-20604/01-04	EGIS PHARMACEUTICALS PLC	HU
Topepsil 200 mg filmom obalené tablety	NL/H/0864/004/DC	21/0377/07-S	EGIS PHARMACEUTICALS PLC	SK
Topepsil 200 mg filmtabletta	NL/H/0863/004/DC	OGYI-T-20604/05-08	EGIS PHARMACEUTICALS PLC	HU
Topepsil 100 mg filmom obalené tablety	NL/H/0864/003	21/0376/07-S	EGIS PHARMACEUTICALS PLC	SK
TOPIBRAIN 100 mg comprimidos recubiertos con película	not available	68666	FERRER INTERNACIONAL, S.A.	ES
TOPIBRAIN 200 mg comprimidos recubiertos con película	not available	68676	FERRER INTERNACIONAL, 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
TOPIBRAIN 25 mg comprimidos recubiertos con película	not available	68664	FERRER INTERNACIONAL, S.A.	ES
TOPIBRAIN 50 mg comprimidos recubiertos con película	not available	68665	FERRER INTERNACIONAL, S.A.	ES
TopiLEK 100, 100 mg, tabletki powlekane	FI/H/0599/003	12684	SANDOZ GMBH	PL
TopiLEK 200, 200 mg, tabletki powlekane	FI/H/0599/004	12685	SANDOZ GMBH	PL
TopiLEK 25, 25 mg, tabletki powlekane	FI/H/0599/001	12680	SANDOZ GMBH	PL
TopiLEK 50, 50 mg, tabletki powlekane	FI/H/0599/002	12683	SANDOZ GMBH	PL
Topilept 100 mg filmomhulde tabletten	NL/H/864/003-004/DC	RVG 34255	EGIS PHARMACEUTICALS PLC	NL
Topilept 200 mg filmomhulde tabletten	NL/H/864/003-004/DC	RVG 34256	EGIS PHARMACEUTICALS PLC	NL
Topilex 100 mg	DK/H/1354/003	21/0402/08-S	G.L. PHARMA GMBH	SK

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
Topilex 100 mg comprimatate filmate	DK/H/1354/003	5531/2013/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 100 mg comprimatate filmate	DK/H/1354/003	5531/2013/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 100 mg comprimatate filmate	DK/H/1354/003	5531/2013/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 100 mg comprimatate filmate	DK/H/1354/003	5531/2013/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 100 mg potahované tablety	DK/H/1354/003	21/456/08-C	G.L. PHARMA GMBH	CZ
Topilex 100 mg- Filmtabletten	DK/H/1354/003	1-27686	G.L. PHARMA GMBH	AT
Topilex 200 mg	DK/H/1354/004	21/0403/08-S	G.L. PHARMA GMBH	SK
Topilex 200 mg comprimatate filmate	DK/H/1354/004	5532/2013/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 200 mg comprimatate filmate	DK/H/1354/004	5532/2013/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO

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
Topilex 200 mg comprimate filmate	DK/H/1354/004	5532/2013/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 200 mg comprimate filmate	DK/H/1354/004	5532/2013/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 200 mg- Filmtabletten	DK/H/1354/004	1-27687	G.L. PHARMA GMBH	AT
Topilex 25 mg	DK/H/1354/001	21/0400/08-S	G.L. PHARMA GMBH	SK
Topilex 25 mg comprimate filmate	DK/H/1354/001	5529/2013/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 25 mg comprimate filmate	DK/H/1354/001	5529/2013/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 25 mg comprimate filmate	DK/H/1354/001	5529/2013/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 25 mg comprimate filmate	DK/H/1354/001	5529/2013/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 25 mg potahované tablety	DK/H/1354/001	21/454/08-C	G.L. PHARMA GMBH	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
Topilex 25 mg- Filmtabletten	DK/H/1354/001	1-27684	G.L. PHARMA GMBH	AT
Topilex 50 mg	DK/H/1354/002	21/0401/08-S	G.L. PHARMA GMBH	SK
Topilex 50 mg comprimatate filmate	DK/H/1354/002	5530/2013/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 50 mg comprimatate filmate	DK/H/1354/002	5530/2013/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 50 mg comprimatate filmate	DK/H/1354/002	5530/2013/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 50 mg comprimatate filmate	DK/H/1354/002	5530/2013/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Topilex 50 mg potahované tablety	DK/H/1354/002	21/455/08-C	G.L. PHARMA GMBH	CZ
Topilex 50 mg- Filmtabletten	DK/H/1354/002	1-27685	G.L. PHARMA GMBH	AT
Topimark 100 mg filmom obalené tablety	DE/H/0984/003	21/0187/08-S	GLENMARK PHARMACEUTICALS S.R.O.	SK

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
Topimark 100 mg potahované tablety	DE/H/0984/003	21/245/08-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Topimark 200 mg filmom obalené tablety	DE/H/0984/004	21/0188/08-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Topimark 25 mg filmom obalené tablety	DE/H/0984/001	21/0185/08-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Topimark 25 mg potahované tablety	DE/H/0984/001	21/243/08-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Topimark 50 mg filmom obalené tablety	DE/H/0984/002	21/0186/08-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
TOPIMARK 50 MG POTAHOVANÉ TABLETY	DE/H/0984/002	21/244/08-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Topimax 100 mg filmdragerade tabletter	SE/H/0110/003	12439	JANSSEN-CILAG AB	SE
Topimax 100 mg filmovertrukne tabletter	SE/H/0110/003	17269	JANSSEN-CILAG A/S	DK
Topimax 100 mg filmuhúðaðar töflur	SE/H/0110/003	950035	JANSSEN-CILAG AB	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
Topimax 100 mg tabletter, filmdrasjerte	SE/H/0110/003	95-792	JANSSEN-CILAG A/S	NO
Topimax 100 mg tabletti, kalvopäällysteinen	SE/H/0110/003	11953	JANSSEN-CILAG OY	FI
Topimax 15 mg hörð hylki	SE/H/0110/007	980529	JANSSEN-CILAG AB	IS
Topimax 15 mg kapseli, kova	SE/H/0110/007	16128	JANSSEN-CILAG OY	FI
Topimax 15 mg kapslar, hårda	SE/H/0110/007	14348	JANSSEN-CILAG AB	SE
Topimax 15 mg kapsler, harde	SE/H/0110/007	98/3715	JANSSEN-CILAG A/S	NO
Topimax 15 mg kapsler, hårde	SE/H/0110/007	19589	JANSSEN-CILAG A/S	DK
Topimax 200 mg filmdragerade tabletter	SE/H/0110/004	12440	JANSSEN-CILAG AB	SE
Topimax 200 mg filmovertrukne tabletter	SE/H/0110/004	17270	JANSSEN-CILAG 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
Topimax 200 mg filmuhúðaðar töflur	SE/H/0110/004	950036	JANSSEN-CILAG AB	IS
Topimax 200 mg tabletter, filmdrasjerte	SE/H/0110/004	95-793	JANSSEN-CILAG A/S	NO
Topimax 200 mg tabletti, kalvopäällysteinen	SE/H/0110/004	11954	JANSSEN-CILAG OY	FI
Topimax 25 mg filmdragerade tabletter	SE/H/0110/001	12437	JANSSEN-CILAG AB	SE
Topimax 25 mg filmovertrukne tabletter	SE/H/0110/001	17267	JANSSEN-CILAG A/S	DK
Topimax 25 mg filmuhúðaðar töflur	SE/H/0110/001	950033	JANSSEN-CILAG AB	IS
Topimax 25 mg hörð hylki	SE/H/0110/008	980530	JANSSEN-CILAG AB	IS
Topimax 25 mg kapseli, kova	SE/H/0110/008	16129	JANSSEN-CILAG OY	FI
Topimax 25 mg kapslar, hárda	SE/H/0110/008	14349	JANSSEN-CILAG AB	SE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topimax 25 mg kapsler, harde	SE/H/0110/008	98/3716	JANSSEN-CILAG A/S	NO
Topimax 25 mg kapsler, hårde	SE/H/0110/008	19590	JANSSEN-CILAG A/S	DK
Topimax 25 mg tabletter, filmdrasjerte	SE/H/0110/001	95-790	JANSSEN-CILAG A/S	NO
Topimax 25 mg tabletti, kalvopäällysteinen	SE/H/0110/001	11951	JANSSEN-CILAG OY	FI
Topimax 50 mg filmdragerade tabletter	SE/H/0110/002	12438	JANSSEN-CILAG AB	SE
Topimax 50 mg filmovertrukne tabletter	SE/H/0110/002	17268	JANSSEN-CILAG A/S	DK
Topimax 50 mg filmuhúðaðar töflur	SE/H/0110/002	950034	JANSSEN-CILAG AB	IS
Topimax 50 mg hörð hylki	SE/H/0110/009	980531	JANSSEN-CILAG AB	IS
Topimax 50 mg kapseli, kova	SE/H/0110/009	16130	JANSSEN-CILAG OY	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topimax 50 mg kapslar, hårda	SE/H/0110/009	14350	JANSSEN-CILAG AB	SE
Topimax 50 mg kapslar, harde	SE/H/0110/009	98/3717	JANSSEN-CILAG A/S	NO
Topimax 50 mg kapslar, hårde	SE/H/0110/009	19591	JANSSEN-CILAG A/S	DK
Topimax 50 mg tabletter, filmdrasjerte	SE/H/0110/002	95-791	JANSSEN-CILAG A/S	NO
Topiramaat 100 mg PCH, filmomhulde tabletten	NO/H/0136/003	RVG 101530	PHARMACHEMIE B.V	NL
Topiramaat 200 mg PCH, filmomhulde tabletten	NO/H/0136/004	RVG 101533	PHARMACHEMIE B.V	NL
Topiramaat 25 mg PCH, filmomhulde tabletten	NO/H/0136/001	RVG 101527	PHARMACHEMIE B.V	NL
Topiramaat 50 mg PCH, filmomhulde tabletten	NO/H/0136/002	RVG 101529	PHARMACHEMIE B.V	NL
Topiramaat Accord 100 mg filmomhulde tabletten	UK/H/1438/003	RVG 102980	ACCORD HEALTHCARE LIMITED	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
Topiramaat Accord 200 mg filmomhulde tabletten	UK/H/1438/004	RVG 102981	ACCORD HEALTHCARE LIMITED	NL
Topiramaat Accord 25 mg filmomhulde tabletten	UK/H/1438/001	RVG 102978	ACCORD HEALTHCARE LIMITED	NL
Topiramaat Accord 50 mg filmomhulde tabletten	UK/H/1438/002	RVG 102979	ACCORD HEALTHCARE LIMITED	NL
Topiramaat Apotex 100 mg, filmomhulde tabletten	NL/H/1231/003	RVG 33206	APOTEX EUROPE B.V.	NL
Topiramaat Apotex 200 mg, filmomhulde tabletten	NL/H/1231/004	RVG 33207	APOTEX EUROPE B.V.	NL
Topiramaat Apotex 25 mg, filmomhulde tabletten	NL/H/1231/001	RVG 33204	APOTEX EUROPE B.V.	NL
Topiramaat Apotex 50 mg, filmomhulde tabletten	NL/H/1231/002	RVG 33205	APOTEX EUROPE B.V.	NL
Topiramaat Auro 100 mg, filmomhulde tabletten	PT/H/0231/003	RVG 104817	AUROBINDO PHARMA B.V.	NL
Topiramaat Auro 200 mg, filmomhulde tabletten	PT/H/0231/004	RVG 104818	AUROBINDO PHARMA B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramaat Auro 25 mg, filmomhulde tabletten	PT/H/0231/001	RVG 104812	AUROBINDO PHARMA B.V.	NL
Topiramaat Auro 50 mg, filmomhulde tabletten	PT/H/0231/002	RVG 104816	AUROBINDO PHARMA B.V.	NL
Topiramaat Aurobindo 100 mg filmomhulde tabletten	not available	RVG 105864	AUROBINDO PHARMA B.V.	NL
Topiramaat Aurobindo 200 mg filmomhulde tabletten	not available	RVG 105865	AUROBINDO PHARMA B.V.	NL
Topiramaat Aurobindo 25 mg filmomhulde tabletten	not available	RVG 105862	AUROBINDO PHARMA B.V.	NL
Topiramaat Aurobindo 50 mg filmomhulde tabletten	not available	RVG 105863	AUROBINDO PHARMA B.V.	NL
Topiramaat CF 100 mg, filmomhulde tabletten	NL/H/0947/003	RVG 34571	CENTRAFARM B.V.	NL
Topiramaat CF 200 mg, filmomhulde tabletten	NL/H/0947/004	RVG 34572	CENTRAFARM B.V.	NL
Topiramaat CF 25 mg, filmomhulde tabletten	NL/H/0947/001	RVG 34569	CENTRAFARM B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramaat CF 50 mg, filmomhulde tabletten	NL/H/0947/002	RVG 34570	CENTRAFARM B.V.	NL
Topiramaat Desitin 100 mg filmomhulde tabletten	NL/H/0876/003	RVG 34263	DESITIN ARZNEIMITTEL GMBH	NL
Topiramaat Desitin 200 mg filmomhulde tabletten	NL/H/0876/004	RVG 34264	DESITIN ARZNEIMITTEL GMBH	NL
Topiramaat Desitin 25 mg filmomhulde tabletten	NL/H/1202/001	RVG 101285	DESITIN ARZNEIMITTEL GMBH	NL
Topiramaat Desitin 50 mg filmomhulde tabletten	NL/H/1202/002	RVG 101286	DESITIN ARZNEIMITTEL GMBH	NL
Topiramaat Glenmark 100 mg, filmomhulde tabletten	DE/H/0984/003	RVG 103744	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Topiramaat Glenmark 200 mg, filmomhulde tabletten	DE/H/0984/004	RVG 103745	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Topiramaat Glenmark 25 mg, filmomhulde tabletten	DE/H/0984/001	RVG 103742	GLENMARK PHARMACEUTICALS EUROPE LIMITED	NL
Topiramaat Glenmark 50 mg, filmomhulde tabletten	DE/H/0984/002	RVG 103743	GLENMARK PHARMACEUTICALS EUROPE LIMITED	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
Topiramaat Mylan 100 mg, filmomhulde tabletten	NL/H/0717/003	RVG 33420	MYLAN B.V.	NL
Topiramaat Mylan 200 mg, filmomhulde tabletten	NL/H/0717/004	RVG 33421	MYLAN B.V.	NL
Topiramaat Mylan 25 mg, filmomhulde tabletten	NL/H/0717/001	RVG 33418	MYLAN B.V.	NL
Topiramaat Mylan 50 mg, filmomhulde tabletten	NL/H/0717/002	RVG 33419	MYLAN B.V.	NL
Topiramaat Sandoz tablet 100 mg, filmomhulde tabletten	FI/H/0599/003	RVG 33977	SANDOZ B.V.	NL
Topiramaat Sandoz tablet 25 mg, filmomhulde tabletten	FI/H/0599/001	RVG 33976	SANDOZ B.V.	NL
Topiramaat Torrent 100 mg filmomhulde tabletten	NL/H/1667/001-005	RVG 104612	TORRENT PHARMA GMBH	NL
Topiramaat Torrent 200 mg filmomhulde tabletten	NL/H/1667/001-005	RVG 104613	TORRENT PHARMA GMBH	NL
Topiramaat Torrent 25 mg filmomhulde tabletten	NL/H/1667/001-005	RVG 104609	TORRENT PHARMA GMBH	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramaat Torrent 400 mg filmomhulde tabletten	NL/H/1667/001-005	RVG 104614	TORRENT PHARMA GMBH	NL
Topiramaat Torrent 50 mg filmomhulde tabletten	NL/H/1667/001-005	RVG 104611	TORRENT PHARMA GMBH	NL
Topiramat - 1 A Pharma® 100 mg Filmdabletten	DE/H/3334/003	67659.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 15 mg Hartkapseln	DE/H/3334/005	67654.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 25 mg Filmdabletten	DE/H/3334/001	67657.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 25 mg Filmdabletten	DE/H/3334/004	67660.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 25 mg Hartkapseln	DE/H/3334/006	67655.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 50 mg Filmdabletten	DE/H/3334/002	67658.00.00	1 A PHARMA GMBH	DE
Topiramat - 1 A Pharma® 50 mg Hartkapseln	DE/H/3334/007	67656.00.00	1 A 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
Topiramat Sandoz 100 mg filmdrasjerte tabletter	FI/H/0599/003	06-4082	SANDOZ A/S	NO
Topiramat Sandoz 200 mg filmdrasjerte tabletter	FI/H/0599/004	06-4083	SANDOZ A/S	NO
Topiramat Sandoz 25 mg filmdrasjerte tabletter	FI/H/0599/001	06-4080	SANDOZ A/S	NO
Topiramat Sandoz 50 mg filmdrasjerte tabletter	FI/H/0599/002	06-4081	SANDOZ A/S	NO
Topiramat "Actavis", filmovertukne tabletter	PT/H/0231/002	44904	ACTAVIS GROUP PTC EHF.	DK
Topiramat "Actavis", filmovertukne tabletter	PT/H/0231/001	44903	ACTAVIS GROUP PTC EHF.	DK
Topiramat "Actavis", filmovertukne tabletter	PT/H/0231/004	44906	ACTAVIS GROUP PTC EHF.	DK
Topiramat "Actavis", filmovertukne tabletter	PT/H/0231/003	44905	ACTAVIS GROUP PTC EHF.	DK
Topiramat "Amneal", filmovertukne tabletter	UK/H/1166/001	41723	AMNEAL PHARMA EUROPE LIMITED	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
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/002	41724	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/003	41725	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/004	41726	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/001	41723	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/002	41724	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/003	41725	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Amneal", filmovertrukne tabletter	UK/H/1166/004	41726	AMNEAL PHARMA EUROPE LIMITED	DK
Topiramat "Hexal", filmovertrukne tabletter	DE/H/3333/001	40209	HEXAL A/S	DK
Topiramat "Hexal", filmovertrukne tabletter	DE/H/3333/002	40210	HEXAL 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
Topiramat "Hexal", filmovertrukne tabletter	DE/H/3333/003	40211	HEXAL A/S	DK
Topiramat "Hexal", filmovertrukne tabletter	DE/H/3333/004	40212	HEXAL A/S	DK
Topiramat "Orion", filmovertrukne tabletter	FI/H/0800/001	42108	ORION CORPORATION	DK
Topiramat "Orion", filmovertrukne tabletter	FI/H/0800/002	42109	ORION CORPORATION	DK
Topiramat "Orion", filmovertrukne tabletter	FI/H/0800/003	42110	ORION CORPORATION	DK
Topiramat "Orion", filmovertrukne tabletter	FI/H/0800/004	42111	ORION CORPORATION	DK
Topiramat 1A Farma 100 mg filmdragerade tabletter	FI/H/0599/003	23824	1A FARMA A/S	SE
Topiramat 1A Farma 200 mg filmdragerade tabletter	FI/H/0599/004	23825	1A FARMA A/S	SE
Topiramat 1A Farma 25 mg filmdragerade tabletter	FI/H/0599/001	23822	1A FARMA A/S	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat 1A Farma 50 mg filmdragerade tabletter	FI/H/0599/002	23823	1A FARMA A/S	SE
Topiramat 1A Pharma 100 mg - Filmtabletten	DE/H/3334/003	1-27614	1A PHARMA GMBH	AT
Topiramat 1A Pharma 200 mg - Filmtabletten	DE/H/3334/004	1-27615	1A PHARMA GMBH	AT
Topiramat 1A Pharma 25 mg - Filmtabletten	DE/H/3334/001	1-27612	1A PHARMA GMBH	AT
Topiramat 1A Pharma 50 mg - Filmtabletten	DE/H/3334/002	1-27613	1A PHARMA GMBH	AT
Topiramat Actavis 100 mg comprimate filmate	815/2008/01-23	815/2008/01-23	ACTAVIS GROUP HF.	RO
TOPIRAMAT ACTAVIS 100 MG FILMDRAGERADE TABLETTER	PT/H/0231/003	42362	ACTAVIS GROUP PTC EHF.	SE
Topiramat Actavis 100 mg Filmtabletten	PT/H/0231/003	1-28639	ACTAVIS GROUP PTC EHF.	AT
Topiramat Actavis 100 mg filmuhúðaðar töflur	PT/H/0231/003	IS/1/09/069/03	ACTAVIS GROUP PTC EHF.	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
TOPIRAMAT ACTAVIS 100 MG POTAHOVANÉ TABLETY	PT/H/0231/003	21/607/09-C	ACTAVIS GROUP PTC EHF.	CZ
TOPIRAMAT ACTAVIS 200 MG FILMDRAGERADE TABLETTER	PT/H/0231/004	42363	ACTAVIS GROUP PTC EHF.	SE
Topiramát Actavis 200 mg Filmtabletten	PT/H/0231/004	1-28640	ACTAVIS GROUP PTC EHF.	AT
Topiramát Actavis 200 mg filmhúðaðar töflur	PT/H/0231/004	IS/1/09/069/04	ACTAVIS GROUP PTC EHF.	IS
Topiramát Actavis 25 mg comprimate filmate	813/2008/01-21	813/2008/01-21	ACTAVIS GROUP HF.	RO
Topiramát Actavis 25 mg filmdragerade tabletter	PT/H/0231/001	42360	ACTAVIS GROUP PTC EHF.	SE
Topiramát Actavis 25 mg Filmtabletten	PT/H/0231/001	1-28637	ACTAVIS GROUP PTC EHF.	AT
Topiramát Actavis 25 mg filmhúðaðar töflur	PT/H/0231/001	IS/1/09/069/01	ACTAVIS GROUP PTC EHF.	IS
TOPIRAMAT ACTAVIS 25 MG POTAHOVANÉ TABLETY	PT/H/0231/001	21/605/09-C	ACTAVIS GROUP PTC EHF.	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
Topiramat Actavis 50 mg comprimate filmate	814/2008/01-21	814/2008/01-21	ACTAVIS GROUP HF.	RO
Topiramat Actavis 50 mg film-coated tablets	PT/H/0231/002	42361	ACTAVIS GROUP PTC EHF.	SE
Topiramat Actavis 50 mg Filmtabletten	PT/H/0231/002	1-28638	ACTAVIS GROUP PTC EHF.	AT
Topiramat Actavis 50 mg filmuhúðaðar töflur	PT/H/0231/002	IS/1/09/069/02	ACTAVIS GROUP PTC EHF.	IS
TOPIRAMAT ACTAVIS 50 MG POTAHOVANÉ TABLETY	PT/H/0231/002	21/606/09-C	ACTAVIS GROUP PTC EHF.	CZ
Topiramat AL 100 mg Filmtabletten	not available	67235.00.00	ALIUD PHARMA GMBH	DE
Topiramat AL 200 mg Filmtabletten	not available	67236.00.00	ALIUD PHARMA GMBH	DE
Topiramat AL 25 mg Filmtabletten	not available	67233.00.00	ALIUD PHARMA GMBH	DE
Topiramat AL 50 mg Filmtabletten	not available	67234.00.00	ALIUD 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
Topiramat AL Migräne 25 mg Filmtabletten	not available	76480.00.00	ALIUD PHARMA GMBH	DE
Topiramat AL Migräne 50 mg Filmtabletten	not available	76481.00.00	ALIUD PHARMA GMBH	DE
Topiramat Aurobindo 100 mg comprimate filmate	NL/H/2916/003	7386/2015/01	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 100 mg comprimate filmate	NL/H/2916/003	7386/2015/02	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 100 mg Filmtabletten	not available	82082.00.00	AUROBINDO PHARMA GMBH	DE
Topiramat Aurobindo 200 mg comprimate filmate	NL/H/2916/004	7387/2015/01	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 200 mg comprimate filmate	NL/H/2916/004	7387/2015/02	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 200 mg Filmtabletten	not available	82083.00.00	AUROBINDO PHARMA GMBH	DE
Topiramat Aurobindo 25 mg comprimate filmate	NL/H/2916/001	7384/2015/01	AUROBINDO PHARMA (MALTA) LIMITED	RO

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
Topiramat Aurobindo 25 mg comprimate filmate	NL/H/2916/001	7384/2015/02	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 25 mg Filmtabletten	not available	82080.00.00	AUROBINDO PHARMA GMBH	DE
Topiramat Aurobindo 50 mg comprimate filmate	NL/H/2916/002	7385/2015/01	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 50 mg comprimate filmate	NL/H/2916/002	7385/2015/02	AUROBINDO PHARMA (MALTA) LIMITED	RO
Topiramat Aurobindo 50 mg Filmtabletten	not available	82081.00.00	AUROBINDO PHARMA GMBH	DE
Topiramat Bluefish 100 mg Filmtabletten	AT/H/0612/003	1-29028	BLUEFISH PHARMACEUTICALS AB	AT
Topiramat Bluefish 100 mg Filmtabletten	AT/H/0612/003	74682.00.00	BLUEFISH PHARMACEUTICALS AB	DE
Topiramat Bluefish 100 mg filmuhúðaðar töflur	AT/H/0612/003	IS/1/13/031/03	BLUEFISH PHARMACEUTICALS AB	IS
Topiramat Bluefish 100 mg tabletter, filmdrasjerte	AT/H/0612/003	08-6138	BLUEFISH PHARMACEUTICALS AB	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
Topiramat Bluefish 200 mg Filmdabletten	AT/H/0612/004	1-29030	BLUEFISH PHARMACEUTICALS AB	AT
Topiramat Bluefish 200 mg Filmdabletten	AT/H/0612/004	74683.00.00	BLUEFISH PHARMACEUTICALS AB	DE
Topiramat Bluefish 200 mg filmuhúðaðar töflur	AT/H/0612/004	IS/1/13/031/04	BLUEFISH PHARMACEUTICALS AB	IS
Topiramat Bluefish 200 mg tabletter, filmdrasjerte	AT/H/0612/004	08-6139	BLUEFISH PHARMACEUTICALS AB	NO
Topiramat Bluefish 25 mg Filmdabletten	AT/H/0612/001	1-29026	BLUEFISH PHARMACEUTICALS AB	AT
Topiramat Bluefish 25 mg Filmdabletten	AT/H/0612/001	74680.00.00	BLUEFISH PHARMACEUTICALS AB	DE
Topiramat Bluefish 25 mg filmuhúðaðar töflur	AT/H/0612/001	IS/1/13/031/01	BLUEFISH PHARMACEUTICALS AB	IS
Topiramat Bluefish 25 mg tabletter, filmdrasjerte	AT/H/0612/001	08-6136	BLUEFISH PHARMACEUTICALS AB	NO
Topiramat Bluefish 50 mg Filmdabletten	AT/H/0612/002	1-29027	BLUEFISH PHARMACEUTICALS AB	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
Topiramat Bluefish 50 mg Filmtabletten	AT/H/0612/002	74681.00.00	BLUEFISH PHARMACEUTICALS AB	DE
Topiramat Bluefish 50 mg filmuhúðaðar töflur	AT/H/0612/002	IS/1/13/031/02	BLUEFISH PHARMACEUTICALS AB	IS
Topiramat Bluefish 50 mg tabletter, filmdrasjerte	AT/H/0612/002	08-6137	BLUEFISH PHARMACEUTICALS AB	NO
Topiramat Bluefish, 100 mg, tabletki powlekane	AT/H/0612/003	16741	BLUEFISH PHARMACEUTICALS AB	PL
Topiramat Bluefish, 200 mg, tabletki powlekane	AT/H/0612/004	16742	BLUEFISH PHARMACEUTICALS AB	PL
Topiramat Bluefish, 25 mg, tabletki powlekane	AT/H/0612/001	16740	BLUEFISH PHARMACEUTICALS AB	PL
Topiramat Bluefish, 50 mg, tabletki powlekane	AT/H/0612/002	15314	BLUEFISH PHARMACEUTICALS AB	PL
Topiramat Desitin 100 mg Filmtabletten	NL/H/0876/003	66556.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Topiramat Desitin 25 mg Filmtabletten	NL/H/1202/001	71490.00.00	DESITIN 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
Topiramat Desitin 50 mg Filmtabletten	NL/H/1202/002	71491.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Topiramat Desitin quadro 200 mg Filmtabletten	NL/H/0876/004	66557.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Topiramat Egis 100 mg comprimate filmate	NL/H/864/003/E/001	6654/2014/09-12	EGIS PHARMACEUTICALS PLC	RO
Topiramat Egis 200 mg comprimate filmate	NL/H/864/004/E/001	6655/2014/09-12	EGIS PHARMACEUTICALS PLC	RO
Topiramat G.L. 100 mg - Filmtabletten	not available	1-27806	G.L. PHARMA GMBH	AT
Topiramat G.L. 200 mg - Filmtabletten	not available	1-27807	G.L. PHARMA GMBH	AT
Topiramat G.L. 25 mg - Filmtabletten	not available	1-27804	G.L. PHARMA GMBH	AT
Topiramat G.L. 50 mg - Filmtabletten	not available	1-27805	G.L. PHARMA GMBH	AT
Topiramat Glenmark 100 mg Filmtabletten	DE/H/0984/003	68192.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	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
Topiramat Glenmark 200 mg Filmtabletten	DE/H/0984/004	68193.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Topiramat Glenmark 25 mg Filmtabletten	DE/H/0984/001	68190.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Topiramat Glenmark 50 mg Filmtabletten	DE/H/0984/002	68191.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Topiramat Heumann 100 mg Filmtabletten	NL/H/1667/003/DC	77518.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Topiramat Heumann 200 mg Filmtabletten	NL/H/1667/004/DC	77519.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Topiramat Heumann 25 mg Filmtabletten	NL/H/1667/001/DC	77516.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Topiramat Heumann 400 mg Filmtabletten	NL/H/1667/005/DC	77520.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Topiramat Heumann 50 mg Filmtabletten	NL/H/1667/002/DC	77517.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Topiramat HEXAL® 100 mg Filmtabletten	DE/H/3333/003	67664.00.00	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
Topiramat HEXAL® 15 mg Hartkapseln	DE/H/3333/005	67666.00.00	HEXAL AG	DE
Topiramat HEXAL® 200 mg Filmtabletten	DE/H/3333/004	67665.00.00	HEXAL AG	DE
Topiramat HEXAL® 25 mg Filmtabletten	DE/H/3333/001	67662.00.00	HEXAL AG	DE
Topiramat HEXAL® 25 mg Hartkapseln	DE/H/3333/006	67667.00.00	HEXAL AG	DE
Topiramat HEXAL® 50 mg Filmtabletten	DE/H/3333/002	67663.00.00	HEXAL AG	DE
Topiramat HEXAL® 50 mg Hartkapseln	DE/H/3333/007	67668.00.00	HEXAL AG	DE
Topiramat Migräne STADA® 100 mg Filmtabletten	not available	67239.00.00	STADAPHARM GMBH	DE
Topiramat Migräne STADA® 25 mg Filmtabletten	not available	67237.00.00	STADAPHARM GMBH	DE
Topiramat Migräne STADA® 50 mg Filmtabletten	not available	67238.00.00	STADAPHARM 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
Topiramat Mylan 100 mg potahované tablety	NL/H/0717/003	21/262/07-C	GENERICS [UK] LIMITED	CZ
Topiramat Mylan 25 mg potahované tablety	NL/H/0717/001	21/260/07-C	GENERICS [UK] LIMITED	CZ
Topiramat Mylan 50 mg potahované tablety	NL/H/0717/002	21/261/07-C	GENERICS [UK] LIMITED	CZ
Topiramat Orion 100 mg apvalkotās tabletes	FI/H/0800/003	08-0285	ORION CORPORATION	LV
Topiramat Orion 100 mg filmdragerade tabletter	FI/H/800/003	24167	ORION CORPORATION	FI
Topiramat Orion 100 mg filmdragerade tabletter	FI/H/0800/003	26075	ORION CORPORATION	SE
Topiramat Orion 100 mg plėvele dengtos tabletės	FI/H/0800/003	LT/1/08/1322/005	ORION CORPORATION	LT
Topiramat Orion 100 mg plėvele dengtos tabletės	FI/H/0800/003	LT/1/08/1322/006	ORION CORPORATION	LT
Topiramat Orion 100 mg tabletti, kalvopäällysteinen	FI/H/0800/003	24167	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat Orion 200 mg apvalkotās tabletes	FI/H/0800/004	08-0286	ORION CORPORATION	LV
Topiramat Orion 200 mg filmdragerade tabletter	FI/H/800/004	24168	ORION CORPORATION	FI
Topiramat Orion 200 mg filmdragerade tabletter	FI/H/0800/004	26076	ORION CORPORATION	SE
Topiramat Orion 200 mg plēvele dengtos tabletēs	FI/H/0800/004	LT/1/08/1322/007	ORION CORPORATION	LT
Topiramat Orion 200 mg plēvele dengtos tabletēs	FI/H/0800/004	LT/1/08/1322/008	ORION CORPORATION	LT
Topiramat Orion 200 mg tabletti, kalvopäällysteinen	FI/H/0800/004	24168	ORION CORPORATION	FI
Topiramat Orion 25 mg apvalkotās tabletes	FI/H/0800/001	08-0283	ORION CORPORATION	LV
Topiramat Orion 25 mg filmdragerade tabletter	FI/H/800/001	24165	ORION CORPORATION	FI
Topiramat Orion 25 mg filmdragerade tabletter	FI/H/0800/001	26073	ORION CORPORATION	SE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat Orion 25 mg plėvele dengtos tabletės	FI/H/0800/001	LT/1/08/1322/001	ORION CORPORATION	LT
Topiramat Orion 25 mg plėvele dengtos tabletės	FI/H/0800/001	LT/1/08/1322/002	ORION CORPORATION	LT
Topiramat Orion 25 mg tabletti, kalvopäälysteinen	FI/H/0800/001	24165	ORION CORPORATION	FI
Topiramat Orion 50 mg apvalkotās tabletes	FI/H/0800/002	08-0284	ORION CORPORATION	LV
Topiramat Orion 50 mg filmdragerade tabletter	FI/H/800/002	24166	ORION CORPORATION	FI
Topiramat Orion 50 mg filmdragerade tabletter	FI/H/0800/002	26074	ORION CORPORATION	SE
Topiramat Orion 50 mg plėvele dengtos tabletės	FI/H/0800/002	LT/1/08/1322/003	ORION CORPORATION	LT
Topiramat Orion 50 mg plėvele dengtos tabletės	FI/H/0800/002	LT/1/08/1322/004	ORION CORPORATION	LT
Topiramat Orion 50 mg tabletti, kalvopäälysteinen	FI/H/0800/002	24166	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat Pharmathen 100 mg filmovertukne tabletter	DK/H/0930/003	38894	PHARMATHEN S.A.	DK
Topiramat Pharmathen 200 mg filmovertukne tabletter	DK/H/0930/004	38895	PHARMATHEN S.A.	DK
Topiramat Pharmathen 25 mg filmovertukne tabletter	DK/H/0930/001	38892	PHARMATHEN S.A.	DK
Topiramat Pharmathen 50 mg filmovertukne tabletter	DK/H/0930/002	38893	PHARMATHEN S.A.	DK
Topiramat ratiopharm 100 mg filmdragerade tabletter	DE/H/2064/004	19869	RATIOPHARM GMBH	FI
Topiramat ratiopharm 100 mg filmdragerade tabletter	DE/H/2064/004	23320	RATIOPHARM GMBH	SE
TOPIRAMAT RATIOPHARM 100 mg kalvopäällysteinen tabletti	DE/H/2064/004	19869	RATIOPHARM GMBH	FI
Topiramat ratiopharm 200 mg filmdragerade tabletter	DE/H/2064/005	19870	RATIOPHARM GMBH	FI
Topiramat ratiopharm 200 mg filmdragerade tabletter	DE/H/2064/005	23321	RATIOPHARM GMBH	SE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
TOPIRAMAT RATIOPHARM 200 mg kalvopäällysteinen tabletti	DE/H/2064/005	19870	RATIOPHARM GMBH	FI
Topiramate ratiopharm 25 mg filmdragerade tabletter	DE/H/2064/002	19867	RATIOPHARM GMBH	FI
Topiramate ratiopharm 25 mg filmdragerade tabletter	DE/H/2064/002	23318	RATIOPHARM GMBH	SE
Topiramate ratiopharm 50 mg filmdragerade tabletter	DE/H/2064/003	19868	RATIOPHARM GMBH	FI
Topiramate ratiopharm 50 mg filmdragerade tabletter	DE/H/2064/003	23319	RATIOPHARM GMBH	SE
TOPIRAMAT RATIOPHARM 50 mg kalvopäällysteinen tabletti	DE/H/2064/003	19868	RATIOPHARM GMBH	FI
Topiramate Sandoz 100 mg - Filmtabletten	FI/H/0599/003	1-26702	SANDOZ GMBH	AT
Topiramate Sandoz 100 mg potahované tablety	FI/H/0599/003	21/403/06-C	SANDOZ S.R.O.	CZ
Topiramate Sandoz 100 mg tabletti, kalvopäällysteinen	FI/H/0599/003	19676	SANDOZ A/S	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat Sandoz 15 mg kapseli, kova	FI/H/0599/005	20303	SANDOZ A/S	FI
Topiramat Sandoz 200 mg - Filmtabletten	FI/H/0599/004	1-26703	SANDOZ GMBH	AT
Topiramat Sandoz 200 mg tabletti, kalvopäällysteinen	FI/H/0599/004	19677	SANDOZ A/S	FI
Topiramat Sandoz 25 mg - Filmtabletten	FI/H/0599/001	1-26697	SANDOZ GMBH	AT
Topiramat Sandoz 25 mg kapseli, kova	FI/H/0599/006	20304	SANDOZ A/S	FI
Topiramat Sandoz 25 mg tabletti, kalvopäällysteinen	FI/H/0599/001	19674	SANDOZ A/S	FI
Topiramat Sandoz 25 mg potahované tablety	FI/H/0599/001	21/401/06-C	SANDOZ S.R.O.	CZ
Topiramat Sandoz 50 mg - Filmtabletten	FI/H/0599/002	1-26699	SANDOZ GMBH	AT
Topiramat Sandoz 50 mg kapseli, kova	FI/H/0599/007	20305	SANDOZ A/S	FI

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramat Sandoz 50 mg tabletti, kalvopäällysteinen	FI/H/0599/002	19675	SANDOZ A/S	FI
Topiramat Sandoz 50 mg potahované tablety	FI/H/0599/002	21/402/06-C	SANDOZ S.R.O.	CZ
Topiramat STADA 100 mg Filmtabletten	DE/H/0889/003	1-27335	STADA ARZNEIMITTEL GMBH	AT
Topiramat STADA 100 mg Filmtabletten	DE/H/0889/003	67243.00.00	STADAPHARM GMBH	DE
Topiramat STADA 200 mg Filmtabletten	DE/H/0889/004	1-27334	STADA ARZNEIMITTEL GMBH	AT
Topiramat STADA 200 mg Filmtabletten	DE/H/0889/004	67244.00.00	STADAPHARM GMBH	DE
Topiramat STADA 25 mg Filmtabletten	DE/H/0889/001	1-27332	STADA ARZNEIMITTEL GMBH	AT
Topiramat STADA 25 mg Filmtabletten	DE/H/0889/001	67241.00.00	STADAPHARM GMBH	DE
Topiramat STADA 50 mg Filmtabletten	DE/H/0889/002	1-27333	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
Topiramat STADA 50 mg Filmtabletten	DE/H/0889/002	67242.00.00	STADAPHARM GMBH	DE
Topiramat Teva 100 mg filmdrasjerte tabletter	NO/H/0136/003	05-3168	TEVA SWEDEN AB	NO
Topiramat Teva 200 mg filmdrasjerte tabletter	NO/H/0136/004	05-3169	TEVA SWEDEN AB	NO
Topiramat Teva 25 mg filmdrasjerte tabletter	NO/H/0136/001	05-3166	TEVA SWEDEN AB	NO
Topiramat Teva 300 mg filmdrasjerte tabletter	NO/H/0136/005	05-3170	TEVA SWEDEN AB	NO
Topiramat Teva 400 mg filmdrasjerte tabletter	NO/H/0136/006	05-3171	TEVA SWEDEN AB	NO
Topiramat Teva 50 mg filmdrasjerte tabletter	NO/H/0136/002	05-3167	TEVA SWEDEN AB	NO
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	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
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	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
Topiramat Winthrop 100 mg Filmtabletten	DE/H/1031/003	68676.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	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
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 200 mg Filmtabletten	DE/H/1031/004	68677.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	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
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 25 mg Filmtabletten	DE/H/1031/001	68674.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	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
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	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
Topiramat Winthrop 50 mg Filmtabletten	DE/H/1031/002	68675.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Topiramat-CT 200 mg Filmtabletten	DE/H/2065/005	65339.00.00	ABZ-PHARMA GMBH	DE
Topiramat-CT 25 mg Filmtabletten	DE/H/2065/002	65336.00.00	ABZ-PHARMA GMBH	DE
Topiramat-CT 50 mg Filmtabletten	DE/H/2065/003	65337.00.00	ABZ-PHARMA GMBH	DE
Topiramate / Actavis 100mg film-coated tablets	PT/H/0231/003	37168/24-05-2012	ACTAVIS GROUP PTC EHF.	GR
Topiramate / Actavis 200mg film-coated tablets	PT/H/0231/004	37162/24-05-2012	ACTAVIS GROUP PTC EHF.	GR
Topiramate / Actavis 25mg film-coated tablets	PT/H/0231/001	58673/11/24-05-2012	ACTAVIS GROUP PTC EHF.	GR
Topiramate / Actavis 50mg film-coated tablets	PT/H/0231/002	37163/24-05-2012	ACTAVIS GROUP PTC EHF.	GR
Topiramate 100 mg film-coated tablets	NL/H/2916/003	MA807/08703	AUROBINDO PHARMA (MALTA) LIMITED	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramate 100 mg Film-coated Tablets	not available	PL 36390/0071	CIPLA (EU) LIMITED	UK
Topiramate 100 mg Film-coated tablets	PT/H/0199/003	PL 20117/0166	MORNINGSIDE HEALTHCARE LTD	UK
Topiramate 200 mg film-coated tablets	NL/H/2916/004	MA807/08704	AUROBINDO PHARMA (MALTA) LIMITED	MT
Topiramate 200 mg Film-coated Tablets	not available	PL 36390/0072	CIPLA (EU) LIMITED	UK
Topiramate 200 mg Film-coated tablets	PT/H/0199/004	PL 20117/0167	MORNINGSIDE HEALTHCARE LTD	UK
Topiramate 25 mg film-coated tablets	NL/H/2916/001	MA807/08701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Topiramate 25 mg Film-coated Tablets	not available	PL 36390/0069	CIPLA (EU) LIMITED	UK
Topiramate 25 mg Film-coated tablets	PT/H/0199/001	PL 20117/0164	MORNINGSIDE HEALTHCARE LTD	UK
Topiramate 300 mg Film-coated Tablets	NO/H/0136/005	PL 00289/1057	TEVA UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramate 400 mg Film-coated Tablets	NO/H/0136/006	PL 00289/1058	TEVA UK LIMITED	UK
Topiramate 50 mg film-coated tablets	NL/H/2916/002	MA807/08702	AUROBINDO PHARMA (MALTA) LIMITED	MT
Topiramate 50 mg Film-coated Tablets	not available	PL 36390/0070	CIPLA (EU) LIMITED	UK
Topiramate 50 mg Film-coated tablets	PT/H/0199/002	PL 20117/0165	MORNINGSIDE HEALTHCARE LTD	UK
Topiramate Accord 100 mg film-coated tablets	UK/H/1438/003	PL 20075/ 0147	ACCORD HEALTHCARE LIMITED	UK
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/034	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/035	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/036	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/037	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/038	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/039	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/040	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/041	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/042	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/043	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/045	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/046	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/047	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/048	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/033	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 100 mg plêvele dengtos tabletès	UK/H/1438/003	LT/1/10/1962/044	ACCORD HEALTHCARE LIMITED	LT
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 585 181 4 0	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 554 3 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 553 7 7	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 559 5 7	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 557 2 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 585 183 7 9	ACCORD HEALTHCARE FRANCE SAS	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
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 560 3 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 558 9 6	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 585 182 0 1	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 100 mg, comprimé pelliculé	not available	34009 274 556 6 7	ACCORD HEALTHCARE FRANCE SAS	FR
Topiramate Accord 200 mg film-coated tablets	UK/H/1438/004	PL 20075/0148	ACCORD HEALTHCARE LIMITED	UK
Topiramate Accord 200 mg plevle dengtos tabletės	UK/H/1438/004	LT/1/10/1962/056	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plėvele dengtos tabletės	UK/H/1438/004	LT/1/10/1962/050	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plėvele dengtos tabletės	UK/H/1438/004	LT/1/10/1962/051	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plėvele dengtos tabletės	UK/H/1438/004	LT/1/10/1962/052	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/053	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/054	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/055	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/057	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/058	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/059	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/061	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/062	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/063	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/064	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/049	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 200 mg plêvele dengtos tabletès	UK/H/1438/004	LT/1/10/1962/060	ACCORD HEALTHCARE LIMITED	LT
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 585 178 3 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 546 0 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 585 177 7 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 551 4 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 550 8 7	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 545 4 7	ACCORD HEALTHCARE FRANCE SAS	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
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 552 0 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 547 7 6	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 585 180 8 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 200 mg, comprimé pelliculé	not available	34009 274 548 3 7	ACCORD HEALTHCARE FRANCE SAS	FR
Topiramate Accord 25 mg film-coated tablets	UK/H/1438/001	PL 20075/ 0145	ACCORD HEALTHCARE LIMITED	UK
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/002	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/003	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/004	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/005	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/006	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/007	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/008	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/009	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/010	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/011	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/013	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/014	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletés	UK/H/1438/001	LT/1/10/1962/015	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/016	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/001	LT/1/10/1962/001	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 25 mg plévele dengtos tabletės	UK/H/1438/01-04/DC	LT/1/10/1962/012	ACCORD HEALTHCARE LIMITED	LT
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 571 5 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 585 190 3 1	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 585 189 5 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 569 0 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 575 0 0	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 572 1 0	ACCORD HEALTHCARE FRANCE SAS	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
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 574 4 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 585 188 9 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 573 8 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 25 mg, comprimé pelliculé	not available	34009 274 570 9 8	ACCORD HEALTHCARE FRANCE SAS	FR
Topiramate Accord 50 mg film-coated tablets	UK/H/1438/002	PL 20075/ 0146	ACCORD HEALTHCARE LIMITED	UK
Topiramate Accord 50 mg plevele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/019	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plevele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/020	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plevele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/021	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plėvele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/018	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/022	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/023	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/024	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/025	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/026	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/027	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/029	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/030	ACCORD HEALTHCARE LIMITED	LT
Topiramate Accord 50 mg plévele dengtos tabletès	UK/H/1438/002	LT/1/10/1962/031	ACCORD HEALTHCARE LIMITED	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
Topiramate Accord 50 mg plévele dengtos tabletés	UK/H/1438/002	LT/1/10/1962/032	ACCORD HEALTHCARE LIMITED	LT
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 565 5 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 563 2 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 567 8 7	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 566 1 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 564 9 7	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 562 6 8	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 585 187 2 0	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 274 568 4 8	ACCORD HEALTHCARE FRANCE SAS	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
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 585 186 6 9	ACCORD HEALTHCARE FRANCE SAS	FR
TOPIRAMATE ACCORD 50 mg, comprimé pelliculé	not available	34009 585 184 3 0	ACCORD HEALTHCARE FRANCE SAS	FR
Topiramate Actavis 100mg Film-Coated Tablets	not available	PL 30306/0187	ACTAVIS GROUP PTC EHF.	UK
Topiramate Actavis 15mg Capsules, hard	UK/H/1579/001	MA628/13601	ACTAVIS GROUP PTC EHF.	MT
Topiramate Actavis 15mg Capsules, hard	UK/H/1579/001	PL 30306/0541	ACTAVIS GROUP PTC EHF.	UK
Topiramate Actavis 200mg Film-Coated Tablets	not available	PL 30306/0188	ACTAVIS GROUP PTC EHF.	UK
Topiramate Actavis 25mg Capsules, hard	UK/H/1579/002	MA628/13602	ACTAVIS GROUP PTC EHF.	MT
Topiramate Actavis 25mg Capsules, hard	UK/H/1579/002	PL 30306/0542	ACTAVIS GROUP PTC EHF.	UK
Topiramate Actavis 25mg Film-Coated Tablets	not available	PL 30306/0185	ACTAVIS GROUP PTC EHF.	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
Topiramate Actavis 50mg Capsules, hard	UK/H/1015/003	MA628/13701	ACTAVIS GROUP PTC EHF.	MT
Topiramate Actavis 50mg Capsules, hard	UK/H/1015/003	PL 30306/0543	ACTAVIS GROUP PTC EHF.	UK
Topiramate Actavis 50mg Film-Coated Tablets	not available	PL 30306/0186	ACTAVIS GROUP PTC EHF.	UK
Topiramate Alvogen 100 mg filmuhúðaðar töflur	not available	IS/1/08/145/03	ALVOGEN EHF	IS
Topiramate Alvogen 25 mg filmuhúðaðar töflur	not available	IS/1/08/145/01	ALVOGEN EHF	IS
Topiramate Alvogen 50 mg filmuhúðaðar töflur	not available	IS/1/08/145/02	ALVOGEN EHF	IS
Topiramate Amneal 100 mg film-coated tablets	UK/H/1166/003	PL 42357/0129	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 100 mg film-coated tablets	UK/H/1166/003	PL 42357/0129	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 200 mg film-coated tablets	UK/H/1166/004	PL 42357/0130	AMNEAL PHARMA EUROPE 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
Topiramate Amneal 200 mg film-coated tablets	UK/H/1166/004	PL 42357/0130	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 25 mg film-coated tablets	UK/H/1166/001	PL 42357/0131	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 25 mg film-coated tablets	UK/H/1166/001	PL 42357/0131	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 50 mg film-coated tablets	UK/H/1166/002	PL 42357/0132	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Amneal 50 mg film-coated tablets	UK/H/1166/002	PL 42357/0132	AMNEAL PHARMA EUROPE LIMITED	UK
Topiramate Aptil 100mg Film-Coated Tablets	-	PL 40378/0078	TORRENT PHARMA LTD.	UK
Topiramate Aptil 200mg Film-Coated Tablets	-	PL 40378/0079	TORRENT PHARMA LTD.	UK
Topiramate Aptil 25mg Film-Coated Tablets	-	PL 40378/0076	TORRENT PHARMA LTD.	UK
Topiramate Aptil 50mg Film-Coated Tablets	-	PL 40378/0077	TORRENT PHARMA 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
TOPIRAMATE ARROW 100 mg, comprimé pelliculé	FR/H/0578/002	33226	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 100 mg, comprimé pelliculé	FR/H/0578/002	33226	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 200 mg, comprimé pelliculé	FR/H/0601/004	33227	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 200 mg, comprimé pelliculé	FR/H/0601/004	33227	ARROW GENERIQUES	FR
Topiramate Arrow 25 mg, gélule	UK/H/1579/002	NL36258	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 50 mg, comprimé pelliculé	FR/H/0601/002	33225	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 50 mg, comprimé pelliculé	FR/H/0601/002	33225	ARROW GENERIQUES	FR
TOPIRAMATE ARROW 50 mg, gélule	UK/H/1015/003	33789	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 100 mg, comprimé pelliculé	not available	32592	ARROW GENERIQUES	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
TOPIRAMATE ARROW GENERIQUES 100 mg, comprimé pelliculé	not available	NL 32592	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 100 mg, comprimé pelliculé	not available	32592	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 100 mg, comprimé pelliculé	not available	NL 32592	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 200 mg, comprimé pelliculé	not available	32593	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 200 mg, comprimé pelliculé	not available	NL 32592	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 200 mg, comprimé pelliculé	not available	NL 32592	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 50 mg, comprimé pelliculé	not available	32591	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 50 mg, comprimé pelliculé	not available	NL 32591	ARROW GENERIQUES	FR
TOPIRAMATE ARROW GENERIQUES 50 mg, comprimé pelliculé	not available	32591	ARROW GENERIQUES	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
TOPIRAMATE ARROW GENERIQUES 50 mg, comprimé pelliculé	not available	NL 32591	ARROW GENERIQUES	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056327	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056907	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056785	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056495	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056617	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056556	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938057157	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938057096	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
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938057218	BIOGARAN	FR
TOPIRAMATE BIOGARAN 100 mg, comprimé pelliculé	not available	3400938056846	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938057386	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400957105518	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400957105747	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400957105457	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938057447	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938057966	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938058048	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
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938057508	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400938057737	BIOGARAN	FR
TOPIRAMATE BIOGARAN 200 mg, comprimé pelliculé	not available	3400957105686	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052183	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052244	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052473	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052534	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052763	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938053135	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
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052305	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938053074	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052824	BIOGARAN	FR
TOPIRAMATE BIOGARAN 50 mg, comprimé pelliculé	not available	3400938052992	BIOGARAN	FR
Topiramate EG 25 mg comprimés pelliculés	DE/H/0889/001	BE324685	EUROGENERICS SA	BE
Topiramate EG 100 mg comprimés pelliculés	DE/H/0889/003	BE 324703	EUROGENERICS N.V./S.A.	BE
Topiramate EG 100 mg filmomhulde tabletten	DE/H/0889/003	BE 324703	EUROGENERICS N.V./S.A.	BE
Topiramate EG 100 mg Filmtabletten	DE/H/0889/003	BE 324703	EUROGENERICS N.V./S.A.	BE
TOPIRAMATE EG 100 mg, comprimé pelliculé	not available	NL34066	EG LABO - LABORATOIRES EUROGENERICS	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
Topiramate EG 200 mg comprimés pelliculés	DE/H/0889/004	BE 324712	EUROGENERICS N.V./S.A.	BE
Topiramate EG 200 mg filmomhulde tabletten	DE/H/0889/004	BE 324712	EUROGENERICS N.V./S.A.	BE
Topiramate EG 200 mg Filmtabletten	DE/H/0889/004	BE 324712	EUROGENERICS N.V./S.A.	BE
TOPIRAMATE EG 200 mg, comprimé pelliculé	not available	NL34067	EG LABO - LABORATOIRES EUROGENERICS	FR
Topiramate EG 25 mg filmomhulde tabletten	DE/H/0889/001	BE 324685	EUROGENERICS SA	BE
Topiramate EG 25 mg Filmtabletten	DE/H/0889/001	BE 324685	EUROGENERICS SA	BE
Topiramate EG 50 mg comprimés pelliculés	DE/H/0889/002	BE324694	EUROGENERICS SA	BE
Topiramate EG 50 mg filmomhulde tabletten	DE/H/0889/002	BE324694	EUROGENERICS SA	BE
Topiramate EG 50 mg Filmtabletten	DE/H/0889/002	BE 324694	EUROGENERICS SA	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
TOPIRAMATE EG 50 mg, comprimé pelliculé	not available	NL34065	EG LABO - LABORATOIRES EUROGENERICS	FR
Topiramate Glenmark 100mg film-coated tablets	DE/H/0984/003	PL 25258/0062	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Topiramate Glenmark 200mg film-coated tablets	DE/H/0984/004	PL 25258/0063	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
TOPIRAMATE GLENMARK 25MG FILM-COATED TABLETS	DE/H/0984/001	PL 25258/0060	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
TOPIRAMATE GLENMARK 50MG FILM-COATED TABLETS	DE/H/0984/002	PL 25258/0061	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Topiramate Milpharm 100 mg film-coated tablets	NL/H/2916/003	PL 16363/0411	MILPHARM LIMITED	UK
Topiramate Milpharm 200 mg film-coated tablets	NL/H/2916/004	PL 16363/0412	MILPHARM LIMITED	UK
Topiramate Milpharm 25 mg film-coated tablets	NL/H/2916/001	PL 16363/0409	MILPHARM LIMITED	UK
Topiramate Milpharm 50 mg film-coated tablets	NL/H/2916/002	PL 16363/0410	MILPHARM 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
Topiramate Mylan 100 mg Film-coated Tablets	NL/H/0717/003	PL 04569/0784	MYLAN B.V.	UK
Topiramate Mylan 100 mg, comprimé pelliculé	not available	NL 32 944	MYLAN S.A.S	FR
Topiramate Mylan 200 mg Film-coated Tablets	NL/H/0717/004	PL 04569/0785	GENERICS [UK] LIMITED	UK
Topiramate Mylan 200 mg, comprimé pelliculé	not available	NL 32 945	MYLAN S.A.S	FR
Topiramate Mylan 25 mg Film-coated Tablets	NL/H/0717/001	PL 04569/0782	GENERICS [UK] LIMITED	UK
Topiramate Mylan 50 mg Film-coated Tablets	NL/H/0717/002	PL 04569-0783	GENERICS [UK] LIMITED	UK
Topiramate Mylan 50 mg, comprimé pelliculé	not available	NL 32 943	MYLAN S.A.S	FR
TOPIRAMATE OPENING PHARMA 100 mg, comprimé pelliculé	not available	389 782-9	TORRENT PHARMA GMBH	FR
TOPIRAMATE OPENING PHARMA 200 mg, comprimé pelliculé	not available	389 834-9	TORRENT PHARMA GMBH	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
TOPIRAMATE OPENING PHARMA 25 mg, comprimé pelliculé	not available	389 248-2	TORRENT PHARMA GMBH	FR
TOPIRAMATE OPENING PHARMA 50mg, comprimé pelliculé	not available	389 220-0	TORRENT PHARMA GMBH	FR
Topiramate Orion, 100 mg õhukese polümeerikattega tabletid	FI/H/0800/003	600708	ORION CORPORATION	EE
Topiramate Orion, 200 mg õhukese polümeerikattega tabletid	FI/H/0800/004	601008	ORION CORPORATION	EE
Topiramate Orion, 25 mg õhukese polümeerikattega tabletid	FI/H/0800/001	600808	ORION CORPORATION	EE
Topiramate Orion, 50 mg õhukese polümeerikattega tabletid	FI/H/0800/002	600908	ORION CORPORATION	EE
Topiramate Sandoz 100 mg Film-coated Tablets	FI/H/0599/003	PL 04416/0725	SANDOZ LTD	UK
Topiramate Sandoz 100 mg filmomhulde tabletten	FI/H/0599/003	BE 294147	SANDOZ N.V.	BE
Topiramate Sandoz 100 mg filmomhulde tabletten	FI/H/0599/003	BE 294156	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
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 169 5 5	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 170 3 7	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 172 6 6	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 173 2 7	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 176 1 7	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 177 8 5	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 174 9 5	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 175 5 6	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 178 4 6	SANDOZ	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
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 179 0 7	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 180 9 6	SANDOZ	FR
TOPIRAMATE Sandoz 100 mg, comprimé pelliculé	not available	34009 390 181 5 7	SANDOZ	FR
Topiramate Sandoz 15 mg Capsules, Hard	FI/H/0599/005	PL 04416/0720	SANDOZ LTD	UK
Topiramate Sandoz 200 mg film-coated Tablets	FI/H/0599/004	PL 04416/0726	SANDOZ LTD	UK
Topiramate Sandoz 200 mg filmomhulde tabletten	FI/H/0599/004	BE 294174	SANDOZ N.V.	BE
Topiramate Sandoz 200 mg filmomhulde tabletten	FI/H/0599/004	BE 294165	SANDOZ N.V.	BE
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 186 7 6	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 187 3 7	SANDOZ	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
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 182 1 8	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 183 8 6	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 184 4 7	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 185 0 8	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 189 6 6	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 190 4 8	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 191 0 9	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 192 7 7	SANDOZ	FR
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 193 3 8	SANDOZ	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
TOPIRAMATE Sandoz 200 mg, comprimé pelliculé	not available	34009 390 195 6 7	SANDOZ	FR
Topiramate Sandoz 25 mg Capsules, Hard	FI/H/0599/006	PL 04416/0721	SANDOZ LTD	UK
Topiramate Sandoz 25 mg Film-coated Tablets	FI/H/0599/001	PL 04416/0723	SANDOZ LTD	UK
Topiramate Sandoz 25 mg filmomhulde tabletten	FI/H/0599/001	BE 293973	SANDOZ N.V.	BE
Topiramate Sandoz 25 mg filmomhulde tabletten	FI/H/0599/001	BE 293946	SANDOZ N.V.	BE
Topiramate Sandoz 50 mg Capsules, Hard	FI/H/0599/007	PL 04416/0722	SANDOZ LTD	UK
Topiramate Sandoz 50 mg Film-coated Tablets	FI/H/0599/002	PL 04416/0724	SANDOZ LTD	UK
Topiramate Sandoz 50 mg filmomhulde tabletten	FI/H/0599/002	BE 293982	SANDOZ N.V.	BE
Topiramate Sandoz 50 mg filmomhulde tabletten	FI/H/0599/002	BE 294131	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
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 139 9 6	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 140 7 8	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 138 2 8	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 141 3 9	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 143 6 8	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 389 144 2 9	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 346 9 8	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 347 5 9	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 349 8 8	SANDOZ	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
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 348 1 0	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 351 2 1	SANDOZ	FR
TOPIRAMATE Sandoz 50 mg, comprimé pelliculé	not available	34009 394 350 6 0	SANDOZ	FR
Topiramate Tecnimedé 100 mg potahované tablety	PT/H/0198/003	21/700/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimedé 100 mg potahované tablety	PT/H/0198/003	21/700/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimedé 200 mg potahované tablety	PT/H/0198/004	21/701/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimedé 200 mg potahované tablety	PT/H/0198/004	21/701/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimedé 25 mg potahované tablety	PT/H/0198/001	21/698/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimedé 25 mg potahované tablety	PT/H/0198/001	21/698/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
Topiramate Tecnimed 50 mg potahované tablety	PT/H/0198/002	21/699/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Tecnimed 50 mg potahované tablety	PT/H/0198/002	21/699/09-C	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	CZ
Topiramate Teva 100mg film-coated tablets	NO/H/0136/003	PL 00289/1055	TEVA UK LIMITED	UK
TOPIRAMATE TEVA 200 mg, comprimé pelliculé sécable	NO/H/0136/005	NL35133	TEVA SANTÉ	FR
Topiramate Teva 200mg film-coated tablets	NO/H/0136/005	PL 00289/1056	TEVA UK LIMITED	UK
Topiramate Teva 25mg film-coated tablets	NO/H/0136/001	PL 00289/1053	TEVA UK LIMITED	UK
Topiramate Teva 50mg film-coated tablets	NO/H/0136/002	PL 00289/1054	TEVA UK LIMITED	UK
Topiramate Torrent 100 mg plévele dengtos tabletés	NL/H/1667/001-005	LT/1/10/2101/2	TORRENT PHARMA GMBH	LT
Topiramate Torrent 25 mg plévele dengtos tabletés	NL/H/1667/001-005	LT/1/10/2101/001-012	TORRENT PHARMA GMBH	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
Topiramate Torrent 400 mg plévele dengtos tabletés	NL/H/1667/001-005	LT/1/10/2101/049-060	TORRENT PHARMA GMBH	LT
Topiramate Torrent 50 mg plévele dengtos tabletés	NL/H/1667/001-005	LT/1/10/2101/013-024	TORRENT PHARMA GMBH	LT
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 384 7 3	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 385 3 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 387 6 3	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 388 2 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 389 9 2	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 100 mg, comprimé pelliculé	not available	34009 391 390 7 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 391 3 5	ZYDUS FRANCE	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
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 393 6 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 394 2 5	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 395 9 3	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 396 5 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 200 mg, comprimé pelliculé	not available	34009 391 397 1 5	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 25 mg, gélule	not available	34009 391 492 4 0	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 25 mg, gélule	not available	34009 391 493 0 1	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 25 mg, gélule	not available	34009 391 713 0 2	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 25 mg, gélule	not available	34009 391 714 7 0	ZYDUS FRANCE	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
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 391 380 1 5	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 391 381 8 3	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 391 382 4 4	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 391 383 0 5	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 574 346 7 0	ZYDUS FRANCE	FR
TOPIRAMATE ZYDUS 50 mg, comprimé pelliculé	not available	34009 574 347 3 1	ZYDUS FRANCE	FR
Topiramate/Mylan 100 mg Επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0717/003	9621 / 15 / 02-06-2016	MYLAN S.A.S	GR
Topiramate/Mylan 25 mg Επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0717/001	18654 / 15 / 02-06-2016	MYLAN S.A.S	GR
Topiramate/Mylan 50 mg Επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0717/003	11353 / 15 / 02-06-2016	MYLAN S.A.S	GR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramate/Teva 200 mg Επικαλυμμένα με λεπτό υμένιο δισκία	NO/H/0136/004	36237/26-5-2015	TEVA PHARMA B.V.	GR
Topiramate/Teva 25 mg Επικαλυμμένα με λεπτό υμένιο δισκία	NO/H/0136/001	36234/26-5-2015	TEVA PHARMA B.V.	GR
TopiramateAccord 50 mg plėvele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/028	ACCORD HEALTHCARE LIMITED	LT
TopiramateAccord 50 mg plėvele dengtos tabletės	UK/H/1438/002	LT/1/10/1962/017	ACCORD HEALTHCARE LIMITED	LT
Topiramat-Hormosan 100 mg Filmtabletten	not available	64821.00.00	HORMOSAN PHARMA GMBH	DE
Topiramat-Hormosan 200 mg Filmtabletten	not available	64822.00.00	HORMOSAN PHARMA GMBH	DE
Topiramat-Hormosan 25 mg Filmtabletten	not available	64819.00.00	HORMOSAN PHARMA GMBH	DE
Topiramat-Hormosan 50 mg Filmtabletten	not available	64820.00.00	HORMOSAN PHARMA GMBH	DE
Topiramat-Janssen 100 mg Filmtabletten	DE/H/2655/003	57000.02.00	JANSSEN-CILAG 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
Topiramat-Janssen 200 mg Filmtabletten	DE/H/2655/004	57000.03.00	JANSSEN-CILAG GMBH	DE
Topiramat-Janssen 25 mg Filmtabletten	DE/H/2655/001	57000.00.00	JANSSEN-CILAG GMBH	DE
Topiramat-Janssen 50 mg Filmtabletten	DE/H/2655/002	57000.01.00	JANSSEN-CILAG GMBH	DE
Topiramat-neuraxpharm 100 mg Filmtabletten	not available	82078.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Topiramat-neuraxpharm 200 mg Filmtabletten	not available	82079.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Topiramat-neuraxpharm 25 mg Filmtabletten	not available	82076.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Topiramat-neuraxpharm 50 mg Filmtabletten	not available	82077.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143330	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143342	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143355	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143367	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143379	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143381	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143393	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143405	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143417	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143429	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143431	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143443	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143456	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143468	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143470	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 100 mg compresse rivestite con film	UK/H/1438/003	041143482	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143494	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143506	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143518	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143520	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143532	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143544	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143557	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143569	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143571	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143583	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143595	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143607	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143619	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143621	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143633	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 200 mg compresse rivestite con film	UK/H/1438/004	041143645	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143013	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143025	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143037	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143049	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143052	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143063	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143076	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143088	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143090	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143102	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143114	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143126	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143138	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143140	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143153	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 25 mg compresse rivestite con film	UK/H/1438/001	041143165	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143177	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143278	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143189	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143191	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143203	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143215	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143227	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143239	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143241	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143254	ACCORD HEALTHCARE LIMITED	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
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143266	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143280	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143292	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143304	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143316	ACCORD HEALTHCARE LIMITED	IT
Topiramato Accord Healthcare 50 mg compresse rivestite con film	UK/H/1438/002	041143328	ACCORD HEALTHCARE LIMITED	IT
Topiramato Almus 100 mg comprimidos revestidos por película	not available	5078142	ALMUS, 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
Topiramato Almus 100 mg comprimidos revestidos por película	not available	5078159	ALMUS, LDA	PT
Topiramato Almus 200 mg comprimidos revestidos por película	not available	5078167	ALMUS, LDA	PT
Topiramato Almus 200 mg comprimidos revestidos por película	not available	5078175	ALMUS, LDA	PT
Topiramato Almus 25 mg comprimidos revestidos por película	not available	5078100	ALMUS, LDA	PT
Topiramato Almus 25 mg comprimidos revestidos por película	not available	5078118	ALMUS, LDA	PT
Topiramato Almus 50 mg comprimidos revestidos por película	not available	5078126	ALMUS, LDA	PT
Topiramato Almus 50 mg comprimidos revestidos por película	not available	5078134	ALMUS, LDA	PT
Topiramato Alter 100 mg comprimidos recubiertos con película EFG	not available	69.882	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 100 mg comprimidos recubiertos con película EFG	not available	69.882	LABORATORIOS ALTER, 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
Topiramato Alter 100 mg comprimidos revestidos por película	PT/H/320/01-04/MR	5014303	ALTER, S.A.	PT
Topiramato Alter 100 mg comprimidos revestidos por película	PT/H/0320/003	5014279	ALTER, S.A.	PT
Topiramato Alter 100 mg comprimidos revestidos por película	PT/H/0320/003	5014261	ALTER, S.A.	PT
Topiramato Alter 200 mg comprimidos recubiertos con película EFG	not available	69.883	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 200 mg comprimidos recubiertos con película EFG	not available	69.883	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 200 mg comprimidos revestidos por película	PT/H/320/01-04/MR	5014337	ALTER, S.A.	PT
Topiramato Alter 200 mg comprimidos revestidos por película	PT/H/0320/004	5014329	ALTER, S.A.	PT
Topiramato Alter 200 mg comprimidos revestidos por película	PT/H/0320/004	5014311	ALTER, S.A.	PT
Topiramato Alter 25 mg comprimidos recubiertos con película EFG	not available	69.880	LABORATORIOS ALTER, 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
Topiramato Alter 25 mg comprimidos recubiertos con película EFG	not available	69.880	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 25 mg comprimidos revestidos por película	PT/H/320/01-04/MR	5014204	ALTER, S.A.	PT
Topiramato Alter 25 mg comprimidos revestidos por película	PT/H/0320/001	5014220	ALTER, S.A.	PT
Topiramato Alter 25 mg comprimidos revestidos por película	PT/H/0320/001	5014212	ALTER, S.A.	PT
Topiramato Alter 25 mg comprimidos revestidos por película	PT/H/0320/001	5014220	ALTER, S.A.	PT
Topiramato Alter 25 mg comprimidos revestidos por película	PT/H/0320/001	5014212	ALTER, S.A.	PT
Topiramato Alter 50 mg comprimidos recubiertos con película EFG	not available	69.881	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 50 mg comprimidos recubiertos con película EFG	not available	69.881	LABORATORIOS ALTER, S.A.	ES
Topiramato Alter 50 mg comprimidos revestidos por película	PT/H/320/01-04/MR	5014253	ALTER, 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
Topiramato Alter 50 mg comprimidos revestidos por película	PT/H/0320/002	5014246	ALTER, S.A.	PT
Topiramato Alter 50 mg comprimidos revestidos por película	PT/H/0320/002	5014238	ALTER, S.A.	PT
Topiramato Amneal 100 mg comprimidos recubiertos con película EFG	UK/H/1166/003	73.002	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 100 mg comprimidos recubiertos con película EFG	UK/H/1166/003	73.002	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 200 mg comprimidos recubiertos con película EFG	UK/H/1166/004	72.999	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 200 mg comprimidos recubiertos con película EFG	UK/H/1166/004	72.999	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 25 mg comprimidos recubiertos con película EFG	UK/H/1166/001	72.998	AMNEAL PHARMA EUROPE LIMITED	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
Topiramato Amneal 25 mg comprimidos recubiertos con película EFG	UK/H/1166/001	72.998	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 50 mg comprimidos recubiertos con película EFG	UK/H/1166/002	73.000	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Amneal 50 mg comprimidos recubiertos con película EFG	UK/H/1166/002	73.000	AMNEAL PHARMA EUROPE LIMITED	ES
Topiramato Aurobindo 100 mg compresse rivestite con film	NL/H/2916/003	043081052	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 100 mg compresse rivestite con film	NL/H/2916/003	043081064	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 100 mg comprimidos recubiertos con película EFG	NL/H/2916/003	79249	LABORATORIOS AUROBINDO S.L.U.	ES
Topiramato Aurobindo 100 mg comprimidos revestidos por película	not available	5281241	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 100 mg comprimidos revestidos por película	not available	5281258	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
Topiramato Aurobindo 100 mg comprimidos revestidos por película	not available	5281266	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 100 mg comprimidos revestidos por película	not available	5281274	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 200 mg compresse rivestite con film	NL/H/2916/004	043081076	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 200 mg compresse rivestite con film	NL/H/2916/004	043081088	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 200 mg comprimidos recubiertos con película EFG	NL/H/2916/004	79250	LABORATORIOS AUROBINDO S.L.U.	ES
Topiramato Aurobindo 200 mg comprimidos revestidos por película	not available	5281308	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 200 mg comprimidos revestidos por película	not available	5281316	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 200 mg comprimidos revestidos por película	not available	5281324	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 200 mg comprimidos revestidos por película	not available	5281332	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
Topiramato Aurobindo 25 mg compresse rivestite con film	NL/H/2916/001	043081013	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 25 mg compresse rivestite con film	NL/H/2916/001	043081025	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 25 mg comprimidos recubiertos con película EFG	NL/H/2916/001	79247	LABORATORIOS AUROBINDO S.L.U.	ES
Topiramato Aurobindo 25 mg comprimidos revestidos por película	not available	5281142	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 25 mg comprimidos revestidos por película	not available	5281159	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 25 mg comprimidos revestidos por película	not available	5281167	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 25 mg comprimidos revestidos por película	not available	5281175	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 50 mg compresse rivestite con film	NL/H/2916/002	043081037	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Topiramato Aurobindo 50 mg compresse rivestite con film	NL/H/2916/002	043081049	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Topiramato Aurobindo 50 mg comprimidos recubiertos con película EFG	NL/H/2916/002	79248	LABORATORIOS AUROBINDO S.L.U.	ES
Topiramato Aurobindo 50 mg comprimidos revestidos por película	not available	5281209	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 50 mg comprimidos revestidos por película	not available	5281217	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 50 mg comprimidos revestidos por película	not available	5281225	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurobindo 50 mg comprimidos revestidos por película	not available	5281233	AUROBINDO PHARMA (PORTUGAL), UNIPessoal LDA	PT
Topiramato Aurovitás	PT/H/0231/003	5034772	ACTAVIS GROUP PTC EHF.	PT
Topiramato Aurovitás 200 mg Comprimidos revestidos por película	PT/H/0231/004	5035001	ACTAVIS GROUP PTC EHF.	PT
Topiramato Aurovitás 25 mg Comprimidos revestidos por película	PT/H/0231/001	5034475	ACTAVIS GROUP PTC EHF.	PT
Topiramato Aurovitás 50 mg Comprimidos revestidos por película	PT/H/0231/002	5034632	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
Topiramato Aurovitas Spain 100 mg comprimidos recubiertos con película EFG	not available	69.969	AUROVITAS SPAIN,S.A.U.	ES
Topiramato Aurovitas Spain 200 mg comprimidos recubiertos con película EFG	not available	69.970	AUROVITAS SPAIN,S.A.U.	ES
Topiramato Aurovitas Spain 25 mg comprimidos recubiertos con película EFG	not available	69.967	AUROVITAS SPAIN,S.A.U.	ES
Topiramato Aurovitas Spain 50 mg comprimidos recubiertos con película EFG	not available	69.968	AUROVITAS SPAIN,S.A.U.	ES
Topiramato Azevedos 100 mg comprimidos revestidos por película	not available	5122866	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 100 mg comprimidos revestidos por película	not available	5122775	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 100 mg comprimidos revestidos por película	not available	5122874	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 100 mg comprimidos revestidos por película	not available	5122767	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
Topiramato Azevedos 200 mg comprimidos revestidos por película	not available	5122916	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 200 mg comprimidos revestidos por película	not available	5122817	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 200 mg comprimidos revestidos por película	not available	5122908	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 200 mg comprimidos revestidos por película	not available	5122809	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 25 mg comprimidos revestidos por película	not available	5122825	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 25 mg comprimidos revestidos por película	not available	5122734	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 25 mg comprimidos revestidos por película	not available	5122833	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 25 mg comprimidos revestidos por película	not available	5122726	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 50 mg comprimidos revestidos por película	not available	5122841	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
Topiramato Azevedos 50 mg comprimidos revestidos por película	not available	5122759	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 50 mg comprimidos revestidos por película	not available	5122858	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Azevedos 50 mg comprimidos revestidos por película	not available	5122742	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Topiramato Bluefish 100 mg comprimidos recubiertos con película EFG	AT/H/0612/003	74580	BLUEFISH PHARMACEUTICALS AB	ES
Topiramato Bluefish 100 mg Comprimidos revestidos por película	AT/H/0612/003	5274717	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 100 mg Comprimidos revestidos por película	AT/H/0612/003	5274709	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 100 mg Comprimidos revestidos por película	AT/H/0612/003	5274667	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 100 mg Comprimidos revestidos por película	AT/H/0612/003	5274659	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 100 mg Comprimidos revestidos por película	AT/H/0612/003	5274675	BLUEFISH PHARMACEUTICALS AB	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
Topiramato Bluefish 200 mg comprimidos recubiertos con película EFG	AT/H/0612/004	73470	BLUEFISH PHARMACEUTICALS AB	ES
Topiramato Bluefish 200 mg Comprimidos revestidos por película	AT/H/0612/004	5274048	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 200 mg Comprimidos revestidos por película	AT/H/0612/004	5274758	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 200 mg Comprimidos revestidos por película	AT/H/0612/004	5274733	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 200 mg Comprimidos revestidos por película	AT/H/0612/004	5274725	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 200 mg Comprimidos revestidos por película	AT/H/0612/004	5274741	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 25 mg comprimidos recubiertos con película EFG	AT/H/0612/001	74579	BLUEFISH PHARMACEUTICALS AB	ES
Topiramato Bluefish 25 mg Comprimidos revestidos por película	AT/H/0612/001	5274576	BLUEFISH PHARMACEUTICALS AB	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
Topiramato Bluefish 25 mg Comprimidos revestidos por película	AT/H/0612/001	5274568	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 25 mg Comprimidos revestidos por película	AT/H/0612/001	5274543	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 25 mg Comprimidos revestidos por película	AT/H/0612/001	5274535	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 25 mg Comprimidos revestidos por película	AT/H/0612/001	5274550	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 50 mg comprimidos recubiertos con película EFG	AT/H/0612/002	74604	BLUEFISH PHARMACEUTICALS AB	ES
Topiramato Bluefish 50 mg Comprimidos revestidos por película	AT/H/0612/002	5274642	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 50 mg Comprimidos revestidos por película	AT/H/0612/002	5274634	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 50 mg Comprimidos revestidos por película	AT/H/0612/002	5274618	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluefish 50 mg Comprimidos revestidos por película	AT/H/0612/002	5274600	BLUEFISH PHARMACEUTICALS AB	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
Topiramato Bluefish 50 mg Comprimidos revestidos por película	AT/H/0612/002	5274626	BLUEFISH PHARMACEUTICALS AB	PT
Topiramato Bluepharma 100 mg comprimidos revestidos por película	not available	5061700	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 100 mg comprimidos revestidos por película	not available	5061718	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 200 mg comprimidos revestidos por película	not available	5061734	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 200 mg comprimidos revestidos por película	not available	5061726	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 25 mg comprimidos revestidos por película	not available	5061650	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 25 mg comprimidos revestidos por película	not available	5061627	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 25 mg comprimidos revestidos por película	not available	5061635	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 25 mg comprimidos revestidos por película	not available	5061643	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, 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
Topiramato Bluepharma 50 mg comprimidos revestidos por película	not available	5061676	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Bluepharma 50 mg comprimidos revestidos por película	not available	5061668	BLUEPHARMA GENÉRICOS - COMÉRCIO DE MEDICAMENTOS, S.A.	PT
Topiramato Ciclum 100 mg comprimidos revestidos por película	not available	5566112	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 100 mg comprimidos revestidos por película	not available	12/H/0216/003	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 100 mg comprimidos revestidos por película	not available	12/H/0216/003	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 200 mg comprimidos revestidos por película	not available	5566120	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 200 mg comprimidos revestidos por película	not available	12/H/0216/004	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 200 mg comprimidos revestidos por película	not available	12/H/0216/004	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 25 mg comprimidos revestidos por película	not available	5566062	CICLUM FARMA 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
Topiramato Ciclum 25 mg comprimidos revestidos por película	not available	5566070	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 25 mg comprimidos revestidos por película	not available	12/H/0216/001	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 50 mg comprimidos revestidos por película	not available	5566104	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 50 mg comprimidos revestidos por película	not available	12/H/0216/002	CICLUM FARMA UNIPessoal LDA.	PT
Topiramato Ciclum 50 mg comprimidos revestidos por película	not available	12/H/0216/002	CICLUM FARMA UNIPessoal LDA.	PT
TOPIRAMATO CINFA 100 MG COMPRIMIDOS RECUBIERTOS CON PELÍCULA EFG	not available	69137	LABORATORIOS CINFA, S.A.	ES
Topiramato Cinfa 100 mg Comprimidos revestidos por película	not available	5128079	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 100 mg Comprimidos revestidos por película	not available	5128103	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 100 mg Comprimidos revestidos por película	not available	5128111	CINFA PORTUGAL, LDA.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramato Cinfa 100 mg Comprimidos revestidos por película	not available	5128129	CINFA PORTUGAL, LDA.	PT
TOPIRAMATO CINFA 200 MG COMPRIMIDOS RECUBIERTOS CON PELÍCULA EFG	not available	69138	LABORATORIOS CINFA, S.A.	ES
TOPIRAMATO CINFA 25 MG COMPRIMIDOS RECUBIERTOS CON PELÍCULA EFG	not available	69135	LABORATORIOS CINFA, S.A.	ES
Topiramato Cinfa 25 mg Comprimidos revestidos por película	not available	5127972	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 25 mg Comprimidos revestidos por película	not available	5128004	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 25 mg Comprimidos revestidos por película	not available	5128012	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 25 mg Comprimidos revestidos por película	not available	5128020	CINFA PORTUGAL, LDA.	PT
TOPIRAMATO CINFA 50 MG COMPRIMIDOS RECUBIERTOS CON PELÍCULA EFG	not available	69136	LABORATORIOS CINFA, 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
Topiramato Cinfa 50 mg Comprimidos revestidos por película	not available	5128038	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 50 mg Comprimidos revestidos por película	not available	5128046	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 50 mg Comprimidos revestidos por película	not available	5128053	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfa 50 mg Comprimidos revestidos por película	not available	5128061	CINFA PORTUGAL, LDA.	PT
Topiramato Cinfamed 100 mg comprimidos recubiertos con película EFG	not available	74.211	LABORATORIOS CINFA, S.A.	ES
Topiramato Cinfamed 200 mg comprimidos recubiertos con película EFG	not available	74.212	LABORATORIOS CINFA, S.A.	ES
Topiramato Cinfamed 25 mg comprimidos recubiertos con película EFG	not available	74.206	LABORATORIOS CINFA, S.A.	ES
Topiramato Cinfamed 50 mg comprimidos recubiertos con película EFG	not available	74.210	LABORATORIOS CINFA, 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
Topiramato Clindonim 100 mg Comprimidos revestidos por película	PT/H/0199/003	5078043	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 100 mg Comprimidos revestidos por película	PT/H/0199/003	5078050	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 200 mg Comprimidos revestidos por película	PT/H/0199/004	5078068	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 200 mg Comprimidos revestidos por película	PT/H/0199/004	5078076	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 25 mg Comprimidos revestidos por película	PT/H/0199/001	5078001	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 25 mg Comprimidos revestidos por película	PT/H/0199/001	5078019	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 50 mg Comprimidos revestidos por película	PT/H/0199/002	5078027	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Clindonim 50 mg Comprimidos revestidos por película	PT/H/0199/002	5078035	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
TOPIRAMATO DOC Generici 100 mg, compresse rivestite con film	NL/H/1231/003	038340117	DOC GENERICI S.R.L.	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
TOPIRAMATO DOC Generici 100 mg, compresse rivestite con film	NL/H/1231/003	038340129	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 200 mg, compresse rivestite con film	NL/H/1231/004	038340168	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 200 mg, compresse rivestite con film	NL/H/1231/004	038340170	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 25 mg, compresse rivestite con film	NL/H/1231/001	038340016	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 25 mg, compresse rivestite con film	NL/H/1231/001	038340028	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 50 mg, compresse rivestite con film	NL/H/1231/002	038340067	DOC GENERICI S.R.L.	IT
TOPIRAMATO DOC Generici 50 mg, compresse rivestite con film	NL/H/1231/002	038340079	DOC GENERICI S.R.L.	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
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314163	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314151	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314175	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314148	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314199	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314187	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314377	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314391	EG SPA	IT
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314403	EG SPA	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
TOPIRAMATO EG 100 mg compresse rivestite con film	DE/H/0889/003	038314136	EG SPA	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314213	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314237	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314225	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314249	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314415	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314427	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314441	EG S.P.A.	IT
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314439	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
TOPIRAMATO EG 200 mg compresse rivestite con film	DE/H/0889/004	038314201	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314050	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314023	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314062	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314047	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314035	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314252	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314264	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314290	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
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314276	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314288	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314302	EG S.P.A.	IT
TOPIRAMATO EG 25 mg compresse rivestite con film	DE/H/0889/001	038314011	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314112	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314098	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314100	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314124	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314086	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
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314365	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314340	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314314	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314338	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314353	EG S.P.A.	IT
TOPIRAMATO EG 50 mg compresse rivestite con film	DE/H/0889/002	038314074	EG S.P.A.	IT
Topiramato Farmalider 100 mg comprimidos recubiertos con película EFG	not available	68.687	FARMALIDER, S.A.	ES
Topiramato Farmalider 200 mg comprimidos recubiertos con película EFG	not available	68.688	FARMALIDER, 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
Topiramato Farmalider 25 mg comprimidos recubiertos con película EFG	not available	68.685	FARMALIDER, S.A.	ES
Topiramato Farmalider 50 mg comprimidos recubiertos con película EFG	not available	68.686	FARMALIDER, S.A.	ES
Topiramato Farnoz 100 mg comprimido revestido por película	not available	5078241	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farnoz 100 mg comprimido revestido por película	not available	5078258	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farnoz 200 mg comprimido revestido por película	not available	5078266	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farnoz 200 mg comprimido revestido por película	not available	5078274	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farnoz 25 mg comprimido revestido por película	not available	5078209	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farnoz 25 mg comprimido revestido por película	not available	5078217	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, 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
Topiramato Farmoz 50 mg comprimido revestido por película	not available	5078225	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Farmoz 50 mg comprimido revestido por película	not available	5078233	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077417	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077425	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077433	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077441	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077458	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077466	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077474	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
Topiramato Genedec 100 mg comprimidos revestidos por película	not available	5077409	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077516	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077524	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077532	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077540	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077557	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077565	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077573	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 200 mg comprimidos revestidos por película	not available	5077508	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
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077219	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077227	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077235	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077243	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077250	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077268	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077276	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 25 mg comprimidos revestidos por película	not available	5077201	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077318	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
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077326	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077334	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077342	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077359	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077367	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077375	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Genedec 50 mg comprimidos revestidos por película	not available	5077300	GENEDEC - MEDICAMENTOS GENÉRICOS, LDA.	PT
Topiramato Generis 100 mg comprimidos, 100 mg, comprimidos revestidos por película	not available	5849591	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 100 mg comprimidos, 100 mg, comprimidos	not available	5849690	GENERIS FARMACÊUTICA, S.A.	PT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
revestidos por película				
Topiramato Generis 100 mg comprimidos, 100 mg, comprimidos revestidos por película	not available	5849393	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 100 mg comprimidos, 100 mg, comprimidos revestidos por película	not available	5849492	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 200 mg comprimidos, 200 mg, comprimidos revestidos por película	not available	5849997	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 200 mg comprimidos, 200 mg, comprimidos revestidos por película	not available	5850094	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 200 mg comprimidos, 200 mg, comprimidos revestidos por película	not available	5849799	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 200 mg comprimidos, 200 mg, comprimidos revestidos por película	not available	5849898	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
Topiramato Generis 25 mg comprimidos, 25 mg, comprimidos revestidos por película	not available	5848791	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 25 mg comprimidos, 25 mg, comprimidos revestidos por película	not available	5848890	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 25 mg comprimidos, 25 mg, comprimidos revestidos por película	not available	5848593	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 25 mg comprimidos, 25 mg, comprimidos revestidos por película	not available	5848692	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 50 mg comprimidos, 50 mg, comprimidos revestidos por película	not available	5849195	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 50 mg comprimidos, 50 mg, comprimidos revestidos por película	not available	5849294	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Generis 50 mg comprimidos, 50 mg, comprimidos revestidos por película	not available	5848999	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
Topiramato Generis 50 mg comprimidos, 50 mg, comprimidos revestidos por película	not available	5849096	GENERIS FARMACÊUTICA, S.A.	PT
TOPIRAMATO GP 100 mg comprimidos revestidos por película	not available	5116074	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 100 mg comprimidos revestidos por película	not available	5116108	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 200 mg comprimidos revestidos por película	not available	5116116	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 200 mg comprimidos revestidos por película	not available	5116124	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 25 mg comprimidos revestidos por película	not available	5116033	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 25 mg comprimidos revestidos por película	not available	5116041	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 50 mg comprimidos revestidos por película	not available	5116058	GP – GENÉRICOS PORTUGUESES, LDA.	PT
TOPIRAMATO GP 50 mg comprimidos revestidos por película	not available	5116066	GP – GENÉRICOS PORTUGUESES, 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
Topiramato KERN PHARMA 100 mg comprimidos recubiertos con película EFG	not available	68.723	KERN PHARMA, S.L.	ES
Topiramato KERN PHARMA 200 mg comprimidos recubiertos con película EFG	not available	68.683	KERN PHARMA, S.L.	ES
Topiramato KERN PHARMA 25 mg comprimidos recubiertos con película EFG	not available	68.681	KERN PHARMA, S.L.	ES
Topiramato KERN PHARMA 50 mg comprimidos recubiertos con película EFG	not available	68.682	KERN PHARMA, S.L.	ES
Topiramato Labesfal 100 mg comprimidos revestidos por película	not available	5103338	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 100 mg comprimidos revestidos por película	not available	5103346	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 100 mg comprimidos revestidos por película	not available	5103437	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 100 mg comprimidos revestidos por película	not available	5103445	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
Topiramato Labesfal 200 mg comprimidos revestidos por película	not available	5103353	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 200 mg comprimidos revestidos por película	not available	5103452	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 200 mg comprimidos revestidos por película	not available	5103353	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 25 mg comprimidos revestidos por película	not available	5103270	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 25 mg comprimidos revestidos por película	not available	5103304	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 25 mg comprimidos revestidos por película	not available	5103379	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 25 mg comprimidos revestidos por película	not available	5103403	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 50 mg comprimidos revestidos por película	not available	5103312	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 50 mg comprimidos revestidos por película	not available	5103320	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
Topiramato Labesfal 50 mg comprimidos revestidos por película	not available	5103411	GENERIS FARMACÊUTICA, S.A.	PT
Topiramato Labesfal 50 mg comprimidos revestidos por película	not available	5103429	GENERIS FARMACÊUTICA, S.A.	PT
TOPIRAMATO Mylan 100 mg comprimidos recubiertos con película EFG	not available	69036	MYLAN PHARMACEUTICALS S.L.	ES
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040647	MYLAN, LDA	PT
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040654	MYLAN, LDA	PT
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040621	MYLAN, LDA	PT
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040639	MYLAN, LDA	PT
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040662	MYLAN, LDA	PT
Topiramato Mylan 100 mg comprimidos revestidos por película	not available	5040670	MYLAN, 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
TOPIRAMATO Mylan 200 mg comprimidos recubiertos con película EFG	not available	69037	MYLAN PHARMACEUTICALS S.L.	ES
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040720	MYLAN, LDA	PT
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040738	MYLAN, LDA	PT
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040704	MYLAN, LDA	PT
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040712	MYLAN, LDA	PT
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040746	MYLAN, LDA	PT
Topiramato Mylan 200 mg comprimidos revestidos por película	not available	5040753	MYLAN, LDA	PT
TOPIRAMATO Mylan 25 mg comprimidos recubiertos con película EFG	not available	69034	MYLAN PHARMACEUTICALS S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040506	MYLAN, LDA	PT
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040514	MYLAN, LDA	PT
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040464	MYLAN, LDA	PT
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040472	MYLAN, LDA	PT
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040522	MYLAN, LDA	PT
Topiramato Mylan 25 mg comprimidos revestidos por película	not available	5040530	MYLAN, LDA	PT
TOPIRAMATO Mylan 50 mg comprimidos recubiertos con película EFG	not available	69035	MYLAN PHARMACEUTICALS S.L.	ES
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040548	MYLAN, LDA	PT
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040555	MYLAN, 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
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040563	MYLAN, LDA	PT
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040571	MYLAN, LDA	PT
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040605	MYLAN, LDA	PT
Topiramato Mylan 50 mg comprimidos revestidos por película	not available	5040613	MYLAN, LDA	PT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755218/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755220/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755232/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755244/M	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
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755257/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755269/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755271/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755283/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755295/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 100 mg compresse rivestite con film	NL/H/0717/003	037755307/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755319/M	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
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755321/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755333/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755345/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755358/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755360/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755372/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755384/M	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
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755396/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 200 mg compresse rivestite con film	NL/H/0717/004	037755408/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755016/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755028/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755030/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755042/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755055/M	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
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755067/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755079/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755081/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755093/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 25 mg compresse rivestite con film	NL/H/0717/001	037755105/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755117/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755129/M	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
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755131/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755143/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755156/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755168/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755170/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755182/M	MYLAN S.P.A.	IT
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755194/M	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
Topiramato Mylan Generics 50 mg compresse rivestite con film	NL/H/0717/002	037755206/M	MYLAN S.P.A.	IT
Topiramato Normon 100 mg comprimidos recubiertos con película EFG	not available	70765	LABORATORIOS NORMON, S.A.	ES
Topiramato NORMON 100 mg comprimidos revestidos por película	not available	5129473	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 100 mg comprimidos revestidos por película	not available	5129515	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 100 mg comprimidos revestidos por película	not available	5129507	LABORATÓRIOS NORMON, S.A.	PT
Topiramato Normon 200 mg comprimidos recubiertos con película EFG	not available	70766	LABORATORIOS NORMON, S.A.	ES
Topiramato NORMON 200 mg comprimidos revestidos por película	not available	5129523	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 200 mg comprimidos revestidos por película	not available	5129549	LABORATÓRIOS NORMON, 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
Topiramato NORMON 200 mg comprimidos revestidos por película	not available	5129531	LABORATÓRIOS NORMON, S.A.	PT
Topiramato Normon 25 mg comprimidos recubiertos con película EFG	not available	70753	LABORATORIOS NORMON, S.A.	ES
Topiramato NORMON 25 mg comprimidos revestidos por película	not available	5128137	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 25 mg comprimidos revestidos por película	not available	5129432	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 25 mg comprimidos revestidos por película	not available	5129424	LABORATÓRIOS NORMON, S.A.	PT
Topiramato Normon 50 mg comprimidos recubiertos con película EFG	not available	70764	LABORATORIOS NORMON, S.A.	ES
Topiramato NORMON 50 mg comprimidos revestidos por película	not available	5129440	LABORATÓRIOS NORMON, S.A.	PT
Topiramato NORMON 50 mg comprimidos revestidos por película	not available	5129465	LABORATÓRIOS NORMON, 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
Topiramato NORMON 50 mg comprimidos revestidos por película	not available	5129457	LABORATÓRIOS NORMON, S.A.	PT
Topiramato Pensa Pharma 100 mg comprimidos recubiertos con película EFG	not available	69.425	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 100 mg comprimidos recubiertos con película EFG	not available	69.425	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 200 mg comprimidos recubiertos con película EFG	not available	69.426	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 200 mg comprimidos recubiertos con película EFG	not available	69.426	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 25 mg comprimidos recubiertos con película EFG	not available	69.423	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 25 mg comprimidos recubiertos con película EFG	not available	69.423	PENSA PHARMA, 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
Topiramato Pensa Pharma 50 mg comprimidos recubiertos con película EFG	not available	69.424	PENSA PHARMA, S.A	ES
Topiramato Pensa Pharma 50 mg comprimidos recubiertos con película EFG	not available	69.424	PENSA PHARMA, S.A	ES
Topiramato Pharma Combix 100 mg comprimidos recubiertos con película EFG	not available	73.765	LABORATORIOS COMBIX, S.L.U.	ES
Topiramato Pharma Combix 200 mg comprimidos recubiertos con película EFG	not available	73.761	LABORATORIOS COMBIX, S.L.U.	ES
Topiramato Pharma Combix 50 mg comprimidos recubiertos con película EFG	not available	73.763	LABORATORIOS COMBIX, S.L.U.	ES
Topiramato Qualigen 100 mg comprimidos recubiertos con película EFG	DK/H/1354/003	70890	QUALIGEN, S.L.	ES
Topiramato Qualigen 200 mg comprimidos recubiertos con película EFG	DK/H/1354/004	70.891	QUALIGEN, S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramato Qualigen 25 mg comprimidos recubiertos con película EFG	DK/H/1354/001	70.868	QUALIGEN, S.L.	ES
Topiramato Qualigen 50 mg comprimidos recubiertos con película EFG	DK/H/1354/002	70.889	QUALIGEN, S.L.	ES
Topiramato ratiopharm 100 mg comprimidos recubiertos con película EFG	DE/H/2064/004	67991	RATIOPHARM ESPANA SA	ES
Topiramato ratiopharm 100 mg comprimidos revestidos por película	not available	5423843	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato ratiopharm 100 mg comprimidos revestidos por película	not available	5423850	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato ratiopharm 200 mg comprimidos recubiertos con película EFG	DE/H/2064/005	67992	RATIOPHARM ESPANA SA	ES
Topiramato ratiopharm 25 mg comprimidos recubiertos con película EFG	DE/H/2064/002	67.989	RATIOPHARM ESPANA SA	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
Topiramato ratiopharm 25 mg comprimidos revestidos por película	not available	5423819	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato ratiopharm 25 mg comprimidos revestidos por película	not available	5423801	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato ratiopharm 50 mg comprimidos recubiertos con película EFG	DE/H/2064/003	67990	RATIOPHARM ESPANA SA	ES
Topiramato ratiopharm 50 mg comprimidos revestidos por película	not available	5423827	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato ratiopharm 50 mg comprimidos revestidos por película	not available	5423835	RATIOPHARM-COMERCIO E INDUSTRIA DE PRODUTOS FARMACEUTICOS LDA	PT
Topiramato Ritisca 100 mg comprimidos revestidos por película	not available	5655279	AUROVITAS UNIPessoal, LDA.	PT
Topiramato Ritisca 200 mg comprimidos revestidos por película	not available	5655303	AUROVITAS UNIPessoal, LDA.	PT
Topiramato Ritisca 25 mg comprimidos revestidos por película	not available	5655246	AUROVITAS UNIPessoal, LDA.	PT
Topiramato Ritisca 25 mg comprimidos revestidos por película	not available	5655253	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
Topiramato Ritisca 50 mg comprimidos revestidos por película	not available	5655261	AUROVITAS UNIPessoal, LDA.	PT
TOPIRAMATO SANDOZ 100 MG COMPRIMIDOS	FI/H/0599/003	5267646	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 100 MG COMPRIMIDOS	FI/H/0599/003	5937388	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 100 MG COMPRIMIDOS	FI/H/0599/003	5937487	SANDOZ FARMACÊUTICA LDA.	PT
Topiramato Sandoz 200 mg comprimidos recubiertos con película EFG	not available	69356	SANDOZ FARMACÊUTICA, S.A.	ES
TOPIRAMATO SANDOZ 25 MG CÁPSULAS	FI/H/0599/006	5937982	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 25 MG CÁPSULAS	FI/H/0599/006	5938089	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 25 MG COMPRIMIDOS	FI/H/0599/001	5937081	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 25 MG COMPRIMIDOS	FI/H/0599/001	5936984	SANDOZ FARMACÊUTICA 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
TOPIRAMATO SANDOZ 25 MG COMPRIMIDOS	FI/H/0599/001	5267620	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 25 MG COMPRIMIDOS	FI/H/0599/001	5267612	SANDOZ FARMACÊUTICA LDA.	PT
Topiramato Sandoz 25 mg comprimidos recubiertos con película EFG	not available	69178	SANDOZ FARMACÊUTICA, S.A.	ES
TOPIRAMATO SANDOZ 50 MG CÁPSULAS	FI/H/0599/007	5938188	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 50 MG CÁPSULAS	FI/H/0599/007	5938287	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 50 MG COMPRIMIDOS	FI/H/0599/002	5267638	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 50 MG COMPRIMIDOS	FI/H/0599/002	5937180	SANDOZ FARMACÊUTICA LDA.	PT
TOPIRAMATO SANDOZ 50 MG COMPRIMIDOS	FI/H/0599/002	5937289	SANDOZ FARMACÊUTICA LDA.	PT
Topiramato Sandoz 50 mg comprimidos recubiertos con película EFG	not available	69179	SANDOZ FARMACÊUTICA, 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
Topiramato Stada 100 mg comprimidos	not available	04/H/0107/003	STADA LDA.	PT
Topiramato Stada 100 mg comprimidos	not available	5694781	STADA LDA.	PT
Topiramato Stada 100 mg comprimidos	not available	5694682	STADA LDA.	PT
Topiramato STADA 100 mg comprimidos recubiertos con película EFG	not available	68.593	LABORATORIO STADA, S.L.	ES
Topiramato Stada 200 mg comprimidos	not available	5694880	STADA LDA.	PT
Topiramato Stada 200 mg comprimidos	not available	5694989	STADA LDA.	PT
Topiramato STADA 200 mg comprimidos recubiertos con película EFG	not available	68.594	LABORATORIO STADA, S.L.	ES
Topiramato Stada 25 mg comprimidos	not available	5694286	STADA LDA.	PT
Topiramato Stada 25 mg comprimidos	not available	5694385	STADA 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
Topiramato STADA 25 mg comprimidos recubiertos con película EFG	not available	68.591	LABORATORIO STADA, S.L.	ES
Topiramato Stada 50 mg comprimidos	not available	5694484	STADA LDA.	PT
Topiramato Stada 50 mg comprimidos	not available	5694583	STADA LDA.	PT
Topiramato STADA 50 mg comprimidos recubiertos con película EFG	not available	68.592	LABORATORIO STADA, S.L.	ES
TOPIRAMATO TARBIS 100 mg comprimidos recubiertos con película EFG	not available	69887	TARBIS FARMA, S.L.	ES
TOPIRAMATO TARBIS 200 mg comprimidos recubiertos con película EFG	not available	69.888	TARBIS FARMA, S.L.	ES
TOPIRAMATO TARBIS 25 mg comprimidos recubiertos con película EFG	not available	69.884	TARBIS FARMA, S.L.	ES
TOPIRAMATO TARBIS 50 mg comprimidos recubiertos con película EFG	not available	69.889	TARBIS FARMA, S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramato Tecnigen 100 mg compresse rivestite con film	PT/H/0199/003	039226055	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato Tecnigen 100 mg compresse rivestite con film	PT/H/0199/003	039226067	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato TecniGen 100 mg comprimidos recubiertos con película EFG	not available	70646	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Topiramato Tecnigen 200 mg compresse rivestite con film	PT/H/0199/004	039226081	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato Tecnigen 200 mg compresse rivestite con film	PT/H/0199/004	039226079	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato TecniGen 200 mg comprimidos recubiertos con película EFG	not available	70647	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Topiramato Tecnigen 25 mg compresse rivestite con film	PT/H/0199/001	039226016	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato Tecnigen 25 mg compresse rivestite con film	PT/H/0199/001	039226028	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
Topiramato TechniGen 25 mg comprimidos recubiertos con película EFG	not available	70644	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Topiramato Technigen 50 mg compresse rivestite con film	PT/H/0199/002	039226042	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato Technigen 50 mg compresse rivestite con film	PT/H/0199/002	039226030	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	IT
Topiramato TechniGen 50 mg comprimidos recubiertos con película EFG	not available	70645	TECNIMEDE ESPAÑA IND. FCA., S.A.	ES
Topiramato Technimede 100 mg Comprimidos revestidos por película	PT/H/0198/003	5079074	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Technimede 100 mg Comprimidos revestidos por película	PT/H/0198/003	5079108	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Technimede 200 mg Comprimidos revestidos por película	PT/H/0198/004	5079116	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Technimede 200 mg Comprimidos revestidos por película	PT/H/0198/004	5079124	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	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
Topiramato Tecnimed 25 mg Comprimidos revestidos por película	PT/H/0198/001	5078407	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Tecnimed 25 mg Comprimidos revestidos por película	PT/H/0198/001	5078415	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Tecnimed 50 mg Comprimidos revestidos por película	PT/H/0198/002	5079058	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Tecnimed 50 mg Comprimidos revestidos por película	PT/H/0198/002	5079066	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974299	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974224	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974275	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974325	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974248	TEVA ITALIA S.R.L.	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
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974263	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974301	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974287	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974236	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974313	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg compresse rivestite con film	NO/H/0136/003	038974251	TEVA ITALIA S.R.L.	IT
Topiramato Teva 100 mg comprimidos recubiertos con película EFG	not available	68.932	TEVA PHARMA S.L.U	ES
Topiramato Teva 100 mg comprimidos revestidos por película	NO/H/0136/003	5108543	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 100 mg comprimidos revestidos por película	NO/H/0136/003	5108535	TEVA PHARMA – PRODUTOS FARMACÊUTICOS 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
Topiramato Teva 100 mg comprimidos revestidos por película	NO/H/0136/003	5108550	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974402	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974390	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974414	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974388	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974426	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974352	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974364	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974438	TEVA ITALIA S.R.L.	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
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974376	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974337	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg compresse rivestite con film	NO/H/0136/004	038974349	TEVA ITALIA S.R.L.	IT
Topiramato Teva 200 mg comprimidos recubiertos con película EFG	not available	68.933	TEVA PHARMA S.L.U	ES
Topiramato Teva 200 mg comprimidos revestidos por película	NO/H/0136/004	5108576	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 200 mg comprimidos revestidos por película	NO/H/0136/004	5108600	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 200 mg comprimidos revestidos por película	NO/H/0136/004	5108568	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974010	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974059	TEVA ITALIA S.R.L.	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
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974022	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/013/001	038974073	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974109	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974085	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974046	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974097	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974034	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg compresse rivestite con film	NO/H/0136/001	038974061	TEVA ITALIA S.R.L.	IT
Topiramato Teva 25 mg comprimidos recubiertos con película EFG	not available	68.930	TEVA PHARMA S.L.U	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
Topiramato Teva 25 mg comprimidos revestidos por película	NO/H/0136/001	5108469	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 25 mg comprimidos revestidos por película	NO/H/0136/001	5108451	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 25 mg comprimidos revestidos por película	NO/H/0136/001	5108477	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974198	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974162	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974186	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974147	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974150	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974123	TEVA ITALIA S.R.L.	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
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974200	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974174	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974111	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974135	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg compresse rivestite con film	NO/H/0136/002	038974212	TEVA ITALIA S.R.L.	IT
Topiramato Teva 50 mg comprimidos recubiertos con película EFG	not available	68.931	TEVA PHARMA S.L.U	ES
Topiramato Teva 50 mg comprimidos revestidos por película	NO/H/0136/002	5108519	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 50 mg comprimidos revestidos por película	NO/H/0136/002	5108527	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Topiramato Teva 50 mg comprimidos revestidos por película	NO/H/0136/002	5108501	TEVA PHARMA – PRODUTOS FARMACÊUTICOS 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
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5857784	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5857883	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5010970	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5011002	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5857784	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5857883	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5010970	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 100 mg comprimidos revestidos por película	not available	5011002	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5858188	TOLIFE - PRODUTOS FARMACÊUTICOS, 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
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5858287	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5011010	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5011028	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5858188	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5858287	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5011010	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 200 mg comprimidos revestidos por película	not available	5011028	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5856984	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5857081	TOLIFE - PRODUTOS FARMACÊUTICOS, 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
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5010939	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5010947	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5856984	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5857081	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5010939	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 25 mg comprimidos revestidos por película	not available	5010947	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5857388	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5857487	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5010954	TOLIFE - PRODUTOS FARMACÊUTICOS, 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
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5010962	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5857388	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5857487	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5010954	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato toLife 50 mg comprimidos revestidos por película	not available	5010962	TOLIFE - PRODUTOS FARMACÊUTICOS, S.A.	PT
Topiramato Tomix 100 mg comprimido revestido por película	not available	5078340	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 100 mg comprimido revestido por película	not available	5078357	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 200 mg comprimido revestido por película	not available	5078365	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 200 mg comprimido revestido por película	not available	5078373	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL 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
Topiramato Tomix 25 mg comprimido revestido por película	not available	5078308	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 25 mg comprimido revestido por película	not available	5078316	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 50 mg comprimido revestido por película	not available	5078324	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Tomix 50 mg comprimido revestido por película	not available	5078332	PENTAFARMA - SOCIEDADE TECNICO-MEDICINAL S.A.	PT
Topiramato Urlabs 100 mg comprimidos recubiertos con película EFG	PT/H/0320/003	76385	ARISTO PHARMA IBERIA, S.L.	ES
Topiramato Urlabs 200 mg comprimidos recubiertos con película EFG	PT/H/320/004	76386	ARISTO PHARMA IBERIA, S.L.	ES
Topiramato Urlabs 25mg comprimidos recubiertos con película EFG	PT/H/320/001	74678	ARISTO PHARMA IBERIA, S.L.	ES
Topiramato Urlabs 50mg comprimidos recubiertos con película EFG	PT/H/320/002	76388	ARISTO PHARMA IBERIA, S.L.	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041058	AXONE, LDA.	PT
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041041	AXONE, LDA.	PT
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041017	AXONE, LDA.	PT
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041009	AXONE, LDA.	PT
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041033	AXONE, LDA.	PT
Topiramato Wynn 100 mg Comprimidos revestidos por película	not available	5041025	AXONE, LDA.	PT
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041124	AXONE, LDA.	PT
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041074	AXONE, LDA.	PT
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041066	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
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041116	AXONE, LDA.	PT
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041108	AXONE, LDA.	PT
Topiramato Wynn 200 mg Comprimidos revestidos por película	not available	5041132	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040910	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040902	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040779	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040878	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040803	AXONE, LDA.	PT
Topiramato Wynn 25 mg Comprimidos revestidos por película	not available	5040761	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
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040977	AXONE, LDA.	PT
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040969	AXONE, LDA.	PT
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040936	AXONE, LDA.	PT
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040928	AXONE, LDA.	PT
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040951	AXONE, LDA.	PT
Topiramato Wynn 50 mg Comprimidos revestidos por película	not available	5040944	AXONE, LDA.	PT
Topiramato Zentiva 100 mg compresse rivestite con film	DE/H/1031/003	038475253	ZENTIVA ITALIA SRL	IT
Topiramato Zentiva 100 mg comprimidos recubiertos con película EFG	UK/H/1483/003	73.224	ZENTIVA, K.S.	ES
Topiramato Zentiva 100 mg comprimidos recubiertos con película	UK/H/1483/003	73.224	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
EFG				
Topiramato Zentiva 200 mg compresse rivestite con film	DE/H/1031/004	038475354	ZENTIVA ITALIA SRL	IT
Topiramato Zentiva 200 mg comprimidos recubiertos con película EFG	UK/H/1483/004	73.225	ZENTIVA, K.S.	ES
Topiramato Zentiva 200 mg comprimidos recubiertos con película EFG	UK/H/1483/004	73.225	ZENTIVA, K.S.	ES
Topiramato Zentiva 25 mg compresse rivestite con film	DE/H/1031/001	038475051	ZENTIVA ITALIA SRL	IT
Topiramato Zentiva 25 mg comprimidos recubiertos con película EFG	UK/H/1483/001	73.222	ZENTIVA, K.S.	ES
Topiramato Zentiva 25 mg comprimidos recubiertos con película EFG	UK/H/1483/001	73.222	ZENTIVA, K.S.	ES
Topiramato Zentiva 50 mg compresse rivestite con film	DE/H/1031/002	038475152	ZENTIVA ITALIA SRL	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
Topiramato Zentiva 50 mg comprimidos recubiertos con película EFG	UK/H/1483/002	73.223	ZENTIVA, K.S.	ES
Topiramato Zentiva 50 mg comprimidos recubiertos con película EFG	UK/H/1483/002	73.223	ZENTIVA, K.S.	ES
Topiramat-ratiopharm 100 mg Filmtabletten	DE/H/2064/004	1-26517	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Topiramat-ratiopharm 200 mg Filmtabletten	DE/H/2064/005	1-26518	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Topiramat-ratiopharm 25 mg Filmtabletten	DE/H/2064/002	1-26515	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Topiramat-ratiopharm 50 mg Filmtabletten	DE/H/2064/003	1-26516	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Topiramat-ratiopharm® 100 mg Filmtabletten	DE/H/2064/003	65333.00.00	RATIOPHARM GMBH	DE
Topiramat-ratiopharm® 200 mg Filmtabletten	DE/H/2064/005	65334.00.00	RATIOPHARM GMBH	DE
Topiramat-ratiopharm® 25 mg Filmtabletten	DE/H/2064/002	65331.00.00	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
Topiramat-ratiopharm® 50 mg Filmtabletten	DE/H/2064/003	65332.00.00	RATIOPHARM GMBH	DE
Topiramat-TEVA® 100 mg Filmtabletten	NO/H/0136/003	71473.00.00	TEVA GMBH	DE
Topiramat-TEVA® 25 mg Filmtabletten	NO/H/0136/001	71471.00.00	TEVA GMBH	DE
Topiramat-TEVA® 50 mg Filmtabletten	NO/H/0136/002	71472.00.00	TEVA GMBH	DE
Topiratore, filmovertrukne tabletter	NL/H/0947/003	40370	STADA ARZNEIMITTEL AG	DK
Topiratore, filmovertrukne tabletter	NL/H/0947/001	40368	STADA ARZNEIMITTEL AG	DK
Topiratore, filmovertrukne tabletter	NL/H/0947/002	40369	STADA ARZNEIMITTEL AG	DK
Topiratore, filmovertrukne tabletter	NL/H/0947/004	40371	STADA ARZNEIMITTEL AG	DK
Topit 100 mg Film-Coated Tablets	FI/H/0599/003	PA0711/111/003	ROWEX 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
Topit 200 mg Film-Coated Tablets	FI/H/0599/004	PA 711/111/4	ROWEX LTD	IE
Topit 25 mg Film-Coated Tablets	FI/H/0599/001	PA 0711/111/001	ROWEX LTD	IE
Topit 50 mg Film-Coated Tablets	FI/H/0599/002	PA 0711/111/002	ROWEX LTD	IE
Toramat 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/003	20244	PHARMATHEN S.A.	CY
Toramat 200 mg επιζυαζιμυζα με ζεπτζ ζμζζζζ δζςζζα	DK/H/0930/004	20245	PHARMATHEN S.A.	CY
Toramat 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/004	40327/18-6-2008	PHARMATHEN S.A.	GR
Toramat 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/001	20242	PHARMATHEN S.A.	CY
Toramat 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/001	40324/18-6-2008	PHARMATHEN S.A.	GR
Toramat 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/002	20243	PHARMATHEN S.A.	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
Toramat 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/002	40325/18-6-2008	PHARMATHEN S.A.	GR
Toramat, 100 mg, tabletki powlekane	DE/H/0984/003	14937	GLENMARK PHARMACEUTICALS S.R.O.	PL
Toramat, 100mg επικαλυμμένα με λεπτό υμένιο δισκία	DK/H/0930/003	40326/18-6-2008	PHARMATHEN S.A.	GR
Toramat, 200 mg, tabletki powlekane	DE/H/0984/004	14938	GLENMARK PHARMACEUTICALS S.R.O.	PL
Toramat, 25 mg, tabletki powlekane	DE/H/0984/001	14935	GLENMARK PHARMACEUTICALS S.R.O.	PL
Toramat, 50 mg, tabletki powlekane	DE/H/0984/002	14936	GLENMARK PHARMACEUTICALS S.R.O.	PL
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/22	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/09	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/17	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/11	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/14	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/16	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/12	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/06	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/07	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/18	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/08	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/21	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/01	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/13	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/03	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/15	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/02	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/19	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/20	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/04	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/05	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/10	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 100 mg comprimate filmate	DE/H/0984/003	6196/2014/23	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/05	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/02	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/01	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/03	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/06	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/04	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/08	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/11	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/07	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/16	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/19	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/20	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/09	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/14	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/08	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/17	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/12	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/10	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/21	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/13	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/22	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/18	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 25 mg comprimate filmate	DE/H/0984/001	6194/2014/15	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/22	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/15	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/14	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/13	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/17	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/16	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/12	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/06	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/10	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/11	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/21	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/19	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/20	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/09	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/07	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/03	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/04	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/18	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/01	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/05	GLENMARK PHARMACEUTICALS S.R.O.	RO

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
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/08	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/02	GLENMARK PHARMACEUTICALS S.R.O.	RO
TORLEPTA 50 mg comprimate filmate	DE/H/0984/002	6195/2014/23	GLENMARK PHARMACEUTICALS S.R.O.	RO
ТАЛОПАМ 100 МГ ФИЛМИРАНИ ТАБЛЕТКИ	not available	20070027	ACTAVIS EAD	BG
ТАЛОПАМ 25 МГ ФИЛМИРАНИ ТАБЛЕТКИ	not available	20070025	ACTAVIS EAD	BG
ТАЛОПАМ 50 МГ ФИЛМИРАНИ ТАБЛЕТКИ	not available	20070026	ACTAVIS EAD	BG
Топилекс 100 mg филмирани таблетки	DK/H/1354/03	20090009	G.L. PHARMA GMBH	BG
Топилекс 25 mg филмирани таблетки	DK/H/1354/01	20090007	G.L. PHARMA GMBH	BG
Топилекс 50 mg филмирани таблетки	DK/H/1354/02	20090008	G.L. PHARMA GMBH	BG

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
Топирамат Акорд 100 mg филмирани таблетки	UK/H/1438/003	20110584	ACCORD HEALTHCARE LIMITED	BG
Топирамат Акорд 200 mg филмирани таблетки	UK/H/1438/004	20110585	ACCORD HEALTHCARE LIMITED	BG
Топирамат Акорд 25 mg филмирани таблетки	UK/H/1438/001	20110582	ACCORD HEALTHCARE LIMITED	BG
Топирамат Акорд 50 mg филмирани таблетки	UK/H/1438/002	20110583	ACCORD HEALTHCARE LIMITED	BG