14 November 2024
EMA/596129/2024
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: oxycodone

Procedure no.: PSUSA/00002254/202404

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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

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
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
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
Orionox Depot 30 mg depottabletter	SE/H/1493/005	12-9223	ORION CORPORATION	NO
Orionox Depot 15 mg depottabletter	SE/H/1493/003	12-9221	ORION CORPORATION	NO
Orionox Depot 60 mg depottabletter	SE/H/1493/007	12-9225	ORION CORPORATION	NO
Orionox Depot, depottabletter	SE/H/1493/003	51490	ORION CORPORATION	DK
Orionox Depot, depottabletter	SE/H/1493/005	51492	ORION CORPORATION	DK
Orionox Depot, depottabletter	SE/H/1493/007	51494	ORION CORPORATION	DK
Oxycan uno 10 mg Retardtabletten	DE/H/3655/001	88009.00.00	HORMOSAN PHARMA GMBH	DE
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

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 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 Abboxia 1 mg/ml injeksjons-/infusjonsvæske, oppløsning	FI/H/1123/01	21-14094	ABBOXIA AB	NO
Oxycodone Abboxia 1 mg/ml injektio-/infuusioneste, liuos	FI/H/1123/001	39429	ABBOXIA AB	FI
Oxycodone Abboxia 1 mg/ml injektions-/infusionsvätska, lösning	FI/H/1123/001	39429	ABBOXIA AB	FI
Oxycodone Abboxia 1 mg/ml injektions-/infusionsvätska, lösning	FI/H/1123/001	62150	ABBOXIA AB	SE
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927785832	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786723	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786891	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786082	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
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927787263	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786433	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786952	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786204	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927787034	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786143	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786372	BIOGARAN	FR
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération prolongée	FR/H/0621/003/DC	3400927786662	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927788505	BIOGARAN	FR
OXYCODONE BIOGARAN	FR/H/0621/005/DC	3400927789564	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 30 mg, comprimé pelliculé à libération prolongée				
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927788963	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927788673	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789045	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927788734	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789274	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789625	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789106	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789335	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé	FR/H/0621/005/DC	3400927789854	BIOGARAN	FR

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pelliculé à libération prolongée				
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789793	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791284	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791925	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791574	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927792465	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927792175	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791635	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927792007	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération	FR/H/0621/007/DC	3400927791406	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
prolongée				
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927792526	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791345	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927791864	BIOGARAN	FR
OXYCODONE BIOGARAN LP 60 mg, comprimé pelliculé à libération prolongée	FR/H/0621/007/DC	3400927792236	BIOGARAN	FR
Oxycodone Depot Orion 15 mg depottabletter	SE/H/1493/03	48495	ORION CORPORATION	SE
Oxycodone Depot Orion 30 mg depottabletter	SE/H/1493/05	48497	ORION CORPORATION	SE
Oxycodone Depot Orion 60 mg depottabletter	SE/H/1493/07	48499	ORION CORPORATION	SE
OXYCODONE EG LP 15 mg, comprimé pelliculé à libération prolongée	DE/H/3641/003	NL42844	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 30 mg, comprimé pelliculé à libération prolongée	DE/H/3641/005	NL42846	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 60 mg, comprimé pelliculé à libération prolongée	DE/H/3641/007	NL42848	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE RENAUDIN 1 mg/mL, solution pour perfusion	not available	34009 301 637 9 5	LABORATOIRE RENAUDIN	FR
OXYCODONE RENAUDIN 1	not available	34009 301 638 0 1	LABORATOIRE RENAUDIN	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
mg/mL, solution pour 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

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
Oxycodon-HCl AL 15 mg Retardtabletten	DE/H/3860/003	89918.00.00	ALIUD PHARMA GMBH	DE
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;	53004.03.00	KRUGMANN GMBH (GERMANY)	DE

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	DE/H/1026/004; DE/H/			
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 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
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

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
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/0112/001-004/MR	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/01/01201/003	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 produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-118192252-02	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-118192252-03	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,	DE/H/0366/001-003/MR	BE253197	MUNDIPHARMA COMM VA	BE

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comprimés à libération prolongée				
OXYCONTIN 10 mg, comprimés à libération prolongée	DE/H/0366/001-003/MR	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 depottabletter	not available	07-4847	MUNDIPHARMA AS	NO
OxyContin 160 mg, tabletten met verlengde afgifte	not available	RVG 100825	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 20 mg comprimidos de liberación prolongada	IE/H/0112/001-004/MR	63447	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 20 mg depottabletter	not available	94-2769	MUNDIPHARMA AS	NO
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

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 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	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 produljenim oslobađanjem	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	HR-H-188823708-05	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-188823708-06	MUNDIPHARMA GES.M.B.H	HR
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/0366/001-003/MR	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 20mg	not available	65/258/00-C	MUNDIPHARMA GES.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 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/0112/001-004/MR	63448	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 40 mg depottabletter	not available	94-2770	MUNDIPHARMA AS	NO
OxyContin 40 mg prolonged release tablets	not available	18610	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 40 mg prolonged release tablets	IE/H/0112/001-004/MR	PA 1688/005/004	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/009	MUNDIPHARMA GES.M.B.H	SI
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/0366/001-003/MR	2009010138	MUNDIPHARMA COMM VA	LU
OxyContin 40 mg, tabletten met verlengde afgifte	DE/H/0366/001-003/MR	BE253276	MUNDIPHARMA COMM VA	BE
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	not available	02-1153	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 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	2009010135	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/0112/001-004/MR	63449	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 80 mg depottabletter	not available	01-2425	MUNDIPHARMA AS	NO
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 tablete s podaljšanim sproščanjem	not available	H/01/01201/012	MUNDIPHARMA GES.M.B.H	SI
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	DE/H/0366/004	BE253303	MUNDIPHARMA COMM VA	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
afgifte				
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 80mg	not available	65/260/00-C	MUNDIPHARMA GES.M.B.H	CZ
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/0366/001-003/MR	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/0366/001-003/MR	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, depottabletter 10 mg	not available	16892	MUNDIPHARMA A/S	DK
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

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 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 354 210 9 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 209 0 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 208 4 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 207 8 7	MUNDIPHARMA SAS	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 211 5 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 206 1 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 604 5 2	MUNDIPHARMA SAS	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 603 9 1	MUNDIPHARMA SAS	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à	not available	34009 384 602 2 3	MUNDIPHARMA SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
libération prolongée				
OXYCONTIN LP 15 mg, comprimé pelliculé à libération prolongée	not available	34009 384 586 7 1	MUNDIPHARMA SAS	FR
OXYCONTIN LP 15 mg, comprimé pelliculé à libération prolongée	not available	34009 384 585 0 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 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 216 7 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 214 4 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 213 8 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 217 3 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 212 1 0	MUNDIPHARMA SAS	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à libération prolongée	not available	34009 384 590 4 3	MUNDIPHARMA SAS	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à libération prolongée	not available	34009 384 589 6 1	MUNDIPHARMA SAS	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à	not available	34009 384 587 3 2	MUNDIPHARMA SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
libération prolongée				
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 223 3 0	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 220 4 0	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 221 0 1	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 222 7 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 225 6 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 219 6 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 905 7 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 904 0 0	MUNDIPHARMA SAS	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 903 4 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 599 1 3	MUNDIPHARMA SAS	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 601 6 2	MUNDIPHARMA SAS	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à	not available	34009 384 598 5 2	MUNDIPHARMA SAS	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
libération prolongée				
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 296 0 5	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 295 4 4	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 228 5 9	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 227 9 8	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 294 8 3	MUNDIPHARMA SAS	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 226 2 0	MUNDIPHARMA SAS	FR
OxyContin retard 10 mg Filmtabletten	IE/H/0112/001-004/MR	1-23358	MUNDIPHARMA GES.M.B.H	AT
OxyContin retard 20 mg Filmtabletten	IE/H/0112/001-004/MR	1-23359	MUNDIPHARMA GES.M.B.H	AT
OxyContin retard 40 mg Filmtabletten	IE/H/0112/001-004/MR	1-23360	MUNDIPHARMA GES.M.B.H	AT
OxyContin retard 5 mg Filmtabletten	IE/H/0112/005	1-26041	MUNDIPHARMA GES.M.B.H	AT
OxyContin retard 80 mg Filmtabletten	IE/H/0112/001-004/MR	1-23361	MUNDIPHARMA GES.M.B.H	AT
OxyContin, 10 mg depottablett	not available	13830	MUNDIPHARMA AB	SE
OxyContin, 10 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	14525	MUNDIPHARMA A/S	PL
OxyContin, 20 mg depottablett	not available	13831	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
OxyContin, 20 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	14526	MUNDIPHARMA A/S	PL
OxyContin, 40 mg depottablett	not available	13832	MUNDIPHARMA AB	SE
OxyContin, 40 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	14527	MUNDIPHARMA A/S	PL
OxyContin, 5 mg depottablett	not available	18349	MUNDIPHARMA AB	SE
OxyContin, 5 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	14524	MUNDIPHARMA A/S	PL
OxyContin, 80 mg depottablett	not available	15982	MUNDIPHARMA AB	SE
OxyContin, 80 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/002; DE/H/1026/003; DE/H/1026/004; DE/H/	14528	MUNDIPHARMA A/S	PL
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435038	MUNDIPHARMA PHARMACEUTICALS SRL	IT
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

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® 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 compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435141	MUNDIPHARMA PHARMACEUTICALS SRL	IT
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

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® 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 compresse a rilascio prolungato	IE/H/0112/001-004/MR	34435230	MUNDIPHARMA PHARMACEUTICALS SRL	IT
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
Oxycorion Depot 15 mg depottabletit	SE/H/1493/03	31051	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycorion Depot 15 mg depottabletter	SE/H/1493/003	31051	ORION CORPORATION	FI
Oxycorion Depot 30 mg depottabletit	SE/H/1493/005	31053	ORION CORPORATION	FI
Oxycorion Depot 30 mg depottabletter	SE/H/1493/005	31053	ORION CORPORATION	FI
Oxycorion Depot 60 mg depottabletit	SE/H/1493/07	31055	ORION CORPORATION	FI
Oxycorion Depot 60 mg depottabletter	SE/H/1493/007	31055	ORION CORPORATION	FI
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 Schmelztabletten	DE/H/0366/007-008	68387.00.00	MUNDIPHARMA GMBH	DE
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	DE/H/0366/015	57886.00.00	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
Injektionslösung/Konzentrat zur Herstellung einer Injektions-/Infusionslösung				
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 cápsulas duras	IE/H/0139/003-005/MR	66575	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 10 mg capsules	not available	20306	MUNDIPHARMA PHARMACEUTICALS LTD	CY
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 361 899 9 7	MUNDIPHARMA SAS	FR
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 solución inyectable y para perfusión	IE/H/0139/006	71243	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 10 mg/ml injeksjonsvæske/infusjonsvæske, oppløsning	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
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 stungulyf, lausn, stungulyfs-/innrennslispykkni, lausn	DE/H/0366/015	IS/1/09/016/01	MUNDIPHARMA A/S	IS
OxyNorm 10 mg/ml, injektio-/infuusioneste, liuos	not available	17150	MUNDIPHARMA OY	FI
OxyNorm 10 mg/ml, injektionsvätska, lösning	not available	18256	MUNDIPHARMA AB	SE
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, solution for injection or infusion	not available	20303	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 10 mg/ml, solution for injection or infusion	IE/H/0139/006	PA 1688/006/001	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
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,	DE/H/0366/015	BE339814	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
solution injectable/solution à diluer injectable ou perfusion				
OXYNORM 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	BE339823	MUNDIPHARMA COMM VA	BE
OXYNORM 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	2009080027	MUNDIPHARMA COMM VA	LU
OxyNorm 20 mg cápsulas duras	IE/H/0139/003-005/MR	66576	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 20 mg capsules	not available	20305	MUNDIPHARMA PHARMACEUTICALS LTD	CY
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 361 905 9 7	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
OxyNorm 20 mg, harde capsules	not available	RVG 27511	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm 5 mg cápsulas duras	IE/H/0139/003-005/MR	66574	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 5 mg capsules	not available	20304	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 5 mg capsules	IE/H/0139/003-005/MR	PA 1688/006/004	MUNDIPHARMA	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
			PHARMACEUTICALS LIMITED	
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, gélule	not available	34009 361 895 3 9	MUNDIPHARMA SAS	FR
OXYNORM 5 mg, gélule	not available	34009 361 893 0 0	MUNDIPHARMA SAS	FR
OXYNORM 5 mg, gélule	not available	34009 362 419 0 9	MUNDIPHARMA SAS	FR
OXYNORM 5 mg, gélule	not available	34009 361 894 7 8	MUNDIPHARMA SAS	FR
OXYNORM 5 mg, gélule	not available	34009 362 418 4 8	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 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
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/0139/001-002/MR	66614	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm concentrate 10 mg/ml	not available	20973	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm concentrate 10mg/ml oral solution	IE/H/0139/001-002/MR	PA 1688/006/003	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

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 Dispersa 10 mg, munndreifitöflur	DE/H/0366/007-008	IS/1/07/015/02	MUNDIPHARMA A/S	IS
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/0366/007-008	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 10 mg	not available	40538	MUNDIPHARMA A/S	DK
OxyNorm Dispersa, smeltetabletter 20 mg	not available	40539	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 10 mg	not available	31295	MUNDIPHARMA A/S	DK
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/0366/007-008	2008110045	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
OxyNorm Instant 10 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319076	MUNDIPHARMA BV	BE
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/0366/007-008	2008110043	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 20 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319085	MUNDIPHARMA BV	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 BV	BE
OxyNorm Instant 5 mg, comprimés orodispersibles	DE/H/0366/007-008	2008110044	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 5 mg, orodispergeerbare tabletten	DE/H/0366/007-008	BE319067	MUNDIPHARMA BV	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/0139/001-002/MR	PA 1688/006/002	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm liquid 5mg/5ml oral solution	not available	20972	MUNDIPHARMA PHARMACEUTICALS LTD	CY
Oxynorm, hårde kapsler 5 mg	not available	31294	MUNDIPHARMA A/S	DK
Oxynorm, injektions- og infusionsvæske, opløsning	not available	33361	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
10 mg/ml				
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
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
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