08 February 2024
EMA/80368/2024
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: octreotide

Procedure no.: PSUSA/00002201/202306

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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
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100110	LIFEPHARMA SPA	IT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100122	LIFEPHARMA SPA	IT
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 112909	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	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 0,1 mg/ml solution injectable en seringue préremplie	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 0,5 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 0,05 mg/ml solution injectable en seringue préremplie	IT/H/0152/001	BE443746	CHEMI S.P.A.	BE
Siroctid 0,05 mg/ml	IT/H/0152/001	BE443746	CHEMI S.P.A.	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
Injektionslösung in einer Fertigspritze				
Siroctid 0,1 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/002	BE443755	CHEMI S.P.A.	BE
Siroctid 0,5 mg /ml Injektionslösung in einer Fertigspritze	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
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
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100045	LIFEPHARMA SPA	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100058	LIFEPHARMA SPA	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa	IT/H/0152/002	039100060	LIFEPHARMA SPA	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
preriempita				
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 112909	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 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

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
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 seringue préremplie	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 0,1 mg /ml	IT/H/0152/002	BE443755	CHEMI S.P.A.	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
Injektionslösung in einer Fertigspritze				
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
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/0153/001	34009 395 353 9 5	CHEMI S.P.A.	FR
Siroctid 0,05 mg/ml, solution	IT/H/0153/001	34009 395 354 5 6	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
injectable ou solution à diluer pour perfusion en seringue preremplie				
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/0153/001	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 seringue préremplie	IT/H/0153/002	34009 395 384 1 9	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 395 385 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	IT/H/0153/003	34009 395 423 7 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
pour perfusion en seringue préremplie				
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
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 iniettabile in siringa preriempita	IT/H/0153/002	039101047	CHEMI S.P.A.	IT
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	IT/H/0153/001	1-28518	CHEMI S.P.A.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Injektionslösung in einer Fertigspritze				
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
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	XI
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	78135.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	XI
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	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 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	78135.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	XI
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	XI
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	XI
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	XI
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	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
solution for injection in prefilled syringe				
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	78135.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	XI
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	78135.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 50 Mikrogramm/ml Injektionslösung	DE/H/6972/001	78135.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 100 Mikrogramm/ml Injektionslösung	DE/H/6972/002	78136.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
Octreotid-hameln 500 Mikrogramm/ml Injektionslösung	DE/H/6972/003	78137.00.00	CHEMI S.P.A.	DE
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	XI
OCTREOTIDE 50 micrograms/ml solution for injection in prefilled syringe	DE/H/6972/001	PL 04465/0005	CHEMI S.P.A.	XI
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
OCTREOTIDE 100 micrograms/ml, solution for injection in prefilled syringe	DE/H/6972/002	PL 04465/0006	CHEMI S.P.A.	XI
OCTREOTIDE 500 micrograms	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	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
/ml, solution for injection in prefilled syringe				
OCTREOTIDE 500 micrograms /ml, solution for injection in prefilled syringe	DE/H/6972/003	PL 04465/0007	CHEMI S.P.A.	XI
Siroctid 0,05 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue preremplie	IT/H/0153/001	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/0153/001	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/0153/001	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 seringue préremplie	IT/H/0153/002	34009 395 384 1 9	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 395 385 8 7	CHEMI S.P.A.	FR
SIROCTID® 0,1 mg/ml, solution injectable ou solution à diluer pour perfusion en seringue	IT/H/0153/002	34009 395 386 4 8	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
préremplie				
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
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

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
Siroctid 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0153/002	039101047	CHEMI S.P.A.	IT
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
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	DE/H/5095/006	6870/2014/01	NOVARTIS PHARMA GMBH	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
pulbere şi solvent pentru suspensie injectabilă				
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	DE/H/5095/005	1998020008	NOVARTIS PHARMA N.V.	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
Repeatable 10 mg, Pulver und Lösungsmittel zur Herstellung einer Injektionssuspension				
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 Herstellung einer Injektionssuspension	DE/H/5095/006	1998020007	NOVARTIS PHARMA N.V.	LU
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

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
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
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	DE/H/5095/007	18079	NOVARTIS IRELAND	CY

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
και διαλύτης για ενέσιμο εναιώρημα			LIMITED	
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 0,1 mg/ml, injekční/infuzní roztok	DE/H/5095/002	56/183/90-B/C	NOVARTIS S.R.O.,	CZ
Sandostatin, 100 mikrogrammi/ml, süste-/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

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 por és oldószer szuszpenziós injekcióhoz	DE/H/5095/005	OGYI-T-1723/03	NOVARTIS HUNGÁRIA KFT.	HU
Sandostatin LAR 10 mg pulveris un ŧ。ŀ₹ŀɀdinŀ》ŀ）s injekciju suspensijas pagatavoŧ。anai	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.	HU
Sandostatin LAR 20 mg pulveris un ŧ。ŀ₹ŀɀdinŀ》ŀ）s injekciju suspensijas pagatavoŧ。anai	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.	HU
Sandostatin LAR 30 mg pulveris un ŧ。ŀ₹ŀɀdinŀ》ŀ）s injekciju suspensijas pagatavoŧ。anai	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 0,1 mg/ ml, Διάλυμα για ένεση / έγχυση	DE/H/5095/002	197980101	NOVARTIS (HELLAS) S.A.C.I.	GR
SANDOSTATIN 0,5 mg/ml,	DE/H/5095/003	197980401	NOVARTIS (HELLAS)	GR

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
Διάλυμα για ένεση / έγχυση			S.A.C.I.	
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, 20mg or 30mg powder and solvent for suspension for injection	DE/H/5095/005	PL 23860/0033	NOVARTIS IRELAND LIMITED	XI
Sandostatin®LAR® 10 mg, 20mg or 30mg powder and solvent for suspension for injection	DE/H/5095/006	PL 23860/0034	NOVARTIS IRELAND LIMITED	XI
Sandostatin®LAR® 10 mg, 20mg or 30mg powder and solvent for suspension for injection	DE/H/5095/007	PL 23860/0035	NOVARTIS IRELAND LIMITED	XI
Sandostatin® 100 microgram/1 ml, solution for injection/infusion	DE/H/5095/002	PL 23860/0031	NOVARTIS IRELAND LIMITED	XI
Sandostatin® 50 microgram/1 ml, solution for injection/infusion	DE/H/5095/001	PL 23860/0030	NOVARTIS IRELAND LIMITED	XI
Sandostatin® 500 microgram/1 ml, solution for injection/infusion	DE/H/5095/003	PL 23860/0032	NOVARTIS IRELAND LIMITED	XI
SANDOSTATIN LAR 10 mg polvo y disolvente para suspensión inyectable	DE/H/5095/005	62.139	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN LAR 20 mg polvo y disolvente para suspensión inyectable	DE/H/5095/006	62.140	NOVARTIS FARMACÉUTICA S.A.	ES
SANDOSTATIN LAR 30 mg polvo y disolvente para suspensión inyectable	DE/H/5095/007	62.141	NOVARTIS FARMACÉUTICA S.A.	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
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
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	DE/H/5095/007	4595	NOVARTIS POLAND SP. Z O. O.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
wstrzykiwań				
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® 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 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

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
Sandostatine LAR 30 mg, poeder en oplosmiddel voor suspensie voor injectie	DE/H/5095/007	RVG 18237	NOVARTIS PHARMA B.V.	NL
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
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, Injektionslösung/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	DE/H/5095/003	2000116137	NOVARTIS PHARMA N.V.	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
Mikrogramm/1 ml, Injektionslösung/Infusionslösung				
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 za suspenzijo za injiciranje	DE/H/5095/006	H/92/01395/005	NOVARTIS PHARMA GMBH	SI
Sandostatin LAR 30 mg prašek in vehikel za suspenzijo za injiciranje	DE/H/5095/007	H/92/01395/006	NOVARTIS PHARMA GMBH	SI
Sandostatin 0,1 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 0,1 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 injiciranje/infundiranje	DE/H/5095/003	H/92/01395/020	NOVARTIS PHARMA GMBH	SI
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	DE/H/5095/003	H/92/01395/022	NOVARTIS PHARMA GMBH	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
za injiciranje/infundiranje				
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
SANDOSTATINE L.P. 20 mg, poudre et solvant pour suspension injectable	DE/H/5095/006	34009 300 214 8 4	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 30 mg, poudre et solvant pour suspension injectable	DE/H/5095/007	34009 300 215 4 5	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE L.P. 10 mg poudre et solvant pour suspension injectable	DE/H/5095/005	34009 300 214 7 7	NOVARTIS PHARMA S.A.S.	FR
Sandostatin 100 mikrogramų/ml injekcinis ar infuzinis tirpalas	DE/H/5095/002	LT/1/94/1033/001	SIA NOVARTIS BALTICS	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
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 100 mikrogram/ml injektions-/infusionsvätska, lösning	DE/H/5095/002	10983	NOVARTIS SVERIGE AB	SE
Sandostatin 50 mikrogram/ml injektions-/infusionsvätska, lösning	DE/H/5095/001	10982	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
Sandostatin 500 mikrogram/ml injektions-/infusionsvätska, lösning	DE/H/5095/003	11294	NOVARTIS SVERIGE AB	SE
Sandostatin LAR 20 mg pulver och vätska till injektionsvätska, suspension	DE/H/5095/006	14058	NOVARTIS SVERIGE AB	SE
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
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	34009 342 442 7 8	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE 50 microgrammes/1 mL, solution injectable/pour perfusion	DE/H/5095/001	34009 342 441 0 0	NOVARTIS PHARMA S.A.S.	FR
SANDOSTATINE 500	DE/H/5095/003	34009 342 443 3 9	NOVARTIS PHARMA S.A.S.	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
microgrammes/1 mL, solution injectable/pour perfusion				
Sandostatin® 100 µg, Injektionslösung/Infusionslösung	DE/H/5095/002	29423.01.00	NOVARTIS PHARMA GMBH	DE
Sandostatin® 500 µg, Injektionslösung/Infusionslösung	DE/H/5095/003	29423.02.00	NOVARTIS PHARMA GMBH	DE
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 23860/0031	NOVARTIS IRELAND LIMITED	XI
Octreotide® 50 microgram /1 ml, solution for injection/infusion	DE/H/5095/001	PL 23860/0030	NOVARTIS IRELAND LIMITED	XI
Octreotide® 500 microgram/1 ml, solution for injection/ infusion	DE/H/5095/003	PL 23860/0032	NOVARTIS IRELAND LIMITED	XI
Sandostatin 100 míkróg/ml stungulyf/innrennslislyf, 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 míkróg/ml stungulyf/innrennslislyf, lausn	DE/H/5095/001	870235	NOVARTIS HEALTHCARE A/S	IS
Sandostatin LAR 10 mg	DE/H/5095/005	950129	NOVARTIS HEALTHCARE A/S	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
stungulyfsstofn og leysir, dreifa				
Sandostatin 100 mikrog/ml injeksjons-/infusjonsvæske, oppløsning	DE/H/5095/002	7556	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
SANDOSTATIN 500 microgram/1 ml, solution for injection/infusion	DE/H/5095/003	MA1249/00601	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	DE/H/5095/007	PA0896/028/006	NOVARTIS IRELAND	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
powder and solvent for suspension for injection			LIMITED	
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 20 mg prašak i otapalo za suspenziju za injekciju	DE/H/5095/006	HR-H-791598560-01	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 30 mg prašak 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
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

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 0,5 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 20 mg prašak i otapalo za suspenziju za injekciju	DE/H/5095/006	HR-H-791598560-02	NOVARTIS HRVATSKA D.O.O.	HR
Sandostatin LAR 30 mg prašak 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
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	not available	027104342	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
iniettabile				
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 solvente per sospensione iniettabile	not available	027104102	NOVARTIS FARMA S.P.A.	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100110	LIFEPHARMA SPA	IT
Treoject 500 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/003	039100122	LIFEPHARMA SPA	IT
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 112909	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	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 0,1 mg/ml solution injectable en seringue préremplie	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 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	BE443764	CHEMI S.P.A.	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
Siroctid 0,5 mg/ml solution injectable en seringue préremplie	IT/H/0152/003	BE443764	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 Injektionslösung in einer Fertigspritze	IT/H/0152/001	BE443746	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 Injektionslösung in einer Fertigspritze	IT/H/0152/003	BE443764	CHEMI S.P.A.	BE
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
Treoject 100 microgrammi/ml,	IT/H/0152/002	039100045	LIFEPHARMA SPA	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				
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100058	LIFEPHARMA SPA	IT
Treoject 100 microgrammi/ml, soluzione iniettabile in siringa preriempita	IT/H/0152/002	039100060	LIFEPHARMA SPA	IT
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,1 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/002	RVG 112910	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,5 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/003	RVG 112911	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 112909	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde spuit	IT/H/0152/001	RVG 112909	CHEMI S.P.A.	NL
Siroctid 0,05 mg/ml oplossing voor injectie in een voorgevulde	IT/H/0152/001	BE443746	CHEMI S.P.A.	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
spuit				
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 seringue préremplie	IT/H/0152/002	BE443755	CHEMI S.P.A.	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
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	IT/H/0152/003	BE443764	CHEMI S.P.A.	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
Injektionslösung in einer Fertigspritze				