20 July 2023
EMA/412983/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): lamotrigine

Procedure No. PSUSA/00001825/202211

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Crisomet 100 mg comprimidos masticables/dispersables	not available	64.462	GLAXOSMITHKLINE, S.A.	ES
Crisomet 200 mg comprimidos masticables/dispersables .	not available	64.463	GLAXOSMITHKLINE, S.A.	ES
Crisomet 25 mg comprimidos masticables/dispersables	not available	63.747	GLAXOSMITHKLINE, S.A.	ES
Crisomet 50 mg comprimidos masticables/dispersables .	not available	64.464	GLAXOSMITHKLINE, S.A.	ES
Labileno 100 mg comprimidos masticables/dispersables .	not available	64.943	GLAXOSMITHKLINE S.A.	ES
Labileno 200 mg comprimidos masticables/dispersables .	not available	65.283	GLAXOSMITHKLINE S.A.	ES
Labileno 25 mg comprimidos masticables/dispersables	not available	63.288	GLAXOSMITHKLINE S.A.	ES
Labileno 50 mg comprimidos masticables/dispersables .	not available	65.284	GLAXOSMITHKLINE S.A.	ES
Lambipol 100 mg comprimés à croquer/dispersibles	NL/H/2051/003	26005010029	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 100 mg comprimés à croquer/dispersibles.	NL/H/2051/003	BE 266122	GLAXOSMITHKLINE PHARMACEUTICALS 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
Lambipol 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/003	BE 266122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/003	26005010029	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 100 mg kauw-/dispergeerbare tabletten.	NL/H/2051/003	BE 266122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 100 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/003	RVG 107455	GLAXOSMITHKLINE B.V.	NL
Lambipol 200 mg comprimés à croquer/dispersibles	NL/H/2051/004	BE 266156	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 200 mg comprimés à croquer/dispersibles	NL/H/2051/004	26005010030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 200 mg Kautabletten/Tabletten	NL/H/2051/004	BE 266156	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/004	26005010030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 200 mg kauw-/dispergeerbare tabletten.	NL/H/2051/004	BE 266156	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 200 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/004	RVG 107456	GLAXOSMITHKLINE B.V.	NL
Lambipol 25 mg comprimés à croquer/dispersibles	NL/H/2051/001	BE 266052	GLAXOSMITHKLINE PHARMACEUTICALS 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
Lambipol 25 mg comprimés à croquer/dispersibles	NL/H/2051/001	26005010026	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg comprimés à croquer/dispersibles.	NL/H/2051/001	BE 266077	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg comprimés à croquer/dispersibles.	NL/H/2051/001	2005010027	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	BE 266052	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	BE 266077	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	26005010026	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	2005010027	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg kauw-/dispergeerbare tabletten.	NL/H/2051/001	BE 266052	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg kauw-/dispergeerbare tabletten.	NL/H/2051/001	BE 266077	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/001	RVG 107451	GLAXOSMITHKLINE B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lambipol 50 mg comprimés à croquer/dispersibles.	NL/H/2051/002	BE 266095	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 50 mg comprimés à croquer/dispersibles.	NL/H/2051/002	26005010028	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/002	BE 266095	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/002	26005010028	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 50 mg kauw-/dispergeerbare tabletten.	NL/H/2051/002	BE 266095	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 50 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/002	RVG 107454	GLAXOSMITHKLINE B.V.	NL
Lamictal	NL/H/1540/003	13904	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1540/004	14790	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1540/002	13903	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1539/001	31464	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1539/003	15831	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1539/002	15830	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1539/004	17951	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal	NL/H/1539/005	15832	GLAXOSMITHKLINE PHARMA 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
Lamictal	NL/H/1539/006	17952	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal µas?µe?a/d?aspe???µe?a d?s??a 2 mg	NL/H/1539/001	2018110	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal 100 mg chewable/dispersible tablets	NL/H/1539/005	PA1077/061/009	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 100 mg chewable/dispersible tablets	not available	PL 00003/0348	THE WELLCOME FOUNDATION LTD	XI
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807078	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807371	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807383	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807395	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807407	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807419	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807421	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807433	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg	NL/H/1540/003	1744/2009/02	GLAXOSMITHKLINE	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
comprimate			(IRELAND) LIMITED	
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/09	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 100 mg comprimés à croquer/dispersibles	NL/H/1539/005	0260/09/10/0609	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 100 mg comprimés à croquer/dispersibles.	NL/H/1539/005	BE 185245	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 100 mg comprimidos	NL/H/1540/003	2253185	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 100 mg comprimidos	NL/H/1540/003	5792742	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 100 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/005	2521987	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 100 mg comprimidos masticables/dispersables.	NL/H/1539/005	61.568	GLAXOSMITHKLINE, S.A.	ES
Lamictal 100 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/005	12323	GLAXOSMITHKLINE OY	FI
Lamictal 100 mg	NL/H/1539/005	33122.03.00	GLAXOSMITHKLINE 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
Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen			CO. KG	
Lamictal 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/005	BE 185245	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/005	0260/09/10/0609	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 100 mg kauw-/dispergeerbare tabletten	NL/H/1539/005	BE 185245	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 100 mg košļājamās/ disperģējamās tabletes	NL/H/1539/005	97-0590	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/029	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/032	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/048	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar	NL/H/1539/005	LT/1/97/1271/030	GLAXOSMITHKLINE TRADING SERVICES	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
disperguojamosios tabletės			LIMITED	
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/031	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/034	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/035	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/047	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/049	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/050	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/051	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 100 mg purutabletti / dispergoituva tabletti	NL/H/1539/005	12323	GLAXOSMITHKLINE OY	FI
Lamictal 100 mg tablete	not available	HR-H-695293019	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/073	GLAXOSMITHKLINE TRADING SERVICES	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			LIMITED	
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/074	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/075	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/076	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/077	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/078	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/079	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/080	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/081	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/109	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/110	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/111	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/112	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/113	GLAXOSMITHKLINE	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			TRADING SERVICES LIMITED	
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/114	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/115	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/116	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/117	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg tablets	NL/H/1540/003	PA1077/061/003	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 100 mg tablets	NL/H/1540/003	MA192/00505	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 100 mg tablets	not available	PL 00003/0274	THE WELLCOME FOUNDATION LTD	XI
Lamictal 100 mg tabletta	NL/H/1540/003	OGYI-T-4094/15	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 100 mg tabletta	NL/H/1540/003	OGYI-T-4094/22	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 100 mg tabletten	NL/H/1540/003	RVG 103366	GLAXOSMITHKLINE B.V.	NL
Lamictal 100 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/005	1-20886	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 100 mg tabletter	NL/H/1540/003	12010	GLAXOSMITHKLINE AB	SE
Lamictal 100 mg tablety	NL/H/1540/003	21/802/92-C/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Lamictal 100 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/005	12792	GLAXOSMITHKLINE 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
Lamictal 100 mg tuggu-/dreifitöflur.	NL/H/1539/005	940291	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 100 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/005	8200	GLAXOSMITHKLINE AS	NO
Lamictal 100 mg žuvacie/dispergovatelné tablety	NL/H/1539/005	21/0405/09-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/031	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/032	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/033	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/034	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/035	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/036	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/037	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/038	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/039	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/145	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/146	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/147	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/148	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/149	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/150	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/151	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/152	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/153	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/154	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/155	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/156	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/157	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvečljive/disperzibilne	NL/H/1539/005	H/96/00871/040	GLAXOSMITHKLINE TRADING SERVICES	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tablete			LIMITED	
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/041	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/042	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/043	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 100 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/005	16178	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
LAMICTAL 100 mg, comprimé dispersible ou à croquer	NL/H/1539/005	NL20298	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 100 mg, närimis/dispergeeruvad tabletid	NL/H/1539/005	426003	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/033	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 2 mg chewable/dispersible tablets	NL/H/1539/001	MA192/00506	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 2 mg chewable/dispersible tablets.	NL/H/1539/001	PA1077/061/005	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 2 mg chewable/dispersible tablets.	not available	PL 00003/0375	THE WELLCOME FOUNDATION LTD	XI
Lamictal 2 mg compresse masticabili/dispersibili	NL/H/1539/001	027807179	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 2 mg comprimate masticabile/dispersabile	NL/H/1539/001	1745/2009/01	GLAXOSMITHKLINE (IRELAND) 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
Lamictal 2 mg comprimés à croquer/dispersibles	NL/H/1539/001	2009100605	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 2 mg comprimés à croquer/dispersibles.	NL/H/1539/001	BE 228611	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/001	3338183	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 2 mg comprimidos masticables/dispersables	NL/H/1539/001	64.391	GLAXOSMITHKLINE S.A.	ES
Lamictal 2 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/001	49825.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 2 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/001	BE 228611	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/001	2009100605	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 2 mg kauw-/dispergeerbare tabletten	NL/H/1539/001	BE 228611	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg purutabletti / dispergoituva tabletti	NL/H/1539/001	23162	GLAXOSMITHKLINE OY	FI
Lamictal 2 mg rágótabletta/diszpergáló dó tabletta	NL/H/1539/001	OGYI-T-4094/18	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 2 mg tuggtabletter /	NL/H/1539/001	23162	GLAXOSMITHKLINE 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
dispergerbara tabletter				
Lamictal 2 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/001	16321	GLAXOSMITHKLINE AB	SE
Lamictal 2 mg tuggu-/dreifitöflur	NL/H/1539/001	IS/1/08/112/01	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 2 mg tyggetabletter/dispergerbare tabletter	NL/H/1539/001	00-2581	GLAXOSMITHKLINE AS	NO
Lamictal 2 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/001	22006	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
LAMICTAL 2 mg, comprimé dispersible ou à croquer	NL/H/1539/001	NL25689	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 200 mg chewable/dispersible tablets	not available	PL 00003/0369	THE WELLCOME FOUNDATION LTD	XI
Lamictal 200 mg chewable/dispersible tablets.	NL/H/1539/006	PA1077/061/010	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807092	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807445	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006/MR	027807458	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807460	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807472	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse	NL/H/1539/006	027807484	GLAXOSMITHKLINE 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
masticabili/dispersibili				
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807496	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg compresse masticabili/dispersibili	NL/H/1539/006	027807508	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 200 mg comprimés à croquer/dispersibles.	NL/H/1539/006	BE185254	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 200 mg comprimés à croquer/dispersibles.	NL/H/1539/006	0260/09/10/0610	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 200 mg comprimidos	NL/H/1540/004	2521581	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 200 mg comprimidos	NL/H/1540/004	5792759	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 200 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/006	4014684	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 200 mg comprimidos masticables/dispersables.	NL/H/1539/006	61.563	GLAXOSMITHKLINE, S.A.	ES
Lamictal 200 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/006	12324	GLAXOSMITHKLINE OY	FI
Lamictal 200 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/006	33122.04.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum	NL/H/1539/006	BE185254	GLAXOSMITHKLINE PHARMACEUTICALS 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
Einnehmen				
Lamictal 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/006	0260/09/10/0610	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 200 mg kauw-/dispergeerbare tabletten	NL/H/1539/006	BE185254	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 200 mg košļājamās/ disperģējamās tabletes	NL/H/1539/006	97-0591	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/036	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/039	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/040	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/042	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/006	LT/1/97/1271/055	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar	NL/H/1539/006	LT/1/97/1271/037	GLAXOSMITHKLINE TRADING SERVICES	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
disperguojamosios tabletės			LIMITED	
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/038	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/041	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/052	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/053	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/054	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/056	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 200 mg purutabletti / dispergoituva tabletti	NL/H/1539/006	12324	GLAXOSMITHKLINE OY	FI
Lamictal 200 mg rágótabletta/diszpergáló dó tabletta	NL/H/1539/006	OGYI-T-4094/17	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 200 mg rágótabletta/diszpergáló dó tabletta	NL/H/1539/006	OGYI-T-4094/23	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/082	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/083	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/084	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/085	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/086	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/118	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/119	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/120	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/121	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/122	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg tablets	NL/H/1540/004	PA1077/061/004	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 200 mg tablets	not available	PL 00003/0297	THE WELLCOME FOUNDATION LTD	XI
Lamictal 200 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/006	1-20216	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 200 mg tabletten.	NL/H/1540/004	RVG 103367	GLAXOSMITHKLINE B.V.	NL
Lamictal 200 mg tabletter	NL/H/1540/004	12011	GLAXOSMITHKLINE 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
Lamictal 200 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/006	13029	GLAXOSMITHKLINE AB	SE
Lamictal 200 mg tuggu-/dreifitöflur.	NL/H/1539/006	960235	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 200 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/006	96-1877	GLAXOSMITHKLINE AS	NO
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/044	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/045	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/168	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/169	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/170	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/161	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvecljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/167	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/046	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/047	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/048	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/049	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/050	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/051	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/052	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/053	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/054	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/055	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/056	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/158	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/159	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/160	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/162	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne	NL/H/1539/006	H/96/00871/163	GLAXOSMITHKLINE TRADING SERVICES	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tablete			LIMITED	
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/164	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/165	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/166	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 200 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/006	18617	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
LAMICTAL 200 mg, comprimé dispersible ou à croquer	NL/H/1539/006	NL21697	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 25 mg chewable/dispersible tablets	not available	PL 00003/0347	THE WELLCOME FOUNDATION LTD	XI
Lamictal 25 mg chewable/dispersible tablets	NL/H/1539/003	PA1077/061/007	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 25 mg chewable/dispersible tablets.	NL/H/1539/003	MA192/00502	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807054	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807268	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807270	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807282	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg	NL/H/1539/003	027807243	GLAXOSMITHKLINE 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
compresse masticabili/dispersibili				
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807256	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/09	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/10	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 25 mg comprimés à croquer/dispersibles	NL/H/1539/003	0260/09/10/0607	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 25 mg comprimés à croquer/dispersibles.	NL/H/1539/003	BE 185227	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 25 mg comprimidos	NL/H/1540/001	5291695	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	5291794	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg	NL/H/1540/001	2252781	GLAXOSMITHKLINE -	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos			PRODUTOS FARMACEUTICOS, LDA	
Lamictal 25 mg comprimidos	NL/H/1540/001	2252880	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	5792650	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	5792668	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	5792676	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	5792700	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/003	2521888	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/003	5291984	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/003	5292081	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos masticables/dispersables	NL/H/1539/003	61.569	GLAXOSMITHKLINE S.A.	ES
Lamictal 25 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/003	12321	GLAXOSMITHKLINE OY	FI
Lamictal 25 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/003	33122.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 25 mg	NL/H/1539/003	BE 185227	GLAXOSMITHKLINE	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
Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen			PHARMACEUTICALS SA	
Lamictal 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/003	0260/09/10/0607	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 25 mg kauw-/dispergeerbare tabletten	NL/H/1539/003	BE 185227	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 25 mg košļājamās/ disperģējamās tabletes	NL/H/1539/003	97-0588	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Lamictal 25 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/003	LT/1/97/1271/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/003	LT/1/97/1271/014	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/003	LT/1/97/1271/015	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/003	LT/1/97/1271/018	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/003	LT/1/97/1271/019	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios	NL/H/1539/003	LT/1/97/1271/004	GLAXOSMITHKLINE TRADING SERVICES 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
tabletės				
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/016	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/017	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 25 mg purutabletti / dispergoituva tabletti	NL/H/1539/003	12321	GLAXOSMITHKLINE OY	FI
Lamictal 25 mg tablete	not available	HR-H-211198060	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/057	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/058	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/059	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/060	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/061	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/062	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/063	GLAXOSMITHKLINE	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			TRADING SERVICES LIMITED	
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/064	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/093	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/094	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/095	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/096	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/097	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/098	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/099	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/100	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg tablets	NL/H/1540/001	PA1077/061/001	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 25 mg tablets	not available	PL 00003/0272	THE WELLCOME FOUNDATION LTD	XI
Lamictal 25 mg tablets.	NL/H/1540/001	MA192/00503	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 25 mg tabletta	NL/H/1540/001	OGYI-T-4094/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 25 mg tabletta	NL/H/1540/001	OGYI-T-4094/20	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 25 mg tabletten	NL/H/1540/001	RVG 103363	GLAXOSMITHKLINE B.V.	NL
Lamictal 25 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/003	1-20875	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 25 mg tabletter	NL/H/1540/001	12008	GLAXOSMITHKLINE AB	SE
Lamictal 25 mg tablety	NL/H/1540/001	21/802/92-A/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Lamictal 25 mg tablety	NL/H/1540/001	21/0802/92-CS	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Lamictal 25 mg tuggtabletter/dispergerbara tabletter.	NL/H/1539/003	12791	GLAXOSMITHKLINE AB	SE
Lamictal 25 mg tuggu-/dreifitöflur	NL/H/1539/003	940290	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 25 mg tyggetabletter/dispergerbare tabletter	NL/H/1539/003	8199	GLAXOSMITHKLINE AS	NO
Lamictal 25 mg žvecljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/013	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvecljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/014	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvecljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/015	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvecljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/017	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvecljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/124	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvecljive/disperzibilne	NL/H/1539/003	H/96/00871/125	GLAXOSMITHKLINE TRADING SERVICES	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tablete			LIMITED	
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/016	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/123	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/126	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/127	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/128	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/129	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/130	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/131	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 25 mg	NL/H/1539/003	16177	GLAXOSMITHKLINE	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
μασώμενα/διασπειρόμενα δισκία			(IRELAND) LIMITED	
LAMICTAL 25 mg, comprimé dispersible ou à croquer	NL/H/1539/003	NL20297	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 25 mg, närimis/dispergeeruvad tabletid	NL/H/1539/003	425403	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Lamictal 5 mg chewable/dispersible tablets	NL/H/1539/002	MA192/00501	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 5 mg chewable/dispersible tablets	not available	PL 00003/0346	THE WELLCOME FOUNDATION LTD	XI
Lamictal 5 mg chewable/dispersible tablets.	NL/H/1539/002	PA1077/061/006	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807066	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807181	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807193	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807205	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807217	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807229	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 5 mg compresse masticabili/dispersibili	NL/H/1539/002	027807231	GLAXOSMITHKLINE 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
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/09	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/10	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/11	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/12	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Lamictal 5 mg comprimate	NL/H/1539/002	1746/2009/13	GLAXOSMITHKLINE (IRELAND) 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
masticabile/dispersabile				
Lamictal 5 mg comprimés à croquer/dispersibles.	NL/H/1539/002	BE 185211	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg comprimés à croquer/dispersibles.	NL/H/1539/002	BE477093	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg comprimés à croquer/dispersibles.	NL/H/1539/002	2009 10 0606	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 5 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/002	2521680	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/002	2521789	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/002	5770029	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg comprimidos masticables/dispersables	NL/H/1539/002	61.561	GLAXOSMITHKLINE S.A.	ES
Lamictal 5 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/002	33122.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 5 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/002	BE 185211	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/002	BE477093	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg	NL/H/1539/002	2009 10 0606	GLAXOSMITHKLINE	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen			PHARMACEUTICALS SA	
Lamictal 5 mg kauw-/dispergeerbare tabletten	NL/H/1539/002	BE 185211	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg kauw-/dispergeerbare tabletten	NL/H/1539/002	BE477093	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 5 mg košlajamas/ dispergejamas tabletes	NL/H/1539/002	97-0587	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/013	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/061	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/063	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/066	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg	NL/H/1539/002	LT/1/97/1271/062	GLAXOSMITHKLINE	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
kramtomosios ar disperguojamosios tabletes			TRADING SERVICES LIMITED	
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/064	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/002	LT/1/97/1271/065	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/043	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 5 mg purutabletti / dispergoituva tabletti	NL/H/1539/002	12320	GLAXOSMITHKLINE OY	FI
Lamictal 5 mg rágótabletta/diszpergáló dó tabletta	NL/H/1539/002	OGYI-T-4094/16	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 5 mg rágótabletta/diszpergáló dó tabletta	NL/H/1539/002	OGYI-T-4094/19	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 5 mg tablete za žvakanje/tablete za	not available	HR-H-287736527	GLAXOSMITHKLINE TRADING SERVICES	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
oralnu suspenziju			LIMITED	
Lamictal 5 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/002	1-20888	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 5 mg tuggtabletter / dispergerbara tabletter	NL/H/1539/002	12320	GLAXOSMITHKLINE OY	FI
Lamictal 5 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/002	12790	GLAXOSMITHKLINE AB	SE
Lamictal 5 mg tuggu-/dreifitöflur.	NL/H/1539/002	940289	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 5 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/002	8198	GLAXOSMITHKLINE AS	NO
Lamictal 5 mg žuvacie/dispergovateľné tablety	NL/H/1539/002	21/0208/98-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/087	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/088	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/092	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/089	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/090	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/091	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 5 mg žvýkací/dispergovatelné tablety	NL/H/1539/002	21/483/96-A/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Lamictal 5 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/002	16176	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
LAMICTAL 5 mg, comprimé dispersible ou à croquer	NL/H/1539/002	NL20296	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 5 mg, närimis/dispergeeruvad tabletid	NL/H/1539/002	425703	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Lamictal 50 mg chewable/dispersible tablets	NL/H/1539/004	PA1077/061/008	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 50 mg chewable/dispersible tablets	not available	PL 00003/0368	THE WELLCOME FOUNDATION LTD	XI
Lamictal 50 mg compresse	NL/H/1539/004	027807080	GLAXOSMITHKLINE 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
masticabili/dispersibili				
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807357	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807369	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807294	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807306	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807318	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807320	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807332	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807344	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg comprimés à croquer/dispersibles	NL/H/1539/004	26009100608	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 50 mg comprimés à croquer/dispersibles.	NL/H/1539/004	BE 185236	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 50 mg comprimidos	NL/H/1540/002	2252989	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos	NL/H/1540/002	2253086	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg	NL/H/1540/002	5291885	GLAXOSMITHKLINE -	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos			PRODUTOS FARMACEUTICOS, LDA	
Lamictal 50 mg comprimidos	NL/H/1540/002	5792718	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos	NL/H/1540/002	5792726	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos	NL/H/1540/002	5792734	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/004	4014585	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/004	5292180	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimidos masticables/dispersables	NL/H/1539/004	61.562	GLAXOSMITHKLINE, S.A.	ES
Lamictal 50 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/004	12322	GLAXOSMITHKLINE OY	FI
Lamictal 50 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/004	33122.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/004	BE 185236	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/004	26009100608	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 50 mg kauw-/dispergeerbare tabletten	NL/H/1539/004	BE 185236	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 50 mg košļājamās/ disperģējamās tabletes	NL/H/1539/004	97-0589	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/020	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/021	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/022	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/023	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/024	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/025	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/026	GLAXOSMITHKLINE TRADING SERVICES 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
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/028	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/044	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/004	LT/1/97/1271/046	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/027	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/045	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Lamictal 50 mg purutabletti / dispergoituva tabletti	NL/H/1539/004	12322	GLAXOSMITHKLINE OY	FI
Lamictal 50 mg tablete	not available	HR-H-052719076	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/065	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/066	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/067	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/068	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/069	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/070	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/071	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/072	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/101	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/102	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/103	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/104	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/105	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/106	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/107	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/108	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg tablets	NL/H/1540/002	PA1077/061/002	GLAXOSMITHKLINE (IRELAND) 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
Lamictal 50 mg tablets	not available	PL 00003/0273	THE WELLCOME FOUNDATION LTD	XI
Lamictal 50 mg tablets.	NL/H/1540/002	MA192/00504	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 50 mg tabletta	NL/H/1540/002	OGYI-T-4094/06	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 50 mg tabletta	NL/H/1540/002	OGYI-T-4094/21	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Lamictal 50 mg tabletten	NL/H/1540/002	RVG 103364	GLAXOSMITHKLINE B.V.	NL
Lamictal 50 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/004	1-20215	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 50 mg tabletter	NL/H/1540/002	12009	GLAXOSMITHKLINE AB	SE
Lamictal 50 mg tablety	NL/H/1540/002	21/802/92-B/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Lamictal 50 mg tablety	NL/H/1540/002	21/0404/09-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Lamictal 50 mg tuggtabletter/dispergerbara tabletter.	NL/H/1539/004	13028	GLAXOSMITHKLINE AB	SE
Lamictal 50 mg tuggu-/dreifitöflur	NL/H/1539/004	960234	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 50 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/004	96-1876	GLAXOSMITHKLINE AS	NO
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/020	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/023	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/026	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg	NL/H/1539/004	H/96/00871/027	GLAXOSMITHKLINE	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
žvecljive/disperzibilne tablete			TRADING SERVICES LIMITED	
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/028	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/029	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/030	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/133	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvecljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/132	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/018	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/019	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/021	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/022	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/024	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/025	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/134	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/135	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/136	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/137	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/138	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/139	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/140	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/141	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/142	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/143	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/144	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Lamictal 50 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/004	18618	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
LAMICTAL 50 mg, comprimé dispersible ou à croquer	NL/H/1539/004	NL21696	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal 50 mg, närimis/dispergeeruvad	NL/H/1539/004	425903	GLAXOSMITHKLINE (IRELAND) LIMITED	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletid.				
Lamictal d?s??a 100 mg.	NL/H/1540/003	2018103	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal Dispers 100 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/005	RVG 19117	GLAXOSMITHKLINE B.V.	NL
Lamictal Dispers 2 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/001	RVG 25344	GLAXOSMITHKLINE B.V.	NL
Lamictal Dispers 200 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/006	RVG 20927	GLAXOSMITHKLINE B.V.	NL
Lamictal Dispers 25 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/003	RVG 19116	GLAXOSMITHKLINE B.V.	NL
Lamictal Dispers 5 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/002	RVG 19115	GLAXOSMITHKLINE B.V.	NL
Lamictal Dispers 50 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/004	RVG 20926	GLAXOSMITHKLINE B.V.	NL
Lamictal δισκία 200 mg.	NL/H/1540/004	2018107	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal δισκία 25 mg	NL/H/1540/001	2018101	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal δισκία 50 mg.	NL/H/1540/002	2018102	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal μασώμενα/διασπειρόμενα δισκία 5 mg	NL/H/1539/002	2018104	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Lamictal, tabletter	NL/H/1540/001	13902	GLAXOSMITHKLINE PHARMA A/S	DK
Lamitrin S, 100 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/005	7873	GLAXOSMITHKLINE (IRELAND) LIMITED	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
Lamitrin S, 2 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/001	9643	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin S, 200 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/006	12342	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin S, 25 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/003	7872	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin S, 5 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/002	7871	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin S, 50 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/004	12341	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin, 100 mg, tabletki	NL/H/1540/003	R/3463	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin, 25 mg, tabletki	NL/H/1540/001	R/3548	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamitrin, 50 mg, tabletki	NL/H/1540/002	R/3462	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Lamotrigin Aristo 100 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	DE/H/0720/005	61971.04.00	ARISTO PHARMA GMBH (ART 57)	DE
Lamotrigin Aristo 200 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	DE/H/0720/006	61971.05.00	ARISTO PHARMA GMBH (ART 57)	DE
Lamotrigin Aristo 25 mg Tabletten zur Herstellung	DE/H/0720/003	61971.02.00	ARISTO PHARMA GMBH (ART 57)	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
einer Suspension zum Einnehmen				
Lamotrigin Aristo 50 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	DE/H/0720/004	61971.03.00	ARISTO PHARMA GMBH (ART 57)	DE
Lamotrigin Desitin 5 mg Tabletten	DE/H/7134/001/DC	7004905.00.00	DESITIN ARZNEIMITTEL GMBH	DE
Lamotrigine Aurobindo 100 mg dispersible tablets	NL/H/2260/004	MA807/03302	AUROBINDO PHARMA (MALTA) LIMITED	MT
Lamotrigine Aurobindo 200 mg dispersible tablets	NL/H/2260/005	MA807/03303	AUROBINDO PHARMA (MALTA) LIMITED	MT
Lamotrigine Aurobindo 50 mg dispersible tablets	NL/H/2260/003	MA807/03301	AUROBINDO PHARMA (MALTA) LIMITED	MT
LAMOTRIGINE EG 200 mg, comprimé dispersible ou à croquer	not available	NL30945	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
Lamotrigine Rudipharm 100 mg Tablets	not available	PL 49565/0111	RUDIPHARM LIMITED	XI
Lamotrigine Rudipharm 200 mg Tablets	not available	PL 49565/0112	RUDIPHARM LIMITED	XI
Lamotrigine Rudipharm 25 mg Tablets	not available	PL 49565/0109	RUDIPHARM LIMITED	XI
Lamotrigine Rudipharm 50 mg Tablets	not available	PL 49565/0110	RUDIPHARM LIMITED	XI
LAMOTRIGINE VIATRIS 100 mg, comprimé dispersible	not available	NL 30024	VIATRIS SANTE	FR
LAMOTRIGINE VIATRIS 200 mg, comprimé dispersible	not available	NL 30025	VIATRIS SANTE	FR
LAMOTRIGINE VIATRIS 25 mg, comprimé dispersible	not available	NL 30421	VIATRIS SANTE	FR
LAMOTRIGINE VIATRIS 50 mg, comprimé	not available	NL 30069	VIATRIS SANTE	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
dispersible				
LAMOTRIX 200 mg tablety	SE/H/0728/004	21/662/07-C	MEDOCHEMIE LTD.	CZ
MEDOTRIGIN 200 mg tablety	SE/H/0728/004	21/0413/07-S	MEDOCHEMIE LTD.	SK
Seizal 100 mg Tablets	not available	022071	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 200 mg Tablets	not available	022072	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 25 mg Tablets	not available	022069	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 50 mg Tablets	not available	022070	DELORBIS PHARMACEUTICALS LTD	CY
Ламиктал 100 mg таблетки	NL/H/1540/003	20011211	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Ламиктал 25 mg таблетки	NL/H/1540/001	2001 1209	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG
Ламиктал 5 mg таблетки за дъвчене/диспергиращи се таблетки	NL/H/1539/002	9700508	GLAXOSMITHKLINE (IRELAND) LIMITED	BG
Ламиктал 50 mg таблетки	NL/H/1540/002	2001 1210	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG