EMA/203553/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): octreotide

Procedure No.: PSUSA/00002201/202006

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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
SANDOSTATIN LAR 10 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/005	6869/2014/01	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN LAR 10 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/005	6869/2014/02	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN LAR 20 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/006	6870/2014/01	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN LAR 20 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/006	6870/2014/02	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN LAR 30 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/007	6871/2014/01	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN LAR 30 mg pulbere şi solvent pentru suspensie injectabilă	DE/H/5095/007	6871/2014/02	NOVARTIS PHARMA GMBH	RO
SANDOSTATIN 100 microgramе/1 ml soluţie injectabilă/perfuzabilă	DE/H/5095/002	4756/2012/01	NOVARTIS PHARMA GMBH	RO
Sandostatin 100 mikrog/ml, injektions-/infusionsvätska, lösning	DE/H/5095/002	10165	NOVARTIS FINLAND OY	FI
Sandostatin 500 mikrog/ml, injektions-/infusionsvätska, lösning	DE/H/5095/003	10166	NOVARTIS FINLAND OY	FI
Sandostatin 100 mikrog/ml, injektio-/infuusioneste, liuos	DE/H/5095/002	10165	NOVARTIS FINLAND OY	FI
Sandostatin 50 mikrog/ml, injektio-/infuusioneste, liuos	DE/H/5095/001	10164	NOVARTIS FINLAND OY	FI
Sandostatin 50 mikrog/ml, injektions-/infusionsvätska, lösning	DE/H/5095/001	10164	NOVARTIS FINLAND OY	FI
Sandostatin 500 mikrog/ml, injektio-/infuusioneste, liuos	DE/H/5095/003	10166	NOVARTIS FINLAND OY	FI
Sandostatine Long Acting Repeatable 10 mg, poudre et solvant pour suspension injectable	DE/H/5095/005	1998020008	NOVARTIS PHARMA N.V.	LU
Sandostatine Long Acting Repeatable 10 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/005	1998020008	NOVARTIS PHARMA N.V.	LU
Sandostatine Long Acting Repeatable 20 mg, poudre et solvant pour suspension injectable	DE/H/5095/006	1998020007	NOVARTIS PHARMA N.V.	LU
Sandostatine Long Acting Repeatable 20 mg, Pulver und Lösungsmittel zur	DE/H/5095/006	1998020007	NOVARTIS PHARMA N.V.	LU

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Herstellung einer Injektionssuspension				
Sandostatine Long Acting Repeatable 30 mg, poudre et solvant pour suspension injectable	DE/H/5095/007	1998020011	NOVARTIS PHARMA N.V.	LU
Sandostatine Long Acting Repeatable 30 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/007	1998020011	NOVARTIS PHARMA N.V.	LU
Sandostatine Long Acting Repeatable 10 mg poudre et solvant pour suspension injectable	DE/H/5095/005	BE191746	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 10 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/005	BE191746	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 20 mg poudre et solvant pour suspension injectable	DE/H/5095/006	BE191737	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 20 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/006	BE191737	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 30 mg poudre et solvant pour suspension injectable	DE/H/5095/007	BE191685	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 30 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/007	BE191685	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 10 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/005	BE191746	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 20 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/006	BE191737	NOVARTIS PHARMA N.V.	BE
Sandostatine Long Acting Repeatable 30 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/007	BE191685	NOVARTIS PHARMA N.V.	BE
САНДОСТАТИН LAR 20 mg прах и разтворител за инжекционна суспензия	DE/H/5095/006	20000366	NOVARTIS PHARMA GMBH	BG

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САНДОСТАТИН LAR 30 MG ПРАХ И РАЗТВОРИТЕЛ ЗА ИНЖЕКЦИОННА СУСПЕНЗИЯ	DE/H/5095/007	20000367	NOVARTIS PHARMA GMBH	BG
САНДОСТАТИН 100 микрограма/1 ml инжекционен/инфузионен разтвор	DE/H/5095/002	20000349	NOVARTIS PHARMA GMBH	BG
SANDOSTATIN LAR 20 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/006	18078	NOVARTIS IRELAND LIMITED	CY
SANDOSTATIN LAR 30 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/007	18079	NOVARTIS IRELAND LIMITED	CY
SANDOSTATIN LAR 10 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/005	18077	NOVARTIS IRELAND LIMITED	CY
SANDOSTATIN LAR 20 mg, prášek a rozpouštědlo pro injekční suspenzi	DE/H/5095/006	56/125/00-C	NOVARTIS S.R.O.,	CZ
SANDOSTATIN LAR 30 mg, prášek a rozpouštědlo pro injekční suspenzi	DE/H/5095/007	56/126/00-C	NOVARTIS S.R.O.,	CZ
Sandostatin LAR, 10 mg süstesuspensiooni pulber ja lahusti	DE/H/5095/005	283099	SIA "NOVARTIS BALTICS"	EE
Sandostatin LAR, 20 mg süstesuspensiooni pulber ja lahusti	DE/H/5095/006	283199	SIA "NOVARTIS BALTICS"	EE
Sandostatin LAR, 30 mg süstesuspensiooni pulber ja lahusti	DE/H/5095/007	283299	SIA "NOVARTIS BALTICS"	EE
SANDOSTATIN 0,05 mg/ml, injekční/infuzní roztok	DE/H/5095/004	56/183/90-A/C	NOVARTIS S.R.O.,	CZ
SANDOSTATIN 0,1 mg/ml, injekční/infuzní roztok	DE/H/5095/002	56/183/90-B/C	NOVARTIS S.R.O.,	CZ
SANDOSTATIN 0,5 mg/ml, injekční/infuzní roztok	DE/H/5095/003	56/183/90-D/C	NOVARTIS S.R.O.,	CZ
Sandostatin, 100 mikrogrammi/ml, süste- või infusioonilahus	DE/H/5095/002	217498	SIA "NOVARTIS BALTICS"	EE
Sandostatin LAR, pulver og solvens til injektionsvæske, suspension	DE/H/5095/005	17059	NOVARTIS HEALTHCARE A/S	DK
Sandostatin LAR, pulver og solvens til injektionsvæske, suspension	DE/H/5095/006	17060	NOVARTIS HEALTHCARE A/S	DK
Sandostatin LAR, pulver og solvens til injektionsvæske, suspension	DE/H/5095/007	17061	NOVARTIS HEALTHCARE A/S	DK
SANDOSTATIN LAR 10 mg por és oldószer	DE/H/5095/005	OGYI-T-1723/03	NOVARTIS HUNGÁRIA 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
szuszpenziós injekcióhoz			PHARMA	
Sandostatin LAR 10 mg pulveris un šķīdinātājs injekciju suspensijas pagatavošanai	DE/H/5095/005	98-0428	SIA "NOVARTIS BALTICS"	LV
SANDOSTATIN LAR 20 mg por és oldószer szuszpenziós injekcióhoz	DE/H/5095/006	OGYI-T-1723/04	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Sandostatin LAR 20 mg pulveris un šķīdinātājs injekciju suspensijas pagatavošanai	DE/H/5095/006	98-0429	SIA "NOVARTIS BALTICS"	LV
SANDOSTATIN LAR 30 mg por és oldószer szuszpenziós injekcióhoz	DE/H/5095/007	OGYI-T-1723/05	NOVARTIS HUNGÁRIA KFT. PHARMA	HU
Sandostatin LAR 30 mg pulveris un šķīdinātājs injekciju suspensijas pagatavošanai	DE/H/5095/007	98-0430	SIA "NOVARTIS BALTICS"	LV
Sandostatin 100 mikrogramm/1 ml oldatos injekció/infúzió	DE/H/5095/002	OGYI-T-1723/01	NOVARTIS HUNGÁRIA KFT.	HU
Sandostatin 100 mikrogrami/ml šķīdums injekcijām/infūzijām	DE/H/5095/002	94-0058	SIA "NOVARTIS BALTICS"	LV
SANDOSTATIN LAR 10 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/005	225670401	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN LAR 20 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/006	225670501	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN LAR 30 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/007	225670601	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN® ενέσιμο διάλυμα ή πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	DE/H/5095/002	197980101	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN® ενέσιμο διάλυμα ή πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	DE/H/5095/003	197980401	NOVARTIS (HELLAS) S.A.C.I.	GR
Sandostatin LAR 10 mg milteliai ir tirpiklis injekcinei suspensijai	DE/H/5095/005	LT/1/94/1033/005	SIA "NOVARTIS BALTICS"	LT
Sandostatin LAR 20 mg milteliai ir tirpiklis injekcinei suspensijai	DE/H/5095/006	LT/1/94/1033/007	SIA "NOVARTIS BALTICS"	LT
Sandostatin LAR 30 mg milteliai ir tirpiklis injekcinei suspensijai	DE/H/5095/007	LT/1/94/1033/009	SIA "NOVARTIS BALTICS"	LT
SANDOSTATIN® LAR® 10 MG POWDER	DE/H/5095/005	PL 00101/0511	NOVARTIS PHARMACEUTICALS UK	UK

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AND SOLVENT FOR SUSPENSION FOR INJECTION			LIMITED	
SANDOSTATIN® LAR® 20 MG POWDER AND SOLVENT FOR SUSPENSION FOR INJECTION	DE/H/5095/006	PL 00101/0512	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
SANDOSTATIN® LAR® 30 MG POWDER AND SOLVENT FOR SUSPENSION FOR INJECTION	DE/H/5095/007	PL 00101/0513	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Sandostatin® 100 microgram/1 ml, solution for injection/infusion	DE/H/5095/002	PL 00101/0213	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Sandostatin® 50 microgram/1 ml, solution for injection/infusion	DE/H/5095/001	PL 00101/0212	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Sandostatin® 500 microgram/1 ml, solution for injection/infusion	DE/H/5095/003	PL 00101/0214	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
SANDOSTATIN LAR 10 mg polvo para suspensión inyectable	DE/H/5095/005	62.139	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN LAR 20mg polvo para suspensión inyectable	DE/H/5095/006	62.140	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN LAR 30 mg polvo para suspensión inyectable	DE/H/5095/007	62.141	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN 100 microgramos/ml solución inyectable y para perfusión	DE/H/5095/002	59.559	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN 50 microgramos/ml solución inyectable y para perfusión	DE/H/5095/001	59.561	NOVARTIS FARMACÉUTICA S.A.	ES
Sandostatin® 50 µg, Injektionslösung/Infusionslösung	DE/H/5095/001	29423.00.00	NOVARTIS PHARMA GMBH	DE
Sandostatina LAR 10 mg pó e veículo para suspensão injetável	DE/H/5095/005	8705145	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Sandostatina LAR 20 mg pó e veículo para suspensão injetável	DE/H/5095/006	8705152	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Sandostatina LAR 30 mg pó e veículo para suspensão injetável	DE/H/5095/007	8705160	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
SANDOSTATIN 100 mikrogramów/1 ml, roztwór do wstrzykiwań / do infuzji	DE/H/5095/002	R/0429	NOVARTIS POLAND SP. Z O. O.	PL
SANDOSTATIN 50 mikrogramów/1 ml, roztwór do wstrzykiwań / do infuzji	DE/H/5095/001	R/0427	NOVARTIS POLAND SP. Z O. O.	PL

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SANDOSTATIN LAR 10 mg proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań	DE/H/5095/005	4597	NOVARTIS POLAND SP. Z O. O.	PL
SANDOSTATIN LAR 20 mg proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań	DE/H/5095/006	4596	NOVARTIS POLAND SP. Z O. O.	PL
SANDOSTATIN LAR 30 mg proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań	DE/H/5095/007	4595	NOVARTIS POLAND SP. Z O. O.	PL
SANDOSTATIN® LAR® 10 mg Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/005	1-22217	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® LAR® 20 mg Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/006	1-22215	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® LAR® 30 mg Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/007	1-22216	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® 50 Mikrogramm/ml Injektions-/Infusionslösung	DE/H/5095/001	1-19101	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® 100 Mikrogramm/ml Injektions-/Infusionslösung	DE/H/5095/002	1-19099	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® 1000 Mikrogram/ 5 ml Injektions-/Infusionslösung	DE/H/5095/004	1-19108	NOVARTIS PHARMA GMBH	AT
SANDOSTATIN® 500 Mikrogramm/ml Injektions-/Infusionslösung	DE/H/5095/003	1-19100	NOVARTIS PHARMA GMBH	AT
Sandostatine 50 microgram/1 ml, oplossing voor injectie/infusie	DE/H/5095/001	RVG 12612	NOVARTIS PHARMA B.V.	NL
Sandostatine 100 microgram/1 ml, oplossing voor injectie/infusie	DE/H/5095/002	RVG 12613	NOVARTIS PHARMA B.V.	NL
Sandostatine 500 microgram/1 ml, oplossing voor injectie/infusie	DE/H/5095/003	RVG 14997	NOVARTIS PHARMA B.V.	NL
Sandostatine LAR 10 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/005	RVG 18235	NOVARTIS PHARMA B.V.	NL
Sandostatine LAR 20 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/006	RVG 18236	NOVARTIS PHARMA B.V.	NL
Sandostatine LAR 30 mg, poeder en	DE/H/5095/007	RVG 18237	NOVARTIS PHARMA B.V.	NL

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oplosmiddel voor suspensie voor injectie				
SANDOSTATIN LAR 20 mg prášok a disperzné prostredie na injekčnú suspenziu	DE/H/5095/006	56/0276/00-S	NOVARTIS SLOVAKIA S.R.O.	SK
SANDOSTATIN LAR 30 mg prášok a disperzné prostredie na injekčnú suspenziu	DE/H/5095/007	56/0277/00-S	NOVARTIS SLOVAKIA S.R.O.	SK
Sandostatina LAR 10 mg polvere e solvente per sospensione iniettabile	DE/H/5095/005	027083082	NOVARTIS FARMA S.P.A.	IT
Sandostatina LAR 20 mg polvere e solvente per sospensione iniettabile	DE/H/5095/006	027083094	NOVARTIS FARMA S.P.A.	IT
Sandostatina LAR 30 mg polvere e solvente per sospensione iniettabile	DE/H/5095/007	027083106	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083017	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083029	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083031	NOVARTIS FARMA S.P.A.	IT
Sandostatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	DE/H/5095/004	027083043	NOVARTIS FARMA S.P.A.	IT
Sandostatine 100 Mikrogramm/1 ml, Injektionlösung/Infusionslösung	DE/H/5095/002	BE141382	NOVARTIS PHARMA N.V.	BE
Sandostatine 500 Mikrogramm/1 ml, Injektionlösung/Infusionslösung	DE/H/5095/003	BE156362	NOVARTIS PHARMA N.V.	BE
Sandostatine 100 microgrammes/1 ml, solution injectable/pour perfusion	DE/H/5095/002	2000116136	NOVARTIS PHARMA N.V.	LU
Sandostatine 100 Mikrogramm/1 ml, Injektionlösung/Infusionslösung	DE/H/5095/002	2000116136	NOVARTIS PHARMA N.V.	LU
Sandostatine 500 microgrammes/1 ml, solution injectable/pour perfusion	DE/H/5095/003	2000116137	NOVARTIS PHARMA N.V.	LU
Sandostatine 500 Mikrogramm/1 ml, Injektionlösung/Infusionslösung	DE/H/5095/003	2000116137	NOVARTIS PHARMA N.V.	LU
Sandostatin LAR 10 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/005	H/92/01395/004	NOVARTIS PHARMA GMBH	SI
Sandostatin LAR 20 mg prašek in vehikel	DE/H/5095/006	H/92/01395/005	NOVARTIS PHARMA GMBH	SI

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za suspenzijo za injiciranje				
Sandostatin LAR 30 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/007	H/92/01395/006	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/001	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/002	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/003	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/007	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/008	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/009	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/010	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/011	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,05 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/001	H/92/01395/012	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/013	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/014	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/015	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/016	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/017	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/002	H/92/01395/018	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/019	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za	DE/H/5095/003	H/92/01395/020	NOVARTIS PHARMA GMBH	SI

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injiciranje/infundiranje				
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/021	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/022	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/023	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,5 mg/ml raztopina za injiciranje/infundiranje	DE/H/5095/003	H/92/01395/024	NOVARTIS PHARMA GMBH	SI
Sandostatin LAR 10 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/005	H/92/01395/025	NOVARTIS PHARMA GMBH	SI
Sandostatin LAR 20 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/006	H/92/01395/026	NOVARTIS PHARMA GMBH	SI
Sandostatin LAR 30 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/007	H/92/01395/027	NOVARTIS PHARMA GMBH	SI
Sandostatina LAR 10 mg polvere e solvente per sospensione iniettabile	DE/H/5095/005	027083322	NOVARTIS FARMA S.P.A.	IT
Sandostatina LAR 20 mg polvere e solvente per sospensione iniettabile	DE/H/5095/006	027083334	NOVARTIS FARMA S.P.A.	IT
Sandostatina LAR 30 mg polvere e solvente per sospensione iniettabile	DE/H/5095/007	027083346	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083169	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083171	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083120	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083132	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083144	NOVARTIS FARMA S.P.A.	IT
Sandostatina 50 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/001	027083157	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083183	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml,	DE/H/5095/002	027083195	NOVARTIS FARMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluzione iniettabile/infusione				
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083221	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083207	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083219	NOVARTIS FARMA S.P.A.	IT
Sandostatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/002	027083233	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083296	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083272	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083284	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083260	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083245	NOVARTIS FARMA S.P.A.	IT
Sandostatina 500 microgrammi/1 ml, soluzione iniettabile/infusione	DE/H/5095/003	027083258	NOVARTIS FARMA S.P.A.	IT
Sandostatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	DE/H/5095/004	027083118	NOVARTIS FARMA S.P.A.	IT
Sandostatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	DE/H/5095/004	027083308	NOVARTIS FARMA S.P.A.	IT
Sandostatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	DE/H/5095/004	027083310	NOVARTIS FARMA S.P.A.	IT
SANDOSTATINE L.P. 20 mg, poudre et solvant pour suspension injectable	DE/H/5095/006	3400930021484	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 30 mg, poudre et solvant pour suspension injectable	DE/H/5095/007	3400930021545	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 10 mg, poudre et solvant pour suspension injectable	DE/H/5095/005	3400930021477	NOVARTIS PHARMA S.A.S.	FR
Sandostatina 50 microgramas/1 ml, solução injetável ou para perfusão	DE/H/5095/001	8705103	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Sandostatina 0,1 mg/1 ml ampolas,	DE/H/5095/002	8705111	NOVARTIS FARMA - PRODUTOS	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
solução injetável (s.c.) ou concentrado para solução para perfusão (perfusão i.v.)			FARMACÊUTICOS S.A.	
Sandostatina 0,5 mg/1 ml ampolas, solução injetável (s.c.) ou concentrado para solução para perfusão (perfusão i.v.)	DE/H/5095/003	8705129	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Sandostatin 100 mikrogramu/ml injekcinis ar infuzinis tirpalas	DE/H/5095/002	LT/1/94/1033/001	SIA "NOVARTIS BALTICS"	LT
Sandostatin LAR 10 mg injektiokuiva-aine ja liuotin, suspensiota varten	DE/H/5095/005	11990	NOVARTIS FINLAND OY	FI
Sandostatin LAR 10 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/005	11990	NOVARTIS FINLAND OY	FI
Sandostatin LAR 20 mg injektiokuiva-aine ja liuotin, suspensiota varten	DE/H/5095/006	11991	NOVARTIS FINLAND OY	FI
Sandostatin LAR 20 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/006	11991	NOVARTIS FINLAND OY	FI
Sandostatin LAR 30 mg injektiokuiva-aine ja liuotin, suspensiota varten	DE/H/5095/007	11992	NOVARTIS FINLAND OY	FI
Sandostatin LAR 30 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/007	11992	NOVARTIS FINLAND OY	FI
Sandostatin, injektions- /infusionsvæske, opløsning	DE/H/5095/001	12987	NOVARTIS HEALTHCARE A/S	DK
Sandostatin, injektions- /infusionsvæske, opløsning	DE/H/5095/002	12988	NOVARTIS HEALTHCARE A/S	DK
SANDOSTATIN 100 mikrogramov/1 ml injekčný/infúzny roztok	DE/H/5095/002	56/0060/17-S	NOVARTIS SLOVAKIA S.R.O.	SK
SANDOSTATIN 500 mikrogramov/1 ml injekcný/infúzny roztok	DE/H/5095/003	56/0061/17-S	NOVARTIS SLOVAKIA S.R.O.	SK
SANDOSTATIN 50 mikrogramov/1 ml injekčný/infúzny roztok	DE/H/5095/001	56/0183/90-CS	NOVARTIS SLOVAKIA S.R.O.	SK
Sandostatin 100 microgram/ml, solution for injection	DE/H/5095/002	10983	NOVARTIS SVERIGE AB	SE
Sandostatin 50 mikrogram/ml injektionsvätska, lösning	DE/H/5095/001	10982	NOVARTIS SVERIGE AB	SE
Sandostatin 500 mikrogram/ml injektionsvätska, lösning	DE/H/5095/003	11294	NOVARTIS SVERIGE AB	SE
Sandostatin LAR 20 mg pulver och vätska	DE/H/5095/006	14058	NOVARTIS SVERIGE 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
till injektionsvätska, suspension				
Sandostatin LAR 30 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/007	14059	NOVARTIS SVERIGE AB	SE
Sandostatin LAR 10 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/005	14057	NOVARTIS SVERIGE AB	SE
SANDOSTATIN 100 μικρογραμμάρια/1 ml, ενέσιμο διάλυμα/διάλυμα προς έγχυση	DE/H/5095/002	18418	NOVARTIS IRELAND LIMITED	CY
SANDOSTATIN 50 μικρογραμμάρια/1 ml, ενέσιμο διάλυμα/διάλυμα προς έγχυση	DE/H/5095/001	18417	NOVARTIS IRELAND LIMITED	CY
Sandostatine 100 microgrammes/1 ml, solution injectable/pour perfusion	DE/H/5095/002	BE141382	NOVARTIS PHARMA N.V.	BE
Sandostatine 100 microgram/1 ml, oplossing voor injectie/infusie	DE/H/5095/002	BE141382	NOVARTIS PHARMA N.V.	BE
Sandostatine 500 microgrammes/1 ml, solution injectable/pour perfusion	DE/H/5095/003	BE156362	NOVARTIS PHARMA N.V.	BE
Sandostatine 500 microgram/1 ml, oplossing voor injectie/infusie	DE/H/5095/003	BE156362	NOVARTIS PHARMA N.V.	BE
SANDOSTATINE 100 microgrammes/1 mL, solution injectable/pour perfusion	DE/H/5095/002	3400934244278	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE 50 microgrammes/1 mL, solution injectable/pour perfusion	DE/H/5095/001	3400934244100	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE 500 microgrammes/1 mL, solution injectable/pour perfusion	DE/H/5095/003	3400934244339	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 10 mg, poudre et solvant pour suspension injectable	DE/H/5095/005	3400936510630	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 20 mg, poudre et solvant pour suspension injectable	DE/H/5095/006	3400936510869	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 30 mg, poudre et solvant pour suspension injectable	DE/H/5095/007	3400936510920	NOVARTIS PHARMA S.A.S.	FR
Sandostatin® 100 µg, Injektionslösung/Infusionslösung	DE/H/5095/002	29423.01.00	NOVARTIS PHARMA GMBH	DE
Sandostatin® 1000 µg, Injektionslösung/Infusionslösung	DE/H/5095/004	29423.03.00	NOVARTIS PHARMA GMBH	DE
Sandostatin® 500 µg, Injektionslösung/Infusionslösung	DE/H/5095/003	29423.02.00	NOVARTIS PHARMA 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
SANDOSTATIN LAR-MONATSDEPOT 10 MG, PULVER UND LÖSUNGSMITTEL ZUR HERSTELLUNG EINER INJEKTIONSSUSPENSION	DE/H/5095/005	43320.00.00	NOVARTIS PHARMA GMBH	DE
SANDOSTATIN LAR-MONATSDEPOT 20 MG, PULVER UND LÖSUNGSMITTEL ZUR HERSTELLUNG EINER INJEKTIONSSUSPENSION	DE/H/5095/006	43320.01.00	NOVARTIS PHARMA GMBH	DE
Sandostatin® LAR®-Monatsdepot 30 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension	DE/H/5095/007	43320.02.00	NOVARTIS PHARMA GMBH	DE
Octreotide® 100 microgram/1 ml, solution for injection/infusion	DE/H/5095/002	PL 00101/0213	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Octreotide® 50 microgram /1 ml, solution for injection/infusion	DE/H/5095/001	PL 00101/0212	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Octreotide® 500 microgram/1 ml, solution for injection/ infusion	DE/H/5095/003	PL 00101/0214	NOVARTIS PHARMACEUTICALS UK LIMITED	UK
Sandostatin 100 µg/ml stungulyf, lausn	DE/H/5095/002	870236	NOVARTIS HEALTHCARE A/S	IS
Sandostatin LAR 20 mg stungulyfsstofn og leysir, dreifa	DE/H/5095/006	950130	NOVARTIS HEALTHCARE A/S	IS
Sandostatin LAR 30 mg stungulyfsstofn og leysir, dreifa	DE/H/5095/007	950131	NOVARTIS HEALTHCARE A/S	IS
Sandostatin 50 µg/ml stungulyf, lausn	DE/H/5095/001	870235	NOVARTIS HEALTHCARE A/S	IS
Sandostatin LAR 10 mg stungulyfsstofn og leysir, dreifa	DE/H/5095/005	950129	NOVARTIS HEALTHCARE A/S	IS
Sandostatin 100 mikrog/ml injeksjons-/infusjonsvæske, oppløsning	DE/H/5095/002	7556	NOVARTIS NORGE AS	NO
Sandostatin 200 mikrog/ml injeksjons-/infusjonsvæske, oppløsning	DE/H/5095/004	7557	NOVARTIS NORGE AS	NO
Sandostatin 50 mikrog/ml injeksjons-/infusjonsvæske, oppløsning	DE/H/5095/001	7555	NOVARTIS NORGE AS	NO
Sandostatin LAR 10 mg pulver og væske til injeksjonsvæske, suspensjon	DE/H/5095/005	95-3602	NOVARTIS NORGE AS	NO
Sandostatin LAR 20 mg pulver og væske til injeksjonsvæske, suspensjon	DE/H/5095/006	04-2502	NOVARTIS NORGE AS	NO
Sandostatin LAR 30 mg pulver og væske til injeksjonsvæske, suspensjon	DE/H/5095/007	04-2503	NOVARTIS NORGE AS	NO

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
SANDOSTATIN 500 microgram/1 ml, solution for injection/infusion	DE/H/5095/003	MA 1249/00601	NOVARTIS IRELAND LIMITED	MT
SANDOSTATIN LAR 10 mg powder and solvent for suspension for injection	DE/H/5095/005	MA1249/00603	NOVARTIS IRELAND LIMITED	MT
SANDOSTATIN LAR 20 mg powder and solvent for suspension for injection	DE/H/5095/006	MA1249/00604	NOVARTIS IRELAND LIMITED	MT
SANDOSTATIN LAR 30 mg powder and solvent for suspension for injection	DE/H/5095/007	MA1249/00605	NOVARTIS IRELAND LIMITED	MT
SANDOSTATIN LAR 20 mg powder and solvent for suspension for injection	DE/H/5095/006	PA0896/028/005	NOVARTIS IRELAND LIMITED	IE
SANDOSTATIN LAR 10 mg powder and solvent for suspension for injection	DE/H/5095/005	PA0896/028/004	NOVARTIS IRELAND LIMITED	IE
SANDOSTATIN LAR 30 mg powder and solvent for suspension for injection	DE/H/5095/007	PA0896/028/006	NOVARTIS IRELAND LIMITED	IE
SANDOSTATIN 100 microgram/1 ml, solution for injection/infusion	DE/H/5095/002	PA0896/028/002	NOVARTIS IRELAND LIMITED	IE
SANDOSTATIN 500 microgram/1 ml, solution for injection/infusion	DE/H/5095/003	PA0896/028/003	NOVARTIS IRELAND LIMITED	IE
SANDOSTATIN 50 microgram/1 ml, solution for injection/infusion	DE/H/5095/001	PA0896/028/001	NOVARTIS IRELAND LIMITED	IE
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 10 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/005	HR-H-910510401-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 20 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/006	HR-H-791598560-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 30 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/007	HR-H-983646054-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-02	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-03	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-04	NOVARTIS HRVATSKA D.O.O.	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
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-05	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-06	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,1 mg/ml otopina za injekciju/infuziju	DE/H/5095/002	HR-H-698681159-07	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-02	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-03	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-04	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-05	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-06	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin 0,5 mg/ml otopina za injekciju/infuziju	DE/H/5095/003	HR-H-463777391-07	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 10 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/005	HR-H-910510401-02	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 20 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/006	HR-H-791598560-02	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 30 mg prasak i otapalo za suspenziju za injekciju	DE/H/5095/007	HR-H-983646054-02	NOVARTIS HRVATSKA D.O.O.	HR
SANDOSTATIN LAR 10 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/005	225670402	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN LAR 20 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/006	225670502	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN LAR 30 mg κόνις και διαλύτης για ενέσιμο εναιώρημα	DE/H/5095/007	225670602	NOVARTIS (HELLAS) S.A.C.I.	GR
Octreotide Chemi 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/001	040625016	CHEMI S.P.A.	IT
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	IT/H/0227/002	PL 04465/0006	CHEMI S.P.A.	UK
Octreotide Chemi 100 microgrammi/ml,	IT/H/0227/002	040625042	CHEMI S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluzione iniettabile in siringa preriempita				
Octreotide Chemi 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/002	040625055	CHEMI S.P.A.	IT
Octreotide Chemi 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/002	040625067	CHEMI S.P.A.	IT
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	IT/H/0227/002	78136.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	IT/H/0227/001	PL 04465/0005	CHEMI S.P.A.	UK
Octreotide Chemi 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/001	040625028	CHEMI S.P.A.	IT
Octreotide Chemi 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/001	040625030	CHEMI S.P.A.	IT
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	IT/H/0227/001	78135.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	IT/H/0227/003	PL 04465/0007	CHEMI S.P.A.	UK
Octreotide Chemi 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/003	040625079	CHEMI S.P.A.	IT
Octreotide Chemi 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/003	040625081	CHEMI S.P.A.	IT
Octreotide Chemi 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0227/003	040625093	CHEMI S.P.A.	IT
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	IT/H/0227/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	IT/H/227/001	78135.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	IT/H/227/001	78135.00.00	CHEMI S.P.A.	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
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	IT/H/227/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	IT/H/227/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	IT/H/227/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	IT/H/227/003	78137.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	IT/H/0227/001	PL 04465/0005	CHEMI S.P.A.	UK
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	IT/H/0227/001	PL 04465/0005	CHEMI S.P.A.	UK
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	IT/H/0227/002	PL 04465/0006	CHEMI S.P.A.	UK
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	IT/H/0227/002	PL 04465/0006	CHEMI S.P.A.	UK
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	IT/H/0227/003	PL 04465/0007	CHEMI S.P.A.	UK
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	IT/H/0227/003	PL 04465/0007	CHEMI S.P.A.	UK
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039100110	LIFEPHARMA SPA	IT
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/153/001	34009 395 353 9 5	CHEMI S.P.A.	FR
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/153/001	34009 395 354 5 6	CHEMI S.P.A.	FR
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/153/001	34009 395 355 1 7	CHEMI S.P.A.	FR
Siroctid 0,05 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/001	1-28518	CHEMI S.P.A.	AT
Siroctid 0,1 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/002	1-28519	CHEMI S.P.A.	AT
SIROCTID® 0,1 mg/ml, solution injectable ou solution à diluer pour perfusion en	IT/H/0153/002	34009 395 384 1 9	CHEMI S.P.A.	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
seringue préremplie				
SIROCTID® 0,1 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue préremplie	IT/H/0153/002	34009 395 385 8 7	CHEMI S.P.A.	FR
SIROCTID® 0,1 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue préremplie	IT/H/0153/002	34009 395 386 4 8	CHEMI S.P.A.	FR
Siroctid 0,5 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/003	1-28520	CHEMI S.P.A.	AT
SIROCTID® 0,5 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue préremplie	IT/H/0153/003	34009 395 422 0 1	CHEMI S.P.A.	FR
SIROCTID® 0,5 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue préremplie	IT/H/0153/003	34009 395 423 7 9	CHEMI S.P.A.	FR
SIROCTID® 0,5 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue préremplie	IT/H/0153/003	34009 395 424 3 0	CHEMI S.P.A.	FR
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039100045	LIFEPHARMA SPA	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039100058	LIFEPHARMA SPA	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039100060	LIFEPHARMA SPA	IT
Siroctid 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/001	039101011	CHEMI S.P.A.	IT
Siroctid 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/001	039101023	CHEMI S.P.A.	IT
Siroctid 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/001	039101035	CHEMI S.P.A.	IT
Siroctid 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/003	039101074	CHEMI S.P.A.	IT
Siroctid 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/003	039101086	CHEMI S.P.A.	IT
Siroctid 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/003	039101098	CHEMI S.P.A.	IT
Siroctid 100 microgrammi/ml, soluzione	IT/H/0153/002	039101047	CHEMI S.P.A.	IT

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iniettabile in siringa preriempita				
Siroctid 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039101050	CHEMI S.P.A.	IT
Siroctid 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039101062	CHEMI S.P.A.	IT
Siroctid 0,05 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/001	1-28518	CHEMI S.P.A.	AT
Siroctid 0,05 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/001	1-28518	CHEMI S.P.A.	AT
Siroctid 0,1 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/002	1-28519	CHEMI S.P.A.	AT
Siroctid 0,1 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/002	1-28519	CHEMI S.P.A.	AT
Siroctid 0,5 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/003	1-28520	CHEMI S.P.A.	AT
Siroctid 0,5 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0153/003	1-28520	CHEMI S.P.A.	AT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100122	LIFEPHARMA SPA	IT
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	112911	CHEMI S.P.A.	NL
Treoject 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/001	039100019	LIFEPHARMA SPA	IT
Treoject 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/001	039100021	LIFEPHARMA SPA	IT
Treoject 50 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/001	039100033	LIFEPHARMA SPA	IT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100072	LIFEPHARMA SPA	IT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100084	LIFEPHARMA SPA	IT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100096	LIFEPHARMA SPA	IT
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	112910	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie	IT/H/0152/002	112910	CHEMI S.P.A.	NL

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in een voorgevulde spuit				
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	112910	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	112911	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	112909	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	112909	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml Injektionslösung in einer Fertigspritze	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml solution injectable en seringue préremplie	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml solution injectable en seringue préremplie	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml solution injectable en seringue préremplie	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml solution injectable en seringue préremplie	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg/ml solution injectable en	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE

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seringue préremplie				
Siroctid 0,1 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,1 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml solution injectable en seringue préremplie	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml solution injectable en seringue préremplie	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg/ml solution injectable en seringue préremplie	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Siroctid 0,5 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
Longastatina LAR 10 mg polvere e solvente per sospensione iniettabile	not available	027104088	NOVARTIS FARMA S.P.A.	IT
Longastatina LAR 10 mg polvere e solvente per sospensione iniettabile	not available	027104328	NOVARTIS FARMA S.P.A.	IT
Longastatina LAR 20 mg polvere e solvente per sospensione iniettabile	not available	027104090	NOVARTIS FARMA S.P.A.	IT
Longastatina LAR 30 mg polvere e solvente per sospensione iniettabile	not available	027104342	NOVARTIS FARMA S.P.A.	IT
Longastatina LAR 20 mg polvere e solvente per sospensione iniettabile	not available	027104330	NOVARTIS FARMA S.P.A.	IT
Longastatina LAR 30 mg polvere e	not available	027104102	NOVARTIS FARMA S.P.A.	IT

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solvente per sospensione iniettabile				
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104025	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104177	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104189	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104191	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104203	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104215	NOVARTIS FARMA S.P.A.	IT
Longastatina 100 microgrammi/1 ml, soluzione iniettabile/infusione	not available	027104227	NOVARTIS FARMA S.P.A.	IT
Longastatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	not available	027104049	NOVARTIS FARMA S.P.A.	IT
Longastatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	not available	027104292	NOVARTIS FARMA S.P.A.	IT
Longastatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	not available	027104304	NOVARTIS FARMA S.P.A.	IT
Longastatina 1000 microgrammi/5 ml, soluzione iniettabile/infusione	not available	027104316	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104013	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104114	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104126	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104138	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104140	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104153	NOVARTIS FARMA S.P.A.	IT
Longastatina 50 microgrammi /1 ml,	not available	027104165	NOVARTIS FARMA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluzione iniettabile/infusione				
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104037	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104239	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104241	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104254	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104266	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104278	NOVARTIS FARMA S.P.A.	IT
Longastatina 500 microgrammi /1 ml, soluzione iniettabile/infusione	not available	027104280	NOVARTIS FARMA S.P.A.	IT