16 October 2025
EMADOC-1700519818-2663923
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): gabapentin

EURD list No. PSUSA/00001499/202502

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
GabaLiquid GeriaSan 50 mg/ml Lösung zum Einnehmen	DE/H/4632/001	96498.00.00	INFECTOPHARM ARZNEIMITTEL UND CONSILIUM GMBH	DE
Gabapentin Activase 100mg Capsules	not available	PL28444/0261	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase 100mg Capsules	not available	PL28444/0261	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase	not available	PL28444/0261	ACTIVASE	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
100mg Capsules			PHARMACEUTICALS LIMITED	
Gabapentin Activase 100mg Capsules	not available	PL28444/0261	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase 100mg Capsules	not available	PL28444/0261	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase 100mg Capsules	not available	PL28444/0261	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase 300mg Capsules	not available	PL28444/0262	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Activase 300mg Capsules	not available	PL28444/0262	ACTIVASE PHARMACEUTICALS LIMITED	XI
Gabapentin Brown & Burk 50 mg/ml sugar free oral solution	not available	PL 25298/0291	BROWN & BURK UK LIMITED	XI
Gabapentin Colonis 50mg/ml Oral Solution	UK/H/6181/001	PL 41344/0028	COLONIS PHARMA LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Glenmark 50 mg/ml Oral Solution	not available	PL 25258/0315	GLENMARK PHARMACEUTICALS EUROPE LIMITED	XI
Gabapentin Medreich 100mg Capsules	not available	PL 21880/0075	MEDREICH PLC	XI
Gabapentin Medreich 100mg Capsules	not available	PL 21880/0075	MEDREICH PLC	XI
Gabapentin Medreich 100mg Capsules	not available	PL 21880/0075	MEDREICH PLC	XI
Gabapentin Medreich 300mg Capsules	not available	PL 21880/0076	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Medreich 400mg Capsules	not available	PL21880/0077	MEDREICH PLC	XI
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Rosemont 50mg/ml Oral Solution	IE/H/1015/001	PA23081/010/001	TAW PHARMA (IRELAND) LTD	IE
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI
Gabapentin Teva 50 mg/ml oral solution.	not available	PL 00289/2154	TEVA UK LIMITED	XI
Gabapentin Thame 50mg/ml Oral Solution	UK/H/7068/001/MR	MA1296/00601	SYRI PHARMA LIMITED	MT
Gabapentin Thame 50mg/ml Oral Solution	UK/H/7068/001/MR	MA1296/00601	SYRI PHARMA LIMITED	MT
Gabapentin Thame 50mg/ml Oral Solution	UK/H/7068/001/MR	MA1296/00601	SYRI PHARMA LIMITED	MT
Gabapentin Thame 50mg/ml Oral Solution	not available	PL 39307/0068	SYRI LIMITED T/A THAME LABORATORIES	XI
Gabapentin Thame 50mg/ml Oral Solution	not available	PL 39307/0068	SYRI LIMITED T/A THAME LABORATORIES	XI
Gabapentin Thame 50mg/ml Oral Solution	not available	PL 39307/0068	SYRI LIMITED T/A THAME LABORATORIES	XI
Neurisol 50 mg/ml mikstur, oppløsning	SE/H/2518/001	23-15937	ORESUND PHARMA APS	NO
Neurisol 50 mg/ml oraaliliuos	SE/H/2518/001	43274	ORESUND PHARMA APS	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
Neurisol 50 mg/ml oral lösning	SE/H/2518/001	65711	ORESUND PHARMA APS	SE
Neurisol, oral opløsning	SE/H/2518/001	70844	ORESUND PHARMA APS	DK
Neurontin 100 mg cápsulas	DE/H/0899/001	5815584	VIATRIS HEALTHCARE LIMITED	PT
Neurontin 100 mg cápsulas	DE/H/0899/001	2433084	VIATRIS HEALTHCARE LDA.	PT
Neurontin 100 mg cápsulas	DE/H/0899/001	2433183	UPJOHN EESV	PT
Neurontin 100 mg capsule rigide	DE/H/0899/001	028740013	VIATRIS PHARMA S.R.L.	IT
NEURONTIN 100 mg cietas kapsulas	DE/H/0899/001	01-0276	UPJOHN EESV	LV
Neurontin 100 mg gélules	DE/H/0899/001	BE181151	VIATRIS HEALTHCARE	BE
Neurontin 100 mg gélules	DE/H/0899/001	2009030266	VIATRIS HEALTHCARE	LU
Neurontin 100 mg hard capsules	DE/H/0899/001	PA 23055/004/001	UPJOHN EESV	IE
Neurontin 100 mg hard capsules	DE/H/0899/001	PL 50622/0046	UPJOHN UK LIMITED	XI
Neurontin 100 mg hårda kapslar	DE/H/0899/001	12072	UPJOHN EESV	SE
Neurontin 100 mg harde capsules	DE/H/0899/001	BE181151	VIATRIS HEALTHCARE	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
Neurontin 100 mg harde kapsler	DE/H/0899/001	00-2995	UPJOHN EESV	NO
Neurontin 100 mg Hartkapseln	DE/H/0899/001	BE181151	VIATRIS HEALTHCARE	BE
Neurontin 100 mg Hartkapseln	DE/H/0899/001	2009030266	VIATRIS HEALTHCARE	LU
Neurontin 100 mg hörð hylki	DE/H/0899/001	IS/1/01/119/01	UPJOHN EESV	IS
Neurontin 100 mg kemény kapszula	DE/H/0899/001	OGYI-T-4966/01	VIATRIS HEALTHCARE LTD	HU
Neurontin 100 mg kemény kapszula	DE/H/0899/001	OGYI-T-4966/02	VIATRIS HEALTHCARE LTD	HU
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/002	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/001	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/008	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/004	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/006	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/005	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/011	UPJOHN EESV	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
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/007	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/009	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/003	UPJOHN EESV	SI
Neurontin 100 mg trde kapsule	DE/H/0899/001	H/99/01105/010	UPJOHN EESV	SI
NEURONTIN 100 mg tvrdé tobolky	DE/H/0899/001	21/461/97-C	UPJOHN EESV	CZ
NEURONTIN 100 mg, gélule	DE/H/0899/001	34009 337 898 6 2	VIATRIS UP	FR
NEURONTIN 100 mg, gélule	DE/H/0899/001	34009 337 896 3 3	VIATRIS UP	FR
NEURONTIN 100, 100 mg, kapsułki twarde	DE/H/0899/001	7692	UPJOHN EESV	PL
Neurontin 100, harde capsules 100 mg	DE/H/0899/001	RVG 22481	VIATRIS HEALTHCARE LTD	NL
Neurontin 300 mg cápsulas	DE/H/0899/002	2433282	VIATRIS HEALTHCARE LDA.	PT
Neurontin 300 mg cápsulas	DE/H/0899/002	3669587	VIATRIS HEALTHCARE LDA.	PT
Neurontin 300 mg cápsulas	DE/H/0899/002	5815683	VIATRIS HEALTHCARE LDA.	PT
Neurontin 300 mg cápsulas duras	DE/H/0899/002	60.621	VIATRIS HEALTHCARE LIMITED	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Neurontin 300 mg capsule rigide	DE/H/0899/002	028740025	VIATRIS PHARMA S.R.L.	IT
NEURONTIN 300 mg cietas kapsulas	DE/H/0899/002	01-0277	UPJOHN EESV	LV
Neurontin 300 mg gélules	DE/H/0899/002	BE181142	VIATRIS HEALTHCARE	BE
Neurontin 300 mg gélules	DE/H/0899/002	2009030267	VIATRIS HEALTHCARE	LU
Neurontin 300 mg hard capsules	DE/H/0899/002	PA 23055/004/002	UPJOHN EESV	IE
Neurontin 300 mg hard capsules	DE/H/0899/002	PL 50622/0047	UPJOHN UK LIMITED	XI
Neurontin 300 mg hårda kapslar	DE/H/0899/002	11662	UPJOHN EESV	FI
Neurontin 300 mg hårda kapslar	DE/H/0899/002	12058	UPJOHN EESV	SE
Neurontin 300 mg harde capsules	DE/H/0899/002	BE181142	VIATRIS HEALTHCARE	BE
Neurontin 300 mg harde kapsler	DE/H/0899/002	8220	UPJOHN EESV	NO
Neurontin 300 mg Hartkapseln	DE/H/0899/002	BE181142	VIATRIS HEALTHCARE	BE
Neurontin 300 mg Hartkapseln	DE/H/0899/002	2009030267	VIATRIS HEALTHCARE	LU
Neurontin 300 mg hörð hylki	DE/H/0899/002	920269	UPJOHN EESV	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Neurontin 300 mg kemény kapszula	DE/H/0899/002	OGYI-T-4966/04	VIATRIS HEALTHCARE LTD	HU
Neurontin 300 mg kemény kapszula	DE/H/0899/002	OGYI-T-4966/03	VIATRIS HEALTHCARE LTD	HU
Neurontin 300 mg kovat kapselit	DE/H/0899/002	11662	UPJOHN EESV	FI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/012	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/022	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/016	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/015	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/020	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/019	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/017	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/018	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/014	UPJOHN EESV	SI
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/013	UPJOHN EESV	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
Neurontin 300 mg trde kapsule	DE/H/0899/002	H/99/01105/021	UPJOHN EESV	SI
Neurontin 300 mg tvrde kapsule	not available	HR-H-198579675	UPJOHN EESV	HR
NEURONTIN 300 mg tvrdé tobolky	DE/H/0899/002	21/462/97-C	UPJOHN EESV	CZ
Neurontin 300 mg καψάκια σκληρά	DE/H/0899/002	17445	VIATRIS HELLAS LTD.	CY
Neurontin 300 mg καψάκια σκληρά	DE/H/0899/002	26441/9-4-2009	VIATRIS HELLAS LTD.	GR
NEURONTIN 300 mg, gélule	DE/H/0899/002	34009 337 901 7 2	VIATRIS UP	FR
NEURONTIN 300 mg, gélule	DE/H/0899/002	34009 337 900 0 4	VIATRIS UP	FR
NEURONTIN 300, 300 mg, kapsułki twarde	DE/H/0899/002	7693	UPJOHN EESV	PL
Neurontin 300, harde capsules 300 mg	DE/H/0899/002	RVG 22482	VIATRIS HEALTHCARE LTD	NL
Neurontin 400 mg cápsulas	DE/H/0899/003	5815782	VIATRIS HEALTHCARE LDA.	PT
Neurontin 400 mg cápsulas	DE/H/0899/003	2433381	VIATRIS HEALTHCARE LDA.	PT
Neurontin 400 mg cápsulas duras	DE/H/0899/003	60.622	VIATRIS HEALTHCARE LIMITED	ES
Neurontin 400 mg capsule rigide	DE/H/0899/003	028740037	VIATRIS PHARMA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
NEURONTIN 400 mg cietas kapsulas	DE/H/0899/003	01-0278	UPJOHN EESV	LV
Neurontin 400 mg gélules	DE/H/0899/003	BE181133	VIATRIS HEALTHCARE	BE
Neurontin 400 mg gélules	DE/H/0899/003	2009030268	VIATRIS HEALTHCARE	LU
Neurontin 400 mg hard capsules	DE/H/0899/003	PA 23055/004/003	UPJOHN EESV	IE
Neurontin 400 mg hard capsules	DE/H/0899/003	PL 50622/0048	UPJOHN UK LIMITED	XI
Neurontin 400 mg hårda kapslar	DE/H/0899/003	11663	UPJOHN EESV	FI
Neurontin 400 mg harde capsules	DE/H/0899/003	BE181133	VIATRIS HEALTHCARE	BE
Neurontin 400 mg harde kapsler	DE/H/0899/003	8221	UPJOHN EESV	NO
Neurontin 400 mg Hartkapseln	DE/H/0899/003	BE181133	VIATRIS HEALTHCARE	BE
Neurontin 400 mg Hartkapseln	DE/H/0899/003	2009030268	VIATRIS HEALTHCARE	LU
Neurontin 400 mg hörð hylki	DE/H/0899/003	920270	UPJOHN EESV	IS
Neurontin 400 mg kemény kapszula	DE/H/0899/003	OGYI-T-4966/05	VIATRIS HEALTHCARE LTD	HU
Neurontin 400 mg kemény kapszula	DE/H/0899/003	OGYI-T-4966/06	VIATRIS HEALTHCARE LTD	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
Neurontin 400 mg kovat kapselit	DE/H/0899/003	11663	UPJOHN EESV	FI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/025	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/027	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/023	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/026	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/029	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/028	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/030	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/033	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/032	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/031	UPJOHN EESV	SI
Neurontin 400 mg trde kapsule	DE/H/0899/003	H/99/01105/024	UPJOHN EESV	SI
Neurontin 400 mg tvrde kapsule	not available	HR-H-227912224	UPJOHN EESV	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
NEURONTIN 400 mg tvrdé tobolky	DE/H/0899/003	21/463/97-C	UPJOHN EESV	CZ
Neurontin 400 mg καψάκια σκληρά	DE/H/0899/003	17446	VIATRIS HELLAS LTD.	CY
Neurontin 400 mg καψάκια σκληρά	DE/H/0899/003	26442/9-4-2009	VIATRIS HELLAS LTD.	GR
NEURONTIN 400 mg, gélule	DE/H/0899/003	34009 337 904 6 2	VIATRIS UP	FR
NEURONTIN 400 mg, gélule	DE/H/0899/003	34009 338 014 4 1	VIATRIS UP	FR
NEURONTIN 400, 400 mg, kapsułki twarde	DE/H/0899/003	7694	UPJOHN EESV	PL
Neurontin 400, harde capsules 400 mg	DE/H/0899/003	RVG 22483	VIATRIS HEALTHCARE LTD	NL
NEURONTIN 600 mg apvalkotas tabletes	DE/H/0899/004	03-0016	UPJOHN EESV	LV
Neurontin 600 mg comprimés pelliculés	DE/H/0899/004	BE369013	VIATRIS HEALTHCARE	BE
Neurontin 600 mg comprimés pelliculés	DE/H/0899/004	2009030269	VIATRIS HEALTHCARE	LU
Neurontin 600 mg comprimidos recubiertos con película	DE/H/0899/004	63.150	VIATRIS HEALTHCARE LIMITED	ES
Neurontin 600 mg comprimidos revestidos por película	DE/H/0899/004	5815881	VIATRIS HEALTHCARE 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
Neurontin 600 mg comprimidos revestidos por película	DE/H/0899/004	2960680	VIATRIS HEALTHCARE LDA.	PT
Neurontin 600 mg film-coated tablets	DE/H/0899/004	PA 23055/004/004	UPJOHN EESV	IE
Neurontin 600 mg film-coated tablets	DE/H/0899/004	PL 50622/0049	UPJOHN UK LIMITED	XI
Neurontin 600 mg filmdragerade tabletter	DE/H/0899/004	14287	UPJOHN EESV	FI
Neurontin 600 mg filmdragerade tabletter	DE/H/0899/004	15369	UPJOHN EESV	SE
Neurontin 600 mg filmdrasjerte tabletter	DE/H/0899/004	99-854	UPJOHN EESV	NO
Neurontin 600 mg filmomhulde tabletten	DE/H/0899/004	BE369013	VIATRIS HEALTHCARE	BE
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/034	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/036	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/049	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/043	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/047	UPJOHN EESV	SI
Neurontin 600 mg	DE/H/0899/004	H/99/01105/053	UPJOHN EESV	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
filmsko obložene tablete				
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/044	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/037	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/038	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/035	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/042	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/040	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/046	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/045	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/051	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/052	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/041	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/050	UPJOHN EESV	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
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/039	UPJOHN EESV	SI
Neurontin 600 mg filmsko obložene tablete	DE/H/0899/004	H/99/01105/048	UPJOHN EESV	SI
Neurontin 600 mg Filmtabletten	DE/H/0899/004	BE369013	VIATRIS HEALTHCARE	BE
Neurontin 600 mg Filmtabletten	DE/H/0899/004	2009030269	VIATRIS HEALTHCARE	LU
Neurontin 600 mg filmuhúðaðar töflur	DE/H/0899/004	990081	UPJOHN EESV	IS
Neurontin 600 mg kalvopäällysteiset tabletit	DE/H/0899/004	14287	UPJOHN EESV	FI
NEURONTIN 600 mg potahované tablety	DE/H/0899/004	21/352/03-C	UPJOHN EESV	CZ
Neurontin 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	DE/H/0899/004	26443/9-4-2009	VIATRIS HELLAS LTD.	GR
NEURONTIN 600 mg, comprimé pelliculé	DE/H/0899/004	34009 359 002 5 8	VIATRIS UP	FR
NEURONTIN 600 mg, comprimé pelliculé	DE/H/0899/004	34009 347 591 0 9	VIATRIS UP	FR
NEURONTIN 600 mg, comprimé pelliculé	DE/H/0899/004	34009 359 001 9 7	VIATRIS UP	FR
NEURONTIN 600 mg, comprimé pelliculé	DE/H/0899/004	34009 347 340 8 3	VIATRIS UP	FR

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NEURONTIN 600 mg, comprimé pelliculé	DE/H/0899/004	34009 300 535 5 3	VIATRIS UP	FR
NEURONTIN 600, 600 mg, tabletki powlekane	DE/H/0899/004	10174	UPJOHN EESV	PL
Neurontin 600, filmomhulde tabletten 600 mg	DE/H/0899/004	RVG 25247	VIATRIS HEALTHCARE LTD	NL
NEURONTIN 800 mg apvalkotās tabletes	DE/H/0899/005	03-0017	UPJOHN EESV	LV
Neurontin 800 mg comprimés pelliculés	DE/H/0899/005	BE369022	VIATRIS HEALTHCARE	BE
Neurontin 800 mg comprimés pelliculés	DE/H/0899/005	2009030270	VIATRIS HEALTHCARE	LU
Neurontin 800 mg comprimidos recubiertos con película	DE/H/0899/005	63.149	VIATRIS HEALTHCARE LIMITED	ES
Neurontin 800 mg comprimidos revestidos por película	DE/H/0899/005	2960789	VIATRIS HEALTHCARE LDA.	PT
Neurontin 800 mg comprimidos revestidos por película	DE/H/0899/005	5815980	VIATRIS HEALTHCARE LDA.	PT
Neurontin 800 mg film-coated tablets	DE/H/0899/005	PA 23055/004/005	UPJOHN EESV	IE
Neurontin 800 mg film-coated tablets	DE/H/0899/005	PL 50622/0050	UPJOHN UK LIMITED	XI
Neurontin 800 mg	DE/H/0899/005	14288	UPJOHN EESV	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
filmdragerade tabletter				
Neurontin 800 mg filmdragerade tabletter	DE/H/0899/005	15370	UPJOHN EESV	SE
Neurontin 800 mg filmdrasjerte tabletter	DE/H/0899/005	99-855	UPJOHN EESV	NO
Neurontin 800 mg filmomhulde tabletten	DE/H/0899/005	BE369022	VIATRIS HEALTHCARE	BE
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/071	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/073	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/072	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/054	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/058	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/069	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/068	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/067	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/062	UPJOHN EESV	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
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/057	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/059	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/070	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/066	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/063	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/061	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/064	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/065	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/055	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/060	UPJOHN EESV	SI
Neurontin 800 mg filmsko obložene tablete	DE/H/0899/005	H/99/01105/056	UPJOHN EESV	SI
Neurontin 800 mg Filmtabletten	DE/H/0899/005	BE369022	VIATRIS HEALTHCARE	BE
Neurontin 800 mg Filmtabletten	DE/H/0899/005	2009030270	VIATRIS HEALTHCARE	LU

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Neurontin 800 mg filmuhúðaðar töflur	DE/H/0899/005	990082	UPJOHN EESV	IS
Neurontin 800 mg kalvopäällysteiset tabletit	DE/H/0899/005	14288	UPJOHN EESV	FI
NEURONTIN 800 mg potahované tablety	DE/H/0899/005	21/353/03-C	UPJOHN EESV	CZ
NEURONTIN 800 mg, comprimé pelliculé	DE/H/0899/005	34009 359 004 8 7	VIATRIS UP	FR
NEURONTIN 800 mg, comprimé pelliculé	DE/H/0899/005	34009 359 003 1 9	VIATRIS UP	FR
NEURONTIN 800 mg, comprimé pelliculé	DE/H/0899/005	34009 300 535 6 0	VIATRIS UP	FR
NEURONTIN 800 mg, comprimé pelliculé	DE/H/0899/005	34009 347 592 7 7	VIATRIS UP	FR
NEURONTIN 800 mg, comprimé pelliculé	DE/H/0899/005	34009 347 342 0 5	VIATRIS UP	FR
NEURONTIN 800, 800 mg, tabletki powlekane	DE/H/0899/005	10175	UPJOHN EESV	PL
Neurontin 800, filmomhulde tabletten 800 mg	DE/H/0899/005	RVG 25248	VIATRIS HEALTHCARE LTD	NL
Neurontin, 300 mg kõvakapsel	DE/H/0899/002	212298	UPJOHN EESV	EE
Neurontin, filmovertrukne tabletter	DE/H/0899/004	32601	UPJOHN EESV	DK

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Neurontin, filmovertrukne tabletter	DE/H/0899/005	32602	UPJOHN EESV	DK
Neurontin, hårde kapsler	DE/H/0899/002	32061	UPJOHN EESV	DK
Neurontin, hårde kapsler	DE/H/0899/003	32062	UPJOHN EESV	DK
Neurontin® 100 mg Hartkapseln	DE/H/0899/001	47275.00.00	VIATRIS PHARMA GMBH	DE
Neurontin® 300 mg - Hartkapseln	DE/H/0899/002	1-20679	UPJOHN EESV	AT
Neurontin® 300 mg Hartkapseln	DE/H/0899/002	47275.01.00	VIATRIS PHARMA GMBH	DE
Neurontin® 400 mg - Hartkapseln	DE/H/0899/003	1-20678	UPJOHN EESV	AT
Neurontin® 400 mg Hartkapseln	DE/H/0899/003	47275.02.00	VIATRIS PHARMA GMBH	DE
Neurontin® 600 mg - Filmtabletten	DE/H/0899/004	1-22737	UPJOHN EESV	AT
Neurontin® 600 mg Filmtabletten	DE/H/0899/004	43728.00.00	VIATRIS PHARMA GMBH	DE
Neurontin® 800 mg - Filmtabletten	DE/H/0899/005	1-22738	UPJOHN EESV	AT
Neurontin® 800 mg Filmtabletten	DE/H/0899/005	43728.01.00	VIATRIS PHARMA GMBH	DE
Неуронтин 300 mg твърди капсули	not available	20000710	UPJOHN EESV	BG
Неуронтин 400 mg	not available	20000711	UPJOHN EESV	BG

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твърди капсули				