09 November 2023
EMA/577152/2023
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: oxycodone

Procedure no.: PSUSA/00002254/202304

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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
Carenoxal 30 mg Retardtabletten	DE/H/3638/005	87923.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 60 mg Retardtabletten	DE/H/3638/007	87925.00.00	ZENTIVA PHARMA GMBH	DE
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI

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Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	H/09/00405/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Enoxy Depot 10 mg depottabletter	SE/H/2082/001	60988	TEVA B.V	SE
Enoxy Depot 20 mg depottabletter	SE/H/2082/002	60989	TEVA B.V	SE
Enoxy Depot 40 mg depottabletter	SE/H/2082/003	60990	TEVA B.V	SE
Enoxy Depot 80 mg depottabletter	SE/H/2082/004	60991	TEVA B.V	SE
Ixyldone 15 mg prolonged-release tablets	NA	PL 20117/0307	MORNINGSIDE HEALTHCARE LTD	XI
Ixyldone 30 mg prolonged-release tablets	NA	PL 20117/0309	MORNINGSIDE HEALTHCARE LTD	XI
Ixyldone 60 mg prolonged-release tablets	NA	PL 20117/0311	MORNINGSIDE HEALTHCARE LTD	XI
Onexila XL 10 mg prolonged-release tablets	UK/H/6610/001	PL 33155/0042	RIVOPHARM UK LTD	XI
Onexila XL 20 mg prolonged-release tablets	UK/H/6610/002	PL 33155/0044	RIVOPHARM UK LTD	XI
Onexila XL 40 mg prolonged-release tablets	UK/H/6610/003	PL 33155/0045	RIVOPHARM UK LTD	XI
Onexila XL 80 mg prolonged-release tablets	UK/H/6610/004	PL 33155/0046	RIVOPHARM UK LTD	XI
Oxycan® uno 10 mg Retardtabletten	DE/H/3655/001	88009.00.00	HORMOSAN PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycan® uno 20 mg Retardtabletten	DE/H/3655/002	88010.00.00	HORMOSAN PHARMA GMBH	DE
Oxycan® uno 40 mg Retardtabletten	DE/H/3655/003	88011.00.00	HORMOSAN PHARMA GMBH	DE
Oxycan® uno 80 mg Retardtabletten	DE/H/3655/004	88012.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon HCl retard CF 15 mg, tabletten met verlengde afgifte	DE/H/3641/003	RVG 112600	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 30 mg, tabletten met verlengde afgifte	DE/H/3641/005	RVG 112602	CENTRAFARM B.V.	NL
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 858 3 2	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 867 2 3	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 868 9 1	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 860 8 2	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 872 6 3	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 864 3 3	BIOGARAN	FR
OXYCODONE BIOGARAN	FR/H/0621/003	34009 277 869 5 2	BIOGARAN	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
LP 15 mg, comprimé pelliculé à libération prolongée				
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 862 0 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 870 3 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 861 4 3	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 863 7 2	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003	34009 277 866 6 2	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 885 0 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 895 6 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 889 6 3	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé	FR/H/0621/005	34009 277 886 7 3	BIOGARAN	FR

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pelliculé à libération prolongée				
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 890 4 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 887 3 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 892 7 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 896 2 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 891 0 6	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 893 3 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789854	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005	34009 277 897 9 3	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération	FR/H/0621/007	34009 277 912 8 4	BIOGARAN	FR

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prolongée				
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 919 2 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 915 7 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 924 6 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 921 7 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 916 3 5	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 920 0 7	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 914 0 6	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 925 2 6	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 913 4 5	BIOGARAN	FR

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OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 918 6 4	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007	34009 277 922 3 6	BIOGARAN	FR
OXYCODONE EG LP 15 mg, comprimé pelliculé à libération prolongée	DE/H/3641/003	NL42844	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
OXYCODONE EG LP 30 mg, comprimé pelliculé à libération prolongée	DE/H/3641/005	NL42846	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
OXYCODONE EG LP 60 mg, comprimé pelliculé à libération prolongée	DE/H/3641/007	NL42848	EG LABO LABORATOIRES EUROGENERICS - DO NOT USE	FR
Oxycodone G.L. Pharma 10 mg prolonged-release tablets	not available	PL 21597/0022	G.L. PHARMA GMBH	XI
Oxycodone G.L. Pharma 20 mg prolonged-release tablets	not available	PL 21597/0023	G.L. PHARMA GMBH	XI
Oxycodone G.L. Pharma 40 mg prolonged-release tablets	not available	PL 21597/0024	G.L. PHARMA GMBH	XI
Oxycodone G.L. Pharma 5 mg prolonged-release tablets	not available	PL 21597/0021	G.L. PHARMA GMBH	XI
Oxycodone G.L. Pharma 80 mg prolonged-release tablets	not available	PL 21597/0025	G.L. PHARMA GMBH	XI
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 301 637 9 5	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour	not available	34009 301 638 0 1	LABORATOIRE RENAUDIN	FR

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perfusion				
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 550 601 1 6	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 550 601 2 3	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 301 905 1 7	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 550 601 4 7	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 301 905 3 1	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 550 601 6 1	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 10 mg/mL, solution pour perfusion	not available	34009 301 905 4 8	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 10 mg/mL, solution pour perfusion	not available	34009 550 600 4 8	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 10 mg/mL, solution pour perfusion	not available	34009 301 905 5 5	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 10 mg/mL, solution pour perfusion	not available	34009 550 600 6 2	LABORATOIRE RENAUDIN	FR
Oxycodon-HCl AbZ 30 mg Retardtabletten	DE/H/3495/001	86471.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 60 mg Retardtabletten	DE/H/3495/002	86472.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AL 15 mg	DE/H/3860/003	89918.00.00	ALIUD PHARMA GMBH	DE

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Retardtabletten				
Oxycodon-HCl AL 30 mg Retardtabletten	DE/H/3860/005	89920.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 60 mg Retardtabletten	DE/H/3860/007	89922.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl beta 1 × täglich 10 mg Retardtabletten	DE/H/3654/001	88005.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 1 x täglich 20 mg Retardtabletten	DE/H/3654/002	88006.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 1 x täglich 40 mg Retardtabletten	DE/H/3654/003	88007.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 1 x täglich 80 mg Retardtabletten	DE/H/3654/004	88008.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 30 mg Retardtabletten	DE/H/0667/004	69806.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 60 mg Retardtabletten	DE/H/0667/005	69807.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Krugmann 10 mg Retardtabletten	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	53004.00.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann 20 mg Retardtabletten	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	53004.01.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann 40 mg Retardtabletten	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	53004.02.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann 5 mg Retardtabletten	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	53004.04.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann 80 mg Retardtabletten	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	53004.03.00	KRUGMANN GMBH (GERMANY)	DE

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Oxycodon-HCl Krugmann akut 10 mg Hartkapseln	not available	58373.00.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann akut 20 mg Hartkapseln	not available	58374.00.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl Krugmann akut 5 mg Hartkapseln	not available	58372.00.00	KRUGMANN GMBH (GERMANY)	DE
Oxycodon-HCl ratiopharm® Uno 10 mg Retardtabletten	SE/H/2082/001	140880	TEVA B.V	AT
Oxycodon-HCl ratiopharm® Uno 20 mg Retardtabletten	SE/H/2082/002	140881	TEVA B.V	AT
Oxycodon-HCl ratiopharm® Uno 40 mg Retardtabletten	SE/H/2082/003	140882	TEVA B.V	AT
Oxycodon-HCl ratiopharm® Uno 80 mg Retardtabletten	SE/H/2082/004	140883	TEVA B.V	AT
Oxycodon-HCL Sandoz 5 mg Retardtabletten	DE/H/1154/001	69179.00.00	HEXAL AG	DE
Oxycodon-HCl Zentiva 30 mg Retardtabletten	DE/H/3476/001	86290.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl Zentiva 60 mg Retardtabletten	DE/H/3476/002	86291.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl-ratiopharm® 30 mg Retardtabletten	DE/H/3475/001	86288.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 60 mg Retardtabletten	DE/H/3475/002	86289.00.00	RATIOPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 15 mg Retardtabletten	DE/H/3641/003	87945.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 30 mg Retardtabletten	DE/H/3641/005	87947.00.00	STADAPHARM GMBH	DE

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Oxycodonhydrochlorid STADA® 60 mg Retardtabletten	DE/H/3641/007	87949.00.00	STADAPHARM GMBH	DE
Oxyconica 15 mg Retardtabletten	DE/H/3644/003	87969.00.00	ACINO AG	DE
Oxyconica 30 mg Retardtabletten	DE/H/3644/005	87971.00.00	ACINO AG	DE
Oxyconica 60 mg Retardtabletten	DE/H/3644/007	87973.00.00	ACINO AG	DE
Oxyconoica 15 mg Retardtabletten	DE/H/3645/003	87977.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Oxyconoica 30 mg Retardtabletten	DE/H/3645/005	87979.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Oxyconoica 60 mg Retardtabletten	DE/H/3645/007	87981.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
OxyContin 10 mg comprimidos de liberación prolongada	IE/H/0112/001	63446	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 10 mg depottabletter	not available	94-2768	MUNDIPHARMA AS	NO
OxyContin 10 mg prolonged release tablets	not available	18608	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 10 mg prolonged release tablets	IE/H/0112/001-004/MR	PA 1688/005/002	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin 10 mg retard filmtabletta	not available	OGYI-T-7166/03	MUNDIPHARMA GES.M.B.H	HU
OxyContin 10 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/001	MUNDIPHARMA GES.M.B.H	SI
OxyContin 10 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/002	MUNDIPHARMA GES.M.B.H	SI
OxyContin 10 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/003	MUNDIPHARMA GES.M.B.H	SI
OxyContin 10 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-01	MUNDIPHARMA GES.M.B.H	HR
OxyContin 10 mg tablete s	DE/H/1026/002;	HR-H-118192252-02	MUNDIPHARMA GES.M.B.H	HR

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produljenim oslobađanjem	DE/H/1026/003; DE/H/1026/004; DE/H/			
OxyContin 10 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-03	MUNDIPHARMA GES.M.B.H	HR
OxyContin 10 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-04	MUNDIPHARMA GES.M.B.H	HR
OxyContin 10 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-05	MUNDIPHARMA GES.M.B.H	HR
OxyContin 10 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-06	MUNDIPHARMA GES.M.B.H	HR
OXYCONTIN 10 mg, comprimés à libération prolongée	DE/H/0366/001-003/MR	BE253197	MUNDIPHARMA COMM VA	BE
OXYCONTIN 10 mg, comprimés à libération prolongée	DE/H/0366/001	2009010136	MUNDIPHARMA COMM VA	LU
OxyContin 10 mg, tabletten met verlengde afgifte	DE/H/0366/001-003/MR	BE253197	MUNDIPHARMA COMM VA	BE
OxyContin 10 mg, tabletten met verlengde afgifte	not available	RVG 22107	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 10 mg, toimeainet prolongeeritult vabastavad tabletid	not available	300500	MUNDIPHARMA GES.M.B.H	EE
OxyContin 10mg	not available	65/257/00-C	MUNDIPHARMA GES.M.B.H	CZ
OxyContin 120 mg depottabletter	not available	07-5181	MUNDIPHARMA AS	NO
OxyContin 120 mg, tabletten met verlengde afgifte	not available	RVG 100824	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 15 mg	not available	07-4847	MUNDIPHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
depottabletter				
OxyContin 160 mg, tabletten met verlengde afgifte	not available	RVG 100825	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 20 mg comprimidos de liberación prolongada	IE/H/0112/002	63447	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 20 mg depottabletter	not available	94-2769	MUNDIPHARMA AS	NO
OxyContin 20 mg depottabletter	not available	13831	MUNDIPHARMA AB	SE
OxyContin 20 mg prolonged release tablets	not available	18609	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 20 mg prolonged release tablets	IE/H/0112/001-004/MR	PA 1688/005/003	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin 20 mg retard filmtabletta	not available	OGYI-T-7166/04	MUNDIPHARMA GES.M.B.H	HU
OxyContin 20 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/004	MUNDIPHARMA GES.M.B.H	SI
OxyContin 20 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/005	MUNDIPHARMA GES.M.B.H	SI
OxyContin 20 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/006	MUNDIPHARMA GES.M.B.H	SI
OxyContin 20 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-01	MUNDIPHARMA GES.M.B.H	HR
OxyContin 20 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-02	MUNDIPHARMA GES.M.B.H	HR
OxyContin 20 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-03	MUNDIPHARMA GES.M.B.H	HR
OxyContin 20 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-04	MUNDIPHARMA GES.M.B.H	HR
OxyContin 20 mg tablete s	DE/H/1026/002;	HR-H-188823708-05	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
produljenim oslobađanjem	DE/H/1026/003; DE/H/1026/004; DE/H/			
OxyContin 20 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-06	MUNDIPHARMA GES.M.B.H	HR
OxyContin 20 mg tablety s prodlouženým uvolňováním	not available	65/258/00-C	MUNDIPHARMA GESELLSCHAFT M.B.H.	CZ
OXYCONTIN 20 mg, comprimés à libération prolongée	DE/H/0366/001-003/MR	BE253242	MUNDIPHARMA COMM VA	BE
OXYCONTIN 20 mg, comprimés à libération prolongée	DE/H/0366/002	2009010137	MUNDIPHARMA COMM VA	LU
OxyContin 20 mg, tabletten met verlengde afgifte	DE/H/0366/001-003/MR	BE253242	MUNDIPHARMA COMM VA	BE
OxyContin 20 mg, tabletten met verlengde afgifte	not available	RVG 22108	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 20 mg, toimeainet prolongeeritult vabastavad tabletid	not available	300600	MUNDIPHARMA GES.M.B.H	EE
OxyContin 30 mg depottabletter	not available	07-4848	MUNDIPHARMA AS	NO
OxyContin 30 mg, tabletten met verlengde afgifte	not available	RVG 100820	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 40 mg comprimidos de liberación prolongada	IE/H/0112/003	63448	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 40 mg depottabletter	not available	94-2770	MUNDIPHARMA AS	NO
OxyContin 40 mg depottabletter	not available	13832	MUNDIPHARMA AB	SE
OxyContin 40 mg	not available	18610	MUNDIPHARMA	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
prolonged release tablets			PHARMACEUTICALS LTD	
OxyContin 40 mg prolonged release tablets	IE/H/0112/001-004/MR	PA 1688/005/004	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin 40 mg retard filmtabletta	not available	OGYI-T-7166/05	MUNDIPHARMA GES.M.B.H	HU
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/007	MUNDIPHARMA GES.M.B.H	SI
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/008	MUNDIPHARMA GES.M.B.H	SI
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/009	MUNDIPHARMA GES.M.B.H	SI
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-01	MUNDIPHARMA GES.M.B.H	HR
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-02	MUNDIPHARMA GES.M.B.H	HR
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-03	MUNDIPHARMA GES.M.B.H	HR
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-04	MUNDIPHARMA GES.M.B.H	HR
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-05	MUNDIPHARMA GES.M.B.H	HR
OxyContin 40 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-849997645-06	MUNDIPHARMA GES.M.B.H	HR
OXYCONTIN 40 mg, comprimés à libération prolongée	DE/H/0366/001-003/MR	BE253276	MUNDIPHARMA COMM VA	BE
OXYCONTIN 40 mg, comprimés à libération prolongée	DE/H/0366/003	2009010138	MUNDIPHARMA COMM VA	LU
OxyContin 40 mg,	DE/H/0366/001-003/MR	BE253276	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
tabletten met verlengde afgifte				
OxyContin 40 mg, tabletten met verlengde afgifte	not available	RVG 22109	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 40 mg, toimeainet prolongeeritult vabastavad tabletid	not available	300700	MUNDIPHARMA GES.M.B.H	EE
OxyContin 40mg	not available	65/259/00-C	MUNDIPHARMA GES.M.B.H	CZ
OxyContin 5 mg comprimidos de liberación prolongada	IE/H/0112/005	68605	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 5 mg depottabletter	not available	02-1153	MUNDIPHARMA AS	NO
OxyContin 5 mg depottabletter	not available	18349	MUNDIPHARMA AB	SE
OxyContin 5 mg prolonged release tablets	not available	20346	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 5 mg prolonged release tablets	IE/H/0112/005	PA 1688/005/001	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OXYCONTIN 5 mg, comprimés à libération prolongée	DE/H/0366/005	BE274942	MUNDIPHARMA COMM VA	BE
OXYCONTIN 5 mg, comprimés à libération prolongée	DE/H/0366/005	1495/05100109	MUNDIPHARMA COMM VA	LU
OxyContin 5 mg, tabletten met verlengde afgifte	DE/H/0366/005	BE274942	MUNDIPHARMA COMM VA	BE
OxyContin 5 mg, tabletten met verlengde afgifte	not available	RVG 27536	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 60 mg depottabletter	not available	07-5180	MUNDIPHARMA AS	NO
OxyContin 80 mg comprimidos de liberación prolongada	IE/H/0112/004	63449	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 80 mg	not available	01-2425	MUNDIPHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
depottabletter				
OxyContin 80 mg depottabletter	not available	15982	MUNDIPHARMA AB	SE
OxyContin 80 mg prolonged release tablets	not available	18611	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 80 mg prolonged release tablets	IE/H/0112/001-004/MR	PA 1688/005/005	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin 80 mg retard filmtabletta	not available	OGYI-T-7166/06	MUNDIPHARMA GES.M.B.H	HU
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/010	MUNDIPHARMA GES.M.B.H	SI
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/011	MUNDIPHARMA GES.M.B.H	SI
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/012	MUNDIPHARMA GES.M.B.H	SI
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-01	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-02	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-03	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-04	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-05	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablete s produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-763441426-06	MUNDIPHARMA GES.M.B.H	HR
OxyContin 80 mg tablety s prodlouženým uvolňováním	not available	65/260/00-C	MUNDIPHARMA GESELLSCHAFT M.B.H.	CZ

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
OXYCONTIN 80 mg, comprimés à libération prolongée	DE/H/0366/004	BE253303	MUNDIPHARMA COMM VA	BE
OXYCONTIN 80 mg, comprimés à libération prolongée	DE/H/0366/004	2009010139	MUNDIPHARMA COMM VA	LU
OxyContin 80 mg, tabletten met verlengde afgifte	DE/H/0366/004	BE253303	MUNDIPHARMA COMM VA	BE
OxyContin 80 mg, tabletten met verlengde afgifte	not available	RVG 22110	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 80 mg, toimeainet prolongeeritult vabastavad tabletid	not available	300800	MUNDIPHARMA GES.M.B.H	EE
OxyContin Depot 10 mg depottabletit	not available	11969	MUNDIPHARMA OY	FI
OxyContin Depot 10 mg, forðatöflur	DE/H/0366/001-003/MR	IS/1/02/047/01	MUNDIPHARMA A/S	IS
OxyContin Depot 20 mg depottabletit	not available	11970	MUNDIPHARMA OY	FI
OxyContin Depot 20 mg, forðatöflur	DE/H/0366/002	IS/1/02/047/02	MUNDIPHARMA A/S	IS
OxyContin Depot 40 mg depottabletit	not available	11971	MUNDIPHARMA OY	FI
OxyContin Depot 40 mg, forðatöflur	DE/H/0366/003	IS/1/02/047/03	MUNDIPHARMA A/S	IS
OxyContin Depot 5 mg depottabletit	not available	16771	MUNDIPHARMA OY	FI
OxyContin Depot 5 mg, forðatöflur	DE/H/0366/005	IS/1/05/021/01	MUNDIPHARMA A/S	IS
OxyContin Depot 80 mg depottabletit	not available	13673	MUNDIPHARMA OY	FI
OxyContin Depot 80 mg, forðatöflur	DE/H/0366/004	IS/1/02/047/04	MUNDIPHARMA A/S	IS
OxyContin Depot,	not available	16892	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
depottabletter 10 mg				
OxyContin Depot, depottabletter 120 mg	not available	42030	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 15 mg	not available	41088	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 20 mg	not available	16893	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 30 mg	not available	41089	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 40 mg	not available	16894	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 5 mg	not available	33548	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 60 mg	not available	42029	MUNDIPHARMA A/S	DK
OxyContin Depot, depottabletter 80 mg	not available	30177	MUNDIPHARMA A/S	DK
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 209 0 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 602 2 3	MUNDIPHARMA SAS	FR
OXYCONTIN LP 15 mg, comprimé pelliculé à libération prolongée	not available	34009 384 584 4 2	MUNDIPHARMA SAS	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 215 0 0	MUNDIPHARMA SAS	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à libération prolongée	not available	34009 384 587 3 2	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 222 7 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 5 mg,	not available	34009 366 903 4 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
comprimé pelliculé à libération prolongée				
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 598 5 2	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 294 8 3	MUNDIPHARMA SAS	FR
OxyContin retard 10 mg Filmtabletten	IE/H/0112/001	1-23358	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyContin retard 20 mg Filmtabletten	IE/H/0112/002	1-23359	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyContin retard 40 mg Filmtabletten	IE/H/0112/003	1-23360	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyContin retard 5 mg Filmtabletten	IE/H/0112/005	1-26041	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyContin retard 80 mg Filmtabletten	IE/H/0112/004	1-23361	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyContin, 10 mg depottablett	not available	13830	MUNDIPHARMA AB	SE
OxyContin, 10 mg, tabletki o przedłu□onym uwalnianiu	DE/H/1026/002	14525	MUNDIPHARMA A/S	PL
OxyContin, 20 mg, tabletki o przedłu□onym uwalnianiu	DE/H/1026/003	14526	MUNDIPHARMA A/S	PL
OxyContin, 40 mg, tabletki o przedłu□onym uwalnianiu	DE/H/1026/004	14527	MUNDIPHARMA A/S	PL
OxyContin, 5 mg, tabletki o przedłu□onym uwalnianiu	DE/H/1026/001	14524	MUNDIPHARMA A/S	PL
OxyContin, 80 mg, tabletki o przedłu□onym uwalnianiu	DE/H/1026/005	14528	MUNDIPHARMA A/S	PL
OxyContin® 10 mg	IE/H/0112/001-004/MR	34435038	MUNDIPHARMA	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
compresse a rilascio prolungato			PHARMACEUTICALS SRL	
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435053	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435040	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435026	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435014	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435065	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435077	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435089	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435091	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435115	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435127	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435103	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg	IE/H/0112/001-004/MR	34435141	MUNDIPHARMA	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
compresse a rilascio prolungato			PHARMACEUTICALS SRL	
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435178	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435180	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435154	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435139	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435166	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435255	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	34435279	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	34435267	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	34435293	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	34435281	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435192	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg	IE/H/0112/001-004/MR	34435230	MUNDIPHARMA	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
compresse a rilascio prolungato			PHARMACEUTICALS SRL	
OxyContin® 80 mg compresse a rilascio prolungato	MAA-OXYCODONE HYDROCHLORIDE PROLONGED-RELEASE TABL	034435242	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435216	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435228	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435204	MUNDIPHARMA PHARMACEUTICALS SRL	IT
Oxygesic 10 mg Retardtabletten	DE/H/0366/001-003/MR	41153.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 20 mg Retardtabletten	DE/H/0366/001-003/MR	41153.01.00	MUNDIPHARMA GMBH	DE
Oxygesic 40 mg Retardtabletten	DE/H/0366/001-003/MR	41153.02.00	MUNDIPHARMA GMBH	DE
Oxygesic 5 mg Retardtabletten	DE/H/0366/005	56594.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 80 mg Retardtabletten	DE/H/0366/004	41153.03.00	MUNDIPHARMA GMBH	DE
Oxygesic akut 10 mg Hartkapseln	not available	58368.00.00	MUNDIPHARMA GMBH	DE
Oxygesic akut 20 mg Hartkapseln	not available	58369.00.00	MUNDIPHARMA GMBH	DE
Oxygesic akut 5 mg Hartkapseln	not available	58367.00.00	MUNDIPHARMA GMBH	DE
Oxygesic Dispersa 10 mg Schmelztabletten	DE/H/0366/007-008	68388.00.00	MUNDIPHARMA GMBH	DE
Oxygesic Dispersa 20 mg Schmelztabletten	DE/H/0366/007-008	68389.00.00	MUNDIPHARMA GMBH	DE
Oxygesic Dispersa 5 mg	DE/H/0366/007-008	68387.00.00	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
Schmelztabletten				
Oxygesic infusio 50 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/0366/014/DC	72147.00.00	MUNDIPHARMA GMBH	DE
Oxygesic injekt 10 mg/ml Injektionslösung/Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/0366/015	57886.00.00	MUNDIPHARMA GMBH	DE
OxyNorm 1 mg/ml mikstur, oppløsning	not available	01-9979	MUNDIPHARMA AS	NO
OxyNorm 1 mg/ml oraaliliuos	not available	14807	MUNDIPHARMA OY	FI
OxyNorm 1 mg/ml oral lösning	not available	15792	MUNDIPHARMA AB	SE
OxyNorm 10 mg	not available	20306	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 10 mg cápsulas	IE/H/0139/004	66575	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 10 mg capsules	IE/H/0139/003-005/MR	PA 1688/006/005	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm 10 mg hårda kapslar	not available	16083	MUNDIPHARMA AB	SE
OxyNorm 10 mg harde kapsler	not available	01-7402	MUNDIPHARMA AS	NO
OxyNorm 10 mg kapseli, kova	not available	15321	MUNDIPHARMA OY	FI
OXYNORM 10 mg, gélule	not available	34009 361 898 2 9	MUNDIPHARMA SAS	FR
OXYNORM 10 mg, gélule	not available	34009 361 897 6 8	MUNDIPHARMA SAS	FR
OXYNORM 10 mg, gélule	not available	34009 361 900 7 8	MUNDIPHARMA SAS	FR
OXYNORM 10 mg, gélule	not available	34009 362 421 5 9	MUNDIPHARMA SAS	FR
OXYNORM 10 mg, gélule	not available	34009 362 420 9 8	MUNDIPHARMA SAS	FR
OxyNorm 10 mg, harde capsules	not available	RVG 27510	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm 10 mg/ml injeksjons-	not available	02-216	MUNDIPHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
/infusjonsvæske, oppløsning				
OxyNorm 10 mg/ml injektionsvätska, lösning	not available	18256	MUNDIPHARMA AB	SE
OxyNorm 10 mg/ml mikstur, oppløsning	not available	01-9980	MUNDIPHARMA AS	NO
OxyNorm 10 mg/ml oraaliliuos	not available	14808	MUNDIPHARMA OY	FI
OxyNorm 10 mg/ml oral lösning	not available	15793	MUNDIPHARMA AB	SE
OxyNorm 10 mg/ml solución inyectable y para perfusión.	IE/H/0139/006	71243	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 10 mg/ml solution for injection or infusion.	IE/H/0139/006	PA1688/006/001	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Oxynorm 10 mg/ml stungulyf, lausn, stungulyfs-/innrennslispykkni, lausn	DE/H/0366/015	IS/1/09/016/01	MUNDIPHARMA A/S	IS
OXYNORM 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	2009080027	MUNDIPHARMA COMM VA	LU
OxyNorm 10 mg/ml, injektio-/infuusioneste, liuos	not available	17150	MUNDIPHARMA OY	FI
OxyNorm 10 mg/ml, oplossing voor injectie	DE/H/0366/015	BE339814	MUNDIPHARMA COMM VA	BE
OxyNorm 10 mg/ml, oplossing voor injectie	DE/H/0366/015	BE339823	MUNDIPHARMA COMM VA	BE
OXYNORM 10 mg/ml, solution buvable	not available	34009 301 318 1 7	MUNDIPHARMA SAS	FR
OXYNORM 10 mg/ml, solution buvable	not available	34009 301 318 2 4	MUNDIPHARMA SAS	FR
OxyNorm 10 mg/ml,	not available	20303	MUNDIPHARMA	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
solution for injection or infusion			PHARMACEUTICALS LTD	
OxyNorm 10 mg/ml, solution for injection or infusion	not available	20303	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OXYNORM 10 mg/ml, solution injectable	not available	34009 392 317 1 6	MUNDIPHARMA SAS	FR
OXYNORM 10 mg/ml, solution injectable	not available	34009 366 915 2 0	MUNDIPHARMA SAS	FR
OXYNORM 10 mg/ml, solution injectable	not available	34009 366 914 6 9	MUNDIPHARMA SAS	FR
OXYNORM 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	BE339814	MUNDIPHARMA COMM VA	BE
OXYNORM 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	BE339823	MUNDIPHARMA COMM VA	BE
OxyNorm 20 mg	not available	20305	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 20 mg cápsulas	IE/H/0139/005	66576	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 20 mg capsules	IE/H/0139/003-005/MR	PA 1688/006/006	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm 20 mg hårda kapslar	not available	16084	MUNDIPHARMA AB	SE
OxyNorm 20 mg harde kapsler	not available	01-7403	MUNDIPHARMA AS	NO
OxyNorm 20 mg kapseli, kova	not available	15322	MUNDIPHARMA OY	FI
OXYNORM 20 mg, gélule	not available	34009 362 423 8 8	MUNDIPHARMA SAS	FR
OXYNORM 20 mg, gélule	not available	34009 361 904 2 9	MUNDIPHARMA SAS	FR
OXYNORM 20 mg, gélule	not available	34009 361 903 6 8	MUNDIPHARMA SAS	FR
OXYNORM 20 mg, gélule	not available	34009 361 901 3 9	MUNDIPHARMA SAS	FR
OXYNORM 20 mg, gélule	not available	34009 362 422 1 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
OxyNorm 20 mg, harde capsules	not available	RVG 27511	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm 5 mg	not available	20304	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 5 mg cápsulas	IE/H/0139/003	66574	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 5 mg capsules	IE/H/0139/003-005/MR	PA 1688/006/004	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm 5 mg hårda kapslar	not available	16454	MUNDIPHARMA AB	SE
OxyNorm 5 mg harde kapsler	not available	01-7401	MUNDIPHARMA AS	NO
OxyNorm 5 mg kapseli, kova	not available	15609	MUNDIPHARMA OY	FI
OXYNORM 5 mg, gélule	not available	34009 361 892 4 9	MUNDIPHARMA SAS	FR
OxyNorm 5 mg, harde capsules	not available	RVG 27509	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm 50 mg/ml injeksjonsvæske/infusjonsvæske, oppløsning	not available	07-5504	MUNDIPHARMA AS	NO
OXYNORM 50 mg/ml, solution à diluer pour perfusion	DE/H/0366/014	2009120027	MUNDIPHARMA COMM VA	LU
OXYNORM 50 mg/ml, solution injectable	not available	34009 387 625 3 2	MUNDIPHARMA SAS	FR
OxyNorm concentrado 10 mg/ml solución oral.	IE/H/0139/002	66.614	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm concentrate 10 mg/ml	not available	20973	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm concentrate 10 mg/ml oral solution	IE/H/0139/001	PA1688/006/003	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 10 mg Orodispersible Tablets	not available	PA 1688/006/008	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 10 mg orodispersible tablets	not available	PL06934/0220	ETHYPHARM	XI
OxyNorm Dispersa 10 mg,	DE/H/0366/007	IS/1/07/015/02	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
munndreifitöflur				
OxyNorm Dispersa 20 mg Orodispersible Tablets	not available	PA 1688/006/009	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 20 mg orodispersible tablets	not available	PL06934/0221	ETHYPHARM	XI
OxyNorm Dispersa 20 mg, munndreifitöflur	DE/H/0366/008	IS/1/07/015/03	MUNDIPHARMA A/S	IS
OxyNorm Dispersa 5 mg orodispersible tablets	not available	PA 1688/006/007	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 5 mg orodispersible tablets	not available	PL06934/0219	ETHYPHARM	XI
OxyNorm Dispersa 5 mg, munndreifitöflur	DE/H/0366/007-008	IS/1/07/015/01	MUNDIPHARMA A/S	IS
OxyNorm Dispersa, smeltetabletter	not available	40539	MUNDIPHARMA A/S	DK
OxyNorm Dispersa, smeltetabletter 10 mg	not available	40538	MUNDIPHARMA A/S	DK
OxyNorm Dispersa, smeltetabletter 5 mg	not available	40537	MUNDIPHARMA A/S	DK
OxyNorm drank 10 mg/ml	not available	RVG 27940	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm drank 5 mg / 5 ml	not available	RVG 27939	MUNDIPHARMA PHARMACEUTICALS BV	NL
Oxynorm hårde kapsler 20 mg	not available	31296	MUNDIPHARMA A/S	DK
OxyNorm injectie 10 mg/ml, oplossing voor injectie of infusie	not available	RVG 29031	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm injectie 50 mg/ml, oplossing voor injectie of infusie	not available	RVG 101605	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm Instant 10 mg, comprimés orodispersibles	DE/H/0366/007	2008110045	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 10 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319076	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
OxyNorm Instant 10 mg, orodispergeerbare tabletten	not available	RVG 34776	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm Instant 10 mg, comprimés orodispersibles	DE/H/0366/007-008	BE319076	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 20 mg, comprimés orodispersibles	DE/H/0366/008	2008110043	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 20 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319085	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 20 mg, orodispergeerbare tabletten	not available	RVG 34777	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm Instant 20 mg, comprimés orodispersibles	DE/H/0366/007-008	BE319085	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 5 mg, comprimés orodispersibles	DE/H/0366/006	2008110044	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 5 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319067	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 5 mg, orodispergeerbare tabletten	not available	RVG 34775	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm Instant 5 mg, comprimés orodispersibles	DE/H/0366/007-008	BE319067	MUNDIPHARMA COMM VA	BE
OxyNorm liquid 1 mg/ml oral solution	IE/H/0139/002	PA1688/006/002	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm liquid 5mg/5ml	not available	20972	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm, 10 mg/ml, roztwór do wstrzykiwań / koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	PL/H/0355/006	15220	MUNDIPHARMA A/S	PL
OxyNorm, hårde kapsler	not available	31294	MUNDIPHARMA A/S	DK
OxyNorm, hårde kapsler	not available	31295	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
Oxynorm, injektions- og infusionsvæske, opløsning 10 mg/ml	not available	33361	MUNDIPHARMA A/S	DK
Oxynorm, oral opløsning 1 mg/ml	not available	31157	MUNDIPHARMA A/S	DK
Oxynorm, oral opløsning 10 mg/ml	not available	31158	MUNDIPHARMA A/S	DK
OxyNorm® 50 mg/ml, solution for injection or infusion	not available	20595	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm® 50 mg/ml, solution for injection or infusion	not available	20595	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm® 50 mg/ml, solution for injection or infusion	not available	PA 1688/006/010	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 426 5 8	MUNDIPHARMA SAS	FR
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 427 1 9	MUNDIPHARMA SAS	FR
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 425 9 7	MUNDIPHARMA SAS	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 429 4 8	MUNDIPHARMA SAS	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 430 2 0	MUNDIPHARMA SAS	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 428 8 7	MUNDIPHARMA SAS	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 424 2 9	MUNDIPHARMA SAS	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 423 6 8	MUNDIPHARMA SAS	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 421 3 9	MUNDIPHARMA SAS	FR
Oxypro 15 mg prolonged-release tablets	not available	PL 48804/0003	RIDGE PHARMA 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
Oxypro 30 mg prolonged-release tablets	not available	PL 48804/0005	RIDGE PHARMA LIMITED	XI
Oxypro 60 mg prolonged-release tablets	not available	PL 48804/0007	RIDGE PHARMA LIMITED	XI
Zarenoxin 15 mg Retardtabletten	DE/H/3642/003	87953.00.00	ACINO AG	DE
Zarenoxin 30 mg Retardtabletten	DE/H/3642/005	87955.00.00	ACINO AG	DE
Zarenoxin 60 mg Retardtabletten	DE/H/3642/007	87957.00.00	ACINO AG	DE