19 September 2018
EMA/752859/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: valproic acid, sodium valproate, valproate pivoxil, valproate semisodium, valpromide, valproate bismuth, calcium valproate, valproate magnesium

Procedure no.: PSUSA/00003090/201801

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Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Convulex 100 mg/ml oldatos injekció	not available	OGYI-T-1112/13	G.L. PHARMA GMBH	HU
Convulex 300 mg maagsapresistente capsules	not available	BE115586	G.L. PHARMA GMBH	BE
Convulex 300 mg maagsapresistente capsules	not available	2003048215	G.L. PHARMA GMBH	LU
Convulex 500 mg maagsapresistente capsules	not available	BE123076	G.L. PHARMA GMBH	BE
Convulex 500 mg maagsapresistente capsules	not available	2003048216	G.L. PHARMA GMBH	LU
DEPAKENE CHRONO	not available	022483111	SANOFI SPA	IT
DEPAKIN	not available	022483010	SANOFI SPA	IT
DEPAKIN	not available	022483034	SANOFI SPA	IT
DEPAKIN	not available	022483061	SANOFI SPA	IT
DEPAKIN	not available	022483147	SANOFI SPA	IT
DEPAKIN	not available	022483150	SANOFI SPA	IT
DEPAKIN	not available	022483162	SANOFI SPA	IT
DEPAKIN	not available	022483174	SANOFI SPA	IT
DEPAKIN	not available	022483186	SANOFI SPA	IT
DEPAKIN	not available	022483198	SANOFI SPA	IT
DEPAKIN	not available	022483200	SANOFI SPA	IT
DEPAKIN	not available	022483212	SANOFI SPA	IT
DEPAKIN	not available	022483224	SANOFI SPA	IT
DEPAKIN	not available	022483236	SANOFI SPA	IT
DEPAKIN	not available	022483248	SANOFI SPA	IT
DEPAKIN	not available	022483251	SANOFI SPA	IT
DEPAKIN	not available	022483O22	SANOFI SPA	IT
DEPAKIN CHRONO	not available	022483109	SANOFI SPA	IT
DEPAKINE	not available	20010250	SANOFI BULGARIA EOOD	BG
DEPAKINE	not available	20010272	SANOFI BULGARIA EOOD	BG
DEPAKINE	not available	9600303	SANOFI BULGARIA EOOD	BG
DEPAKINE	not available	RVG 14996	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE	not available	RVG 14996	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE	not available	RVG 17569	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE	not available	RVG 18153	SANOFI-AVENTIS NETHERLANDS B.V.	NL

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE	not available	RVG 18153	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE	not available	21/0322/94-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
DEPAKINE 100 MG/ML POLVO Y DISOLVENTE PARA SOLUCION INYECTABLE	not available	60352	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 100 MG/ML POR ES OLDOSZER OLDATOS INJEKCIOHOZ	not available	OGYI-T-5527/01	SANOFI-AVENTIS ZRT	HU
DEPAKINE 100 MG/ML POR ES OLDOSZER OLDATOS INJEKCIOHOZ	not available	OGYI-T-5527/05	SANOFI-AVENTIS ZRT	HU
Depakine 200 mg comprimate gastrorezistente	not available	8141/2006/01	SANOFI-AVENTIS FRANCE	RO
DEPAKINE 200 MG COMPRIMIDOS GASTRORRESISTENTES	not available	48.827	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 200 MG COMPRIMIDOS GASTRORRESISTENTES	not available	48.827	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 200 MG COMPRIMIDOS GASTRORRESISTENTES	not available	48827	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	302 929-2	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	330 536-1	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	339 265-0	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	344 448-2	SANOFI-AVENTIS FRANCE	FR

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	344 449-9	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	344 450-7	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	360 923-3	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	360 925-6	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 200 MG, COMPRIME GASTRO RESISTANT	not available	563 915-5	SANOFI-AVENTIS FRANCE	FR
Depakine 200 mg/ml solução oral	not available	9729400	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE 200 MG/ML SOLUCION ORAL	not available	48828	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 200 MG/ML, SOLUTION BUVABLE	not available	302 930-0	SANOFI-AVENTIS FRANCE	FR
Depakine 300 mg/5 ml siroop	not available	BE110923	SANOFI BELGIUM	BE
Depakine 300 mg/5 ml sirop	not available	BE110923	SANOFI BELGIUM	BE
DEPAKINE 300 MG/5 ML SIROP	not available	0029320	SANOFI BELGIUM	LU
Depakine 300 mg/ml drank	not available	BE048316	SANOFI BELGIUM	BE
Depakine 300 mg/ml solution buvable	not available	BE048316	SANOFI BELGIUM	BE
DEPAKINE 300 MG/ML SOLUTION BUVABLE	not available	0029317	SANOFI BELGIUM	LU
DEPAKINE 300 MG/ML-TROPFEN	not available	15.699	SANOFI-AVENTIS GMBH OSTERREICH	AT
Depakine 40 mg/ml xarope	not available	2829091	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE 400 MG/4 ML	not available	21/0674/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Depakine 400 mg/4 ml pó e solvente para solução injetável	not available	8729418	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine 400 mg/4 ml, prášek a rozpouštedlo pro injekcní roztok	not available	21/265/96-C	SANOFI-AVENTIS SRO	CZ
DEPAKINE 400 MG/4 ML, PREPARATION INJECTABLE POUR VOIE I.V.	not available	3400932557691	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 400 MG/4 ML, PREPARATION INJECTABLE POUR VOIE I.V.	not available	554 104-8	SANOFI-AVENTIS FRANCE	FR
Depakine 5,0 g/100 ml, sirup	not available	21/312/99-C	SANOFI-AVENTIS SRO	CZ
DEPAKINE 50 MG/ML SZIRUP	not available	OGYI-T-5527/02	SANOFI-AVENTIS ZRT	HU
DEPAKINE 500 MG COMPRIMIDOS GASTRORRESISTENTES	not available	54.470	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 500 MG COMPRIMIDOS GASTRORRESISTENTES	not available	54.470	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 500 MG COMPRIMIDOS GASTRORRESISTENTES	not available	54470	SANOFI-AVENTIS, S.A.	ES
DEPAKINE 500 mg γαστροανθεκτικά δισκία	not available	41428/07/27-5-2008	SANOFI-AVENTIS AEBE	GR
DEPAKINE 500 mg γαστροανθεκτικά δισκία	not available	41428/07/27-5-2008	SANOFI-AVENTIS AEBE	GR
DEPAKINE 500 mg γαστροανθεκτικά δισκία	not available	41428/07/27-5-2008	SANOFI-AVENTIS AEBE	GR
DEPAKINE 500 mg γαστροανθεκτικά δισκία	not available	41428/07/27-5-2008	SANOFI-AVENTIS AEBE	GR
DEPAKINE 500 mg γαστροανθεκτικά δισκία	not available	41428/07/27-5-2008	SANOFI-AVENTIS AEBE	GR

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DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	319 227-6	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	339 264-4	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	344 451-3	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	344 453-6	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	360 921-0	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 500 MG, COMPRIME GASTRO RESISTANT	not available	360 922-7	SANOFI-AVENTIS FRANCE	FR
Depakine 57,64 mg/ml sirop	not available	6368/2006/01	SANOFI-AVENTIS FRANCE	RO
Depakine 57,64 mg/ml sirop	not available	6368/2006/02	SANOFI-AVENTIS FRANCE	RO
DEPAKINE 57,64 MG/ML, SIROP	not available	326 345-0	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 57,64 MG/ML, SIROP	not available	326 346-7	SANOFI-AVENTIS FRANCE	FR
DEPAKINE 57.64 MG/ML SYRUP	not available	LT/1/94/0973/001	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE 57.64 MG/ML SYRUP	not available	96-0149	SANOFI-AVENTIS LATVIA SIA	LV
DEPAKINE CHRONO	not available	20010812	SANOFI BULGARIA EOOD	BG
DEPAKINE CHRONO	not available	9900385	SANOFI BULGARIA EOOD	BG
DEPAKINE CHRONO	not available	019172	SANOFI-AVENTIS CYPRUS LTD	CY
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR

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DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	41972/10/21-06-2011	SANOFI-AVENTIS AEBE	GR
DEPAKINE CHRONO	not available	R/6943	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO	not available	R/6943	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO	not available	R/6943	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO	not available	R/6944	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO	not available	R/6944	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO	not available	R/6944	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONO 300	not available	RVG 13157	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONO 300	not available	RVG 13157	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONO 300	not available	RVG 13157	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONO 300	not available	RVG 13157	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Depakine Chrono 300 300 mg comprimidos de libertação prolongada	not available	5168307	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 300 300 mg comprimidos de libertação prolongada	not available	5169578	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 300 300 mg comprimidos de libertação prolongada	not available	5493531	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 300 300 mg comprimidos de libertação prolongada	not available	8729335	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 300 300 mg comprimidos de libertação prolongada	not available	8729343	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 300 mg comprimate cu eliberare prelungita	not available	6493/2006/01	SANOFI-AVENTIS FRANCE	RO

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Depakine Chrono 300 mg comprimés à libération prolongée	not available	BE166512	SANOFI BELGIUM	BE
Depakine Chrono 300 mg comprimés à libération prolongée	not available	BE166512	SANOFI BELGIUM	BE
DEPAKINE CHRONO 300 MG COMPRIMÉS À LIBÉRATION PROLONGÉE	not available	0210611	SANOFI BELGIUM	LU
DEPAKINE CHRONO 300 MG COMPRIMÉS À LIBÉRATION PROLONGÉE	not available	0210611	SANOFI BELGIUM	LU
DEPAKINE CHRONO 300 MG FILMSKO OBLOZENE TABLETE S PODALJZANIM SPROSCANJEM	not available	H/00/00450/001	SANOFI-AVENTIS D.O.O.	SI
DEPAKINE CHRONO 300 MG FILMTABLETTA	not available	OGYI-T-5527/03	SANOFI-AVENTIS ZRT	HU
Depakine Chrono 300 mg ilgstošās darbības tabletes	not available	96-0286	SANOFI-AVENTIS LATVIA SIA	LV
DEPAKINE CHRONO 300 MG MODIFIED RELEASE TABLETS	not available	LT/1/94/0818/001	UAB SANOFI-AVENTIS LIETUVA	LT
Depakine Chrono 300 mg tablete s prilagođenim oslobađanjem	not available	HR-H-132091526-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Depakine Chrono 300 mg tabletten met verlengde afgifte	not available	BE166512	SANOFI BELGIUM	BE
Depakine Chrono 300 mg tabletten met verlengde afgifte	not available	BE166512	SANOFI BELGIUM	BE
DEPAKINE CHRONO 300 mg, toimeainet prolongeeritult vabastavad tabletid	not available	151096	SANOFI-AVENTIS ESTONIA OÜ	EE
DEPAKINE CHRONO 300mg SECABLE	not available	21/056/91-A/C	SANOFI-AVENTIS SRO	CZ

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE CHRONO 500	not available	RVG 11775	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONO 500	not available	RVG 11775	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONO 500	not available	RVG 11775	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Depakine Chrono 500 500 mg comprimidos de libertação prolongada	not available	5169602	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 500 500 mg comprimidos de libertação prolongada	not available	5493523	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chrono 500 500 mg comprimidos de libertação prolongada	not available	8729350	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONO 500 MG	not available	21/0056/91-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
DEPAKINE CHRONO 500 MG	not available	21/0056/91-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Depakine Chrono 500 mg comprimate cu eliberare prelungita	not available	1671/2009/01	SANOFI-AVENTIS FRANCE	RO
Depakine Chrono 500 mg comprimés à libération prolongée	not available	BE166521	SANOFI BELGIUM	BE
Depakine Chrono 500 mg comprimés à libération prolongée	not available	BE166521	SANOFI BELGIUM	BE
DEPAKINE CHRONO 500 MG COMPRIMÉS À LIBÉRATION PROLONGÉE	not available	0210686	SANOFI BELGIUM	LU
DEPAKINE CHRONO 500 MG COMPRIMÉS À LIBÉRATION PROLONGÉE	not available	0210686	SANOFI BELGIUM	LU

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE CHRONO 500 MG FILMSKO OBLOZENE TABLETE S PODALJZANIM SPROSCANJEM	not available	H/00/00450/002	SANOFI-AVENTIS D.O.O.	SI
DEPAKINE CHRONO 500 MG FILMTABLETTA	not available	OGYI-T-5527/04	SANOFI-AVENTIS ZRT	HU
Depakine Chrono 500 mg ilgstošās darbības tabletes	not available	96-0324	SANOFI-AVENTIS LATVIA SIA	LV
DEPAKINE CHRONO 500 MG MODIFIED RELEASE TABLETS	not available	LT/1/94/0818/002	UAB SANOFI-AVENTIS LIETUVA	LT
Depakine Chrono 500 mg sécable tablety s řízeným uvolňováním	not available	21/056/91 - B/C	SANOFI-AVENTIS SRO	CZ
Depakine Chrono 500 mg sécable tablety s řízeným uvolňováním	not available	21/056/91 - B/C	SANOFI-AVENTIS SRO	CZ
Depakine Chrono 500 mg tablete s prilagođenim oslobađanjem	not available	HR-H-202764510-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Depakine Chrono 500 mg tabletten met verlengde afgifte	not available	BE166521	SANOFI BELGIUM	BE
Depakine Chrono 500 mg tabletten met verlengde afgifte	not available	BE166521	SANOFI BELGIUM	BE
DEPAKINE CHRONO 500 MG, COMPRIME PELLICULE SECABLE A LIBERATION PROLONGEE	not available	330 180-2	SANOFI-AVENTIS FRANCE	FR
DEPAKINE CHRONO 500 MG, COMPRIME PELLICULE SECABLE A LIBERATION PROLONGEE	not available	364 626-3	SANOFI-AVENTIS FRANCE	FR
DEPAKINE CHRONO 500 MG, COMPRIME PELLICULE SECABLE A LIBERATION PROLONGEE	not available	556 145-3	SANOFI-AVENTIS FRANCE	FR

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE CHRONO 500 MG, COMPRIME PELLICULE SECABLE A LIBERATION PROLONGEE	not available	559 008-7	SANOFI-AVENTIS FRANCE	FR
DEPAKINE CHRONO 500 MG, COMPRIME PELLICULE SECABLE A LIBERATION PROLONGEE	not available	584 894-7	SANOFI-AVENTIS FRANCE	FR
DEPAKINE CHRONO 500 MG, TOIMEAINET PROLONGEERITULT VABASTAVAD TABLETID	not available	151196	SANOFI-AVENTIS ESTONIA OÜ	EE
DEPAKINE CHRONO RETARD 300 MG-FILMTABLETTEN	not available	1-19787	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE CHRONO RETARD 500 MG-FILMTABLETTEN	not available	1-19786	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE CHRONOSPHERE	not available	RVG 30759	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30759	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30760	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30760	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30761	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30761	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30762	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30762	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30763	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	RVG 30763	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE CHRONOSPHERE	not available	11946	SANOFI-AVENTIS SP Z.O.O.	PL

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DEPAKINE CHRONOSPHERE	not available	11946	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11947	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11947	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11948	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11948	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11949	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11949	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11950	SANOFI-AVENTIS SP Z.O.O.	PL
DEPAKINE CHRONOSPHERE	not available	11950	SANOFI-AVENTIS SP Z.O.O.	PL
Depakine Chronosphere 100 mg granulado de libertação modificada	not available	5035837	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 100 mg granulado de libertação modificada	not available	5035845	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 100 mg granulado de libertação modificada	not available	5334586	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 100 mg granulado de libertação modificada	not available	5334685	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 1000 mg granulado de libertação modificada	not available	5035936	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 1000 mg granulado de libertação modificada	not available	5035944	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 1000 mg granulado de libertação modificada	not available	5335385	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 1000 mg granulado de libertação modificada	not available	5335484	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONOSPHERE 1000 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/011	UAB SANOFI-AVENTIS LIETUVA	LT

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DEPAKINE CHRONOSPHERE 1000 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/012	UAB SANOFI-AVENTIS LIETUVA	LT
Depakine Chronosphere 250 mg granulado de libertação modificada	not available	5035852	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 250 mg granulado de libertação modificada	not available	5035860	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 250 mg granulado de libertação modificada	not available	5334784	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 250 mg granulado de libertação modificada	not available	5334883	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONOSPHERE 250 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/005	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE CHRONOSPHERE 250 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/006	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE CHRONOSPHERE 250 MG-RETARDGRANULAT IN BEUTELN	not available	1-25371	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE CHRONOSPHERE 250 MG-RETARDGRANULAT IN BEUTELN	not available	1-25371	SANOFI-AVENTIS GMBH OSTERREICH	AT
Depakine Chronosphere 50 mg, granulado de libertação modificada	not available	5035811	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 50 mg, granulado de libertação modificada	not available	503829	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 50 mg, granulado de libertação modificada	not available	5334388	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Depakine Chronosphere 50 mg, granulado de libertação modificada	not available	5334487	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONOSPHERE 50 MG-RETARDGRANULAT IN BEUTELN	not available	1-25369	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE CHRONOSPHERE 50 MG-RETARDGRANULAT IN BEUTELN	not available	1-25369	SANOFI-AVENTIS GMBH OSTERREICH	AT
Depakine Chronosphere 500 mg granulado de libertação modificada	not available	5035878	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 500 mg granulado de libertação modificada	not available	5035902	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 500 mg granulado de libertação modificada	not available	5334982	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 500 mg granulado de libertação modificada	not available	5335088	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONOSPHERE 500 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/007	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE CHRONOSPHERE 500 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/008	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE CHRONOSPHERE 500 MG-RETARDGRANULAT IN BEUTELN	not available	1-25372	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE CHRONOSPHERE 500 MG-RETARDGRANULAT IN BEUTELN	not available	1-25372	SANOFI-AVENTIS GMBH OSTERREICH	AT
Depakine Chronosphere 750 mg granulado de libertação modificada	not available	5035910	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Depakine Chronosphere 750 mg granulado de libertação modificada	not available	5035928	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 750 mg granulado de libertação modificada	not available	5335187	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Depakine Chronosphere 750 mg granulado de libertação modificada	not available	5335286	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
DEPAKINE CHRONOSPHERE 750 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/009	UAB SANOFI-AVENTIS LIETUVA	LT
DEPAKINE CHRONOSPHERE 750 MG MODIFIED RELEASE GRANULES	not available	LT/1/94/0952/010	UAB SANOFI-AVENTIS LIETUVA	LT
Depakine Crono 300 mg comprimidos de liberación prolongada	not available	60.351	SANOFI-AVENTIS, S.A.	ES
Depakine Crono 300 mg comprimidos de liberación prolongada	not available	60.351	SANOFI-AVENTIS, S.A.	ES
Depakine Crono 300 mg comprimidos de liberación prolongada	not available	60.351	SANOFI-AVENTIS, S.A.	ES
Depakine Crono 500 mg comprimidos de liberación prolongada	not available	60.350	SANOFI-AVENTIS, S.A.	ES
Depakine Crono 500 mg comprimidos de liberación prolongada	not available	60.350	SANOFI-AVENTIS, S.A.	ES
Depakine Crono 500 mg comprimidos de liberación prolongada	not available	60.350	SANOFI-AVENTIS, S.A.	ES
DEPAKINE ENTERIC	not available	RVG 07055	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE ENTERIC	not available	RVG 07055	SANOFI-AVENTIS NETHERLANDS B.V.	NL

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKINE ENTERIC	not available	RVG 07405	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE ENTERIC	not available	RVG 07476	SANOFI-AVENTIS NETHERLANDS B.V.	NL
DEPAKINE ENTERIC	not available	RVG 07476	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Depakine Enteric 150 mg comprimés gastro-résistants	not available	BE110905	SANOFI BELGIUM	BE
Depakine Enteric 150 mg comprimés gastro-résistants	not available	BE110905	SANOFI BELGIUM	BE
DEPAKINE ENTERIC 150 MG COMPRIMÉS GASTRO-RÉSISTANTS	not available	0029334	SANOFI BELGIUM	LU
Depakine Enteric 150 mg maagsapresistente tabletten	not available	BE110905	SANOFI BELGIUM	BE
Depakine Enteric 150 mg maagsapresistente tabletten	not available	BE110905	SANOFI BELGIUM	BE
Depakine Enteric 300 mg comprimés gastro-résistants	not available	BE092775	SANOFI BELGIUM	BE
Depakine Enteric 300 mg comprimés gastro-résistants	not available	BE092775	SANOFI BELGIUM	BE
DEPAKINE ENTERIC 300 MG COMPRIMÉS GASTRO-RÉSISTANTS	not available	0145297	SANOFI BELGIUM	LU
Depakine Enteric 300 mg maagsapresistente tabletten	not available	BE092775	SANOFI BELGIUM	BE
Depakine Enteric 300 mg maagsapresistente tabletten	not available	BE092775	SANOFI BELGIUM	BE
Depakine Enteric 500 mg comprimés gastro-résistants	not available	BE110932	SANOFI BELGIUM	BE
Depakine Enteric 500 mg comprimés gastro-résistants	not available	BE110932	SANOFI BELGIUM	BE
Depakine Enteric 500 mg comprimés gastro-résistants	not available	BE110932	SANOFI BELGIUM	BE
Depakine Enteric 500 mg comprimés gastro-résistants	not available	0029348	SANOFI BELGIUM	LU

Product full name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Depakine Enteric 500 mg maagsapresistente tabletten	not available	BE110932	SANOFI BELGIUM	BE
Depakine Enteric 500 mg maagsapresistente tabletten	not available	BE110932	SANOFI BELGIUM	BE
Depakine Enteric 500 mg maagsapresistente tabletten	not available	BE110932	SANOFI BELGIUM	BE
Depakine I.V. 400 mg/4 ml poeder en oplosmiddel voor oplossing voor injectie	not available	BE163134	SANOFI BELGIUM	BE
Depakine I.V. 400 mg/4ml poudre et solvant pour solution injectable	not available	BE163134	SANOFI BELGIUM	BE
DEPAKINE I.V. 400 MG/ML POUDRE ET SOLVANT POUR SOLUTION INJECTABLE	not available	0196069	SANOFI BELGIUM	LU
DEPAKINE, 288,2 mg/5 ml, syrop	not available	R/3074	SANOFI-AVENTIS FRANCE	PL
DEPAKINE, 400 mg (400 mg/4 ml), proszek i rozpuszczalnik do sporzadzania roztworu do wstrzykiwan	not available	R/7170	SANOFI-AVENTIS FRANCE	PL
DEPAKINE, 400 MG SUSTELAHUSE PULBER JA LAHUSTI	not available	150996	SANOFI-AVENTIS ESTONIA OÜ	EE
DEPAKINE-TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	1-24529	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKINE-TROCKENSTECHAMPULLEN MIT LOSUNGSMITTEL	not available	1-24529	SANOFI-AVENTIS GMBH OSTERREICH	AT
DEPAKOTE 250 mg, comprimé gastro-résistant	not available	328 242-4	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 250 mg, comprimé gastro-résistant	not available	348 762-3	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 250 mg, comprimé gastro-résistant	not available	348 764-6	SANOFI-AVENTIS FRANCE	FR

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPAKOTE 250 mg, comprimé gastro-résistant	not available	561 893-4	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 250MG TABLETS	not available	PL 04425/0199	AVENTIS PHARMA LTD	UK
DEPAKOTE 250MG TABLETS	not available	PL 04425/0199	AVENTIS PHARMA LTD	UK
DEPAKOTE 250MG TABLETS	not available	PL 04425/0199	AVENTIS PHARMA LTD	UK
DEPAKOTE 500 mg, comprimé gastro-résistant	not available	328 241-8	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 500 mg, comprimé gastro-résistant	not available	348 765-2	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 500 mg, comprimé gastro-résistant	not available	348 766-9	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 500 mg, comprimé gastro-résistant	not available	354 442-7	SANOFI-AVENTIS FRANCE	FR
DEPAKOTE 500MG TABLETS	not available	PL 04425/0200	AVENTIS PHARMA LTD	UK
DEPAKOTE 500MG TABLETS	not available	PL 04425/0200	AVENTIS PHARMA LTD	UK
DEPAKOTE 500MG TABLETS	not available	PL 04425/0200	AVENTIS PHARMA LTD	UK
Depamag 100 mg/ml soluzione orale	not available	027107034	ALFASIGMA S.P.A.	IT
Depamag 200 mg compresse gastroresistenti	not available	027107010	ALFASIGMA S.P.A.	IT
Depamag 500 mg compresse gastroresistenti	not available	027107022	ALFASIGMA S.P.A.	IT
DEPAMIDE	not available	023105036	SANOFI SPA	IT
DEPAMIDE	not available	023105048	SANOFI SPA	IT
DEPAMIDE 300 mg, comprimé pelliculé gastro-résistant	not available	320 706-1	SANOFI-AVENTIS FRANCE	FR
DEPAMIDE 300 mg, comprimé pelliculé gastro-résistant	not available	559 000-6	SANOFI-AVENTIS FRANCE	FR
DEPRAKINE	not available	17782	SANOFI-AVENTIS DENMARK A/S	DK
DEPRAKINE	not available	17782	SANOFI-AVENTIS DENMARK A/S	DK
DEPRAKINE	not available	17783	SANOFI-AVENTIS DENMARK A/S	DK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPRAKINE	not available	17783	SANOFI-AVENTIS DENMARK A/S	DK
DEPRAKINE 200 MG/ML TIPAT, LIUOS	not available	12216	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG DEPOTTABLETTI	not available	10266	SANOFI OY	FI
DEPRAKINE 300 MG ENTEROTABLETTI	not available	12218	SANOFI OY	FI
DEPRAKINE 300 MG ENTEROTABLETTI	not available	12218	SANOFI OY	FI
DEPRAKINE 400 MG INJEKTIOKUIVA-AINE JA LIUOTIN, LIUOSTA VARTEN	not available	13405	SANOFI OY	FI
DEPRAKINE 500 MG DEPOTTABLETTI	not available	10267	SANOFI OY	FI
DEPRAKINE 500 MG DEPOTTABLETTI	not available	10267	SANOFI OY	FI
DEPRAKINE 500 MG DEPOTTABLETTI	not available	10267	SANOFI OY	FI
DEPRAKINE 500 MG ENTEROTABLETTI	not available	12219	SANOFI OY	FI
DEPRAKINE 500 MG ENTEROTABLETTI	not available	12219	SANOFI OY	FI
DEPRAKINE 60 MG/ML ORAALILIUOS	not available	12215	SANOFI OY	FI
DEPRAKINE RETARD	not available	13148	SANOFI-AVENTIS DENMARK A/S	DK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DEPRAKINE RETARD	not available	13148	SANOFI-AVENTIS DENMARK A/S	DK
DEPRAKINE RETARD	not available	13230	SANOFI-AVENTIS DENMARK A/S	DK
DEPRAKINE RETARD	not available	13230	SANOFI-AVENTIS DENMARK A/S	DK
Deprakine® 500 mg depottabletti	not available	10267	SANOFI OY	FI
Deprakine® 500 mg depottabletti	not available	10267	SANOFI OY	FI
Diplexil 150, 150mg, Prolonged-release capsules	PT/H/1406/01/MR	PT/H/1406/01/MR	TECNIFAR, INDÚSTRIA TÉCNICA FARMACÊUTICA, SA	PT
EPILIM 100MG CRUSHABLE TABLETS	not available	PA 540/150/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM 100MG CRUSHABLE TABLETS	not available	PA 540/150/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM 100MG CRUSHABLE TABLETS	not available	PL 04425/0317	AVENTIS PHARMA LTD	UK
EPILIM 100MG CRUSHABLE TABLETS	not available	PL 04425/0317	AVENTIS PHARMA LTD	UK
EPILIM 200 GASTRO-RESISTANT TABLETS	not available	PL 04425/0302	AVENTIS PHARMA LTD	UK
EPILIM 200 GASTRO-RESISTANT TABLETS	not available	PL 04425/0302	AVENTIS PHARMA LTD	UK
EPILIM 400MG POWDER AND SOLVENT FOR SOLUTION FOR INJECTION/INFUSION	not available	PL 04425/0685	AVENTIS PHARMA LTD	UK
EPILIM 400MG POWDER AND SOLVENT FOR SOLUTION FOR INJECTION/INFUSION	not available	PL 04425/0685	AVENTIS PHARMA LTD	UK
EPILIM 500 GASTRO-RESISTANT TABLETS	not available	PL 04425/0303	AVENTIS PHARMA LTD	UK
EPILIM 500 GASTRO-RESISTANT TABLETS	not available	PL 04425/0303	AVENTIS PHARMA LTD	UK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EPILIM CHRONO 200	not available	MA082/04301	SANOFI MALTA LTD	MT
EPILIM CHRONO 200MG CONTROLLED RELEASE TABLETS	not available	PA 540/150/10	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM CHRONO 200MG CONTROLLED RELEASE TABLETS	not available	PL 04425/0307	AVENTIS PHARMA LTD	UK
EPILIM CHRONO 300	not available	MA 082/04310	SANOFI MALTA LTD	MT
EPILIM CHRONO 300 CONTROLLED RELEASE TABLETS	not available	PA 540/150/11	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM CHRONO 300 CONTROLLED RELEASE TABLETS	not available	PL 04425/0308	AVENTIS PHARMA LTD	UK
EPILIM CHRONO 500	not available	MA082/04302	SANOFI MALTA LTD	MT
EPILIM CHRONO 500MG CONTROLLED RELEASE TABLETS	not available	PA 540/150/12	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM CHRONO 500MG CONTROLLED RELEASE TABLETS	not available	PL 04425/0309	AVENTIS PHARMA LTD	UK
Epilim Chronosphere 100mg prolonged-release granules	not available	PA 540/150/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Chronosphere 100mg prolonged-release granules	not available	PA 540/150/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Chronosphere 250mg prolonged-release granules	not available	PA 540/150/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Chronosphere 250mg prolonged-release granules	not available	PA 540/150/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Chronosphere 500mg prolonged-release granules	not available	PA 540/150/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Chronosphere 500mg prolonged-release granules	not available	PA 540/150/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM CHRONOSPHERE MR 1000MG MODIFIED RELEASE GRANULES	not available	PL 04425/0316	AVENTIS PHARMA LTD	UK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EPILIM CHRONOSPHERE MR 1000MG MODIFIED RELEASE GRANULES	not available	PL 04425/0316	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 100MG MODIFIED RELEASE GRANULES	not available	PL 04425/0312	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 100MG MODIFIED RELEASE GRANULES	not available	PL 04425/0312	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 250MG MODIFIED RELEASE GRANULES	not available	PL 04425/0313	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 250MG MODIFIED RELEASE GRANULES	not available	PL 04425/0313	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 500MG MODIFIED RELEASE GRANULES	not available	PL 04425/0314	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 500MG MODIFIED RELEASE GRANULES	not available	PL 04425/0314	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 50MG MODIFIED RELEASE GRANULES	not available	PL 04425/0310	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 50MG MODIFIED RELEASE GRANULES	not available	PL 04425/0310	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 750MG MODIFIED RELEASE GRANULES	not available	PL 04425/0315	AVENTIS PHARMA LTD	UK
EPILIM CHRONOSPHERE MR 750MG MODIFIED RELEASE GRANULES	not available	PL 04425/0315	AVENTIS PHARMA LTD	UK
EPILIM ENTERIC 200MG GASTRO-RESISTANT COATED TABLETS	not available	PA 540/150/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
EPILIM ENTERIC 200MG GASTRO-RESISTANT COATED TABLETS	not available	PA 540/150/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM ENTERIC 500MG GASTRO-RESISTANT COATED TABLETS	not available	PA 540/150/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM ENTERIC 500MG GASTRO-RESISTANT COATED TABLETS	not available	PA 540/150/3	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM INTRAVENOUS	not available	PA 540/150/13	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM INTRAVENOUS	not available	PA 540/150/13	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM INTRAVENOUS	not available	MA082/04303	SANOFI MALTA LTD	MT
EPILIM LIQUID	not available	082/04311	SANOFI MALTA LTD	MT
EPILIM LIQUID	not available	PL 04425/0300	AVENTIS PHARMA LTD	UK
Epilim Liquid 200 mg/5ml Oral Solution	not available	PA 540/150/14	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Liquid 200 mg/5ml Oral Solution	not available	PA 540/150/14	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
EPILIM SYRUP	not available	PL 04425/0301	AVENTIS PHARMA LTD	UK
EPILIM SYRUP	not available	PL 04425/0301	AVENTIS PHARMA LTD	UK
Epilim Syrup 200mg/5ml Oral Solution	not available	PA 540/150/15	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Epilim Syrup 200mg/5ml Oral Solution	not available	PA 540/150/15	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Episenta® 1000 mg Prolonged-release Granules	not available	PL 14040/0027	DESITIN ARZNEIMITTEL GMBH	UK
Episenta® 150 mg Prolonged-release Capsule	not available	PL 14040/0024	DESITIN ARZNEIMITTEL GMBH	UK
Episenta® 300 mg Prolonged-release Capsule	not available	PL 14040/0025	DESITIN ARZNEIMITTEL GMBH	UK
Episenta® 500 mg Prolonged-release Granules	not available	PL 14040/0026	DESITIN ARZNEIMITTEL GMBH	UK
Episenta® solution for injection	not available	PL 14040/0028	DESITIN ARZNEIMITTEL GMBH	UK

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ERGENYL 100 MG ENTEROTABLETTER	not available	12828	SANOFI AB	SE
ERGENYL 100 MG ENTEROTABLETTER	not available	12828	SANOFI AB	SE
Ergenyl 150 mg, magensaftresistente Filmtabletten	not available	378.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 150 mg, magensaftresistente Filmtabletten	not available	378.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 150 mg, magensaftresistente Filmtabletten	not available	378.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 150 mg, magensaftresistente Filmtabletten	not available	378.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 150 mg, magensaftresistente Filmtabletten	not available	378.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ERGENYL 200 MG/ML ORALA DROPPAR, LOSNING	not available	12827	SANOFI AB	SE
ERGENYL 300 MG ENTEROTABLETTER	not available	12829	SANOFI AB	SE
ERGENYL 500 MG ENTEROTABLETTER	not available	12830	SANOFI AB	SE
Ergenyl 500 mg, magensaftresistente Filmtabletten	not available	378.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 500 mg, magensaftresistente Filmtabletten	not available	378.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 500 mg, magensaftresistente Filmtabletten	not available	378.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl 500 mg, magensaftresistente Filmtabletten	not available	378.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ergenyl 500 mg, magensaftresistente Filmtabletten	not available	378.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ERGENYL 60 MG/ML ORAL LOSNING	not available	12831	SANOFI AB	SE
ERGENYL INTRAVENOS	not available	32555.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ERGENYL INTRAVENOS	not available	32555.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ERGENYL INTRAVENOS	not available	32555.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ERGENYL PULVER OCH VATSKA TILL INJEKTIONSVATSKA, LOSNING	not available	14282	SANOFI AB	SE
ERGENYL RETARD 100 MG DEPOTGRANULAT, DOSPASE	not available	20985	SANOFI AB	SE
ERGENYL RETARD 100 MG DEPOTGRANULAT, DOSPASE	not available	20985	SANOFI AB	SE
ERGENYL RETARD 100 MG DEPOTGRANULAT, DOSPASE	not available	20985	SANOFI AB	SE
ERGENYL RETARD 250 MG DEPOTGRANULAT, DOSPASE	not available	20986	SANOFI AB	SE
ERGENYL RETARD 250 MG DEPOTGRANULAT, DOSPASE	not available	20986	SANOFI AB	SE
ERGENYL RETARD 250 MG DEPOTGRANULAT, DOSPASE	not available	20986	SANOFI AB	SE
ERGENYL RETARD 300 MG DEPOTTABLETTER	not available	13043	SANOFI AB	SE
ERGENYL RETARD 300 MG DEPOTTABLETTER	not available	13043	SANOFI AB	SE
ERGENYL RETARD 500 MG DEPOTGRANULAT, DOSPASE	not available	20987	SANOFI AB	SE
ERGENYL RETARD 500 MG DEPOTGRANULAT, DOSPASE	not available	20987	SANOFI AB	SE
ERGENYL RETARD 500 MG DEPOTGRANULAT, DOSPASE	not available	20987	SANOFI AB	SE

Product full name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ERGENYL RETARD 500 MG DEPOTTABLETTER	not available	13044	SANOFI AB	SE
ERGENYL RETARD 500 MG DEPOTTABLETTER	not available	13044	SANOFI AB	SE
ERGENYL VIAL	not available	32556.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl vial 400 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	not available	32556.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Ergenyl vial 400 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung	not available	32556.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
MICROPAKINE L.P. 250 MG, GRANULES A LIBERATION PROLONGEE	not available	365 512-1	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 250 MG, GRANULES A LIBERATION PROLONGEE	not available	565 742-0	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 750 mg, granulés à libération prolongée en sachet-dose	not available	365 514-4	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 750 mg, granulés à libération prolongée en sachet-dose	not available	565 744-3	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 100 MG, GRANULES À LIBERATION PROLONGEE EN SACHET-DOSE	not available	365 511-5	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 100 MG, GRANULES À LIBERATION PROLONGEE EN SACHET-DOSE	not available	565 741-4	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 1000 MG , GRANULES A LIBERATION PROLONGEE EN SACHET-DOSE	not available	365 517-3	SANOFI-AVENTIS FRANCE	FR

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MICROPAKINE L.P. 1000 MG , GRANULES A LIBERATION PROLONGEE EN SACHET-DOSE	not available	565 746-6	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 500 MG, GRANULES A LIBERATION PROLONGEE EN SACHET-DOSE	not available	365 513-8	SANOFI-AVENTIS FRANCE	FR
MICROPAKINE L.P. 500 MG, GRANULES A LIBERATION PROLONGEE EN SACHET-DOSE	not available	565 743-7	SANOFI-AVENTIS FRANCE	FR
Orfiril 100 mg/ml injektioneste, liuos	FI/H/0127/001	12593	DESITIN ARZNEIMITTEL GMBH	FI
Orfiril 100 mg/ml Injektionslösung	not available	39415.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Orfiril 100 mg/ml Injektionslösung	FI/H/0127/001	50248.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Orfiril 100 mg/ml, injeksjonsvæske, oppløsning	not available	96-2608	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril 100 mg/ml, oplossing voor injectie	not available	RVG 24465	PHARMACHEMIE B.V	NL
Orfiril CR 1000 mg, granulaat met gereguleerde afgifte	not available	RVG 24464	PHARMACHEMIE B.V	NL
Orfiril CR 300 mg, capsules met gereguleerde afgifte	not available	RVG 24462	PHARMACHEMIE B.V	NL
Orfiril CR 500 mg, granulaat met gereguleerde afgifte	not available	RVG 24463	PHARMACHEMIE B.V	NL
Orfiril long 1000 mg depotgranulat i endosebeholder	not available	98-2477	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long 1000 mg depotgranulat i endosebeholder	not available	98-2477	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long 1000 mg, Retard-Minitabletten	not available	57471.01.01	DESITIN ARZNEIMITTEL GMBH	DE

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Orfiril long 150 mg depotkapsel, hård	not available	13188	DESITIN ARZNEIMITTEL GMBH	SE
Orfiril long 150 mg depotkapseli, kova	not available	13214	DESITIN ARZNEIMITTEL GMBH	FI
Orfiril long 150 mg, Hartkapseln, retardiert	not available	57471.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Orfiril long 150 mg, toimeainet prolongeeritult vabastavad kõvakapslid	not available	326200	DESITIN ARZNEIMITTEL GMBH	EE
Orfiril long 300 mg depotkapsel, hård	not available	13189	DESITIN ARZNEIMITTEL GMBH	SE
Orfiril long 300 mg depotkapseli, kova	not available	13215	DESITIN ARZNEIMITTEL GMBH	FI
Orfiril long 300 mg, Hartkapseln, retardiert	not available	57471.01.00	DESITIN ARZNEIMITTEL GMBH	DE
Orfiril long 300 mg, toimeainet prolongeeritult vabastavad kõvakapslid	EE/H/0104/001	326300	DESITIN ARZNEIMITTEL GMBH	EE
Orfiril long 500 mg depotgranulat	not available	14577	DESITIN ARZNEIMITTEL GMBH	SE
Orfiril long 500 mg depotgranulat i endosebeholder	not available	98-2476	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long 500 mg depotgranulat i endosebeholder	not available	98-2476	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long 500 mg depotrakeet	not available	13216	DESITIN ARZNEIMITTEL GMBH	FI
Orfiril long 500 mg, Retard-Minitabletten	not available	57471.00.01	DESITIN ARZNEIMITTEL GMBH	DE
Orfiril long 500 mg, toimeainet prolongeeritult vabastavad graanulid	EE/H/0104/002	326400	DESITIN ARZNEIMITTEL GMBH	EE
Orfiril long depotkapsler	not available	96-1965	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long depotkapsler	not available	96-1965	DESITIN ARZNEIMITTEL GMBH	NO

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Orfiril long depotkapsler	not available	96-1966	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long depotkapsler	not available	96-1966	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril long, Depotgranulat	not available	19055	DESITIN ARZNEIMITTEL GMBH	DK
Orfiril long, Depotgranulat	not available	19056	DESITIN ARZNEIMITTEL GMBH	DK
Orfiril long, Depotkapsler, hårde	not available	18324	DESITIN ARZNEIMITTEL GMBH	DK
Orfiril long, Depotkapsler, hårde	not available	18325	DESITIN ARZNEIMITTEL GMBH	DK
Orfiril retard 300 mg depottablett	not available	7866	DESITIN ARZNEIMITTEL GMBH	NO
Orfiril retard 300 mg forðatöflur	not available	920075	DESITIN ARZNEIMITTEL GMBH	IS
Orfiril retard, Depottabletter	not available	14700	DESITIN ARZNEIMITTEL GMBH	DK
Orfiril, Injektionsvæske, opløsning	not available	18520	DESITIN ARZNEIMITTEL GMBH	DK
Sodium Valproate 100mg/ml Solution for Injection or Infusion	UK/H/5168/01/DC	MA154/10201	WOCKHARDT UK LTD	MT
Sodium Valproate 100mg/ml Solution for Injection or Infusion	UK/H/6313/001	PL 12762/0529	MERCURY PHARMACEUTICALS LTD.	UK
Sodium Valproate 100mg/ml Solution for Injection or Infusion	UK/H/5168/001	PL 29831/0506	WOCKHARDT UK LTD	UK
Valproát chrono Sandoz 300 mg tablety s predĺženým uvoľňovaním	NL/H/0736/001	21/0267/06-S	SANDOZ PHARMACEUTICALS D.D.	SK
Valproát chrono Sandoz 300 mg tablety s predĺženým uvoľňovaním	NL/H/0736/001	21/0267/06-S	SANDOZ PHARMACEUTICALS D.D.	SK

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Valproát chrono Sandoz 300 mg tablety s predĺženým uvoľňovaním	NL/H/0736/001	21/0267/06-S	SANDOZ PHARMACEUTICALS D.D.	SK
VALPROATE DE SODIUM AGUETTANT 400 mg/4 ml, solution injectable	not available	378 574.0	LABORATOIRE AGUETTANT	FR
VALPROATE DE SODIUM AGUETTANT 400 mg/4 ml, solution injectable	not available	378 574.0	LABORATOIRE AGUETTANT	FR
VALPROATE DE SODIUM AGUETTANT 400 mg/4 ml, solution injectable	not available	378 575.7	LABORATOIRE AGUETTANT	FR
VALPROATE DE SODIUM AGUETTANT 400 mg/4 ml, solution injectable	not available	378 575.7	LABORATOIRE AGUETTANT	FR
VALPROATE DE SODIUM ZENTIVA 200 MG, COMPRIME GASTRO-RESISTANT	not available	34009 361 072 7 4	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 200 MG, COMPRIME GASTRO-RESISTANT	not available	34009 564 609 5 3	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 200 mg, comprimé gastro-résistant	not available	34009 352 104 7 0	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 200 mg, comprimé gastro-résistant	not available	34009 361 071 0 6	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 200 mg, comprimé gastro-résistant	not available	34009 367 385 7 7	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 200 mg/ml, solution buvable	not available	367 388-6	SANOFI-AVENTIS FRANCE	FR

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VALPROATE DE SODIUM ZENTIVA 500 MG, COMPRIME GASTRO-RESISTANT	not available	361 073-3	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 500 MG, COMPRIME GASTRO-RESISTANT	not available	361 075-6	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA 500 MG, COMPRIME GASTRO-RESISTANT	not available	367 386-3	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA LP 500 mg, comprimé pelliculé sécable à libération prolongée	not available	369 346-9	SANOFI-AVENTIS FRANCE	FR
VALPROATE DE SODIUM ZENTIVA LP 500 mg, comprimé pelliculé sécable à libération prolongée	not available	567 053-8	SANOFI-AVENTIS FRANCE	FR
Valproat-neuraxpharm 60 mg/ml Lösung zum Einnehmen	DE/H/3620/001	87742.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE