26 October 2017
EMA/716726/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): naratriptan

Procedure No.: PSUSA/00002126/201702

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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
FORMIGRAN® 2,5 mg Filmtabletten	not available	41308.00.00	GLAXOSMITHKLINE CONSUMER HEALTHCARE GMBH & CO. KG	DE
Naragran	not available	18644	GLAXOSMITHKLINE PHARMA A/S	DK
NARAMIG 2,5 mg comprimés pelliculés	SE/H/0131/001	BE185053	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
NARAMIG 2,5 mg comprimés pelliculés	SE/H/0131/001	260/09/02/0196	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	5803689	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	2576080	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	2576189	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	2711588	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	2711687	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimido revestido por película	SE/H/0131/001	2575983	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
Naramig 2,5 mg comprimidos recubiertos con película	SE/H/0131/001	61.828	GLAXO S.A.	ES
Naramig 2,5 mg filmdragerade tabletter	SE/H/0131/001	12783	GLAXOSMITHKLINE OY	FI
NARAMIG 2,5 mg filmomhulde tabletten	SE/H/0131/001	BE185053	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
NARAMIG 2,5 mg filmomhulde tabletten	SE/H/0131/001	260/09/02/0196	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Naramig 2,5 mg filmsko obložene tablete	not available	H/98/01081/001	GLAXOSMITHKLINE D.O.O.	SI
Naramig 2,5 mg filmsko obložene tablete	not available	H/98/01081/002	GLAXOSMITHKLINE D.O.O.	SI
Naramig 2,5 mg Filmtabletten	SE/H/0131/001	BE185053	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Naramig 2,5 mg Filmtabletten	SE/H/0131/001	260/09/02/0196	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Naramig 2,5 mg kalvopäällysteinen tabletti	SE/H/0131/001	12783	GLAXOSMITHKLINE OY	FI
NARAMIG 2,5 mg plėvele	not available	LT/1/98/1508/001	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
dengtos tabletės				
Naramig 2,5 mg potahované tablety	not available	33/531/99-C	GLAXO GROUP LIMITED	CZ
Naramig 2,5 mg tablett, filmdrasjert	not available	96-3619	GLAXOSMITHKLINE AS	NO
NARAMIG 2,5 mg, comprime pellicule	SE/H/0131/001	NL22647	LABORATOIRE GLAXOSMITHKLINE	FR
Naramig 2,5mg επικαλυμμένα με λεπτό υμένιο δισκίο.	SE/H/0131/001	2355601	GLAXOSMITHKLINE AEBE	GR
Naramig 2,5mg, filmdragerad tablett	SE/H/0131/001	13382	GLAXOSMITHKLINE AB	SE
Naramig 2.5 mg film-coated tablets	not available	PL 10949/0273	GLAXO WELLCOME UK LTD TRADING AS GLAXOSMITHKLINE UK	UK
Naramig, 2,5 mg, tabletki powlekane	not available	10614	GLAXOSMITHKLINE EXPORT LTD	PL
Naramig, filmomhulde tabletten 2,5 mg.	SE/H/0131/001	RVG 21444	GLAXOSMITHKLINE BV	NL
Naramig® 2,5 mg Filmtabletten	SE/H/0131/001	40706.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE