22 June 2023
EMA/356967/2023
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: morphine, morphine / cyclizine

Procedure no.: PSUSA/00010549/202210

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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
Actimorph 1 mg Orodispersible tablets	DE/H/6657/001	PL 06934/0245	ETHYPHARM	XI
Actimorph 10 mg Orodispersible tablets	DE/H/6657/004	PL 06934/0248	ETHYPHARM	XI
Actimorph 2.5 mg Orodispersible tablets	DE/H/6657/002	PL 06934/0246	ETHYPHARM	XI
Actimorph 20 mg Orodispersible tablets	DE/H/6657/005	PL 06934/0249	ETHYPHARM	XI
Actimorph 30 mg Orodispersible tablets	DE/H/6657/006	PL 06934/0250	ETHYPHARM	XI
Actimorph 5 mg Orodispersible tablets	DE/H/6657/003	PL 06934/0247	ETHYPHARM	XI
ACTISKENAN 1 mg, comprimé orodispersible	DE/H/6657/001	34009 302 333 1 3	ETHYPHARM	FR
ACTISKENAN 10 mg, comprimé orodispersible	DE/H/6657/004	34009 302 333 5 1	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 896 3 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 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

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 349 906 9 4	ETHYPHARM	FR
ACTISKENAN 10 mg, gélule	not available	34009 349 907 5 5	ETHYPHARM	FR
ACTISKENAN 2,5 mg, comprimé orodispersible	DE/H/6657/002	34009 302 333 3 7	ETHYPHARM	FR
ACTISKENAN 20 mg, comprimé orodispersible	DE/H/6657/005	34009 302 333 6 8	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

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 349 913 5 6	ETHYPHARM	FR
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, comprimé orodispersible	DE/H/6657/006	34009 302 333 7 5	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 922 4 7	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 349 923 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

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 349 920 1 8	ETHYPHARM	FR
ACTISKENAN 30 mg, gélule	not available	34009 349 921 8 6	ETHYPHARM	FR
ACTISKENAN 5 mg, comprimé orodispersible	DE/H/6657/003	34009 302 333 4 4	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

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 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	DE
Capros 100 mg Hartkapsel, retardiert	not available	30177.03.00	ETHYPHARM	DE
Capros 30 mg Hartkapsel, retardiert	not available	30177.01.00	ETHYPHARM	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 60 mg Hartkapsel, retardiert	not available	30177.02.00	ETHYPHARM	DE
Capros akut 1 mg Schmelztabletten	DE/H/6657/001	7000173.00.00	ETHYPHARM	DE
Capros akut 10 mg Kapseln	not available	57760.00.00	ETHYPHARM	DE
Capros akut 10 mg Schmelztabletten	DE/H/6657/004	7000176.00.00	ETHYPHARM	DE
Capros akut 2,5 mg Schmelztabletten	DE/H/6657/002	7000174.00.00	ETHYPHARM	DE
Capros akut 20 mg Kapseln	not available	57761.00.00	ETHYPHARM	DE
Capros akut 20 mg Schmelztabletten	DE/H/6657/005	7000177.00.00	ETHYPHARM	DE
Capros akut 30 mg Kapseln	not available	57762.00.00	ETHYPHARM	DE
Capros akut 30 mg Schmelztabletten	DE/H/6657/006	7000178.00.00	ETHYPHARM	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 5 mg Kapseln	not available	57759.00.00	ETHYPHARM	DE
Capros akut 5 mg Schmelztabletten	DE/H/6657/003	7000175.00.00	ETHYPHARM	DE
Contalgin 10 mg forðatöflur.	not available	839001	MUNDIPHARMA A/S	IS
Contalgin 10 mg forðatöflur.	not available	839001	MUNDIPHARMA A/S	IS
Contalgin 100 mg forðatöflur.	not available	843416	MUNDIPHARMA A/S	IS
Contalgin 100 mg forðatöflur.	not available	843416	MUNDIPHARMA A/S	IS
Contalgin 200 mg forðatöflur.	not available	920094	MUNDIPHARMA A/S	IS
Contalgin 200 mg forðatöflur.	not available	920094	MUNDIPHARMA A/S	IS
Contalgin 30 mg forðatöflur.	not available	833071	MUNDIPHARMA 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
Contalgin 30 mg forðatöflur.	not available	833071	MUNDIPHARMA A/S	IS
Contalgin 5 mg forðatöflur.	not available	930012	MUNDIPHARMA A/S	IS
Contalgin 5 mg forðatöflur.	not available	930012	MUNDIPHARMA A/S	IS
Contalgin 60 mg forðatöflur.	not available	930290	MUNDIPHARMA A/S	IS
Contalgin 60 mg forðatöflur.	not available	930290	MUNDIPHARMA A/S	IS
Contalgin Uno 120 mg hörð forðahylki.	not available	960023	MUNDIPHARMA A/S	IS
Contalgin Uno 120 mg hörð forðahylki.	not available	960023	MUNDIPHARMA A/S	IS
Contalgin Uno 150 mg hörð forðahylki.	not available	960024	MUNDIPHARMA A/S	IS
Contalgin Uno 150 mg hörð forðahylki.	not available	960024	MUNDIPHARMA 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
Contalgin Uno 200 mg hörð forðahylki.	not available	960025	MUNDIPHARMA A/S	IS
Contalgin Uno 200 mg hörð forðahylki.	not available	960025	MUNDIPHARMA A/S	IS
Contalgin Uno 30 mg hörð forðahylki.	not available	960020	MUNDIPHARMA A/S	IS
Contalgin Uno 30 mg hörð forðahylki.	not available	960020	MUNDIPHARMA A/S	IS
Contalgin Uno 60 mg hörð forðahylki.	not available	960021	MUNDIPHARMA A/S	IS
Contalgin Uno 60 mg hörð forðahylki.	not available	960021	MUNDIPHARMA A/S	IS
Contalgin Uno 90 mg hörð forðahylki.	not available	960022	MUNDIPHARMA A/S	IS
Contalgin Uno 90 mg hörð forðahylki.	not available	960022	MUNDIPHARMA A/S	IS
Contalgin Uno, hårde depotkapsler	not available	17797	MUNDIPHARMA 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
Contalgin Uno, hårde depotkapsler	not available	17798	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17799	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17800	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17801	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17802	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17797	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17798	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17799	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17800	MUNDIPHARMA 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
Contalgin Uno, hårde depotkapsler	not available	17801	MUNDIPHARMA A/S	DK
Contalgin Uno, hårde depotkapsler	not available	17802	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	15421	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	10801	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	11021	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	13515	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	11641	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	14491	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	15421	MUNDIPHARMA 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
Contalgin, depottabletter	not available	10801	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	11021	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	13515	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	11641	MUNDIPHARMA A/S	DK
Contalgin, depottabletter	not available	14491	MUNDIPHARMA A/S	DK
Cyclimorph 10 Solution for Injection	not available	PA 1142/3/1	AMDIPHARM LIMITED	IE
Cyclimorph 15 Solution for Injection	not available	PA 1142/3/2	AMDIPHARM LIMITED	IE
Cyclimorph-10 Injection 1 ml Ampoules	not available	PL 20072/0007	AMDIPHARM UK LIMITED	XI
Cyclimorph-15 Injection	not available	PL 20072/0008	AMDIPHARM UK LIMITED	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
Cyclizine Tartrate 50mg/ml and Morphine Tartrate 10mg/ml Injection	not available	PL 20072/0007	AMDIPHARM UK LIMITED	XI
Cyclizine Tartrate 50mg/ml and Morphine Tartrate 15mg/ml Injection	not available	PL 20072/0008	AMDIPHARM UK LIMITED	XI
Dolcontin 10 mg depottabletter	not available	7070	MUNDIPHARMA AS	NO
Dolcontin 10 mg depottabletter	not available	7070	MUNDIPHARMA AS	NO
Dolcontin 10 mg depottabletter	not available	10847	MUNDIPHARMA AB	SE
Dolcontin 100 mg depottabletter	not available	7072	MUNDIPHARMA AS	NO
Dolcontin 100 mg depottabletter	not available	7072	MUNDIPHARMA AS	NO
Dolcontin 100 mg depottabletter	not available	10849	MUNDIPHARMA 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
Dolcontin 200 mg depottabletter	not available	7912	MUNDIPHARMA AS	NO
Dolcontin 200 mg depottabletter	not available	7912	MUNDIPHARMA AS	NO
Dolcontin 200 mg depottabletter	not available	11694	MUNDIPHARMA AB	SE
Dolcontin 30 mg depottabletter	not available	7071	MUNDIPHARMA AS	NO
Dolcontin 30 mg depottabletter	not available	7071	MUNDIPHARMA AS	NO
Dolcontin 30 mg depottabletter	not available	10848	MUNDIPHARMA AB	SE
Dolcontin 5 mg depottabletter	not available	7940	MUNDIPHARMA AS	NO
Dolcontin 5 mg depottabletter	not available	7940	MUNDIPHARMA AS	NO
Dolcontin 5 mg depottabletter	not available	11832	MUNDIPHARMA 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
Dolcontin 60 mg depottabletter	not available	7561	MUNDIPHARMA AS	NO
Dolcontin 60 mg depottabletter	not available	7561	MUNDIPHARMA AS	NO
Dolcontin 60 mg depottabletter	not available	11076	MUNDIPHARMA AB	SE
Dolcontin Unotard 120 mg depotkapseli, kova	not available	12864	MUNDIPHARMA AB	SE
Dolcontin Unotard 150 mg depotkapseli, kova	not available	12865	MUNDIPHARMA AB	SE
Dolcontin Unotard 200 mg depotkapseli, kova	not available	12866	MUNDIPHARMA AB	SE
Dolcontin Unotard 30 mg depotkapseli, kova	not available	12861	MUNDIPHARMA AB	SE
Dolcontin Unotard 60 mg depotkapseli, kova	not available	12862	MUNDIPHARMA AB	SE
Dolcontin Unotard 90 mg depotkapseli, kova	not available	12863	MUNDIPHARMA 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
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/001	KALCEKS	LT
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/002	KALCEKS	LT
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/003	KALCEKS	LT
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/004	KALCEKS	LT
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/005	KALCEKS	LT
Maracex 20 mg/ml injekcinis ar infuzinis tirpalas	EE/H/0246/001	LT/1/18/4219/006	KALCEKS	LT
Maracex 20 mg/ml injekční/infuzní roztok	EE/H/0246/001	65/525/18-C	KALCEKS	CZ
Maracex 20 mg/ml injekčný/infúzny roztok	EE/H/0246/001	65/0162/19-S	KALCEKS	SK
Maracex 20 mg/ml oplossing voor injectie/infusie	EE/H/0246/001	RVG 121082	KALCEKS	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
Maracex 20 mg/ml šķīdums injekcijām/infūzijām	EE/H/0246/001	18-0066	KALCEKS	LV
Maracex 20 mg/ml solución inyectable y para perfusión	EE/H/0246/001	84129	KALCEKS	ES
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/01	KALCEKS	RO
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/02	KALCEKS	RO
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/03	KALCEKS	RO
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/04	KALCEKS	RO
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/05	KALCEKS	RO
Maracex 20 mg/ml soluție injectabilă/perfuzabilă	EE/H/0246/001	10808/2018/06	KALCEKS	RO
Maracex 20 mg/ml solution for injection/infusion	EE/H/0246/001	MA1207/00101	KALCEKS	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Maracex 20 mg/ml solution for injection/infusion	EE/H/0246/001	MA1207/00102	KALCEKS	MT
Maracex 20 mg/ml solution for injection/infusion	EE/H/0246/001	MA1207/00103	KALCEKS	MT
Maracex 20 mg/ml süste-/infusioonilahus	EE/H/0246/001	962518	KALCEKS	EE
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/01	KALCEKS	HU
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/02	KALCEKS	HU
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/03	KALCEKS	HU
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/04	KALCEKS	HU
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/05	KALCEKS	HU
Maracex 20 mg/ml oldatos injekció vagy infúzió	EE/H/0246/001	OGYI-T-23989/06	KALCEKS	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
Maracex, 20 mg/ml, roztwór do wstrzykiwań/do infuzji	EE/H/0246/001	24750	KALCEKS	PL
M-long 10 mg, Hartkapseln, retardiert	not available	30173.00.00	ETHYPHARM	DE
M-long 100 mg, Hartkapseln, retardiert	not available	30173.03.00	ETHYPHARM	DE
M-long 30 mg, Hartkapseln, retardiert	not available	30173.01.00	ETHYPHARM	DE
M-long 60 mg, Hartkapseln, retardiert	not available	30173.02.00	ETHYPHARM	DE
Morapid 10 mg Filmtabletten	not available	1-20439	MUNDIPHARMA GES.M.B.H	AT
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
Morapid 20 mg Filmtabletten	not available	1-20444	MUNDIPHARMA GES.M.B.H	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
Morfin Alternova 5 mg tabletter	not available	59121	ALTERNOVA A/S	SE
Morfina B. Braun 1 mg/ml solución inyectable	not available	82748	B.BRAUN MEDICAL, S.A.	ES
Morfina B. Braun 1 mg/ml solución inyectable	not available	82748	B.BRAUN MEDICAL, S.A.	ES
Morfina B. Braun 20 mg/ml solución inyectable en vial	not available	82749	B.BRAUN MEDICAL, S.A.	ES
Morfina B. Braun 20 mg/ml solución inyectable en vial	not available	82749	B.BRAUN MEDICAL, S.A.	ES
Morfina B. Braun 40 mg/ml solución inyectable	not available	82750	B.BRAUN MEDICAL, S.A.	ES
Morfina SERRA 10 mg/ml solución inyectable	not available	37473	LABORATORIOS SERRA PAMIES S.A.	ES
Morfina SERRA 20 mg/ml solución inyectable	not available	37472	LABORATORIOS SERRA PAMIES S.A.	ES
Morfina Solfato SUN 1 mg/ml - soluzione per infusione in siringa preriempita	DE/H/6814/001	049154014	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	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
Morfina Solfato SUN 1 mg/ml - soluzione per infusione in siringa preriempita	DE/H/6814/001	049154026	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Morfina Solfato SUN 2 mg/ml - soluzione per infusione in siringa preriempita	DE/H/6814/002	049154040	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Morfina Solfato SUN 2 mg/ml - soluzione per infusione in siringa preriempita	DE/H/6814/002	049154038	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Morfina Sun 1 mg/ml solución para perfusión en jeringa precargada	DE/H/6814/001	86587	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Morfina Sun 2 mg/ml solución para perfusión en jeringa precargada	DE/H/6814/002	86588	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Morphgesic SR 10 mg Tablets	not available	PL 20072/0231	AMDIPHARM UK LIMITED	XI
Morphgesic SR 100 mg Tablets	not available	PL 20072/0234	AMDIPHARM UK LIMITED	XI
Morphgesic SR 30 mg Tablets	not available	PL 20072/0232	AMDIPHARM UK LIMITED	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
Morphgesic SR 60 mg Tablets	not available	PL 20072/0233	AMDIPHARM UK LIMITED	XI
Morphin Aristo 20 mg Retardtabletten	not available	94198.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Morphin Aristo 20 mg Retardtabletten	not available	94198.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Morphin Aristo 45 mg Retardtabletten	not available	94199.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Morphin Aristo 45 mg Retardtabletten	not available	94199.00.00	ARISTO PHARMA GMBH (ART 57)	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® 10 mg, Injektionslösung	not available	6108826.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Morphin Merck® 100 mg, Infusionslösung	not available	6108826.02.00	MERCK HEALTHCARE GERMANY 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
Morphin Merck® 20 mg Injektionslösung	not available	6108826.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Morphin Merck® Tropfen 0,5%	not available	31264.00.00	MERCK HEALTHCARE GERMANY GMBH	DE
Morphin Merck® Tropfen 2%	not available	31264.01.00	MERCK HEALTHCARE GERMANY GMBH	DE
Morphine "Orion", injektionsvæske, opløsning	NO/H/0258/002	57511	ORION CORPORATION	DK
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 075 5 3	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 077 8 2	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 074 9 2	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 365 6 3	COOPERATION PHARMACEUTIQUE FRANCAISE	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
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 366 2 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 076 1 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 565 882 7 5	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 939 2 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 101 6 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 102 2 5	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 940 0 6	COOPERATION PHARMACEUTIQUE FRANCAISE	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
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 941 7 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 103 9 3	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 369 104 5 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 942 3 5	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) COOPER 10 mg/ml, solution injectable	not available	34009 566 944 6 4	COOPERATION PHARMACEUTIQUE FRANCAISE	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 10 mg/ml, solution injectable	not available	34009 340 213 0 5	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

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
MORPHINE (CHLORHYDRATE) LAVOISIER 10 mg/ml, solution injectable	not available	34009 342 912 3 4	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 10 mg/ml, solution injectable	not available	34009 560 276 1 3	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 20 mg/ml, solution injectable	not available	34009 344 109 3 2	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 20 mg/ml, solution injectable	not available	34009 553 533 2 4	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 20 mg/ml, solution injectable	not available	34009 342 914 6 3	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (CHLORHYDRATE) LAVOISIER 20 mg/ml, solution injectable	not available	34009 560 278 4 2	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (SULFATE) LAVOISIER 1 mg/ml, solution injectable	not available	34009 356 206 9 9	LABORATOIRES CHAIX ET DU MARAIS	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
MORPHINE (SULFATE) LAVOISIER 50 mg/ml, solution injectable	not available	34009 356 199 2 1	LABORATOIRES CHAIX ET DU MARAIS	FR
MORPHINE (SULFATE) LAVOISIER 50 mg/ml, solution injectable	not available	34009 356 201 7 0	LABORATOIRES CHAIX ET DU MARAIS	FR
Morphine 1 mg/ml solution for infusion in pre-filled syringe	DE/H/6814/001	PL 31750/0195	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	XI
Morphine 2 mg/ml solution for infusion in pre-filled syringe	DE/H/6814/002	PL 31750/0196	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	XI
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 mg/ml injektionsvätska, lösning	NO/H/0258/002	34067	ORION CORPORATION	FI
Morphine Sulfate Injection 10 mg in 1 ml	not available	PL 17907/0597	BRISTOL LABORATORIES LIMITED	XI
Morphine Sulfate Injection 15 mg in 1 ml	not available	PL 17907/0598	BRISTOL LABORATORIES LIMITED	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
Morphine Sulfate Injection 30 mg in 1 ml	not available	PL 17907/0599	BRISTOL LABORATORIES LIMITED	XI
Morphine Sulfate Injection BP Minijet™	not available	PL 03265/0037	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 10 mg prolonged-release tablets	not available	19746	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 10 mg prolonged-release tablets	not available	19746	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 100 mg prolonged-release tablets	not available	19748	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 100	not available	19748	MUNDIPHARMA PHARMACEUTICALS LTD	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
mg prolonged-release tablets				
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 30 mg prolonged-release tablets	not available	19745	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 30 mg prolonged-release tablets	not available	19745	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 60 mg prolonged-release tablets	not available	19747	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Morphine sulfate prolonged-release tablets (Q12h) MST® CONTINUS® 60 mg prolonged-release tablets	not available	19747	MUNDIPHARMA PHARMACEUTICALS LTD	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
Morphine sulfate Sevredol 10 mg film-coated tablets	not available	19750	MUNDIPHARMA GES.M.B.H	CY
Morphine sulfate Sevredol 10 mg film-coated tablets	not available	19750	MUNDIPHARMA GES.M.B.H	CY
Morphine sulfate Sevredol 20 mg film-coated tablets	not available	19749	MUNDIPHARMA GES.M.B.H	CY
Morphine sulfate Sevredol 20 mg film-coated tablets	not available	19749	MUNDIPHARMA GES.M.B.H	CY
Morphine Sulphate 10mg/ml Solution for Injection	not available	PA 73/20/1	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 1mg/5ml Solution for Injection	not available	PA 73/20/3	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 30 mg/ml Solution for Injection	not available	PA 73/20/4	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Sulphate 60 mg/ml Solution for Injection	not available	PA 73/20/8	MERCURY PHARMACEUTICALS (IRELAND) LTD.	IE
Morphine Unimedic 1 mg/ml injektionsvätska, lösning	SE/H/1657/001	56117	UNIMEDIC PHARMA 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
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 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 libération prolongée	not available	34009 555 597 8 8	MUNDIPHARMA SAS	FR
MOSCONTIN 100 mg, comprimé enrobé à libération prolongée	not available	34009 328 700 2 8	MUNDIPHARMA SAS	FR
MOSCONTIN 100 mg, comprimé enrobé à libération prolongée	not available	34009 555 597 8 8	MUNDIPHARMA SAS	FR
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

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
MOSCONTIN 30 mg, comprimé enrobé à libération prolongée	not available	34009 328 698 8 6	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 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 L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 343 006-6 0	MUNDIPHARMA SAS	FR
MOSCONTIN L.P. 200 mg, comprimé pelliculé à libération prolongée	not available	34009 343 006-6 0	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
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 558 273 9 9	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	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	RVG 11205	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 10 mg, tabletten met verlengde afgifte	not available	RVG 11205	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 100 mg, comprimés à libération prolongée	not available	2005078273	MUNDIPHARMA COMM VA	LU
MS Contin 100 mg, tabletten met verlengde afgifte	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
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 100 mg, tabletten met verlengde afgifte	not available	RVG 11208	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 15 mg, tabletten met verlengde afgifte	not available	RVG 16674	MUNDIPHARMA PHARMACEUTICALS BV	NL
MS Contin 15 mg, tabletten met verlengde afgifte	not available	RVG 16674	MUNDIPHARMA PHARMACEUTICALS BV	NL
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	BE 134041	MUNDIPHARMA COMM VA	BE
MS Contin 30 mg, tabletten met verlengde afgifte	not available	RVG 11206	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, tabletten met verlengde afgifte	not available	RVG 11206	MUNDIPHARMA PHARMACEUTICALS BV	NL
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	1495/05078272	MUNDIPHARMA COMM VA	LU
MS Contin 60 mg, tabletten met verlengde afgifte	not available	BE 134057	MUNDIPHARMA COMM VA	BE
MS Contin 60 mg, tabletten met verlengde afgifte	not available	BE 134057	MUNDIPHARMA COMM VA	BE
MS Contin 60 mg, tabletten met verlengde afgifte	not available	RVG 11207	MUNDIPHARMA PHARMACEUTICALS BV	NL
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

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 Direct 10 mg, omhulde tabletten	not available	BE192482	MUNDIPHARMA COMM VA	BE
MS Direct 10 mg, omhulde tabletten	not available	BE192482	MUNDIPHARMA COMM VA	BE
MST 10 mg Mundipharma Retardtabletten	not available	4034.01.00	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 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 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

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 30 mg Mundipharma Retardtabletten	not available	4034.00.00	MUNDIPHARMA GMBH	DE
MST 60 mg Mundipharma Retardtabletten	not available	4034.02.00	MUNDIPHARMA GMBH	DE
MST 60 mg Mundipharma Retardtabletten	not available	4034.02.00	MUNDIPHARMA GMBH	DE
MST CONTINUS 10 mg comprimidos de liberación prolongada	not available	57.898	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 10 mg comprimidos de liberación prolongada	not available	57.898	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 10 mg prolonged- release tablets	not available	PA 1688/004/002	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 tablete s produljenim oslobađanjem	not available	HR-H-863692829	MUNDIPHARMA GES.M.B.H	HR
MST Continus 10 mg tablete s produljenim oslobađanjem	not available	HR-H-863692829	MUNDIPHARMA GES.M.B.H	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
MST CONTINUS 100 mg comprimidos de liberación prolongada	not available	57.900	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 100 mg comprimidos de liberación prolongada	not available	57.900	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 100 mg prolonged-release tablets	not available	PA 1688/004/006	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	MUNDIPHARMA GES.M.B.H	HR
MST Continus 100 mg tablete s produljenim oslobađanjem	not available	HR-H-448781299	MUNDIPHARMA GES.M.B.H	HR
MST CONTINUS 15 mg comprimidos de liberación prolongada	not available	61.081	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 15 mg comprimidos de liberación prolongada	not available	61.081	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 15 mg Prolonged-release tablets	not available	PA 1688/004/003	MUNDIPHARMA PHARMACEUTICALS 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
MST CONTINUS 200 mg comprimidos de liberación prolongada	not available	61.909	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 30 mg comprimidos de liberación prolongada	not available	57.897	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 30 mg Prolonged-release tablets	not available	PA 1688/004/004	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	MUNDIPHARMA GES.M.B.H	HR
MST Continus 30 mg tablete s produljenim oslobađanjem	not available	HR-H-113395944	MUNDIPHARMA GES.M.B.H	HR
MST Continus 30 mg tablety s riadeným uvoľňovaním	not available	65/0259/13-S	MUNDIPHARMA GES.M.B.H	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
MST Continus 30 mg tablety s riadeným uvoľňovaním	not available	65/0259/13-S	MUNDIPHARMA GES.M.B.H	SK
MST CONTINUS 5 mg comprimidos de liberación prolongada	not available	61.080	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 5 mg comprimidos de liberación prolongada	not available	61.080	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 5 mg prolonged-release tablets	not available	PA 1688/004/001	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
MST CONTINUS 60 mg comprimidos de liberación prolongada	not available	57.899	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 60 mg comprimidos de liberación prolongada	not available	57.899	MUNDIPHARMA PHARMACEUTICALS SL	ES
MST CONTINUS 60 mg prolonged-release tablets	not available	PA 1688/004/005	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	MUNDIPHARMA GES.M.B.H	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
MST Continus 60 mg tablete s produljenim oslobađanjem	not available	HR-H-734333555	MUNDIPHARMA GES.M.B.H	HR
MST Continus 60 mg tablety s riadeným uvoľňovaním	not available	65/0257/13-S	MUNDIPHARMA GES.M.B.H	SK
MST Continus 60 mg tablety s riadeným uvoľňovaním	not available	65/0258/13-S	MUNDIPHARMA GES.M.B.H	SK
MST Continus 60 mg tablety s riadeným uvoľňovaním	not available	65/0257/13-S	MUNDIPHARMA GES.M.B.H	SK
MST Continus 60 mg tablety s riadeným uvoľňovaním	not available	65/0258/13-S	MUNDIPHARMA GES.M.B.H	SK
MST Continus, 10 mg tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4762	MUNDIPHARMA A/S	PL
MST Continus, 10 mg tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4762	MUNDIPHARMA A/S	PL
MST Continus, 100 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4765	MUNDIPHARMA 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
MST Continus, 100 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4765	MUNDIPHARMA A/S	PL
MST Continus, 200 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4766	MUNDIPHARMA A/S	PL
MST Continus, 200 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4766	MUNDIPHARMA A/S	PL
MST Continus, 30 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4763	MUNDIPHARMA A/S	PL
MST Continus, 30 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4763	MUNDIPHARMA A/S	PL
MST Continus, 60 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4764	MUNDIPHARMA A/S	PL
MST Continus, 60 mg Tabletki powlekane o zmodyfikowanym uwalnianiu	not available	4764	MUNDIPHARMA 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
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 20mg/ml	not available	PA 2256/004/002	GLENWOOD GMBH	IE
Oramorph Oral Solution 10 mg/5 ml	not available	PA 2256/004/001	GLENWOOD GMBH	IE
Rhotard Morphine SR 10 mg Tablets	not available	PL 20072/0231	AMDIPHARM UK LIMITED	XI
Rhotard Morphine SR 30 mg Tablets	not available	PL 20072/0232	AMDIPHARM UK LIMITED	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
Rhotard Morphine SR 60 mg Tablets	not available	PL 20072/0233	AMDIPHARM UK LIMITED	XI
Rhotard Morphine Tablets SR 100mg.	not available	PL 20072/0234	AMDIPHARM UK LIMITED	XI
SEVREDOL 10 mg comprimate filmate	not available	9264/2016/01	MUNDIPHARMA GES.M.B.H	RO
Sevredol 10 mg comprimidos recubiertos con película.	not available	59.656	MUNDIPHARMA PHARMACEUTICALS SL	ES
Sevredol 10 mg film-coated tablets	not available	PA 1688/009/001	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Sevredol 10 mg filmom obalené tablety	not available	65/0302/98-S	MUNDIPHARMA GES.M.B.H	SK
Sevredol 10 mg filmom obalené tablety	not available	65/0302/98-S	MUNDIPHARMA GES.M.B.H	SK
Sevredol 10 mg filmom obložene tablete	not available	HR-H-915182609	MUNDIPHARMA GES.M.B.H	HR
Sevredol 10 mg filmom obložene tablete	not available	HR-H-915182609	MUNDIPHARMA GES.M.B.H	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
Sevredol 10 mg filmsko obložene tablete	not available	H/01/01416/001	MUNDIPHARMA GES.M.B.H	SI
Sevredol 10 mg filmsko obložene tablete	not available	H/01/01416/002	MUNDIPHARMA GES.M.B.H	SI
Sevredol 10 mg Filmtabletten	not available	25932.00.00	MUNDIPHARMA GMBH	DE
Sevredol 10 mg Filmtabletten	not available	25932.00.00	MUNDIPHARMA GMBH	DE
SEVREDOL 10 mg, comprimé pelliculé sécable	not available	34009 334 799 7 8	MUNDIPHARMA SAS	FR
SEVREDOL 10 mg, comprimé pelliculé sécable	not available	34009 334 799 7 8	MUNDIPHARMA SAS	FR
SEVREDOL 20 mg comprimate filmate	not available	9265/2016/01-03	MUNDIPHARMA GES.M.B.H	RO
SEVREDOL 20 mg comprimidos recubiertos con película	not available	59.655	MUNDIPHARMA PHARMACEUTICALS SL	ES
Sevredol 20 mg film-coated tablets	not available	PA 1688/009/002	MUNDIPHARMA PHARMACEUTICALS 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
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmom obložene tablete	not available	HR-H-932315368	MUNDIPHARMA GES.M.B.H	HR
Sevredol 20 mg filmsko obložene tablete	not available	H/01/01416/003	MUNDIPHARMA GES.M.B.H	SI
Sevredol 20 mg filmsko obložene tablete	not available	H/01/01416/004	MUNDIPHARMA GES.M.B.H	SI
Sevredol 20 mg Filmtabletten	not available	25932.01.00	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
Sevredol 20 mg Filmtabletten	not available	25932.01.00	MUNDIPHARMA GMBH	DE
SEVREDOL 20 mg, comprimé pelliculé sécable	not available	34009 334 800 5 9	MUNDIPHARMA SAS	FR
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 20mg	not available	65/300/99-C	MUNDIPHARMA GES.M.B.H	CZ
Sevredol 50 mg film-coated tablets	not available	PA 1688/009/003	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Sevredol, 20 mg, tabletki powlekane	not available	8827	MUNDIPHARMA A/S	PL
Sevredol, 20 mg, tabletki powlekane	not available	8827	MUNDIPHARMA A/S	PL
SKENAN L.P. 10 mg, microgranules à libération prolongée en gélule	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
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. 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 100 mg Hartkapseln, retardiert	not available	31512.01.00	MUNDIPHARMA GMBH	DE
Substitol 120 mg trde kapsule s podaljšanim sproščanjem	not available	H/04/01467/001	MUNDIPHARMA GESELLSCHAFT M.B.H.	SI
Substitol 200 mg Hartkapseln, retardiert	not available	31512.00.00	MUNDIPHARMA GMBH	DE
Substitol 200 mg trde kapsule s podaljšanim sproščanjem	not available	H/04/01467/002	MUNDIPHARMA GESELLSCHAFT M.B.H.	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
Substitol 30 mg Hartkapseln, retardiert	not available	31512.04.00	MUNDIPHARMA GMBH	DE
Substitol 60 mg Hartkapseln, retardiert	not available	31512.02.00	MUNDIPHARMA GMBH	DE
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
TWICE 10 mg Capsule rigide a rilascio prolungato	not available	033484015	ETHYPHARM	IT
TWICE 100 mg Capsule rigide a rilascio prolungato	not available	033484041	ETHYPHARM	IT
TWICE 30 mg Capsule rigide a rilascio prolungato	not available	033484027	ETHYPHARM	IT
TWICE 60 mg Capsule rigide a rilascio prolungato	not available	033484039	ETHYPHARM	IT
ZOMORPH 100mg capsules	not available	PL 06934/0185	ETHYPHARM	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
ZOMORPH 10mg capsules	not available	PL 06934/0182	ETHYPHARM	XI
ZOMORPH 200mg capsules	not available	PL 06934/0186	ETHYPHARM	XI
ZOMORPH 30mg capsules	not available	PL 06934/0183	ETHYPHARM	XI
ZOMORPH 60mg capsules	not available	PL 06934/0184	ETHYPHARM	XI
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