27 June 2018
EMA/524132/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: morphine morphine-cyclizine

Procedure no.: PSUSA/00010549/201710

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

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
ACTISKENAN 10 mg, gélule	not available	34009 349 896 3 6	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 898 6 5	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 899 2 6	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 904 6 5	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 561 958 9 3	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 905 2 6	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 908 1 6	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 900 0 7	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 901 7 5	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 902 3 6	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 906 9 4	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 907 5 5	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 912 9 5	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 910 6 6	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 914 1 7	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 915 8 5	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 911 2 7	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 913 5 6	ETHYPHARM	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
ACTISKENAN 20 mg, gélule	not available	34009 349 916 4 6	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 918 7 5	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 917 0 7	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 909 8 4	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 349 919 3 6	ETHYPHARM	FR
ACTISKENAN 20 mg, gélule	not available	34009 561 959 5 4	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 922 4 7	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 923 0 8	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 924 7 6	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 925 3 7	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 927 6 6	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 928 2 7	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 930 7 7	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 931 3 8	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 561 960 3 6	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 929 9 5	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 920 1 8	ETHYPHARM	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
ACTISKENAN 30 mg, gélule	not available	34009 349 921 8 6	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 510 8 4	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 512 0 6	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 514 3 5	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 516 6 4	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 518 9 3	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 519 5 4	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 894 0 7	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 895 7 5	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 561 957 2 5	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 513 7 4	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 511 4 5	ETHYPHARM	FR
ACTISKENAN 5 mg, gélule	not available	34009 349 517 2 5	ETHYPHARM	FR
Capros 10 mg Hartkapsel, retardiert	not available	30177.00.00	ETHYPHARM GMBH	DE
Capros 100 mg Hartkapsel, retardiert	not available	30177.03.00	ETHYPHARM GMBH	DE
Capros 30 mg Hartkapsel, retardiert	not available	30177.01.00	ETHYPHARM GMBH	DE
Capros 60 mg Hartkapsel, retardiert	not available	30177.02.00	ETHYPHARM 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
Capros akut 10 mg Kapseln	not available	57760.00.00	ETHYPHARM	DE
Capros akut 20 mg Kapseln	not available	57761.00.00	ETHYPHARM	DE
Capros akut 30 mg Kapseln	not available	57762.00.00	ETHYPHARM	DE
Capros akut 5 mg Kapseln	not available	57759.00.00	ETHYPHARM	DE
Contalgin 10 mg forðatöflur	not available	839001	PFIZER APS	IS
Contalgin 100 mg forðatöflur	not available	843416	PFIZER APS	IS
Contalgin 200 mg forðatöflur	not available	920094	PFIZER APS	IS
Contalgin 30 mg forðatöflur	not available	833071	PFIZER APS	IS
Contalgin 5 mg forðatöflur	not available	930012	PFIZER APS	IS
Contalgin 60 mg forðatöflur	not available	930290	PFIZER APS	IS
Contalgin Uno 120 mg hörð forðahylki	not available	960023	PFIZER APS	IS
Contalgin Uno 150 mg hörð forðahylki	not available	960024	PFIZER APS	IS
Contalgin Uno 200 mg hörð forðahylki	not available	960025	PFIZER APS	IS
Contalgin Uno 30 mg hörð forðahylki	not available	960020	PFIZER APS	IS
Contalgin Uno 60 mg hörð forðahylki	not available	960021	PFIZER APS	IS
Contalgin Uno 90 mg hörð forðahylki	not available	960022	PFIZER APS	IS
Contalgin Uno, depotkapsler, hårde	not available	17800	PFIZER APS	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
Contalgin Uno, depotkapsler, hårde	not available	17799	PFIZER APS	DK
Contalgin Uno, depotkapsler, hårde	not available	17801	PFIZER APS	DK
Contalgin Uno, depotkapsler, hårde	not available	17802	PFIZER APS	DK
Contalgin Uno, depotkapsler, hårde	not available	17798	PFIZER APS	DK
Contalgin Uno, depotkapsler, hårde	not available	17797	PFIZER APS	DK
Contalgin, depotgranulat til oral suspension, enkeltdosisbeholder	UK/H/0110/004	18031	PFIZER APS	DK
Contalgin, depotgranulat til oral suspension, enkeltdosisbeholder	UK/H/0110/002	18029	PFIZER APS	DK
Contalgin, depotgranulat til oral suspension, enkeltdosisbeholder	UK/H/0110/001	18028	PFIZER APS	DK
Contalgin, depotgranulat til oral suspension, enkeltdosisbeholder	UK/H/0110/003	18030	PFIZER APS	DK
Contalgin, depotgranulat til oral suspension, enkeltdosisbeholder	UK/H/0110/005	18032	PFIZER APS	DK
Contalgin, depottabletter	not available	10801	PFIZER APS	DK
Contalgin, depottabletter	not available	11641	PFIZER APS	DK
Contalgin, depottabletter	not available	11021	PFIZER APS	DK
Contalgin, depottabletter	not available	13515	PFIZER APS	DK
Contalgin, depottabletter	not available	15421	PFIZER APS	DK
Contalgin, depottabletter	not available	14491	PFIZER APS	DK
Cyclimorph 10 Solution for Injection	not available	PA1142/003/001	AMDIPHARM LIMITED	IE
Cyclimorph 15 Solution	not available	PA1142/003/002	AMDIPHARM LIMITED	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
for Injection				
Cyclimorph-10 Injection 1 ml Ampoules	not available	PL 20072/0007	AMDIPHARM UK LIMITED	UK
Cyclimorph-15 Injection	not available	PL 20072/0008	AMDIPHARM UK LIMITED	UK
Cyclizine Tartrate 50mg/ml and Morphine Tartrate 10mg/ml Injection	not available	PL 20072/0007	AMDIPHARM UK LIMITED	UK
Cyclizine Tartrate 50mg/ml and Morphine Tartrate 15mg/ml Injection	not available	PL 20072/0008	AMDIPHARM UK LIMITED	UK
Dolcontin 10 mg depottabletter	not available	7070	PFIZER AS	NO
Dolcontin 10 mg depottabletter	not available	10847	PFIZER AB	SE
Dolcontin 10 mg depottabletti	not available	9109	MUNDIPHARMA OY	FI
Dolcontin 100 mg depotgranulat till oral suspension	not available	12875	PFIZER AB	SE
Dolcontin 100 mg depotrakeet oraalisuspensiota varten	not available	12450	MUNDIPHARMA OY	FI
Dolcontin 100 mg depottabletter	not available	7072	PFIZER AS	NO
Dolcontin 100 mg depottabletter	not available	10849	PFIZER AB	SE
Dolcontin 100 mg depottabletti	not available	9829	MUNDIPHARMA OY	FI
Dolcontin 20 mg depotgranulat til mikstur, suspensjon i dosepose	not available	95-3811	PFIZER AS	NO
Dolcontin 20 mg depotgranulat till oral	not available	12872	PFIZER 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
suspension				
Dolcontin 20 mg depotrakeet oraalisuspensiota varten	not available	12447	MUNDIPHARMA OY	FI
Dolcontin 200 mg depotrakeet oraalisuspensiota varten	not available	12451	MUNDIPHARMA OY	FI
Dolcontin 200 mg depottabletter	not available	7912	PFIZER AS	NO
Dolcontin 200 mg depottabletter	not available	11694	PFIZER AB	SE
Dolcontin 200 mg depottabletti	not available	11089	MUNDIPHARMA OY	FI
Dolcontin 30 mg depotgranulat till oral suspension	not available	12873	PFIZER AB	SE
Dolcontin 30 mg depotrakeet oraalisuspensiota varten	not available	12448	MUNDIPHARMA OY	FI
Dolcontin 30 mg depottabletter	not available	7071	PFIZER AS	NO
Dolcontin 30 mg depottabletter	not available	10848	PFIZER AB	SE
Dolcontin 30 mg depottabletti	not available	9110	MUNDIPHARMA OY	FI
Dolcontin 5 mg depottabletter	not available	7940	PFIZER AS	NO
Dolcontin 5 mg depottabletter	not available	11832	PFIZER AB	SE
Dolcontin 60 mg depotgranulat till oral suspension	not available	12874	PFIZER AB	SE
Dolcontin 60 mg depotrakeet oraalisuspensiota varten	not available	12449	MUNDIPHARMA OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dolcontin 60 mg depottabletter	not available	7561	PFIZER AS	NO
Dolcontin 60 mg depottabletter	not available	11076	PFIZER AB	SE
Dolcontin 60 mg depottabletti	not available	10539	MUNDIPHARMA OY	FI
Dolcontin Unotard 120 mg depotkapsel, hård	not available	12864	PFIZER AB	SE
Dolcontin Unotard 120 mg depotkapseli, kova	not available	12518	MUNDIPHARMA OY	FI
Dolcontin Unotard 150 mg depotkapsel, hård	not available	12865	PFIZER AB	SE
Dolcontin Unotard 150 mg depotkapseli, kova	not available	12519	MUNDIPHARMA OY	FI
Dolcontin Unotard 200 mg depotkapsel, hård	not available	12866	PFIZER AB	SE
Dolcontin Unotard 200 mg depotkapseli, kova	not available	12520	MUNDIPHARMA OY	FI
Dolcontin Unotard 30 mg depotkapsel, hård	not available	12861	PFIZER AB	SE
Dolcontin Unotard 30 mg depotkapseli, kova	not available	12515	MUNDIPHARMA OY	FI
Dolcontin Unotard 60 mg depotkapsel, hård	not available	12862	PFIZER AB	SE
Dolcontin Unotard 60 mg depotkapseli, kova	not available	12516	MUNDIPHARMA OY	FI
Dolcontin Unotard 90 mg depotkapsel, hård	not available	12863	PFIZER AB	SE
Dolcontin Unotard 90 mg depotkapseli, kova	not available	12517	MUNDIPHARMA OY	FI
DOLQ 20 mg comprimidos efervescentes	not available	68892	ARAFARMA GROUP, S.A	ES
Kapanol 20, capsules	not available	RVG 17210	GLAXOSMITHKLINE 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
met gereguleerde afgifte 20 mg				
Kapanol 50, capsules met gereguleerde afgifte 50 mg	not available	RVG 17211	GLAXOSMITHKLINE B.V.	NL
M-long 10 mg, Hartkapseln, retardiert	not available	30173.00.00	GRÜNENTHAL GMBH	DE
M-long 100 mg, Hartkapseln, retardiert	not available	30173.03.00	GRÜNENTHAL GMBH	DE
M-long 30 mg, Hartkapseln, retardiert	not available	30173.01.00	GRÜNENTHAL GMBH	DE
M-long 60 mg, Hartkapseln, retardiert	not available	30173.02.00	GRÜNENTHAL GMBH	DE
Morapid 10 mg Filmtabletten	not available	1-20439	MUNDIPHARMA GES.M.B.H	AT
Morapid 20 mg Filmtabletten	not available	1-20444	MUNDIPHARMA GES.M.B.H	AT
Morph - 1 A Pharma 20 mg Retardtabletten	not available	94190.00.00	1 A PHARMA GMBH	DE
Morph - 1 A Pharma 45 mg Retardtabletten	not available	94191.00.00	1 A PHARMA GMBH	DE
Morphin HEXAL® 20 mg Retardtabletten	not available	88049.00.00	HEXAL AG	DE
Morphin HEXAL® 45 mg Retardtabletten	not available	88050.00.00	HEXAL AG	DE
Morphin Merck ® 100 mg, Infusionslösung Zur Anwendung bei Säuglingen, Kindern und Erwachsenen	not available	6108826.02.00	MERCK SERONO GMBH	DE
Morphin Merck® 10 mg, Injektionslösung Zur Anwendung bei Kindern und Erwachsenen	not available	6108826.00.00	MERCK SERONO GMBH	DE
Morphin Merck® 20 mg,	not available	6108826.01.00	MERCK SERONO 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
Injektionslösung Zur Anwendung bei Kindern und Erwachsenen				
Morphin Merck® Tropfen 0,5%	not available	31264.00.00	MERCK SERONO GMBH	DE
Morphin Merck® Tropfen 2%	not available	31264.01.00	MERCK SERONO GMBH	DE
Morphine "Orion", injektionsvæske, opløsning	NO/H/0258/002	57511	ORION CORPORATION	DK
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	NL 19143-3400936907782	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 10 mg/ml, solution injectable	not available	34009 553 532 6 3	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 10 mg/ml, solution injectable	not available	34009 553 532 6 3	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (SULFATE) LAVOISIER 1 mg/ml, solution injectable	not available	NL24641-3400935620699	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (SULFATE) LAVOISIER 50 mg/ml, solution injectable	not available	NL24418-3400935620170	LABORATOIRES CHAIX ET DU MARAIS	FR
Morphine Orion 20 mg/ml injeksjonsvæske, oppløsning	NO/H/0258/002	16-11113	ORION CORPORATION	NO
Morphine Orion 20 mg/ml injektioneste, liuos	NO/H/0258/002	34067	ORION CORPORATION	FI
Morphine Orion 20	NO/H/0258/002	34067	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/ml injektionsvätska, lösning				
Morphine Sulfate Injection 10 mg in 1 ml	not available	PL 17907/0597	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Morphine Sulfate Injection 15 mg in 1 ml	not available	PL 17907/0598	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Morphine Sulfate Injection 30 mg in 1 ml	not available	PL 17907/0599	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Morphine Sulfate Injection BP MinijetTM	not available	PL 03265/0037	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	UK
Morphine Sulphate 10mg/ml Solution for Injection	not available	PA0073/020/001	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 1mg/5ml Solution for Injection	not available	PA0073/020/003	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 30 mg/ml Solution for Injection	not available	PA0073/020/004	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 60 mg/ml Solution for Injection	not available	PA0073/020/008	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphinsulfat Auto-Injector, 10 mg, Injektionslösung in Fertigpen	not available	58602.00.00	MERIDIAN MEDICAL TECHNOLOGIES LTD.	DE
MOSCONTIN 10 mg, comprimé enrobé à libération prolongée	not available	34009 555 594 9 8	MUNDIPHARMA SAS	FR
MOSCONTIN 10 mg, comprimé enrobé à libération prolongée	not available	34009 328 697 1 8	MUNDIPHARMA SAS	FR
MOSCONTIN 100 mg, comprimé enrobé à	not available	34009 555 597 8 8	MUNDIPHARMA SAS	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
libération prolongée				
MOSCONTIN 100 mg, comprimé enrobé à libération prolongée	not available	34009 328 700 2 8	MUNDIPHARMA SAS	FR
MOSCONTIN 30 mg, comprimé enrobé à libération prolongée	not available	34009 555 595 5 9	MUNDIPHARMA SAS	FR
MOSCONTIN 30 mg, comprimé enrobé à libération prolongée	not available	34009 328 698 8 6	MUNDIPHARMA SAS	FR
MOSCONTIN 60 mg, comprimé enrobé à libération prolongée	not available	34009 555 596 1 0	MUNDIPHARMA SAS	FR
MOSCONTIN 60 mg, comprimé enrobé à libération prolongée	not available	34009 328 699 4 7	MUNDIPHARMA SAS	FR
MOSCONTIN LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 558 273 9 9	MUNDIPHARMA SAS	FR
MOSCONTIN LP 200 mg, comprimé pelliculé à libération prolongée	not available	34009 343 006-6 0	MUNDIPHARMA SAS	FR
MS Contin 10 mg, comprimés à libération prolongée	not available	BE 134032	MUNDIPHARMA COMM VA	BE
MS Contin 10 mg, comprimés à libération prolongée	not available	1495/05078269	MUNDIPHARMA COMM VA	LU
MS Contin 10 mg, tabletten met verlengde afgifte	not available	BE 134032	MUNDIPHARMA COMM VA	BE
MS Contin 10 mg, tabletten met verlengde afgifte	not available	RVG 11205	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 100 mg,	not available	BE 134066	MUNDIPHARMA COMM VA	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimés à libération prolongée				
MS Contin 100 mg, comprimés à libération prolongée	not available	1495/05078273	MUNDIPHARMA COMM VA	LU
MS Contin 100 mg, tabletten met verlengde afgifte	not available	BE 134066	MUNDIPHARMA COMM VA	BE
MS Contin 100 mg, tabletten met verlengde afgifte	not available	RVG 11208	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 15 mg, comprimés à libération prolongée	not available	BE 175874	MUNDIPHARMA COMM VA	BE
MS Contin 15 mg, comprimés à libération prolongée	not available	1495/05078270	MUNDIPHARMA COMM VA	LU
MS Contin 15 mg, tabletten met verlengde afgifte	not available	BE 175874	MUNDIPHARMA COMM VA	BE
MS Contin 15 mg, tabletten met verlengde afgifte	not available	RVG 16674	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 200 mg, comprimés à libération prolongée	not available	BE 168646	MUNDIPHARMA COMM VA	BE
MS Contin 200 mg, comprimés à libération prolongée	not available	1495/05078274	MUNDIPHARMA COMM VA	LU
MS Contin 200 mg, tabletten met verlengde afgifte	not available	BE 168646	MUNDIPHARMA COMM VA	BE
MS Contin 200 mg, tabletten met verlengde afgifte	not available	RVG 15376	MUNDIPHARMA PHARMACEUTICALS BV	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
MS Contin 30 mg, comprimés à libération prolongée	not available	BE 134041	MUNDIPHARMA COMM VA	BE
MS Contin 30 mg, comprimés à libération prolongée	not available	1495/05078271	MUNDIPHARMA COMM VA	LU
MS Contin 30 mg, tabletten met verlengde afgifte	not available	BE 134041	MUNDIPHARMA COMM VA	BE
MS Contin 30 mg, tabletten met verlengde afgifte	not available	RVG 11206	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 5 mg, comprimés à libération prolongée	not available	BE 175865	MUNDIPHARMA COMM VA	BE
MS Contin 5 mg, comprimés à libération prolongée	not available	1495/05078268	MUNDIPHARMA COMM VA	LU
MS Contin 5 mg, tabletten met verlengde afgifte	not available	BE 175865	MUNDIPHARMA COMM VA	BE
MS Contin 5 mg, tabletten met verlengde afgifte	not available	RVG 16673	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 5 mg, tabletten met verlengde afgifte	not available	RVG 16673	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 60 mg, comprimés à libération prolongée	not available	BE 134057	MUNDIPHARMA COMM VA	BE
MS Contin 60 mg, comprimés à libération prolongée	not available	1495/05078272	MUNDIPHARMA COMM VA	LU
MS Contin 60 mg, tabletten met verlengde	not available	BE 134057	MUNDIPHARMA COMM VA	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
afgifte				
MS Contin 60 mg, tabletten met verlengde afgifte	not available	RVG 11207	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Direct 10 mg, comprimés enrobés	not available	1495/02/12/7018	MUNDIPHARMA COMM VA	LU
MS Direct 10 mg, omhulde tabletten	not available	BE192482	MUNDIPHARMA COMM VA	BE
MS Direct 10 mg, comprimés enrobés	not available	BE192482	MUNDIPHARMA COMM VA	BE
MS Direct 20 mg, comprimés enrobés	not available	1495/02/12/7019	MUNDIPHARMA COMM VA	LU
MS Direct 20 mg, omhulde tabletten	not available	BE192473	MUNDIPHARMA COMM VA	BE
MS Direct 20 mg, comprimés enrobés	not available	BE192473	MUNDIPHARMA COMM VA	BE
MSR 10 mg Mundipharma® Zäpfchen	not available	30760.00.00	MUNDIPHARMA GMBH	DE
MSR 20 mg Mundipharma® Zäpfchen	not available	30760.01.00	MUNDIPHARMA GMBH	DE
MSR 30 mg Mundipharma® Zäpfchen	not available	30760.02.00	MUNDIPHARMA GMBH	DE
MST 10 mg Mundipharma® Retardtabletten	not available	4034.01.00	MUNDIPHARMA GMBH	DE
MST 100 mg Mundipharma® Retardtabletten	not available	4034.03.00	MUNDIPHARMA GMBH	DE
MST 200 mg Mundipharma® Retardtabletten	not available	26813.00.00	MUNDIPHARMA GMBH	DE
MST 30 mg Mundipharma® Retardtabletten	not available	4034.00.00	MUNDIPHARMA GMBH	DE
MST 60 mg	not available	4034.02.00	MUNDIPHARMA 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
Mundipharma® Retardtabletten				
MST CONTINUS ® 100 mg comprimidos de liberación prolongada	not available	57.900	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 10 mg prolonged release tablets	not available	MA198/00201	NAPP PHARMACEUTICALS LTD	MT
MST CONTINUS 10 mg prolonged-release tablets	not available	PA 1688/4/2	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST Continus 10 mg retard filmtabletta	not available	OGYI-T-2187/01	MUNDIPHARMA GES.M.B.H	HU
MST Continus 10 mg	not available	65/0119/91-C/S	MUNDIPHARMA GES.M.B.H	SK
MST Continus 100 mg	not available	65/0257/13-S	MUNDIPHARMA GES.M.B.H	SK
MST CONTINUS 100 mg Prolonged-release Tablets	not available	PA 1688/4/6	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST Continus 100 mg retard filmtabletta	not available	OGYI-T-2187/05	MUNDIPHARMA GES.M.B.H	HU
MST Continus 100 mg tablete s produljenim oslobađanjem	not available	HR-H-448781299	MEDIS ADRIA D.O.O.	HR
MST CONTINUS 15 mg prolonged-release tablets	not available	PA 1688/4/3	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST Continus 30 mg	not available	65/0259/13-S	MUNDIPHARMA GES.M.B.H	SK
MST CONTINUS 30 mg prolonged release tablets	not available	MA198/00202	NAPP PHARMACEUTICALS LTD	MT
MST CONTINUS 30 mg prolonged-release tablets	not available	PA 1688/4/4	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST Continus 30 mg retard filmtabletta	not available	OGYI-T-2187/03	MUNDIPHARMA GES.M.B.H	HU
MST Continus 30 mg tablete s produljenim oslobađanjem	not available	HR-H-113395944	MEDIS ADRIA D.O.O.	HR
MST CONTINUS 5 mg	not available	PA 1688/4/1	MUNDIPHARMA	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
prolonged-release tablets			PHARMACEUTICALS LIMITED	
MST CONTINUS 60 mg prolonged release tablets	not available	MA198/00203	NAPP PHARMACEUTICALS LTD	MT
MST CONTINUS 60 mg Prolonged-release tablets	not available	PA 1688/4/5	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST Continus 60 mg retard filmtabletta	not available	OGYI-T-2187/04	MUNDIPHARMA GES.M.B.H	HU
MST Continus 60 mg tablete s produljenim oslobađanjem	not available	HR-H-734333555	MEDIS ADRIA D.O.O.	HR
MST Continus 60 mg	not available	65/0258/13-S	MUNDIPHARMA GES.M.B.H	SK
MST CONTINUS suspension 100 mg	UK/H/0110/004	PL 16950/0033	NAPP PHARMACEUTICALS LTD	UK
MST CONTINUS suspension 20 mg	UK/H/0110/001	PL 16950/0030	NAPP PHARMACEUTICALS LTD	UK
MST CONTINUS suspension 200 mg	UK/H/0110/005	PL 16950/0034	NAPP PHARMACEUTICALS LTD	UK
MST CONTINUS suspension 30 mg	UK/H/0110/002	PL 16950/0031	NAPP PHARMACEUTICALS LTD	UK
MST CONTINUS suspension 60 mg	UK/H/0110/003	PL 16950/0032	NAPP PHARMACEUTICALS LTD	UK
MST CONTINUS tablets 10 mg	not available	19746	MUNDIPHARMA PHARMACEUTICALS LTD	CY
MST CONTINUS tablets 100 mg	not available	19748	MUNDIPHARMA PHARMACEUTICALS LTD	CY
MST CONTINUS tablets 30 mg	not available	19745	MUNDIPHARMA PHARMACEUTICALS LTD	CY
MST CONTINUS tablets 60 mg	not available	19747	MUNDIPHARMA PHARMACEUTICALS LTD	CY
MST Continus, 100 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4765	NORPHARMA A/S	PL
MST Continus, 10mg	not available	4762	NORPHARMA A/S	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
tabletki powlekane o zmodyfikowanym uwalnianiu				
MST Continus, 200 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4766	NORPHARMA A/S	PL
MST Continus, 30 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4763	NORPHARMA A/S	PL
MST Continus, 60 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4764	NORPHARMA A/S	PL
MST CONTINUS® 10 mg comprimidos de liberación prolongada	not available	57.898	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS® 15 mg comprimidos de liberación prolongada	not available	61.081	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS® 200 mg comprimidos de liberación prolongada	not available	61.909	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS® 30 mg comprimidos de liberación prolongada	not available	57.897	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS® 5 mg comprimidos de liberación prolongada	not available	61.080	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS® 60 mg comprimidos de liberación prolongada	not available	57.899	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST® 100 mg Retard-Granulat	not available	30679.03.00	MUNDIPHARMA 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
MST® 20 mg Retard-Granulat	not available	30679.00.00	MUNDIPHARMA GMBH	DE
MST® 200 mg Retard-Granulat	not available	30679.04.00	MUNDIPHARMA GMBH	DE
MST® 30 mg Retard-Granulat	not available	30679.01.00	MUNDIPHARMA GMBH	DE
MST® 60 mg Retard-Granulat	not available	30679.02.00	MUNDIPHARMA GMBH	DE
Mundidol retard 10 mg Filmtabletten	not available	1-18003	MUNDIPHARMA GES.M.B.H	AT
Mundidol retard 100 mg Filmtabletten	not available	1-18376	MUNDIPHARMA GES.M.B.H	AT
Mundidol retard 200 mg Filmtabletten	not available	1-19435	MUNDIPHARMA GES.M.B.H	AT
Mundidol retard 30 mg Filmtabletten	not available	1-18004	MUNDIPHARMA GES.M.B.H	AT
Mundidol retard 60 mg Filmtabletten	not available	1-18375	MUNDIPHARMA GES.M.B.H	AT
Oramorph Concentrated Oral Solution 20 mg/ml	not available	PL 0015/0125	BOEHRINGER INGELHEIM LTD.	UK
Oramorph Concentrated Oral Solution 20mg/ml	not available	PA 7/44/2	BOEHRINGER INGELHEIM LTD.	IE
Oramorph Oral Solution 10 mg/5 ml	not available	PA 7/44/1	BOEHRINGER INGELHEIM LTD.	IE
Oramorph Oral Solution 10 mg/5 ml	not available	PL 0015/0122	BOEHRINGER INGELHEIM LTD.	UK
Sevredol 10 mg	not available	65/0302/98-S	MUNDIPHARMA GES.M.B.H	SK
Sevredol 10 mg filmom obložene tablete	not available	HR-H-915182609	MEDIS ADRIA D.O.O.	HR
Sevredol 10 mg filmsko obložene tablete	not available	H/01/01416/001	MEDIS, D.O.O.	SI
Sevredol 10 mg filmsko obložene tablete	not available	H/01/01416/002	MEDIS, D.O.O.	SI
SEVREDOL 10 mg,	not available	34009 334 799 7 8	MUNDIPHARMA SAS	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
comprimé pelliculé sécable				
SEVREDOL 10mg	not available	65/299/99-C	MUNDIPHARMA GES.M.B.H	CZ
Sevredol 20 mg	not available	65/0256/13-S	MUNDIPHARMA GES.M.B.H	SK
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MEDIS ADRIA D.O.O.	HR
Sevredol 20 mg filmsko obložene tablete	not available	H/01/01416/003	MEDIS, D.O.O.	SI
Sevredol 20 mg filmsko obložene tablete	not available	H/01/01416/004	MEDIS, D.O.O.	SI
SEVREDOL 20 mg, comprimé pelliculé sécable	not available	34009 334 800 5 9	MUNDIPHARMA SAS	FR
SEVREDOL 20mg	not available	65/300/99-C	MUNDIPHARMA GES.M.B.H	CZ
Sevredol tablets 10 mg	not available	19750	MUNDIPHARMA PHARMACEUTICALS LTD	CY
SEVREDOL tablets 20 mg	not available	19749	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Sevredol, 10 mg, tabletki powlekane	not available	8826	NORPHARMA A/S	PL
Sevredol, 20 mg, tabletki powlekane	not available	8827	NORPHARMA A/S	PL
Sevredol® 10 mg comprimidos recubiertos con película	not available	59.656	MUNDIPHARMA PHARMACEUTICALS SL	ES
Sevredol® 10 mg Filmtabletten	not available	25932.00.00	MUNDIPHARMA GMBH	DE
Sevredol® 20 mg comprimidos recubiertos con película	not available	59.655	MUNDIPHARMA PHARMACEUTICALS SL	ES
Sevredol® 20 mg Filmtabletten	not available	25932.01.00	MUNDIPHARMA GMBH	DE
SKENAN L.P. 10 mg, microgranules à	not available	34009 333 235 2 3	ETHYPHARM	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
libération prolongée en gélule				
SKENAN L.P. 100 mg, microgranules à libération prolongée en gélule	not available	34009 333 238 1 3	ETHYPHARM	FR
SKENAN L.P. 200 mg, microgranules à libération prolongée en gélule	not available	34009 340 538 7 0	ETHYPHARM	FR
SKENAN L.P. 200 mg, microgranules à libération prolongée en gélule	not available	34009 340 537 0 2	ETHYPHARM	FR
SKENAN L.P. 200 mg, microgranules à libération prolongée en gélule	not available	34009 340 539 3 1	ETHYPHARM	FR
SKENAN L.P. 30 mg, microgranules à libération prolongée en gélule	not available	34009 333 236 9 1	ETHYPHARM	FR
SKENAN L.P. 60 mg, microgranules à libération prolongée en gélule	not available	34009 333 237 5 2	ETHYPHARM	FR
Substitol 120 mg trde kapsule s podaljšanim sproščanjem	not available	H/04/01467/001	MEDIS, D.O.O.	SI
Substitol 200 mg trde kapsule s podaljšanim sproščanjem	not available	H/04/01467/002	MEDIS, D.O.O.	SI
Substitol 30 mg Hartkapseln, retardiert	not available	31512.04.00	MUNDIPHARMA GMBH	DE
Substitol 60 mg	not available	31512.02.00	MUNDIPHARMA 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
Hartkapseln, retardiert				
Substitol retard 120 mg Kapseln	not available	1-22749	MUNDIPHARMA GES.M.B.H	AT
Substitol retard 200 mg Kapseln	not available	1-22750	MUNDIPHARMA GES.M.B.H	AT
Substitol® 100 mg Hartkapseln, retardiert	not available	31512.01.00	MUNDIPHARMA GMBH	DE
Substitol® 200 mg Hartkapseln, retardiert	not available	31512.00.00	MUNDIPHARMA GMBH	DE
TWICE 10 mg Capsule rigide a rilascio prolungato	not available	033484015	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TWICE 100 mg Capsule rigide a rilascio prolungato	not available	033484041	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TWICE 30 mg Capsule rigide a rilascio prolungato	not available	033484027	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
TWICE 60 mg Capsule rigide a rilascio prolungato	not available	033484039	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
Zomorph 10 mg cápsulas duras de liberación prolongada	not available	60.125	ETHYPHARM	ES
Zomorph 100 mg cápsulas duras de liberación prolongada	not available	60.122	ETHYPHARM	ES
Zomorph 30 mg cápsulas duras de liberación prolongada	not available	60.124	ETHYPHARM	ES
Zomorph 60 mg cápsulas	not available	60.123	ETHYPHARM	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
duras de liberación prolongada				
ZOMORPH capsules 100mg	not available	PL 06934/0185	ETHYPHARM	UK
ZOMORPH capsules 10mg	not available	PL 06934/0182	ETHYPHARM	UK
ZOMORPH capsules 200mg	not available	PL 06934/0186	ETHYPHARM	UK
ZOMORPH capsules 30mg	not available	PL 06934/0183	ETHYPHARM	UK
ZOMORPH Capsules 60mg	not available	PL 06934/0184	ETHYPHARM	UK
МСТ КОНТИНУС 10 mg таблетки с изменено освобождаване	not available	20000384	MUNDIPHARMA GES.M.B.H	BG
МСТ КОНТИНУС 100 mg таблетки с изменено освобождаване	not available	20000387	MUNDIPHARMA GES.M.B.H	BG
МСТ КОНТИНУС 30 mg таблетки с изменено освобождаване	not available	20000385	MUNDIPHARMA GES.M.B.H	BG
МСТ КОНТИНУС 60 mg таблетки с изменено освобождаване	not available	20000386	MUNDIPHARMA GES.M.B.H	BG
Субститол 120 mg капсули с удължено освобождаване	not available	20030413	MUNDIPHARMA GESELLSCHAFT M.B.H.	BG
Субститол 200 mg капсули с удължено освобождаване	not available	20030414	MUNDIPHARMA GESELLSCHAFT M.B.H.	BG