25 July 2018
EMA/600046/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: lamotrigine

Procedure no.: PSUSA/00001825/201711

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Seizal 25 mg Tablets	not available	22069	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 50 mg Tablets	not available	22070	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 100 mg Tablets	not available	22071	DELORBIS PHARMACEUTICALS LTD	CY
Seizal 200 mg Tablets	not available	22072	DELORBIS PHARMACEUTICALS LTD	CY
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/057	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/058	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/059	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/060	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/061	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/062	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/063	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg tablete	NL/H/1540/001	H/96/00871/064	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/065	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/066	GLAXOSMITHKLINE D.O.O.	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/96/00871/067	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/068	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/069	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/070	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/071	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg tablete	NL/H/1540/002	H/96/00871/072	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/073	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/074	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/075	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/076	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/077	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/078	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/079	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg tablete	NL/H/1540/003	H/96/00871/080	GLAXOSMITHKLINE D.O.O.	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 tablete	NL/H/1540/003	H/96/00871/081	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/082	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/083	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/084	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/085	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg tablete	NL/H/1540/004	H/96/00871/086	GLAXOSMITHKLINE D.O.O.	SI
Lamictal δισκία 25 mg	NL/H/1540/001	2018101	GLAXOSMITHKLINE AEBE	GR
Lamictal δισκία 100 mg	NL/H/1540/003	2018103	GLAXOSMITHKLINE AEBE	GR
Lamictal δισκία 200 mg	NL/H/1540/004	2018107	GLAXOSMITHKLINE AEBE	GR
Lamictal δισκία 50 mg	NL/H/1540/002	2018102	GLAXOSMITHKLINE AEBE	GR
Lamictal 25 mg tablets.	NL/H/1540/001	PL 00003/0272	THE WELLCOME FOUNDATION LTD	UK
Ламиктал 25 mg таблетки	NL/H/1540/001	2001 1209	GLAXOSMITHKLINE EOOD	BG
Lamictal, tabletter	NL/H/1540/001	13902	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 25 mg tablets	NL/H/1540/001	PA1077/061/001	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 25 mg tablets.	NL/H/1540/001	MA 192/00503	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 25 mg tabletten	NL/H/1540/001	RVG 103363	GLAXOSMITHKLINE B.V.	NL
Lamitrin, 25 mg, tabletki	NL/H/1540/001	R/3548	GLAXOSMITHKLINE EXPORT LTD	PL
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 comprimidos	NL/H/1540/001	2252781	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg comprimidos	NL/H/1540/001	2252880	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 25 mg tablety	NL/H/1540/001	21/0802/92-C/S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Lamictal 25 mg tabletter.	NL/H/1540/001	12008	GLAXOSMITHKLINE AB	SE
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/01	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/02	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/03	THE WELLCOME FOUNDATION LTD	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 25 mg comprimate	NL/H/1540/001	1742/2009/04	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/05	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/06	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/07	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/08	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/09	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg comprimate	NL/H/1540/001	1742/2009/10	THE WELLCOME FOUNDATION LTD	RO
Lamictal 25 mg tabletta	NL/H/1540/001	OGYI-T-4094/04	GLAXOSMITHKLINE KFT.	HU
Lamictal 25 mg tabletta	NL/H/1540/001	OGYI-T-4094/08	GLAXOSMITHKLINE KFT.	HU
Lamictal 100 mg tablets	NL/H/1540/003	PL 00003/0274	THE WELLCOME FOUNDATION LTD	UK
Ламиктал 100 mg таблетки	NL/H/1540/003	2001 1211	GLAXOSMITHKLINE EOOD	BG
Lamictal 100 mg, tablety	NL/H/1540/003	21/802/92-C/C	THE WELLCOME FOUNDATION LTD	CZ
Lamictal, tabletter	NL/H/1540/003	13904	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 100 mg tabletta	NL/H/1540/003	OGYI-T-4094/10	GLAXOSMITHKLINE KFT.	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 100 mg tabletta	NL/H/1540/003	OGYI-T-4094/15	GLAXOSMITHKLINE KFT.	HU
Lamictal 100 mg tablets	NL/H/1540/003	PA1077/061/003	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 100 mg tablets	NL/H/1540/003	MA 192/00505	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 100 mg tabletten	NL/H/1540/003	RVG 103366	GLAXOSMITHKLINE B.V.	NL
Lamitrin, 100 mg, tabletki	NL/H/1540/003	R/3463	GLAXOSMITHKLINE EXPORT LTD	PL
Lamictal 100 mg comprimidos	NL/H/1540/003	2253185	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/01	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/02	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/03	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/04	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/05	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/06	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/07	THE WELLCOME FOUNDATION LTD	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 100 mg comprimate	NL/H/1540/003	1744/2009/08	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg comprimate	NL/H/1540/003	1744/2009/09	THE WELLCOME FOUNDATION LTD	RO
Lamictal 100 mg tabletter	NL/H/1540/003	12010	GLAXOSMITHKLINE AB	SE
Lamictal 200 mg tablets	NL/H/1540/004	PL 00003/0297	THE WELLCOME FOUNDATION LTD	UK
Lamictal, tabletter	NL/H/1540/004	14790	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 200 mg tablets	NL/H/1540/004	PA1077/061/004	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 200 mg tabletten	NL/H/1540/004	RVG 103367	GLAXOSMITHKLINE B.V.	NL
Lamictal 200 mg comprimidos	NL/H/1540/004	2521581	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 200 mg tabletter	NL/H/1540/004	12011	GLAXOSMITHKLINE AB	SE
Lamictal 50 mg tablets.	NL/H/1540/002	PL 00003/0273	THE WELLCOME FOUNDATION LTD	UK
Ламиктал 50 mg таблетки	NL/H/1540/002	2001 1210	GLAXOSMITHKLINE EOOD	BG
Lamictal 50 mg, tablety	NL/H/1540/002	21/802/92-B/C	THE WELLCOME FOUNDATION LTD	CZ
Lamictal, tabletter	NL/H/1540/002	13903	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 50 mg tabletta	NL/H/1540/002	OGYI-T-4094/06	GLAXOSMITHKLINE KFT.	HU
Lamictal 50 mg tabletta	NL/H/1540/002	OGYI-T-4094/09	GLAXOSMITHKLINE KFT.	HU
Lamictal 50 mg tablets	NL/H/1540/002	PA1077/061/002	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 50 mg tablets.	NL/H/1540/002	MA 192/00504	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 50 mg tabletten	NL/H/1540/002	RVG 103364	GLAXOSMITHKLINE B.V.	NL
Lamitrin, 50 mg, tabletki	NL/H/1540/002	R/3462	GLAXOSMITHKLINE EXPORT LTD	PL
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 comprimidos	NL/H/1540/002	5291885	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/01	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/02	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/03	THE WELLCOME FOUNDATION LTD	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 50 mg comprimate	NL/H/1540/002	1743/2009/04	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/05	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/06	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/07	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/08	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg comprimate	NL/H/1540/002	1743/2009/09	THE WELLCOME FOUNDATION LTD	RO
Lamictal 50 mg tablety	NL/H/1540/002	21/0404/09-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Lamictal 50 mg tabletter.	NL/H/1540/002	12009	GLAXOSMITHKLINE AB	SE
Lamictal 5 mg, žvýkací/dispergovatelné tablety	NL/H/1539/002	21/483/96-A/C	THE WELLCOME FOUNDATION LTD	CZ
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
Lamotrigine Aurobindo 50 mg dispersible tablets	NL/H/2260/003	MA807/03301	AUROBINDO PHARMA (MALTA) LIMITED	MT
Lamotrigine Aurobindo 100 mg dispersible tablets	NL/H/2260/004	MA807/03302	AUROBINDO PHARMA (MALTA) LIMITED	MT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Lamotrigine Aurobindo 200 mg dispersible tablets	NL/H/2260/005	MA807/03303	AUROBINDO PHARMA (MALTA) LIMITED	MT
Lamictal 2 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/001	135577	GLAXOSMITHKLINE PHARMA GMBH.	AT
Ламиктал 2 mg таблетки за дъвчене/диспергиращи се таблетки	NL/H/1539/001	20140091	GLAXOSMITHKLINE EOOD	BG
Lamictal 5 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/002	16176	GLAXO GROUP LIMITED	CY
Lamictal 2 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/001	22006	GLAXO GROUP LIMITED	CY
Lamictal 2 mg tablete za žvakanje/tablete za oralnu suspenziju	NL/H/1539/001	UP/I-530-09/13-01/70	GLAXOSMITHKLINE D.O.O.	HR
Lamictal 25 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/003	2521888	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg tuggtabletter / dispergerbara tabletter	NL/H/1539/002	12320	GLAXOSMITHKLINE OY	FI
Lamictal 2 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/001	23162	GLAXOSMITHKLINE OY	FI
Lamictal 25 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/003	12321	GLAXOSMITHKLINE OY	FI
Lamictal 50 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/004	12322	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
Lamictal 100 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/005	12323	GLAXOSMITHKLINE OY	FI
Lamictal 200 mg dispergerbara tabletter / tuggtabletter	NL/H/1539/006	12324	GLAXOSMITHKLINE OY	FI
Lamictal 5 mg kauw-/dispergeerbare tabletten	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	BE 185211	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 25 mg kauw-/dispergeerbare tabletten	NL/H/1539/003	BE 185227	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/003	BE185227	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 50 mg kauw-/dispergeerbare tabletten	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	BE185236	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 100 mg kauw-/dispergeerbare tabletten	NL/H/1539/005	BE 185245	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
Lamictal 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/005	BE185245	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 200 mg kauw-/dispergeerbare tabletten	NL/H/1539/006	BE 185254	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/006	BE185254	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg kauw-/dispergeerbare tabletten	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	BE 228611	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/001	0260/09/10/0605	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 5 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/002	0260/09/10/0606	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/005	0260/09/10/0609	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 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/006	0260/09/10/0610	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/003	0260/09/10/0607	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/004	260/09/10/0608	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 2 mg compresse masticabili/dispersibili	NL/H/1539/001	027807179	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 2 mg žvecljive/disperzibilne tablete	NL/H/1539/001	H/96/00871/001	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/002	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/003	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvecljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/004	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/005	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/006	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/007	GLAXOSMITHKLINE D.O.O.	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/96/00871/008	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/009	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/010	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/011	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/012	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/013	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/014	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/015	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/016	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 25 mg žvečljive/disperzibilne tablete	NL/H/1539/003	H/96/00871/017	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/018	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/019	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/020	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/021	GLAXOSMITHKLINE D.O.O.	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/96/00871/022	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/023	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/024	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/025	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/026	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/027	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/028	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/029	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 50 mg žvečljive/disperzibilne tablete	NL/H/1539/004	H/96/00871/030	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvecljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/031	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/032	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/033	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/034	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/035	GLAXOSMITHKLINE D.O.O.	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 žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/036	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/037	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/038	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/039	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/040	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/041	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/042	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 100 mg žvečljive/disperzibilne tablete	NL/H/1539/005	H/96/00871/043	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/044	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/045	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/046	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/047	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/048	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/049	GLAXOSMITHKLINE D.O.O.	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/96/00871/050	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/051	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/052	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/053	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/054	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/055	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 200 mg žvečljive/disperzibilne tablete	NL/H/1539/006	H/96/00871/056	GLAXOSMITHKLINE D.O.O.	SI
Lamictal μασώμενα/διασπειρόμενα δισκία 5 mg	NL/H/1539/002	2018104	GLAXOSMITHKLINE AEBE	GR
Lamictal μασώμενα/διασπειρόμενα δισκία 2 mg	NL/H/1539/001	2018110	GLAXOSMITHKLINE AEBE	GR
Lamictal 2 mg chewable/dispersible tablets	NL/H/1539/001	PL 00003/0375	THE WELLCOME FOUNDATION LTD	UK
Lamictal 2 mg comprimés à croquer/dispersibles	NL/H/1539/001	BE 228611	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 2 mg, žvýkací/dispergovatelné tablety	NL/H/1539/001	21/330/01-C	THE WELLCOME FOUNDATION LTD	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/001	31464	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal, 2 mg närimis/dispergeeruvad tabletid	NL/H/1539/001	332800	GLAXO WELLCOME UK LIMITED	EE
Lamictal 2 mg purutabletti / dispergoituva tabletti	NL/H/1539/001	23162	GLAXOSMITHKLINE OY	FI
LAMICTAL 2 mg, comprimé dispersible ou à croquer	NL/H/1539/001	NL25689	LABORATOIRE GLAXOSMITHKLINE	FR
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 tuggu-/dreifitöflur	NL/H/1539/001	IS/1/08/112/01	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 2 mg chewable/dispersible tablets	NL/H/1539/001	PA1077/061/005	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Lamictal 2 mg košļājamās/ disperģējamās tabletes	NL/H/1539/001	02-0401	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 2 mg comprimés à croquer/dispersibles	NL/H/1539/001	0260/09/10/0605	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 2 mg chewable/dispersible tablets	NL/H/1539/001	MA 192/00506	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal Dispers 2 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/001	RVG 25344	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
Lamictal 2 mg tyggetabletter/dispergerbare tabletter	NL/H/1539/001	00-2581	GLAXOSMITHKLINE AS	NO
Lamitrin S, 2 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/001	9643	GLAXOSMITHKLINE EXPORT LTD	PL
Lamictal 2 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/001	3338183	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 2 mg comprimate masticabile/dispersabile	NL/H/1539/001	1745/2009/01	THE WELLCOME FOUNDATION LTD	RO
Lamictal 2 mg žuvacie/dispergovatelné tablety	NL/H/1539/001	21/0150/08-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Lamictal 2 mg comprimidos masticables/dispersables	NL/H/1539/001	64.391	GLAXOSMITHKLINE S.A.	ES
Lamictal 2 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/001	16321	GLAXOSMITHKLINE AB	SE
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/013	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg comprimés à croquer/dispersibles.	NL/H/1539/002	2009 10 0606	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal Dispers 5 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/002	RVG 19115	GLAXOSMITHKLINE BV	NL

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Lamitrin S, 5 mg, tabletki do rozgryzania i żucia / do sporządzania zawiesiny	NL/H/1539/002	7871	GLAXOSMITHKLINE EXPORT LTD	PL
Lamictal 5 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/002	2521680	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg žuvacie/dispergovateľné tablety	NL/H/1539/002	21/0208/98-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Lamictal 5 mg comprimidos masticables/dispersables	NL/H/1539/002	61.561	GLAXOSMITHKLINE S.A.	ES
Lamictal 25 mg chewable/dispersible tablets	NL/H/1539/003	PL 00003/0347	THE WELLCOME FOUNDATION LTD	UK
Lamictal 25 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/003	1-20875	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 25 mg comprimés à croquer/dispersibles	NL/H/1539/003	BE 185227	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 25 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/003	16177	GLAXO GROUP LIMITED	CY
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/003	15831	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 25 mg, närimis/dispergeeruvad tabletid	NL/H/1539/003	425403	GLAXO WELLCOME UK LIMITED	EE
Lamictal 25 mg purutabletti / dispergoituva tabletti	NL/H/1539/003	12321	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
LAMICTAL 25 mg, comprimé dispersible ou à croquer	NL/H/1539/003	NL20297	LABORATOIRE GLAXOSMITHKLINE	FR
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 tuggu-/dreifitöflur	NL/H/1539/003	940290	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 25 mg chewable/dispersible tablets	NL/H/1539/003	PA1077/061/007	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
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	027807130	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg compresse masticabili/dispersibili	NL/H/1539/003	027807142	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 25 mg košļājamās/ disperģējamās tabletes	NL/H/1539/003	97-0588	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/003	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/004	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/005	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/014	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/015	GLAXOSMITHKLINE LIETUVA UAB	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 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/016	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/017	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/018	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/019	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/057	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/003	LT/1/97/1271/058	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 25 mg comprimés à croquer/dispersibles	NL/H/1539/003	0260/09/10/0607	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal 25 mg chewable/dispersible tablets	NL/H/1539/003	MA 192/00502	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal Dispers 25 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/003	RVG 19116	GLAXOSMITHKLINE B.V.	NL
Lamictal 25 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/003	8199	GLAXOSMITHKLINE AS	NO
Lamitrin S, 25 mg, tabletki do rozgryzania i żucia / do sporządzania zawiesiny	NL/H/1539/003	7872	GLAXOSMITHKLINE EXPORT LTD	PL
Lamictal 25 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/003	5291984	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
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 tuggtabletter/dispergerbara tabletter.	NL/H/1539/003	12791	GLAXOSMITHKLINE AB	SE
Lamictal 5 mg chewable/dispersible tablets.	NL/H/1539/002	PL 00003/0346	THE WELLCOME FOUNDATION LTD	UK
Lamictal 5 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/002	1-20888	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 5 mg comprimés à croquer/dispersibles	NL/H/1539/002	BE 185211	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Ламиктал 5 mg таблетки за дъвчене/диспергиращи се таблетки	NL/H/1539/002	970 0508	GLAXO GROUP LIMITED	BG
Lamictal 5 mg, žvýkací/dispergovatelné tablety	NL/H/1539/002	21/483/96-A/C	THE WELLCOME FOUNDATION LTD	CZ
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/002	15830	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 5 mg, närimis/dispergeeruvad tabletid	NL/H/1539/002	425703	GLAXO WELLCOME UK LIMITED	EE
Lamictal 5 mg purutabletti / dispergoituva tabletti	NL/H/1539/002	12320	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
LAMICTAL 5 mg, comprimé dispersible ou à croquer	NL/H/1539/002	NL20296	LABORATOIRE GLAXOSMITHKLINE	FR
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 rágótabletta/diszpergálódó tabletta	NL/H/1539/002	OGYI-T-4094/16	GLAXOSMITHKLINE KFT.	HU
Lamictal 5 mg tuggu-/dreifitöflur	NL/H/1539/002	940289 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
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 košļājamās/ disperģējamās tabletes	NL/H/1539/002	97-0587	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/002	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/043	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/011	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/010	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/009	GLAXOSMITHKLINE LIETUVA UAB	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 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/012	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg chewable/dispersible tablets.	NL/H/1539/002	MA 192/00501	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Lamictal 5 mg tyggetabletter/dispergerbare tabletter	NL/H/1539/002	8198	GLAXOSMITHKLINE AS	NO
Lamictal 5 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/002	2521789	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/01	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/02	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/03	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/04	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/05	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/06	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/07	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/002	12790	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 50 mg chewable/dispersible tablets	NL/H/1539/004	PL 00003/0368	THE WELLCOME FOUNDATION LTD	UK
Lamictal 50 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/004	1-20215	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 50 mg comprimés à croquer/dispersibles	NL/H/1539/004	BE 185236	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 50 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/004	18618	GLAXO GROUP LIMITED	CY
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/004	17951	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 50 mg, närimis/dispergeeruvad tabletid	NL/H/1539/004	425903	GLAXO WELLCOME UK LIMITED	EE
Lamictal 50 mg purutabletti / dispergoituva tabletti	NL/H/1539/004	12322	GLAXOSMITHKLINE OY	FI
LAMICTAL 50 mg, comprimé dispersible ou à croquer	NL/H/1539/004	NL21696	LABORATOIRE GLAXOSMITHKLINE	FR
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 tuggu-/dreifitöflur.	NL/H/1539/004	960234 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 50 mg chewable/dispersible tablets	NL/H/1539/004	PA1077/061/008	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 compresse masticabili/dispersibili	NL/H/1539/004	027807080	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg compresse masticabili/dispersibili	NL/H/1539/004	027807155	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 50 mg košļājamās/ disperģējamās tabletes	NL/H/1539/004	97-0589	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/006	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/020	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/021	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/022	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/023	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/024	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/025	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/026	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/027	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/028	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/044	GLAXOSMITHKLINE LIETUVA UAB	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 tabletės	NL/H/1539/004	LT/1/97/1271/045	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/046	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/004	LT/1/97/1271/059	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 50 mg comprimés à croquer/dispersibles	NL/H/1539/004	260/09/10/0608	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal Dispers 50 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/004	RVG 20926	GLAXOSMITHKLINE B.V.	NL
Lamictal 50 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/004	96-1876	GLAXOSMITHKLINE AS	NO
Lamitrin S, 50 mg, tabletki do rozgryzania i żucia / do sporządzania zawiesiny	NL/H/1539/004	12341	GLAXOSMITHKLINE EXPORT LTD	PL
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 tuggtabletter/dispergerbara tabletter.	NL/H/1539/004	13028	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 chewable/dispersible tablets	NL/H/1539/005	PL 00003/0348	THE WELLCOME FOUNDATION LTD	UK
Lamictal 100 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/005	1-20886	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 100 mg comprimés à croquer/dispersibles	NL/H/1539/005	BE 185245	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 100 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/005	16178	GLAXO GROUP LIMITED	CY
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/005	15832	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 100 mg, närimis/dispergeeruvad tabletid	NL/H/1539/005	426003	GLAXO WELLCOME UK LIMITED	EE
Lamictal 100 mg purutabletti / dispergoituva tabletti	NL/H/1539/005	12323	GLAXOSMITHKLINE OY	FI
LAMICTAL 100 mg, comprimé dispersible ou à croquer	NL/H/1539/005	NL20298	LABORATOIRE GLAXOSMITHKLINE	FR
Lamictal® 100 mg Kautabletten bzw. Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/1539/005	33122.03.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Lamictal 100 mg tuggu-/dreifitöflur	NL/H/1539/005	940291	GLAXOSMITHKLINE PHARMA A/S	IS
Lamictal 100 mg chewable/dispersible tablets	NL/H/1539/005	PA1077/061/009	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 100 mg compresse masticabili/dispersibili	NL/H/1539/005	027807078	GLAXOSMITHKLINE S.P.A.	IT
Lamictal 100 mg košļājamās/ disperģējamās tabletes	NL/H/1539/005	97-0590	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/007	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/005	LT/1/97/1271/029	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/030	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/031	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/032	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/033	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/034	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/035	GLAXOSMITHKLINE LIETUVA UAB	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 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/047	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/048	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/049	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/050	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/005	LT/1/97/1271/051	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 100 mg comprimés à croquer/dispersibles	NL/H/1539/005	0260/09/10/0609	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal Dispers 100 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/005	RVG 19117	GLAXOSMITHKLINE B.V.	NL
Lamictal 100 mg tyggetabletter/dispergerbare tabletter.	NL/H/1539/005	8200	GLAXOSMITHKLINE AS	NO
Lamitrin S, 100 mg, tabletki do rozgryzania i zucia / do sporzadzania zawiesiny	NL/H/1539/005	7873	GLAXOSMITHKLINE EXPORT LTD	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
Lamictal 100 mg comprimidos dispersíveis ou para mastigar	NL/H/1539/005	2521987	GLAXOSMITHKLINE - PRODUTOS FARMACEUTICOS, LDA	PT
Lamictal 100 mg žuvacie/dispergovateľné tablety	NL/H/1539/005	21/0405/09-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Lamictal 100 mg comprimidos masticables/dispersables	NL/H/1539/005	61.568	GLAXOSMITHKLINE S.A.	ES
Lamictal 100 mg tuggtabletter/dispergerbara tabletter	NL/H/1539/005	12792	GLAXOSMITHKLINE AB	SE
Lamictal 200 mg chewable/dispersible tablets	NL/H/1539/006	PL 00003/0369	THE WELLCOME FOUNDATION LTD	UK
Lamictal 200 mg Tabletten (kaubar/suspendierbar)	NL/H/1539/006	1-20216	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 200 mg comprimés à croquer/dispersibles	NL/H/1539/006	BE 185254	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lamictal 200 mg μασώμενα/διασπειρόμενα δισκία	NL/H/1539/006	18617	GLAXO GROUP LIMITED	CY
Lamictal, tyggetabletter/dispergible tabletter	NL/H/1539/006	17952	GLAXOSMITHKLINE PHARMA A/S	DK
Lamictal 200 mg purutabletti / dispergoituva tabletti	NL/H/1539/006	12324	GLAXOSMITHKLINE OY	FI
LAMICTAL 200 mg, comprimé dispersible ou à croquer	NL/H/1539/006	NL21697	LABORATOIRE GLAXOSMITHKLINE	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
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 rágótabletta/diszpergálódó tabletta	NL/H/1539/006	OGYI-T-4094/17	GLAXOSMITHKLINE KFT.	HU
Lamictal 200 mg tuggu-/dreifitöflur	NL/H/1539/006	960235 (IS)	GLAXOSMITHKLINE PHARMA A/S	IS
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 košļājamās/ disperģējamās tabletes	NL/H/1539/006	97-0591	GLAXOSMITHKLINE LATVIA SIA	LV
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/008	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/036	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/037	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/038	GLAXOSMITHKLINE LIETUVA UAB	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 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/039	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/040	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/041	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/042	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/052	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/053	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/054	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/055	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 200 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/006	LT/1/97/1271/056	GLAXOSMITHKLINE LIETUVA UAB	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 200 mg comprimés à croquer/dispersibles	NL/H/1539/006	0260/09/10/0610	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamictal Dispers 200 mg, kauwtabletten/dispergeerbare tabletten	NL/H/1539/006	RVG 20927	GLAXOSMITHKLINE B.V.	NL
Lamictal 200 mg tyggetabletter/dispergerbare tabletter	NL/H/1539/006	96-1877	GLAXOSMITHKLINE AS	NO
Lamitrin S, 200 mg, tabletki do rozgryzania i żucia / do sporządzania zawiesiny	NL/H/1539/006	12342	GLAXOSMITHKLINE EXPORT LTD	PL
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 tuggtabletter/dispergerbara tabletter	NL/H/1539/006	13029	GLAXOSMITHKLINE AB	SE
Lamictal 5 mg comprimés à croquer/dispersibles	NL/H/1539/002	BE477093	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 kauw-/dispergeerbare tabletten	NL/H/1539/002	BE477093	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
Lamictal 2 mg rágótabletta/diszpergálódó tabletta	NL/H/1539/001	OGYI-T-4094/18	GLAXOSMITHKLINE KFT.	HU
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/087	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/088	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/089	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/090	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/091	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg žvečljive/disperzibilne tablete	NL/H/1539/002	H/96/00871/092	GLAXOSMITHKLINE D.O.O.	SI
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/061	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/063	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/066	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/062	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/064	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg kramtomosios ar disperguojamosios tabletės	NL/H/1539/002	LT/1/97/1271/065	GLAXOSMITHKLINE LIETUVA UAB	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 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
Lamictal 2 mg kramtomosios ar disperguojamosios tabletes	NL/H/1539/001	LT/1/97/1271/060	GLAXOSMITHKLINE LIETUVA UAB	LT
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/08	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/09	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/10	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/11	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/12	THE WELLCOME FOUNDATION LTD	RO
Lamictal 5 mg comprimate masticabile/dispersabile	NL/H/1539/002	1746/2009/13	THE WELLCOME FOUNDATION LTD	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
LAMOTRIGINE EG 200 mg, comprimé dispersible ou à croquer	not available	NL30945	EG LABO LABORATOIRES EUROGENERICS	FR
Lamictal 50 mg tablete	not available	HR-H-052719076	GLAXOSMITHKLINE D.O.O.	HR
Labileno 100 mg comprimidos masticables/dispersables	not available	64.943	GLAXOSMITHKLINE S.A.	ES
Crisomet 50 mg comprimidos masticables/dispersables	not available	64.464	GLAXOSMITHKLINE S.A.	ES
Crisomet 100 mg comprimidos masticables/dispersables	not available	64.462	GLAXOSMITHKLINE S.A.	ES
Crisomet 25 mg comprimidos masticables/dispersables	not available	63.747	GLAXOSMITHKLINE S.A.	ES
Crisomet 200 mg comprimidos masticables/dispersables	not available	64.463	GLAXOSMITHKLINE S.A.	ES
Labileno 25 mg comprimidos masticables/dispersables.	not available	63.288	GLAXOSMITHKLINE S.A.	ES
Labileno 200 mg comprimidos masticables/dispersables.	not available	65.283	GLAXOSMITHKLINE S.A.	ES
Labileno 50 mg comprimidos masticables/dispersables	not available	65.284	GLAXOSMITHKLINE S.A.	ES
Lamotrigin GSK 5 mg Tabletten (kaubar/suspendierbar)	not available	1-25285	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamotrigin GSK 25 mg Tabletten (kaubar/suspendierbar)	not available	1-25286	GLAXOSMITHKLINE PHARMA GMBH.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation Number	National Authorisation Number	MAH of product in the Member State	Member State where product is authorised
Lamotrigin GSK 50 mg Tabletten (kaubar/suspendierbar)	not available	1-25287	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamotrigin GSK 100 mg Tabletten (kaubar/suspendierbar)	not available	1-25288	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamotrigin GSK 200 mg Tabletten (kaubar/suspendierbar)	not available	1-25289	GLAXOSMITHKLINE PHARMA GMBH.	AT
Lamictal 25 mg tablete	not available	HR-H-211198060	GLAXOSMITHKLINE D.O.O.	HR
Lamictal 100 mg tablete	not available	HR-H-695293019	GLAXOSMITHKLINE D.O.O.	HR
LAMICTAL 5 mg tablete za žvakanje/tablete za oralnu suspenziju	not available	HR-H-287736527	GLAXOSMITHKLINE D.O.O.	HR
Lamitrin 50 mg tabletta	not available	OGYI-T-8805/04	GLAXOSMITHKLINE KFT.	HU
Lamitrin 50 mg tabletta	not available	OGYI-T-8805/10	GLAXOSMITHKLINE KFT.	HU
Lamitrin 100 mg tabletta	not available	OGYI-T-8805/06	GLAXOSMITHKLINE KFT.	HU
Lamitrin 100 mg tabletta	not available	OGYI-T-8805/11	GLAXOSMITHKLINE KFT.	HU
Lamitrin 200 mg tabletta	not available	OGYI-T-8805/08	GLAXOSMITHKLINE KFT.	HU
Lamitrin 200 mg tabletta	not available	OGYI-T-8805/12	GLAXOSMITHKLINE KFT.	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
Lamitrin 25 mg tabletta	not available	OGYI-T-8805/02	GLAXOSMITHKLINE KFT.	HU
Lamitrin 25 mg tabletta	not available	OGYI-T-8805/09	GLAXOSMITHKLINE KFT.	HU
Lambipol 25 mg kauw-/dispergeerbare tabletten	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 266052	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg kauw-/dispergeerbare tabletten	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	BE 266077	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 50 mg kauw-/dispergeerbare tabletten	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	BE 266095	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 100 mg kauw-/dispergeerbare tabletten	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	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 200 mg kauw-/dispergeerbare 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	BE 266156	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	260/05/01/0026	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/001	260/05/01/0027	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 50 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/002	260/05/01/0028	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 100 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/003	260/05/01/0029	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 200 mg Kautabletten/Tabletten zur Herstellung einer Suspension zum Einnehmen	NL/H/2051/004	260/05/01/0030	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
Lambipol 50 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/002	RVG 107454	GLAXOSMITHKLINE B.V.	NL
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	260/05/01/0028	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 100 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/003	RVG 107455	GLAXOSMITHKLINE B.V.	NL
Lambipol 100 mg comprimés à croquer/dispersibles	NL/H/2051/003	BE 266122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Lambipol 100 mg comprimés à croquer/dispersibles	NL/H/2051/003	260/05/01/0029	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 200 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/004	RVG 107456	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	260/05/01/0030	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg, kauwtabletten/dispergeerbare tabletten	NL/H/2051/001	RVG 107451	GLAXOSMITHKLINE B.V.	NL
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	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	260/05/01/0026	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lambipol 25 mg comprimés à croquer/dispersibles	NL/H/2051/001	260/05/01/0027	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Lamotrigin Aristo® 25 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	DE/H/0720/003	61971.02.00	ARISTO PHARMA GMBH (ART 57)	DE
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 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