22 May 2025
EMA/178301/2025
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): sumatriptan, naproxen / sumatriptan

Procedure No. PSUSA/00002832/202409

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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
Frimig Duo 85 mg/500 mg filmom obalené tablety	FI/H/1108/001	33/0079/24-S	ORION CORPORATION	SK
Frimig Duo 85 mg/500 mg potahované tablety	FI/H/1108/001	33/348/23-C	ORION CORPORATION	CZ
Frimig Duo, 85 mg + 500 mg, tabletki powlekane	FI/H/1108/001	27538	ORION CORPORATION	PL
Imigran 10 mg Nasal Spray, Solution	not available	PA 1077/008/003	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Imigran 10 mg nässpray	NL/H/0115/001	12426	GLAXOSMITHKLINE OY	FI
Imigran 10 mg nässpray, lösning	NL/H/0115/001	13219	GLAXOSMITHKLINE AB	SE
Imigran 10 mg nenäsumute	NL/H/0115/001	12426	GLAXOSMITHKLINE OY	FI
Imigran 10 mg solución para pulverización nasal.	NL/H/0115/001	61.629	GLAXOSMITHKLINE, S.A.	ES
Imigran 10 mg Spray Nasale	NL/H/0115/001	027975123	GLAXOSMITHKLINE S.P.A.	IT
IMIGRAN 10 mg/0,1 ml pršilo za nos, raztopina	not available	H/93/00766/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Imigran 10 Neusspray	NL/H/0115/001	RVG 19469	GLAXOSMITHKLINE B.V.	NL
IMIGRAN 100 mg Compresse rivestite con film	not available	027975059	GLAXOSMITHKLINE S.P.A.	IT
Imigran 100 mg Filmtabletten	not available	25621.00.01	GLAXOSMITHKLINE GMBH & CO. KG	DE
Imigran 100 mg plėvele dengtos tabletės	not available	LT/1/94/0344/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Imigran 100 mg tablett	not available	7896	GLAXOSMITHKLINE AS	NO
Imigran 100 mg tabletta	not available	OGYI-T-5410/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Imigran 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	2026501	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Imigran 12 mg/ml injekčný roztok	not available	33/0266/93-S	GLAXOSMITHKLINE (IRELAND) LIMITED	SK

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
Imigran 12 mg/ml injeksjonsvæske, oppløsning	not available	7895	GLAXOSMITHKLINE AS	NO
Imigran 12 mg/ml injektioneste, liuos	not available	10737	GLAXOSMITHKLINE (IRELAND) LIMITED	FI
Imigran 12 mg/ml injektionsvätska, lösning	not available	10737	GLAXOSMITHKLINE (IRELAND) LIMITED	FI
Imigran 12 mg/ml injektionsvätska, lösning	not available	11435	GLAXOSMITHKLINE AB	SE
Imigran 12 mg/ml stungulyf, lausn.	not available	900125	GLAXOSMITHKLINE PHARMA A/S	IS
Imigran 20 mg Nasal Spray, Solution	not available	PA 1077/008/004	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Imigran 20 mg näs„spray	NL/H/0115/002	12427	GLAXOSMITHKLINE OY	FI
Imigran 20 mg nässpray, lösning	NL/H/0115/002	13220	GLAXOSMITHKLINE AB	SE
Imigran 20 mg nenäsumute	NL/H/0115/002	12427	GLAXOSMITHKLINE OY	FI
Imigran 20 mg solución para pulverización nasal.	NL/H/0115/002	61.630	GLAXOSMITHKLINE, S.A.	ES
Imigran 20 mg Spray Nasale	NL/H/0115/002	027975135	GLAXOSMITHKLINE S.P.A.	IT
Imigran 20 mg Spray Nasale	NL/H/0115/002	027975147	GLAXOSMITHKLINE S.P.A.	IT
Imigran 20 mg nosová aerodisperzia	not available	33/0244/99-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
IMIGRAN 20 mg/0,1 ml pršilo za nos, raztopina	not available	H/93/00766/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Imigran 20 mg/dose, nesespray	not available	95-2920	GLAXOSMITHKLINE AS	NO
Imigran 20 mg/skammt nefúði, lausn.	not available	980420	GLAXOSMITHKLINE PHARMA A/S	IS
Imigran 20 Neusspray	NL/H/0115/002	RVG 19470	GLAXOSMITHKLINE B.V.	NL
IMIGRAN 50 mg Compresse rivestite con film	not available	027975073	GLAXOSMITHKLINE S.P.A.	IT
Imigran 50 mg film coated tablets	not available	MA 192/04001	GLAXOSMITHKLINE (IRELAND) 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
Imigran 50 mg Filmtabletten	not available	31483.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Imigran 50 mg plėvele dengtos tabletės	not available	LT/1/94/0344/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
IMIGRAN 50 mg tablete	not available	HR-H-584958851	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Imigran 50 mg tablett	not available	7991	GLAXOSMITHKLINE AS	NO
Imigran 50 mg tabletta	not available	OGYI-T-5410/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Imigran 50 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	2026504	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Imigran 6 mg injekční roztok v zásobní vložce	not available	33/701/93-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Imigran 6 mg solución inyectable	not available	59.973	GLAXOSMITHKLINE S.A.	ES
Imigran 6 mg/0,5 ml - Spritzampullen	not available	1-19773	GLAXOSMITHKLINE PHARMA GMBH.	AT
IMIGRAN 6 mg/0,5 ml raztopina za injiciranje	not available	H/93/00766/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Imigran 6 mg/0,5 ml solução injetável	not available	8788604	GLAXO WELLCOME FARMACÊUTICA, LDA	PT
IMIGRAN 6 mg/0,5 ml Soluzione iniettabile per uso sottocutaneo	not available	027975061	GLAXOSMITHKLINE S.P.A.	IT
Imigran 6 mg/0,5 mL ενέσιμο διάλυμα	not available	2026502	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Imigran 6 s.c. oplossing voor injectie voor subcutaan gebruik.	not available	RVG 15009	GLAXOSMITHKLINE B.V.	NL
IMIGRAN DR 100 mg comprimate dispersabile	not available	11592/2019/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 100 mg comprimate dispersabile	not available	11592/2019/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 100 mg comprimate dispersabile	not available	11592/2019/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 100 mg comprimate dispersabile	not available	11592/2019/03	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
IMIGRAN DR 100 mg comprimate dispersabile	not available	11592/2019/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 50 mg comprimate dispersabile	not available	11591/2019/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 50 mg comprimate dispersabile	not available	11591/2019/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 50 mg comprimate dispersabile	not available	11591/2019/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR 50 mg comprimate dispersabile	not available	11591/2019/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
IMIGRAN DR50 mg comprimate dispersabile	not available	11591/2019/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Imigran FDT, 100 mg õhukese polümeerikattega tabletid	not available	437604	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Imigran FDT, 50 mg õhukese polümeerikattega tabletid	not available	437504	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Imigran film-coated tablets 50 mg	not available	19508	GLAXOSMITHKLINE (IRELAND) LIMITED	CY
Imigran Ftab 100 mg Film-coated Tablets	not available	PA 1077/008/007	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Imigran FTAB 100, dispergeerbare tabletten 100 mg	not available	RVG 29414	GLAXOSMITHKLINE B.V.	NL
Imigran Ftab 50 mg Film-coated Tablets	not available	PA 1077/008/006	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Imigran FTAB 50, dispergeerbare tabletten 50 mg	not available	RVG 29413	GLAXOSMITHKLINE B.V.	NL
Imigran Juvenil 10 mg/dose, nesespray	not available	95-2919	GLAXOSMITHKLINE AS	NO
Imigran Juvenil 10 mg/skammt nefúði, lausn.	not available	IS/1/05/102/01	GLAXOSMITHKLINE PHARMA A/S	IS
Imigran Nasal 20 mg 20 mg/0,1 ml Lösung	NL/H/115/002	31483.01.01	GLAXOSMITHKLINE GMBH & CO. KG	DE
Imigran Nasal mite 10 mg 10 mg/0,1 ml Lösung	NL/H/0115/001	31483.00.01	GLAXOSMITHKLINE GMBH & CO. KG	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
Imigran Neo 50 mg comprimidos recubiertos con película.	not available	66.406	GLAXOSMITHKLINE, S.A.	ES
Imigran Novum 50 mg filmdragerad tablett	not available	19492	GLAXOSMITHKLINE AB	SE
Imigran oldatos injekció + autoinjektor	not available	OGYI-T-5410/05	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Imigran Radis 100 mg filmdragerad tablett	not available	18198	GLAXOSMITHKLINE OY	FI
Imigran Radis 100 mg kalvopäällysteinen tabletti	not available	18198	GLAXOSMITHKLINE OY	FI
Imigran Radis 100 mg tablett	not available	03-1916	GLAXOSMITHKLINE AS	NO
Imigran Radis 100 mg töflur	not available	IS/1/04/190/02	GLAXOSMITHKLINE PHARMA A/S	IS
Imigran Radis 50 mg filmdragerad tablett	not available	18197	GLAXOSMITHKLINE OY	FI
Imigran Radis 50 mg kalvopäällysteinen tabletti	not available	18197	GLAXOSMITHKLINE OY	FI
Imigran Radis 50 mg tablett	not available	03-1915	GLAXOSMITHKLINE AS	NO
Imigran Radis 50 mg töflur	not available	IS/1/04/190/01	GLAXOSMITHKLINE PHARMA A/S	IS
IMIGRAN SPRINT 100 mg disperzibilne tablete	not available	H/93/00766/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
IMIGRAN SPRINT 100 mg disperzibilne tablete	not available	H/93/00766/005	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Imigran Sprint 100 mg filmtabletta	not available	OGYI-T-10096/08	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
IMIGRAN SPRINT 50 mg disperzibilne tablete	not available	H/93/00766/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
IMIGRAN SPRINT 50 mg disperzibilne tablete	not available	H/93/00766/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
IMIGRAN SPRINT 50 mg disperzibilne tablete	not available	H/93/00766/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Imigran Sprint 50 mg filmtabletta	not available	OGYI-T-10096/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Imigran Sprint, dispergible	not available	34998	GLAXOSMITHKLINE PHARMA	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
tabletter			A/S	
Imigran Sprint, dispergible tabletter	not available	34997	GLAXOSMITHKLINE PHARMA A/S	DK
Imigran T 100 mg plevele dengtos tabletes	not available	LT/1/06/0531/006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Imigran T 100 mg plevele dengtos tabletes	not available	LT/1/06/0531/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Imigran T 50 mg plevele dengtos tabletes	not available	LT/1/06/0531/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Imigran T 50 mg plevele dengtos tabletes	not available	LT/1/06/0531/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Imigran, 100 mg, tabletki powlekane	not available	R/1516	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Imigran, 20 mg/0,1 ml, aerozol do nosa, roztwór	not available	4916	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Imigran, 50 mg, tabletki powlekane	not available	7381	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Imigran, injektionsvæske, opløsning	not available	14113	GLAXOSMITHKLINE PHARMA A/S	DK
Imigran, næsespray, opløsning	NL/H/0115/001	18362	GLAXOSMITHKLINE PHARMA A/S	DK
Imigran, næsespray, opløsning	NL/H/0115/002	18363	GLAXOSMITHKLINE PHARMA A/S	DK
IMIGRANE 10 mg/0,1 ml, solution pour pulvérisation nasale	not available	6 715 333 4	LABORATOIRE GLAXOSMITHKLINE	FR
IMIGRANE 20 mg/0,1 ml, solution pour pulvérisation nasale	not available	6 787 632 8	LABORATOIRE GLAXOSMITHKLINE	FR
IMIGRANE 50 mg, comprimé pelliculé	not available	6 467 603 0	LABORATOIRE GLAXOSMITHKLINE	FR
IMIGRANE 6 mg/0,5 ml, solution injectable pour voie sous-cutanée en seringue pré-remplie	not available	6 948 532 8	LABORATOIRE GLAXOSMITHKLINE	FR
Imigran-Inject 6 mg/0,5 ml Injektionslösung	not available	25623.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Imigran-T 100 mg	not available	25623.00.01	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
Filmtabletten			CO. KG	
Imigran-T 50 mg Filmtabletten	not available	31485.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
IMIJECT 6 mg/0,5 ml, solution injectable pour voie sous-cutanée en seringue pré-remplie	not available	6 6276750	LABORATOIRE GLAXOSMITHKLINE	FR
Imitrex 10 mg Neusspray	NL/H/0115/001	BE 182716	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 10 mg Solution pour pulvérisation nasale.	NL/H/0115/001	BE 182716	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 10 mg Solution pour pulvérisation nasale.	NL/H/0115/001	2007119551	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex 10mg Nasenspray	NL/H/0115/001	BE 182716	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 10mg Nasenspray	NL/H/0115/001	2007119551	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex 20 mg Neusspray	NL/H/0115/002	BE182707	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 20 mg Solution pour pulvérisation nasale.	NL/H/0115/002	BE182707	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 20 mg Solution pour pulvérisation nasale.	NL/H/0115/002	2007119552	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex 20mg Nasenspray	NL/H/0115/002	BE182707	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex 20mg Nasenspray	NL/H/0115/002	2007119552	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex Instant 100 mg	not available	BE279474	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
comprimés dispersibles			PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	
Imitrex Instant 100 mg comprimés dispersibles	not available	2006030005	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex Instant 100 mg dispergeerbare tabletten	not available	BE279474	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex Instant 100 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	BE279474	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex Instant 100 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	2006030005	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex Instant 50 mg comprimés dispersibles	not available	BE279465	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex Instant 50 mg comprimés dispersibles	not available	2006030004	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex Instant 50 mg dispergeerbare tabletten	not available	BE279465	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex Instant 50 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	BE279465	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex Instant 50 mg Tabletten zur Herstellung einer Suspension zum Einnehmen	not available	2006030004	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Imitrex SC 6 mg Injektionslösung	not available	BE157473	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Imitrex SC 6 mg Injektionslösung	not available	2001116611	GLAXOSMITHKLINE PHARMACEUTICALS EN	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
			ABREGE GLAXOSMITHKLINE	
IMITREX SC 6 mg oplossing voor injectie	not available	BE157473	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
IMITREX SC 6 mg solution injectable	not available	BE157473	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
IMITREX SC 6 mg solution injectable	not available	2001116611	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
MigraKind 50 mg Film-Coated Tablets	not available	PL 08553/0693	DR. REDDY'S LABORATORIES (UK) LTD	XI
Migsun 85 mg/500 mg apvalkotās tabletes	FI/H/1108/001	22-0119	ORION CORPORATION	LV
Migsun 85 mg/500 mg film-coated tablets	FI/H/1108/001	PA1327/022/001	ORION CORPORATION	IE
Migsun 85 mg/500 mg õhukese polümeerikattega tabletid	FI/H/1108/001	1066022	ORION CORPORATION	EE
Migsun 85 mg/500 mg plėvele dengtos tabletės	FI/H/1108/001	LT/1/22/4977/001	ORION CORPORATION	LT
Migsun 85 mg/500 mg plėvele dengtos tabletės	FI/H/1108/001	LT/1/22/4977/002	ORION CORPORATION	LT
NOMANESIT 85 mg/500 mg comprimé pelliculé	FI/H/1108/001	34009 302 924 3 3	ORION CORPORATION	FR
NOMANESIT 85 mg/500 mg comprimé pelliculé	FI/H/1108/001	34009 302 924 4 0	ORION CORPORATION	FR
Nomigrin 85 mg/457 mg filmdragerade tabletter	FI/H/1108/001	61618	ORION CORPORATION	SE
Nomigrin 85 mg/500 mg filmdragerade tabletter	FI/H/1108/001	38713	ORION CORPORATION	FI
Nomigrin 85 mg/500 mg filmdrasjerte tabletter	FI/H/1108/001	21-13833	ORION CORPORATION	NO
Nomigrin 85 mg/500 mg filmtabletta	FI/H/1108/001	OGYI-T-24078/01	ORION CORPORATION	HU
Nomigrin 85 mg/500 mg kalvopäällysteiset tabletit	FI/H/1108/001	38713	ORION CORPORATION	FI
Nomigrin, filmovertrukne	FI/H/1108/001	65618	ORION CORPORATION	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
tabletter				
Rosemig 100 mg potahované tablety	not available	33/026/04-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Rosemig 20 mg nosní sprej, roztok	not available	33/149/99-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Rosemig 50 mg potahované tablety	not available	33/025/04-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Rosemig Sprintab 100 mg dispergovatelné tablety	not available	33/196/04-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Rosemig Sprintab 50 mg dispergovatelné tablety	not available	33/195/04-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Sumanet 85 mg/500 mg compresse rivestite con film	FI/H/1108/001	050972013	ORION CORPORATION	IT
Sumanet 85 mg/500 mg compresse rivestite con film	FI/H/1108/001	050972025	ORION CORPORATION	IT
Sumanet 85 mg/500 mg Filmtabletten	FI/H/1108/001	142090	ORION CORPORATION	AT
Sumatriptan 100 mg film-coated tablets.	not available	PL 36687/0366	TORRENT PHARMA (UK) LTD.	XI
Sumatriptan 100 mg tablets	NL/H/0629/002	PL 04416/1045	SANDOZ LTD	XI
Sumatriptan 3 mg/0.5 ml solution for injection in pre-filled pen	NL/H/4660/001	PL 31750/0161	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	XI
Sumatriptan 50 mg film-coated tablets	not available	PL 36687/0365	TORRENT PHARMA (UK) LTD.	XI
Sumatriptan 50 mg tablets	NL/H/0629/001	PL 04416/1044	SANDOZ LTD	XI
Sumatriptan Aurobindo 100 mg tablets	MT/H/0336/002	MA807/10402	AUROBINDO PHARMA (MALTA) LIMITED	MT
Sumatriptan Hormosan 3 mg Injektionslösung im Fertigpen	DE/H/6819/001/DC	7001043.00.00	HORMOSAN PHARMA GMBH	DE
Sumatriptan SUN 3 mg/0,5 ml oplossing voor injectie in voorgevulde pen	NL/H/4660/001	RVG 124476	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Sumatriptán SUN 3 mg/0,5	NL/H/4660/001	85221	SUN PHARMACEUTICAL	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
ml solución inyectable en pluma precargada			INDUSTRIES EUROPE B.V.	
Sumatriptan SUN 3 mg/0,5 ml soluție injectabilă în stilou injector preumplut	NL/H/4660/001	13116/2020/01	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	RO
Sumatriptan SUN 3 mg/0,5 ml soluție injectabilă în stilou injector preumplut	NL/H/4660/001	13116/2020/03	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	RO
Sumatriptan SUN 3 mg/0,5 ml soluție injectabilă în stilou injector preumplut	NL/H/4660/001	13116/2020/02	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	RO
SUMATRIPTAN SUN 3 mg/0,5 ml, solution injectable en seringue préremplie (voie SC)	NL/H/4660/001	34009 302 034 9 1	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
SUMATRIPTAN SUN 3 mg/0,5 ml, solution injectable en seringue préremplie (voie SC)	NL/H/4660/001	34009 302 035 2 1	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
SUMATRIPTAN SUN 3 mg/0,5 ml, solution injectable en seringue préremplie (voie SC)	NL/H/4660/001	34009 302 035 0 7	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
Sumatriptan SUN 6 mg/ml injeksjonsvæske, oppløsning i ferdigfylt penn	NL/H/4660/001	19-12691	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NO
Sumatriptan SUN 6 mg/ml injektionsvätska, lösning i förfylld injektionspenna	NL/H/4660/001	58741	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	SE
Sumatriptan SUN Pharma 3 mg/0,5 ml soluzione iniettabile in penna pre-riempita	NL/H/4660/001	047538020	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Sumatriptan SUN Pharma 3 mg/0,5 ml soluzione iniettabile in penna pre-riempita	NL/H/4660/001	047538032	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Sumatriptan SUN Pharma 3	NL/H/4660/001	047538018	SUN PHARMACEUTICAL	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
mg/0,5 ml soluzione iniettabile in penna pre-riempita			INDUSTRIES EUROPE B.V.	
Sumatriptan SUN, 3 mg/0,5 ml, roztwór do wstrzykiwań we wstrzykiwaczu	NL/H/4660/001	25967	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	PL
Suvexx 85 mg + 500 mg comprimidos revestidos por película	FI/H/1108/001	5872205	ORION CORPORATION	PT
Suvexx 85 mg/500 mg filmomhulde tabletten	FI/H/1108/001	RVG 132188	ORION CORPORATION	NL
Suvexx 85 mg/500 mg επικαλυμμένα με λεπτό υμένιο δισκία	FI/H/1108/001	80278	ORION CORPORATION	GR
Имигран 50 mg филмирани таблетки	not available	9600076	GLAXOSMITHKLINE TRADING SERVICES LIMITED	BG