12 December 2018
EMA/817240/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): oxycodone

Procedure No.: PSUSA/00002254/201804

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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
Abtard 10 mg prolonged-release tablets	UK/H/6614/002	PL 42623/0002	ETHYPHARM UK LTD	UK
Abtard 15 mg prolonged-release tablets	UK/H/6614/003	PL 42623/0003	ETHYPHARM UK LTD	UK
Abtard 20 mg prolonged-release tablets	UK/H/6614/002	PL 42623/0004	ETHYPHARM UK LTD	UK
Abtard 30 mg prolonged-release tablets	UK/H/6614/005	PL 42623/0005	ETHYPHARM UK LTD	UK
Abtard 40 mg prolonged-release tablets	UK/H/6614/006	PL 42623/0006	ETHYPHARM UK LTD	UK
Abtard 5 mg prolonged-release tablets	UK/H/6614/001	PL 42623/0001	ETHYPHARM UK LTD	UK
Abtard 60 mg prolonged-release tablets	UK/H/6614/007	PL 42623/0007	ETHYPHARM UK LTD	UK
Abtard 80 mg prolonged-release tablets	UK/H/6614/008	PL 42623/0008	ETHYPHARM UK LTD	UK
Accordeon, 10 mg, tabletki o przedłużonym uwalnianiu	NL/H/3192/002	19785	ACCORD HEALTHCARE LIMITED	PL
Accordeon, 20 mg, tabletki o przedłużonym uwalnianiu	NL/H/3192/003	19786	ACCORD HEALTHCARE LIMITED	PL
Accordeon, 40 mg, tabletki o przedłużonym uwalnianiu	NL/H/3192/004	19787	ACCORD HEALTHCARE LIMITED	PL
Accordeon, 5 mg, tabletki o przedłużonym uwalnianiu	NL/H/3192/001	19784	ACCORD HEALTHCARE LIMITED	PL
Accordeon, 80 mg, tabletki o przedłużonym uwalnianiu	NL/H/3192/005	19788	ACCORD HEALTHCARE LIMITED	PL
Alivio 10 mg prolonged-release tablets	DE/H/4245/002	PL 17907/0559	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Alivio 20 mg prolonged-release tablets	DE/H/4245/003	PL 17907/0560	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Alivio 30 mg prolonged-	DE/H/4245/004	PL 17907/0561	BRISTOL LABORATORIES	UK

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release tablets			LTD (BERKHAMSTED)	
Alivio 40 mg prolonged-release tablets	DE/H/4245/005	PL 17907/0562	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Alivio 5 mg prolonged-release tablets	DE/H/4245/001	PL 17907/0558	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Alivio 60 mg prolonged-release tablets	DE/H/4245/006	PL 17907/0563	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Alivio 80 mg prolonged-release tablets	DE/H/4245/007	PL 17907/0564	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Candox 10 mg prolonged release tablets	DE/H/1395/002	PL 00289/1710	TEVA UK LIMITED	UK
Candox 20mg prolonged release tablets	DE/H/0790/001	PL 00289/1711	TEVA UK LIMITED	UK
Candox 40mg prolonged release tablets	DE/H/0790/002	PL 00289/1712	TEVA UK LIMITED	UK
Candox 5 mg Prolonged-release Tablets	DE/H/2947/001	PL 00289/1709	TEVA UK LIMITED	UK
Candox 80mg Prolonged-release Tablets	DE/H/0790/003	PL 00289/1713	TEVA UK LIMITED	UK
Carenoxal 10 mg Retardtabletten	DE/H/3638/002	87920.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 15 mg Retardtabletten	DE/H/3638/003	87921.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 20 mg Retardtabletten	DE/H/3638/004	87922.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 30 mg Retardtabletten	DE/H/3638/005	87923.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 40 mg Retardtabletten	DE/H/3638/006	87924.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 5 mg Retardtabletten	DE/H/3638/001	87919.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 60 mg Retardtabletten	DE/H/3638/007	87925.00.00	ZENTIVA PHARMA GMBH	DE
Carenoxal 80 mg Retardtabletten	DE/H/3638/008	87926.00.00	ZENTIVA 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
Carexil 20 mg Prolonged-release Tablets	DE/H/1154/003	PL 04416/0836	SANDOZ LTD	UK
Carexil 5 mg Prolonged-release Tablets	DE/H/1154/001	PL 04416/0834	SANDOZ LTD	UK
Carexil® 10 mg Prolonged-release Tablets	DE/H/1154/002	PL 04416/0835	SANDOZ LTD	UK
Carexil® 40 mg Prolonged-release Tablets	DE/H/1154/004	PL 04416/1304	SANDOZ LTD	UK
Carexil® 80 mg Prolonged-release Tablets	DE/H/1154/006	PL 04416/1305	SANDOZ LTD	UK
Codilek 10 mg tablete s podaljšanim sproščanjem	DE/H/1154/002	5363-I-269/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 10 mg tablete s podaljšanim sproščanjem	DE/H/1154/002	5363-I-270/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 10 mg tablete s podaljšanim sproščanjem	DE/H/1154/002	5363-I-271/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 10 mg tablete s podaljšanim sproščanjem	DE/H/1154/002	5363-I-272/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 10 mg tablete s podaljšanim sproščanjem	DE/H/1154/002	5363-I-273/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 20 mg tablete s podaljšanim sproščanjem	DE/H/1154/003	5363-I-276/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 20 mg tablete s podaljšanim sproščanjem	DE/H/1154/003	5363-I-274/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 20 mg tablete s podaljšanim sproščanjem	DE/H/1154/003	5363-I-275/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 20 mg tablete s podaljšanim sproščanjem	DE/H/1154/003	5363-I-277/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 20 mg tablete s podaljšanim sproščanjem	DE/H/1154/003	5363-I-278/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/068	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 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/069	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/070	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/071	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/072	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/073	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/074	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/076	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/075	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/077	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/078	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/079	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/080	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/081	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/082	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/083	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/084	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 40 mg tablete s podaljšanim sproščanjem	DE/H/1154/004	H/09/00405/067	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s	DE/H/1154/001	5363-I-264/13	LEK PHARMACEUTICALS	SI

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podaljšanim sproščanjem			D.D. LJUBLJANA	
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	5363-I-268/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	5363-I-265/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	5363-I-266/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 5 mg tablete s podaljšanim sproščanjem	DE/H/1154/001	5363-I-267/13	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/086	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/087	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/088	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/089	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/090	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/091	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/092	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/093	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/094	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/095	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/096	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/097	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/098	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI

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Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/099	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/100	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/101	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/102	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 60 mg tablete s podaljšanim sproščanjem	DE/H/1154/005	H/09/00405/085	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/105	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/106	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/107	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/108	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/109	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/110	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/111	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/112	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/113	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/114	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/116	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/118	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s	DE/H/1154/006	H/09/00405/119	LEK PHARMACEUTICALS	SI

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podaljšanim sproščanjem			D.D. LJUBLJANA	
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/120	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/104	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/115	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/117	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codilek 80 mg tablete s podaljšanim sproščanjem	DE/H/1154/006	H/09/00405/103	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Codoxy 10 mg retard tabletta	DE/H/2182/002	OGYI-T-21706/05	G.L. PHARMA GMBH	HU
Codoxy 10 mg retard tabletta	DE/H/2182/002	OGYI-T-21706/06	G.L. PHARMA GMBH	HU
Codoxy 10 mg retard tabletta	DE/H/2182/002	OGYI-T-21706/07	G.L. PHARMA GMBH	HU
Codoxy 10 mg retard tabletta	DE/H/2182/002	OGYI-T-21706/08	G.L. PHARMA GMBH	HU
Codoxy 20 mg retard tabletta	DE/H/2182/003	OGYI-T-21706/09	G.L. PHARMA GMBH	HU
Codoxy 20 mg retard tabletta	DE/H/2182/003	OGYI-T-21706/10	G.L. PHARMA GMBH	HU
Codoxy 20 mg retard tabletta	DE/H/2182/003	OGYI-T-21706/11	G.L. PHARMA GMBH	HU
Codoxy 20 mg retard tabletta	DE/H/2182/003	OGYI-T-21706/12	G.L. PHARMA GMBH	HU
Codoxy 40 mg retard tabletta	DE/H/2182/004	OGYI-T-21706/13	G.L. PHARMA GMBH	HU
Codoxy 40 mg retard tabletta	DE/H/2182/004	OGYI-T-21706/14	G.L. PHARMA GMBH	HU
Codoxy 40 mg retard tabletta	DE/H/2182/004	OGYI-T-21706/15	G.L. PHARMA GMBH	HU
Codoxy 40 mg retard tabletta	DE/H/2182/004	OGYI-T-21706/16	G.L. PHARMA GMBH	HU

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Codoxy 5 mg retard tabletta	DE/H/2182/001	OGYI-T-21706/01	G.L. PHARMA GMBH	HU
Codoxy 5 mg retard tabletta	DE/H/2182/001	OGYI-T-21706/02	G.L. PHARMA GMBH	HU
Codoxy 5 mg retard tabletta	DE/H/2182/001	OGYI-T-21706/03	G.L. PHARMA GMBH	HU
Codoxy 5 mg retard tabletta	DE/H/2182/001	OGYI-T-21706/04	G.L. PHARMA GMBH	HU
Codoxy 80 mg retard tabletta	DE/H/2182/005	OGYI-T-21706/17	G.L. PHARMA GMBH	HU
Codoxy 80 mg retard tabletta	DE/H/2182/005	OGYI-T-21706/18	G.L. PHARMA GMBH	HU
Codoxy 80 mg retard tabletta	DE/H/2182/005	OGYI-T-21706/19	G.L. PHARMA GMBH	HU
Codoxy 80 mg retard tabletta	DE/H/2182/005	OGYI-T-21706/20	G.L. PHARMA GMBH	HU
Contiroxil 10 mg tablety s predĺženým uvoľňovaním	DE/H/1154/002	65/0272/08-S	SANDOZ PHARMACEUTICALS D.D.	SK
Contiroxil 20 mg tablety s predĺženým uvoľňovaním	DE/H/1154/003	65/0273/08-S	SANDOZ PHARMACEUTICALS D.D.	SK
Contiroxil 40 mg tablety s predĺženým uvoľňovaním	DE/H/1154/004	65/0403/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Contiroxil 80 mg tablety s predĺženým uvoľňovaním	DE/H/1154/006	65/0404/12-S	SANDOZ PHARMACEUTICALS D.D.	SK
Contiroxin 40 mg Prolonged-release Tablets	DE/H/0789/002	PL 04416/0754	SANDOZ LTD	UK
Contiroxin 80 mg Prolonged-release Tablets	DE/H/0789/003	PL 04416/0755	SANDOZ LTD	UK
Daloxy 10 mg capsule rigide a rilascio	IS/H/0196/001	042781017	ETHYPHARM	IT

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prolungato				
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781029	ETHYPHARM	IT
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781031	ETHYPHARM	IT
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781043	ETHYPHARM	IT
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781056	ETHYPHARM	IT
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781070	ETHYPHARM	IT
Daloxy 10 mg capsule rigide a rilascio prolungato	IS/H/0196/001	042781068	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781082	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781094	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781106	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781118	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781120	ETHYPHARM	IT
Daloxy 20 mg capsule rigide a rilascio	IS/H/0196/002	042781132	ETHYPHARM	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
prolungato				
Daloxy 20 mg capsule rigide a rilascio prolungato	IS/H/0196/002	042781144	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781157	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781169	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781171	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781183	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781195	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781207	ETHYPHARM	IT
Daloxy 40 mg capsule rigide a rilascio prolungato	IS/H/0196/003	042781219	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781221	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781233	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781245	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio	IS/H/0196/004	042781258	ETHYPHARM	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
prolungato				
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781260	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781272	ETHYPHARM	IT
Daloxy 80 mg capsule rigide a rilascio prolungato	IS/H/0196/004	042781284	ETHYPHARM	IT
Dancex SR 10 mg Prolonged-Release Tablets	DE/H/1155/002	PA0711/143/002	ROWEX LTD	IE
Dancex SR 20 mg Prolonged-Release Tablets	DE/H/1155/003	PA0711/143/003	ROWEX LTD	IE
Dancex SR 40 mg Prolonged-Release Tablets	DE/H/1155/004	PA0711/143/004	ROWEX LTD	IE
Dancex SR 5 mg Prolonged-Release Tablets	DE/H/1155/001	PA0711/143/001	ROWEX LTD	IE
Dancex SR 60 mg Prolonged-Release Tablets	DE/H/1155/005	PA0711/143/005	ROWEX LTD	IE
Dancex SR 80 mg Prolonged-Release Tablets	DE/H/1155/006	PA0711/143/006	ROWEX LTD	IE
Dolanor 10 mg comprimido de libertacao prolongada	DE/H/2183/002	5314919	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5315015	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação	DE/H/2183/002	5315007	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314976	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314968	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314950	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314943	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314935	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314927	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314901	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314877	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314869	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314851	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314844	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação	DE/H/2183/002	5314836	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314828	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314810	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314802	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 10 mg comprimido de libertação prolongada	DE/H/2183/002	5314778	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertacao prolongada	DE/H/2183/003	5315122	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315247	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315239	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315221	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315213	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315205	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315171	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação	DE/H/2183/003	5315163	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315155	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315148	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315130	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315114	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315106	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315072	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315064	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315056	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315049	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315031	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 20 mg comprimido de libertação prolongada	DE/H/2183/003	5315023	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao	DE/H/2183/004	5315478	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315460	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315452	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315445	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315437	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315429	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315411	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315403	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315379	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315361	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315535	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315346	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao	DE/H/2183/004	5315338	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315320	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315312	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315304	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315270	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315262	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 40 mg comprimido de libertacao prolongada	DE/H/2183/004	5315254	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertacao prolongada	DE/H/2183/001	5314554	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertacao prolongada	DE/H/2183/001	5314562	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertacao prolongada	DE/H/2183/001	5314547	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314760	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314752	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação	DE/H/2183/001	5314745	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314737	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314729	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314711	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314703	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314679	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314661	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314653	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314646	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314638	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314620	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314612	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 5 mg comprimido de libertação	DE/H/2183/001	5314604	LANNACHER HEILMITTEL GES.M.B.H.,	PT

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
prolongada				
Dolanor 5 mg comprimido de libertação prolongada	DE/H/2183/001	5314570	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	DE/H/2183/0005	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315700	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315668	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315650	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315643	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315635	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315569	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315544	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315536	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolanor 80 mg comprimido de libertacao prolongada	DE/H/2183/005	5315528	LANNACHER HEILMITTEL GES.M.B.H.,	PT
Dolocodon 10 mg tablety s prodlouženým	CZ/H/0212/002	65/674/10-C	ZENTIVA, K.S.	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
uvolňováním				
Dolocodon 10 mg, toimeainet prolongeeritult vabastavad tabletid	CZ/H/0212/002	707610	ZENTIVA, K.S.	EE
Dolocodon 20 mg tablety s prodlouženým uvolňováním	CZ/H/0212/003	65/675/10-C	ZENTIVA, K.S.	CZ
Dolocodon 20 mg, toimeainet prolongeeritult vabastavad tabletid	CZ/H/0212/003	707810	ZENTIVA, K.S.	EE
Dolocodon 40 mg tablety s prodlouženým uvolňováním	CZ/H/0212/004	65/676/10-C	ZENTIVA, K.S.	CZ
Dolocodon 40 mg, toimeainet prolongeeritult vabastavad tabletid	CZ/H/0212/004	707710	ZENTIVA, K.S.	EE
Dolocodon 5 mg tablety s prodlouženým uvolňováním	CZ/H/0212/001	65/673/10-C	ZENTIVA, K.S.	CZ
Dyxal 10 mg tvrdé tobolky	DE/H/4502/002	65/287/17-C	SANDOZ S.R.O.	CZ
Dyxal 20 mg tvrdé tobolky	DE/H/4502/003	65/288/17-C	SANDOZ S.R.O.	CZ
Dyxal 5 mg tvrdé tobolky	DE/H/4502/001	65/286/17-C	SANDOZ S.R.O.	CZ
Ixyldone 10 mg prolonged-release tablets	DE/H/3639/002	PL 20117/0306	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 15 mg prolonged-release tablets	DE/H/3639/003	PL 20117/307	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 20 mg prolonged-release tablets	DE/H/3639/004	PL 20117/0308	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 30 mg prolonged-release tablets	DE/H/3639/005	PL 20117/0309	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 40 mg	DE/H/3639/006	PL 20117/0310	MORNINGSIDE HEALTHCARE	UK

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			LTD	
Ixyldone 5 mg prolonged-release tablets	DE/H/3639/001	PL 20117/0305	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 60 mg prolonged-release tablets	DE/H/3639/007	PL 20117/0311	MORNINGSIDE HEALTHCARE LTD	UK
Ixyldone 80 mg prolonged-release tablets	DE/H/3639/008	PL 20117/0312	MORNINGSIDE HEALTHCARE LTD	UK
Lenocod 10 mg Retardtabletten	DE/H/3738/002	89062.00.00	ETHYPHARM	DE
Lenocod 20 mg Retardtabletten	DE/H/3738/003	89063.00.00	ETHYPHARM	DE
Lenocod 40 mg Retardtabletten	DE/H/3738/004	89064.00.00	ETHYPHARM	DE
Lenocod 5 mg Retardtabletten	DE/H/3738/001	89061.00.00	ETHYPHARM	DE
Leveraxo 10 mg prolonged-release tablets	DE/H/4246/002	PL 04569/1651	GENERICS [UK] LIMITED	UK
Leveraxo 20 mg prolonged-release tablets	DE/H/4246/003	PL 04569/1652	GENERICS [UK] LIMITED	UK
Leveraxo 30 mg prolonged-release tablets	DE/H/4246/004	PL 04569/1653	GENERICS [UK] LIMITED	UK
Leveraxo 40 mg prolonged-release tablets	DE/H/4246/005	PL 04569/1654	GENERICS [UK] LIMITED	UK
Leveraxo 5 mg prolonged-release tablets	DE/H/4246/001	PL 04569/1650	GENERICS [UK] LIMITED	UK
Leveraxo 60 mg prolonged-release tablets	DE/H/4246/006	PL 04569/1655	GENERICS [UK] LIMITED	UK
Leveraxo 80 mg prolonged-release tablets	DE/H/4246/007	PL 04569/1656	GENERICS [UK] LIMITED	UK
Longtec 10 mg prolonged release tablets	not available	PL 40431/0002	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 120 mg prolonged release tablets	not available	PL 40431/0009	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 15 mg prolonged release tablets	not available	PL 40431/0006	QDEM PHARMACEUTICALS LIMITED	UK

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
Longtec 20 mg prolonged release tablets	not available	PL 40431/0003	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 30 mg prolonged release tablets	not available	PL 40431/0007	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 40 mg prolonged release tablets	not available	PL 40431/0004	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 5 mg prolonged release tablets	not available	PL 40431/0001	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 60 mg prolonged release tablets	not available	PL 40431/0008	QDEM PHARMACEUTICALS LIMITED	UK
Longtec 80 mg prolonged release tablets	not available	PL 40431/0005	QDEM PHARMACEUTICALS LIMITED	UK
Lynlor 10mg Capsules, hard	UK/H/6969/002	PL 30306/0421	ACTAVIS GROUP PTC EHF.	UK
Lynlor 20mg Capsules, hard	UK/H/6969/003	PL 30306/0422	ACTAVIS GROUP PTC EHF.	UK
Lynlor 5mg Capsules, hard	UK/H/6969/001	PL 30306/0420	ACTAVIS GROUP PTC EHF.	UK
Oksikodon Lek 10 mg tablete s podaljšanim sproščanjem	DE/H/1419/002	5363-I-2289/12	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Oksikodon Lek 5 mg tablete s podaljšanim sproščanjem	DE/H/1419/001	5363-I-2288/12	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Oksikodon Vitabalans 10 mg filmsko obložene tablete	SE/H/1119/002	H/12/01153/003	VITABALANS OY	SI
Oksikodon Vitabalans 10 mg filmsko obložene tablete	SE/H/1119/002	H/12/01153/004	VITABALANS OY	SI
Oksikodon Vitabalans 5 mg filmsko obložene tablete	SE/H/1119/001	H/12/01153/001	VITABALANS OY	SI
Oksikodon Vitabalans 5 mg filmsko obložene tablete	SE/H/1119/001	H/12/01153/002	VITABALANS OY	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
Olbete 10 mg comprimidos de libertação prolongada	not available	5600465	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 10 mg comprimidos de libertação prolongada	not available	5600473	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 10 mg comprimidos de libertação prolongada	not available	5600457	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 10 mg comprimidos de libertação prolongada	not available	5600440	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 20 mg comprimidos de libertação prolongada	not available	5600507	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 20 mg comprimidos de libertação prolongada	not available	5600515	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 20 mg comprimidos de libertação prolongada	not available	5600531	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 20 mg comprimidos de libertação prolongada	not available	5600523	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 40 mg comprimidos de libertação prolongada	not available	5601307	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 40 mg comprimidos de libertação prolongada	not available	5601315	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 40 mg comprimidos de libertação prolongada	not available	5601265	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 40 mg comprimidos de libertação prolongada	not available	5601273	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT

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
Olbete 5 mg comprimidos de libertação prolongada	not available	5600150	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 5 mg comprimidos de libertação prolongada	not available	5600200	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 5 mg comprimidos de libertação prolongada	not available	5600168	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 5 mg comprimidos de libertação prolongada	not available	5600176	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 80 mg comprimidos de libertação prolongada	not available	5601331	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 80 mg comprimidos de libertação prolongada	not available	5601323	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 80 mg comprimidos de libertação prolongada	not available	5601356	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Olbete 80 mg comprimidos de libertação prolongada	not available	5601349	TECNIMEDE - SOCIEDADE TÉCNICO-MEDICINAL, SA	PT
Onexila XL 10 mg prolonged-release tablets	DE/H/4465/001	PL 33155/0042	RIVOPHARM UK LTD	UK
Onexila XL 20 mg prolonged-release tablets	DE/H/4465/002	PL 33155/0044	RIVOPHARM UK LTD	UK
Onexila XL 40 mg prolonged-release tablets	DE/H/4465/003	PL 33155/0045	RIVOPHARM UK LTD	UK
Onexila XL 80 mg prolonged-release tablets	DE/H/4465/004	PL 33155/0046	RIVOPHARM UK LTD	UK
Orionox 10 mg depottabletter	SE/H/1493/02	12-9220	ORION CORPORATION	NO
Orionox 15 mg depottabletter	SE/H/1493/03	12-9221	ORION CORPORATION	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
Orionox 20 mg depottabletter	SE/H/1493/04	12-9222	ORION CORPORATION	NO
Orionox 30 mg depottabletter	SE/H/1493/05	12-9223	ORION CORPORATION	NO
Orionox 40 mg depottabletter	SE/H/1493/06	12-9224	ORION CORPORATION	NO
Orionox 5 mg depottabletter	SE/H/1493/01	12-9219	ORION CORPORATION	NO
Orionox 60 mg depottabletter	SE/H/1493/07	12-9225	ORION CORPORATION	NO
Orionox 80 mg depottabletter	SE/H/1493/08	12-9226	ORION CORPORATION	NO
Orionox, depottabletter	SE/H/1493/01	51488	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/02	51489	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/03	51490	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/04	51491	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/05	51492	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/06	51493	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/07	51494	ORION CORPORATION	DK
Orionox, depottabletter	SE/H/1493/08	51495	ORION CORPORATION	DK
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318093	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318105	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318117	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318129	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318131	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318143	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318156	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318168	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318170	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318396	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318408	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318410	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318422	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318434	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318446	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318459	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318461	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318473	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318485	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318497	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318509	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 10 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318511	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318182	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318194	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318206	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318218	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318220	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318232	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318244	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318257	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318269	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318523	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318535	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318547	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318550	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318562	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318574	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318586	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318598	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318600	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318612	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318624	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318636	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 20 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318648	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318016	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318028	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318030	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318042	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318055	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318067	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318079	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318081	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318271	AUROBINDO PHARMA (ITALIA) S.R.L.	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
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318283	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318295	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318307	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318319	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318321	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318333	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318345	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318358	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318360	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318372	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Aurobindo 5 mg compresse a rilascio prolungato	DE/H/5356/001-008/DC	042318384	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio	IT/H/558/001	044919013	BRUNO FARMACEUTICI	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
prolungato				
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919025	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919037	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919049	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919052	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919064	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919076	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919088	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio prolungato	IT/H/558/001	044919090	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 10 mg compresse a rilascio	IT/H/558/001	044919102	BRUNO FARMACEUTICI	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
prolungato				
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919114	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919126	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919138	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919140	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919153	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919165	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919177	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919189	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio	IT/H/558/002	044919191	BRUNO FARMACEUTICI	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
prolungato				
Ossicodone Bruno Farmaceutici, 20 mg compresse a rilascio prolungato	IT/H/558/002	044919203	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919215	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919227	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919239	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919241	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919254	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919266	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919278	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio	IT/H/558/003	044919280	BRUNO FARMACEUTICI	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
prolungato				
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919292	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 40 mg compresse a rilascio prolungato	IT/H/558/003	044919304	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919316	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919328	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919330	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919342	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919355	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919367	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio	IT/H/558/004	044919379	BRUNO FARMACEUTICI	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
prolungato				
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919381	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919393	BRUNO FARMACEUTICI	IT
Ossicodone Bruno Farmaceutici, 80 mg compresse a rilascio prolungato	IT/H/558/004	044919405	BRUNO FARMACEUTICI	IT
Ossicodone Ethypharm, 10 mg compresse a rilascio prolungato	DE/H/4080/002	043428022	ETHYPHARM	IT
Ossicodone Ethypharm, 20 mg compresse a rilascio prolungato	DE/H/4080/003	043428034	ETHYPHARM	IT
Ossicodone Ethypharm, 40 mg compresse a rilascio prolungato	DE/H/4080/004	043428046	ETHYPHARM	IT
Ossicodone Ethypharm, 5mg compresse a rilascio prolungato	DE/H/4080/001	043428010	ETHYPHARM	IT
Ossicodone Molteni 10 mg/ml, soluzione iniettabile o per infusione	UK/H/5881/001	043927019	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ossicodone Molteni 10 mg/ml, soluzione iniettabile o per infusione	UK/H/5881/001	043927021	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ossicodone Molteni 10 mg/ml, soluzione iniettabile o per infusione	UK/H/5881/001	043927033	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ossicodone Molteni 10 mg/ml, soluzione	UK/H/5881/001	043927045	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
iniettabile o per infusione			ESERCIZIO S.P.A.	
Ossicodone Molteni 50 mg/ml, soluzione iniettabile o per infusione	UK/H/5881/002	043927058	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
OXANEST® 10 mg/ml - injektioneste, liuos	not available	6646	TAKEDA OY	FI
Oxeltra 10mg Prolonged-Release Tablets	UK/H/6001/02/MR	29831/0631	WOCKHARDT UK LTD	UK
Oxeltra 15mg Prolonged-Release Tablets	UK/H/6001/03/MR	29831/0632	WOCKHARDT UK LTD	UK
Oxeltra 20mg Prolonged-Release Tablets	UK/H/6001/04/MR	29831/0633	WOCKHARDT UK LTD	UK
Oxeltra 30mg Prolonged-Release Tablets	UK/H/6001/05/MR	29831/0634	WOCKHARDT UK LTD	UK
Oxeltra 40mg Prolonged-Release Tablets	UK/H/6001/06/MR	29831/0635	WOCKHARDT UK LTD	UK
Oxeltra 5mg Prolonged-Release Tablets	UK/H/6001/01/MR	29831/0630	WOCKHARDT UK LTD	UK
Oxeltra 60mg Prolonged-Release Tablets	UK/H/6001/07/MR	29831/0636	WOCKHARDT UK LTD	UK
Oxeltra 80mg Prolonged-Release Tablets	UK/H/6001/08/MR	29831/0637	WOCKHARDT UK LTD	UK
Oxicodona Ethypharm 10 mg comprimidos de liberación prolongada	DE/H/4081/002	80003	ETHYPHARM	ES
Oxicodona Ethypharm 20 mg comprimidos de liberación prolongada	DE/H/4081/003	80004	ETHYPHARM	ES
Oxicodona Ethypharm 40 mg comprimidos de liberación prolongada	DE/H/4081/004	80005	ETHYPHARM	ES
Oxicodona Ethypharm 5 mg comprimidos de liberación prolongada	DE/H/4081/001	80002	ETHYPHARM	ES
Oxicodona Mithridatum	not available	5617121	MITHRIDATUM LIMITED	PT

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 comprimidos de libertação prolongada				
Oxicodona Mithridatum 10 mg comprimidos de libertação prolongada	not available	5617154	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 15 mg comprimidos de libertação prolongada	not available	5617329	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 15 mg comprimidos de libertação prolongada	not available	5617337	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 20 mg comprimidos de libertação prolongada	not available	5617238	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 20 mg comprimidos de libertação prolongada	not available	5617261	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 30 mg comprimidos de libertação prolongada	not available	5617022	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 30 mg comprimidos de libertação prolongada	not available	5617048	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 40 mg comprimidos de libertação prolongada	not available	5617600	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 40 mg comprimidos de libertação prolongada	not available	5617626	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 5 mg comprimidos de libertação prolongada	not available	5616834	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 5 mg comprimidos de libertação prolongada	not available	5616867	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum	not available	5616933	MITHRIDATUM LIMITED	PT

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
60 mg comprimidos de libertação prolongada				
Oxicodona Mithridatum 60 mg comprimidos de libertação prolongada	not available	5616958	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 80 mg comprimidos de libertação prolongada	not available	5617477	MITHRIDATUM LIMITED	PT
Oxicodona Mithridatum 80 mg comprimidos de libertação prolongada	not available	5617527	MITHRIDATUM LIMITED	PT
Oxicodona Sandoz 10 mg comprimidos de liberación prolongada EFG	DE/H/1154/002	70687	SANDOZ FARMACÉUTICA, S.A.	ES
Oxicodona Sandoz 20 mg comprimidos de liberación prolongada EFG	DE/H/1154/003	70686	SANDOZ FARMACÉUTICA, S.A.	ES
Oxicodona Sandoz 40 mg comprimidos de liberacion prolongada EFG	DE/H/1154/004	76865	SANDOZ FARMACÉUTICA, S.A.	ES
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/01	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/02	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/04	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/05	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40	DE/H/3293/001	5047/2012/06	S.C. SANDOZ S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimate cu eliberare prelungită				
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/07	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/08	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/09	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/10	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/11	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/12	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/13	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/14	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 40 mg, comprimate cu eliberare prelungită	DE/H/3293/001	5047/2012/03	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/01	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/02	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60	DE/H/3293/002	5048/2012/03	S.C. SANDOZ S.R.L.	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimate cu eliberare prelungită				
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/04	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/05	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/06	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/07	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/08	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/09	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/10	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/11	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/12	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/13	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 60 mg, comprimate cu eliberare prelungită	DE/H/3293/002	5048/2012/14	S.C. SANDOZ S.R.L.	RO
Oxicodona Sandoz 80 mg	DE/H/1154/006	76867	SANDOZ FARMACÉUTICA,	ES

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de liberacion prolongada EFG			S.A.	
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/01	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/02	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/03	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/04	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/05	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/06	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/07	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/08	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/09	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/10	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/11	S.C. SANDOZ S.R.L.	RO

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/12	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/13	S.C. SANDOZ S.R.L.	RO
OXICODONĂ SANDOZ 80 mg, comprimate cu eliberare prelungită	DE/H/3293/003	5049/2012/14	S.C. SANDOZ S.R.L.	RO
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060158	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060160	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060172	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060184	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060196	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060208	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060210	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060222	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060234	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060259	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060261	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060273	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060285	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060246	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060727	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 10 mg compresse a rilascio prolungato	NL/H/3192/002	042060778	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060297	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060309	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060311	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060323	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060335	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060347	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060350	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060362	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060374	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060386	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060398	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060400	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060412	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060424	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060739	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 20 mg compresse a rilascio prolungato	NL/H/3192/003	042060780	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060436	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060448	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060451	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060475	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060487	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060499	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060501	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060513	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060525	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060537	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060549	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060552	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060564	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060463	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060741	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 40 mg compresse a rilascio prolungato	NL/H/3192/004	042060792	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060018	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060020	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060032	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060044	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060057	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060069	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060071	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060083	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060095	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060107	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060119	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060121	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060133	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060145	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060715	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 5 mg compresse a rilascio prolungato	NL/H/3192/001	042060766	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060588	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060590	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060602	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060626	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060638	ACCORD HEALTHCARE LIMITED	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
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060640	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060653	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060665	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060677	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060691	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060703	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060689	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060614	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060754	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Accord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060804	ACCORD HEALTHCARE LIMITED	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263017/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263029/M	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263031/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263043/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263056/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263068/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263070/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263082/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263094/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263106/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263118/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 40 mg compresse a rilascio prolungato	DE/H/3293/001	041263120/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263183/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263207/M	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263219/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263144/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263132/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263169/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263157/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263171/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263195/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263233/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263221/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 60 mg compresse a rilascio prolungato	DE/H/3293/002	041263245/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263258/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263260/M	SANDOZ S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263272/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263284/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263296/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263308/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263322/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263334/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263346/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263310/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263359/M	SANDOZ S.P.A.	IT
Oxicodone Sandoz 80 mg compresse a rilascio prolungato	DE/H/3293/003	041263361/M	SANDOZ S.P.A.	IT
OxicodoneAccord 80 mg compresse a rilascio prolungato	NL/H/3192/005	042060576	ACCORD HEALTHCARE LIMITED	IT
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/024	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s	DE/H/2183/002	H/10/01199/020	LANNACHER HEILMITTEL	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
podaljsanim sproscanjem			GES.M.B.H.,	
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/025	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/026	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/027	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/021	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/028	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/022	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/023	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/029	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/030	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/031	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/032	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/033	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/034	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/035	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/036	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/037	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 10 mg tablete s podaljsanim sproscanjem	DE/H/2183/002	H/10/01199/038	LANNACHER HEILMITTEL GES.M.B.H.,	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/043	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/039	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/044	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/045	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/046	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/040	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/047	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/041	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/042	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/048	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/049	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/050	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/051	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/052	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/053	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/054	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/055	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 20 mg tablete s	DE/H/2183/003	H/10/01199/056	LANNACHER HEILMITTEL	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
podaljšanim sproščanjem			GES.M.B.H.,	
Oxidol 20 mg tablete s podaljšanim sproščanjem	DE/H/2183/003	H/10/01199/057	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/062	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/058	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/063	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/064	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/065	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/059	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/066	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/060	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/061	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/067	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/068	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/069	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/070	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/071	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/072	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/073	LANNACHER HEILMITTEL GES.M.B.H.,	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/074	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/075	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 40 mg tablete s podaljšanim sproščanjem	DE/H/2183/004	H/10/01199/076	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/005	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/001	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/006	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/007	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/008	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/002	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/009	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/003	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/004	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/010	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/011	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/012	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/013	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/014	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s	DE/H/2183/001	H/10/01199/015	LANNACHER HEILMITTEL	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
podaljšanim sproščanjem			GES.M.B.H.,	
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/016	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/017	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/018	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 5 mg tablete s podaljšanim sproščanjem	DE/H/2183/001	H/10/01199/019	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/081	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/077	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/082	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/083	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/084	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/078	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/085	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/079	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/080	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/086	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/087	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/088	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/089	LANNACHER HEILMITTEL GES.M.B.H.,	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/090	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/091	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/092	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/093	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/094	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidol 80 mg tablete s podaljsanim sproscanjem	DE/H/2183/005	H/10/01199/095	LANNACHER HEILMITTEL GES.M.B.H.,	SI
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/05	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/06	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/07	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/08	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/09	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/10	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/11	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/12	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/13	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/14	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/15	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/16	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/17	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/18	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 10 mg comprimate cu eliberare prelungită	DE/H/2182/002	6281/2014/19	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/05	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/06	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/07	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/08	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/09	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/10	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/11	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/12	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/13	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/14	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/15	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/16	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/17	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/18	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 20 mg comprimate cu eliberare prelungită	DE/H/2182/003	6282/2014/19	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/05	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/06	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/07	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/08	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/09	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/10	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/11	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/12	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/13	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/14	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/15	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/16	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/17	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/18	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 40 mg comprimate cu eliberare prelungită	DE/H/2182/004	6283/2014/19	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/05	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/06	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/07	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/08	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/09	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/10	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/11	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/12	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/13	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/14	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/15	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/16	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/17	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/18	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 5 mg comprimate cu eliberare prelungită	DE/H/2182/001	6280/2014/19	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/01	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/02	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/03	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/04	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/05	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/06	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/07	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/08	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/09	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/10	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/11	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/12	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/13	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/14	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/15	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/16	LANNACHER HEILMITTEL GES.M.B.H.,	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/17	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/18	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxidolor 80 mg comprimate cu eliberare prelungită	DE/H/2182/005	6284/2014/19	LANNACHER HEILMITTEL GES.M.B.H.,	RO
Oxikodon Actavis 10 mg kapslar, hårda	SE/H/1226/002	47450	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Actavis 20 mg kapslar, hårda	SE/H/1226/003	47451	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Actavis 5 mg kapslar, hårda	SE/H/1226/001	47449	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Acino 10 mg depottabletter	DE/H/1153/002	25804	ACINO AG	SE
Oxikodon Depot Acino 20 mg depottabletter	DE/H/1153/003	25805	ACINO AG	SE
Oxikodon Depot Acino 40 mg depottabletter	DE/H/1153/004	25806	ACINO AG	SE
Oxikodon Depot Acino 80 mg depottabletter	DE/H/1153/005	25807	ACINO AG	SE
Oxikodon Depot Actavis 10 mg depottabletter	SE/H/1313/002	48600	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Actavis 15 mg depottabletter	SE/H/1313/003	48601	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Actavis 20 mg depottabletter	SE/H/1313/004	48602	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Actavis 30 mg depottabletter	SE/H/1313/005	48603	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Actavis 40 mg depottabletter	SE/H/1313/006	48604	ACTAVIS GROUP PTC EHF.	SE
Oxikodon Depot Actavis 60 mg depottabletter	SE/H/1313/007	48605	ACTAVIS GROUP PTC EHF.	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
Oxikodon Depot Actavis 80 mg depottabletter	SE/H/1313/008	48606	ACTAVIS GROUP PTC EHF.	SE
Oxpian 10 mg Prolonged-release Tablets	SE/H/1549/002	PL 00289/2028	TEVA UK LIMITED	UK
Oxpian 15 mg Prolonged-release Tablets	SE/H/1549/003	PL 00289/2029	TEVA UK LIMITED	UK
Oxpian 20 mg Prolonged-release Tablets	SE/H/1549/004	PL 00289/2030	TEVA UK LIMITED	UK
Oxpian 30 mg Prolonged-release Tablets	SE/H/1549/005	PL 00289/2031	TEVA UK LIMITED	UK
Oxpian 40 mg Prolonged-release Tablets	SE/H/1549/006	PL 00289/2032	TEVA UK LIMITED	UK
Oxpian 5 mg Prolonged-release Tablets	SE/H/1549/001	PL 00289/2027	TEVA UK LIMITED	UK
Oxpian 60 mg Prolonged-release Tablets	SE/H/1549/007	PL 00289/2033	TEVA UK LIMITED	UK
Oxpian 80 mg Prolonged-release Tablets	SE/H/1549/008	PL 00289/2034	TEVA UK LIMITED	UK
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
Oxycan® uno 80 mg Retardtabletten	DE/H/3655/004	88012.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon AL 10 mg Hartkapseln	DE/H/4774/002	98061.00.00	ALIUD PHARMA GMBH	DE
Oxycodon AL 20 mg	DE/H/4774/003	98062.00.00	ALIUD 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
Hartkapseln				
Oxycodon AL 5 mg Hartkapseln	DE/H/4774/001	98060.00.00	ALIUD PHARMA GMBH	DE
Oxycodon Aristo 1 mg/ml Lösung zum Einnehmen	DE/H/4654/001	96749.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon Aristo 10 mg/ml Lösung zum Einnehmen	DE/H/4654/002	96750.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon dura 10 mg Retardtabletten	DE/H/2439/002	1-30540	MYLAN DURA GMBH	AT
Oxycodon dura 20 mg Retardtabletten	DE/H/2439/003	1-30541	MYLAN DURA GMBH	AT
Oxycodon dura 40 mg Retardtabletten	DE/H/2439/004	1-30542	MYLAN DURA GMBH	AT
Oxycodon dura 5 mg Retardtabletten	DE/H/2439/001	1-30539	MYLAN DURA GMBH	AT
Oxycodon dura 80 mg Retardtabletten	DE/H/2439/005	1-30543	MYLAN DURA GMBH	AT
Oxycodon G.L. 10 mg-Retardtabletten	DE/H/2183/002	1-29628	G.L. PHARMA GMBH	AT
Oxycodon G.L. 20 mg-Retardtabletten	DE/H/2183/003	1-29629	G.L. PHARMA GMBH	AT
Oxycodon G.L. 40 mg-Retardtabletten	DE/H/2183/004	1-29630	G.L. PHARMA GMBH	AT
Oxycodon G.L. 5 mg-Retardtabletten	DE/H/2183/001	1-29627	G.L. PHARMA GMBH	AT
Oxycodon G.L. 80 mg-Retardtabletten	DE/H/2183/005	1-29631	G.L. PHARMA GMBH	AT
Oxycodon HCl 10 mg Teva, tabletten met verlengde afgifte	DE/H/1395/002	RVG 101359	TEVA NEDERLAND B.V.	NL
Oxycodon HCl 20 mg Teva, tabletten met verlengde afgifte	DE/H/0790/001	RVG 34224	TEVA NEDERLAND B.V.	NL
Oxycodon HCl 2x täglich	not available	97768.00.00	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
Hormosan 10 mg Retardtabletten				
Oxycodon HCl 2x täglich Hormosan 20 mg Retardtabletten	not available	97769.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon HCl 2x täglich Hormosan 40 mg Retardtabletten	not available	97771.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon HCl 2x täglich Hormosan 5 mg Retardtabletten	not available	97767.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon HCl 2x täglich Hormosan 80 mg Retardtabletten	not available	97773.00.00	HORMOSAN PHARMA GMBH	DE
Oxycodon HCl 40 mg Teva, tabletten met verlengde afgifte	DE/H/0790/002	RVG 34225	TEVA NEDERLAND B.V.	NL
Oxycodon HCl 5 mg Teva, tabletten met verlengde afgifte	DE/H/2947/001	RVG 108257	TEVA NEDERLAND B.V.	NL
Oxycodon HCl 80 mg Teva, tabletten met verlengde afgifte	DE/H/0790/003	RVG 34226	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Accord 10 mg depottabletit	NL/H/3192/002	29744	ACCORD HEALTHCARE LIMITED	FI
Oxycodon HCl Accord 20 mg depottabletit	NL/H/3192/003	29745	ACCORD HEALTHCARE LIMITED	FI
Oxycodon HCl Accord 40 mg depottabletit	NL/H/3192/004	29746	ACCORD HEALTHCARE LIMITED	FI
Oxycodon HCl Accord 5 mg depottabletit	NL/H/3192/001	29743	ACCORD HEALTHCARE LIMITED	FI
Oxycodon HCl Accord 80 mg depottabletit	DE/H/2440/005	29747	ACCORD HEALTHCARE LIMITED	FI
Oxycodon HCl Apotex 10 mg, tablet met verlengde afgifte	NL/H/2521/002	RVG 104272	APOTEX EUROPE B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon HCl Apotex 20 mg, tablet met verlengde afgifte	NL/H/2521/003	RVG 104273	APOTEX EUROPE B.V.	NL
Oxycodon HCl Apotex 40 mg, tablet met verlengde afgifte	NL/H/2521/004	RVG 104274	APOTEX EUROPE B.V.	NL
Oxycodon HCl Apotex 5 mg, tablet met verlengde afgifte	NL/H/2521/001	RVG 104271	APOTEX EUROPE B.V.	NL
Oxycodon HCl Apotex 80 mg, tablet met verlengde afgifte	NL/H/2521/005	RVG 104275	APOTEX EUROPE B.V.	NL
Oxycodon HCl Aristo® akut 10 mg Filmtabletten Wirkstoff: Oxycodonhydrochlorid	not available	88371.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon HCl Aristo® akut 5 mg Filmtabletten Wirkstoff: Oxycodonhydrochlorid	not available	88370.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon HCl Aurobindo 10 mg, capsules hard	NL/H/4429/001-003	RVG 111251	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo 20 mg, capsules hard	NL/H/4429/001-003	RVG 111252	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo 5 mg, capsules hard	NL/H/4429/001-003	RVG 111250	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 10 mg, tabletten met gereguleerde afgifte	DE/H/5356/002	RVG 112717	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 15 mg, tabletten met gereguleerde afgifte	DE/H/5356/003	RVG 112718	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 20 mg, tabletten met gereguleerde afgifte	DE/H/5356/004	RVG 112719	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo	DE/H/5356/005	RVG 112720	AUROBINDO PHARMA B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Retard 30 mg, tabletten met gereguleerde afgifte				
Oxycodon HCl Aurobindo Retard 40 mg, tabletten met gereguleerde afgifte	DE/H/5356/006	RVG 112721	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 5 mg, tabletten met gereguleerde afgifte	DE/H/5356/001	RVG 112716	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 60 mg, tabletten met gereguleerde afgifte	DE/H/5356/007	RVG 112723	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl Aurobindo Retard 80 mg, tabletten met gereguleerde afgifte	DE/H/5356/008	RVG 112724	AUROBINDO PHARMA B.V.	NL
Oxycodon HCl CF 10 mg, harde capsules	DE/H/4774/002	RVG 119738	CENTRAFARM B.V.	NL
Oxycodon HCl CF 20 mg, harde capsules	DE/H/4774/003	RVG 119739	CENTRAFARM B.V.	NL
Oxycodon HCl CF 5 mg, harde capsules	DE/H/4774/001	RVG 119737	CENTRAFARM B.V.	NL
Oxycodon HCl G.L. 10 mg filmomhulde tabletten	DE/H/2669/002	106669	G.L. PHARMA GMBH	NL
Oxycodon HCl G.L. 20 mg filmomhulde tabletten	DE/H/2669/004	RVG 118173	G.L. PHARMA GMBH	NL
Oxycodon HCl G.L. 5 mg filmomhulde tabletten	DE/H/2669/001	106668	G.L. PHARMA GMBH	NL
Oxycodon HCl Lannacher 10 mg forðatöflur	DE/H/2182/002	IS/1/09/045/02	LANNACHER HEILMITTEL GES.M.B.H.,	IS
Oxycodon HCl Lannacher 10 mg Retardtabletten	DE/H/2182/002	76277.00.00	LANNACHER HEILMITTEL GES.M.B.H.,	DE
Oxycodon HCl Lannacher 10 mg tabletten met verlengde afgifte	DE/H/2183/002	RVG 104139	LANNACHER HEILMITTEL GES.M.B.H.,	NL
Oxycodon HCl Lannacher	DE/H/2182/003	IS/1/09/045/03	LANNACHER HEILMITTEL	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
20 mg forðatöflur			GES.M.B.H.,	
Oxycodon HCl Lannacher 20 mg Retardtabletten	DE/H/2182/003	76278.00.00	LANNACHER HEILMITTEL GES.M.B.H.,	DE
Oxycodon HCl Lannacher 20 mg tabletten met verlengde afgifte	DE/H/2183/003	RVG 104140	LANNACHER HEILMITTEL GES.M.B.H.,	NL
Oxycodon HCl Lannacher 40 mg forðatöflur	DE/H/2182/004	IS/1/09/045/04	LANNACHER HEILMITTEL GES.M.B.H.,	IS
Oxycodon HCl Lannacher 40 mg Retardtabletten	DE/H/2182/004	76279.00.00	LANNACHER HEILMITTEL GES.M.B.H.,	DE
Oxycodon HCl Lannacher 40 mg tabletten met verlengde afgifte	DE/H/2183/004	RVG 104141	LANNACHER HEILMITTEL GES.M.B.H.,	NL
Oxycodon HCl Lannacher 5 mg forðatöflur	DE/H/2182/001	IS/1/09/045/01	LANNACHER HEILMITTEL GES.M.B.H.,	IS
Oxycodon HCl Lannacher 5 mg Retardtabletten	DE/H/2182/001	76276.00.00	LANNACHER HEILMITTEL GES.M.B.H.,	DE
Oxycodon HCl Lannacher 5 mg tabletten met verlengde afgifte	DE/H/2183/001	RVG 104137	LANNACHER HEILMITTEL GES.M.B.H.,	NL
Oxycodon HCl Lannacher 80 mg forðatöflur	DE/H/2182/005	IS/1/09/045/05	LANNACHER HEILMITTEL GES.M.B.H.,	IS
Oxycodon HCl Lannacher 80 mg Retardtabletten	DE/H/2182/005	76280.00.00	LANNACHER HEILMITTEL GES.M.B.H.,	DE
Oxycodon HCl Lannacher 80 mg tabletten met verlengde afgifte	DE/H/2183/005	RVG 104142	LANNACHER HEILMITTEL GES.M.B.H.,	NL
Oxycodon HCl retard CF 10 mg, tabletten met verlengde afgifte	DE/H/3641/002	RVG 112599	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 15 mg, tabletten met verlengde afgifte	DE/H/3641/003	RVG 112600	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 20 mg, tabletten met verlengde afgifte	DE/H/3641/004	RVG 112601	CENTRAFARM B.V.	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon HCl retard CF 30 mg, tabletten met verlengde afgifte	DE/H/3641/005	RVG 112602	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 40 mg, tabletten met verlengde afgifte	DE/H/3641/006	RVG 112603	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 5 mg, tabletten met verlengde afgifte	DE/H/3641/001	RVG 112598	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 60 mg, tabletten met verlengde afgifte	DE/H/3641/007	RVG 112604	CENTRAFARM B.V.	NL
Oxycodon HCl retard CF 80 mg, tabletten met verlengde afgifte	DE/H/3641/008	RVG 112605	CENTRAFARM B.V.	NL
Oxycodon HCl Retard Teva 10 mg, tabletten met verlengde afgifte	SE/H/1549/002	RVG 117834	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 15 mg, tabletten met verlengde afgifte	SE/H/1549/003	RVG 117835	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 20 mg, tabletten met verlengde afgifte	SE/H/1549/004	RVG 117836	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 30 mg, tabletten met verlengde afgifte	SE/H/1549/005	RVG 117837	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 40 mg, tabletten met verlengde afgifte	SE/H/1549/006	RVG 117838	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 5 mg, tabletten met verlengde afgifte	SE/H/1549/001	RVG 117833	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Retard Teva 60 mg, tabletten met verlengde afgifte	SE/H/1549/007	RVG 117839	TEVA NEDERLAND B.V.	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon HCl Retard Teva 80 mg, tabletten met verlengde afgifte	SE/H/1549/008	RVG 117840	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Sandoz 10 mg, harde capsules	DE/H/4502/002	RVG 121589	SANDOZ B.V.	NL
Oxycodon HCl Sandoz 20 mg, harde capsules	DE/H/4502/003	RVG 121586	SANDOZ B.V.	NL
Oxycodon HCl Sandoz 5 mg, harde capsules	DE/H/4502/001	RVG 121571	SANDOZ B.V.	NL
OXYCODON HCl SANDOZ RETARD 10 MG, TABLETTEN MET VERLENGDE AFGIFTE	DE/H/1154/002	RVG 100649	SANDOZ B.V.	NL
OXYCODON HCl SANDOZ RETARD 20 MG, TABLETTEN MET VERLENGDE AFGIFTE	DE/H/1154/003	RVG 100654	SANDOZ B.V.	NL
Oxycodon HCl Sandoz retard 40 mg, tabletten met verlengde afgifte	DE/H/1154/004	RVG 109612	SANDOZ B.V.	NL
OXYCODON HCl SANDOZ RETARD 5 MG, TABLETTEN MET VERLENGDE AFGIFTE	DE/H/1154/001	RVG 100644	SANDOZ B.V.	NL
Oxycodon HCl Sandoz retard 60 mg, tabletten met verlengde afgifte	DE/H/1154/005	RVG 109613	SANDOZ B.V.	NL
Oxycodon HCl Sandoz retard 80 mg, tabletten met verlengde afgifte	DE/H/1154/006	RVG 109614	SANDOZ B.V.	NL
Oxycodon HCl Teva 10 mg, capsules, hard	SE/H/1423/002	RVG 115195	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Teva 20 mg, capsules, hard	SE/H/1423/003	RVG 115196	TEVA NEDERLAND B.V.	NL
Oxycodon HCl Teva 5 mg, capsules, hard	SE/H/1423/001	RVG 115194	TEVA NEDERLAND B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OXYCODON LANNACHER 10 MG TABLETY S PRODLOUŽENÝM UVOLNOVÁNÍM	DE/H/2182/002	65/934/10-C	G.L. PHARMA GMBH	CZ
OXYCODON LANNACHER 20 MG TABLETY S PRODLOUŽENÝM UVOLŇOVÁNÍM	DE/H/2182/003	65/935/10-C	G.L. PHARMA GMBH	CZ
OXYCODON LANNACHER 40 MG TABLETY S PRODLOUŽENÝM UVOLŇOVÁNÍM	DE/H/2182/004	65/936/10-C	G.L. PHARMA GMBH	CZ
OXYCODON LANNACHER 80 MG TABLETY S PRODLOUŽENÝM UVOLNOVÁNÍM	DE/H/2182/005	65/937/10-C	G.L. PHARMA GMBH	CZ
Oxycodon PAINBREAK 10 mg Retardtabletten	not available	98655.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 20 mg Retardtabletten	not available	98656.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 30 mg Retardtabletten	not available	98657.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 40 mg Retardtabletten	not available	98658.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 5 mg Retardtabletten	not available	98654.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 60 mg Retardtabletten	not available	98659.00.00	PB PHARMA GMBH	DE
Oxycodon PAINBREAK 80 mg Retardtabletten	not available	98660.00.00	PB PHARMA GMBH	DE
Oxycodon Retard AB 10 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457404	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 10 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457413	AUROBINDO PHARMA B.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon Retard AB 10 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457422	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 10 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457431	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 15 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457457	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 15 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457440	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 15 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457475	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 15 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457466	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 20 mg, tabletten met verlengde afgifte	SE/H/1313/004	BE457511	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 20 mg, tabletten met verlengde afgifte	SE/H/1313/004	BE457484	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 20 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457493	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 20 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457502	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 30 mg, tabletten met verlengde afgifte	SE/H/1313/005	BE457546	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 30 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457555	AUROBINDO PHARMA B.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon Retard AB 30 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457537	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 30 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457520	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 40 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457591	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 40 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457564	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 40 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457582	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 40 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457573	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 5 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457360	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 5 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457395	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 5 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457386	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 5 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457377	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 60 mg, tabletten met verlengde afgifte	SE/H/1313/007	BE457635	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 60 mg, tabletten met verlengde afgifte	SE/H/1313/007	BE457626	AUROBINDO PHARMA B.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon Retard AB 60 mg, tabletten met verlengde afgifte	SE/H/1313/007	BE457617	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 60 mg, tabletten met verlengde afgifte	SE/H/1313/007	BE457600	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 80 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457671	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 80 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457662	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 80 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457653	AUROBINDO PHARMA B.V.	BE
Oxycodon Retard AB 80 mg, tabletten met verlengde afgifte	DE/H/5356/001-008/DC	BE457644	AUROBINDO PHARMA B.V.	BE
Oxycodon Sandoz 10 mg tabletten met verlengde afgifte	DE/H/1154/002	BE 335307	SANDOZ N.V.	BE
Oxycodon Sandoz 10 mg tabletten met verlengde afgifte	DE/H/1154/002	BE 335316	SANDOZ N.V.	BE
Oxycodon Sandoz 20 mg tabletten met verlengde afgifte	DE/H/1154/003	BE 335334	SANDOZ N.V.	BE
Oxycodon Sandoz 20 mg tabletten met verlengde afgifte	DE/H/1154/003	BE 335325	SANDOZ N.V.	BE
Oxycodon Sandoz 40 mg tabletten met verlengde afgifte	DE/H/1154/004	BE427104	SANDOZ N.V.	BE
Oxycodon Sandoz 40 mg tabletten met verlengde afgifte	DE/H/1154/004	BE427113	SANDOZ N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon Sandoz 5 mg tabletten met verlengde afgifte	DE/H/1154/001	BE 335282	SANDOZ N.V.	BE
Oxycodon Sandoz 5 mg tabletten met verlengde afgifte	DE/H/1154/001	BE 335291	SANDOZ N.V.	BE
Oxycodon Sandoz 80 mg tabletten met verlengde afgifte	DE/H/1154/006	BE427156	SANDOZ N.V.	BE
Oxycodon Sandoz 80 mg tabletten met verlengde afgifte	DE/H/1154/006	BE427147	SANDOZ N.V.	BE
Oxycodon Sandoz Retard 10 mg tablety s prodlouženým uvolňováním	DE/H/1419/002	65/244/09-C	SANDOZ S.R.O.	CZ
Oxycodon Sandoz Retard 40 mg tablety s prodlouženým uvolňováním	DE/H/0789/002	65/620/07-C	SANDOZ S.R.O.	CZ
Oxycodon Sandoz Retard 80 mg tablety s prodlouženým uvolňováním	DE/H/0789/003	65/621/07-C	SANDOZ S.R.O.	CZ
Oxycodon Stada, 10 mg, kapsułki, twarde	DE/H/4774/002	24321	STADA ARZNEIMITTEL AG	PL
Oxycodon Stada, 20 mg, kapsułki, twarde	DE/H/4774/003	24322	STADA ARZNEIMITTEL AG	PL
Oxycodon Stada, 5 mg, kapsułki, twarde	DE/H/4774/001	24320	STADA ARZNEIMITTEL AG	PL
Oxycodon Teva 10 mg Retardtabletten	SE/H/1549/001	137130	TEVA B.V	AT
Oxycodon Teva 20 mg Retardtabletten	SE/H/1549/002	137132	TEVA B.V	AT
Oxycodon Teva 30 mg Retardtabletten	SE/H/1549/003	137133	TEVA B.V	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon Teva 40 mg Retardtabletten	SE/H/1549/004	137134	TEVA B.V	AT
Oxycodon Teva 60 mg Retardtabletten	SE/H/1549/005	137135	TEVA B.V	AT
Oxycodon Teva 80 mg Retardtabletten	SE/H/1549/008	137136	TEVA B.V	AT
OXYCODONE MEDAC 10 mg/ml, solution injectable/pour perfusion	UK/H/5881/001	34009 301 033 6 4	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	FR
OXYCODONE MEDAC 10 mg/ml, solution injectable/pour perfusion	UK/H/5881/001	34009 300 922 7 9	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	FR
OXYCODONE MEDAC 10 mg/ml, solution injectable/pour perfusion	UK/H/5881/001	34009 300 922 8 6	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	FR
OXYCODONE MEDAC 10 mg/ml, solution injectable/pour perfusion	UK/H/5881/001	34009 300 922 9 3	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	FR
Oxycodone ”Teva”, hårde kapsler	SE/H/1423/001	54001	TEVA DENMARK A/S	DK
Oxycodone ”Teva”, hårde kapsler	SE/H/1423/002	54002	TEVA DENMARK A/S	DK
Oxycodone ”Teva”, hårde kapsler	SE/H/1423/003	54003	TEVA DENMARK A/S	DK
Oxycodone ”Vitabalans”, filmovertrukne tabletter	SE/H/1119/001	48874	VITABALANS OY	DK
Oxycodone ”Vitabalans”, filmovertrukne tabletter	SE/H/1119/002	48875	VITABALANS OY	DK
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 9 3	ACCORD HEALTHCARE FRANCE 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
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 0 9	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 1 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 2 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 4 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 5 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 6 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 7 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 8 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 3 1	ACCORD HEALTHCARE FRANCE 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
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 4 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 5 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 6 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 8 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 756 9 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 0 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 1 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 2 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 3 9	ACCORD HEALTHCARE FRANCE 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
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 4 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 5 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 6 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 7 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 8 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 757 9 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 0 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 1 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 2 1	ACCORD HEALTHCARE FRANCE 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
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 9 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 0 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 1 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 3 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 4 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 5 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 6 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 7 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 8 2	ACCORD HEALTHCARE FRANCE 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
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 3 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 4 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 5 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 7 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 300 758 8 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 753 8 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 753 9 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 1 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 2 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 3 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP	not available	34009 300 754 4 9	ACCORD HEALTHCARE	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
5 mg, comprimé pelliculé à libération prolongée			FRANCE SAS	
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 5 6	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 6 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 7 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 8 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 754 9 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 0 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 1 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 300 755 2 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 759 9 9	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 1 9	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé	not available	34009 300 760 2 6	ACCORD HEALTHCARE FRANCE 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
pelliculé à libération prolongée				
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 3 3	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 4 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 5 7	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 6 4	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 7 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 8 8	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 760 9 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 761 0 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé	not available	34009 300 761 1 8	ACCORD HEALTHCARE FRANCE 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
pelliculé à libération prolongée				
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 761 2 5	ACCORD HEALTHCARE FRANCE SAS	FR
OXYCODONE ACCORD LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 300 761 3 2	ACCORD HEALTHCARE FRANCE SAS	FR
Oxycodone Actavis 10 mg depottabletit	FI/H/0965/002	31118	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 10 mg harde kapsler	SE/H/1226/002	11-8801	ACTAVIS GROUP PTC EHF.	NO
Oxycodone Actavis 15 mg depottabletit	FI/H/0965/003	31119	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 20 mg depottabletit	FI/H/0965/004	31120	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 20 mg harde kapsler	SE/H/1226/003	11-8802	ACTAVIS GROUP PTC EHF.	NO
OXYCODONE ACTAVIS 30 MG DEPOTTABLETIT	FI/H/0965/005	31121	ACTAVIS GROUP PTC EHF.	FI
OXYCODONE ACTAVIS 40 MG DEPOTTABLETIT	FI/H/0965/006	31122	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 5 mg capsules, hard	SE/H/1226/001	20130044	ACTAVIS GROUP PTC EHF.	BG
Oxycodone Actavis 5 mg depottabletit	FI/H/0965/001	31117	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 5 mg harde kapsler	SE/H/1226/001	11-8800	ACTAVIS GROUP PTC EHF.	NO
Oxycodone Actavis 60 mg depottabletit	FI/H/0965/007	31123	ACTAVIS GROUP PTC EHF.	FI
Oxycodone Actavis 80 mg depottabletit	FI/H/0965/008	31124	ACTAVIS GROUP PTC EHF.	FI
OXYCODONE ARROW 10	not available	NL 42519	ARROW GENERIQUES	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, gélule				
OXYCODONE ARROW 20 mg, gélule	not available	NL 42520	ARROW GENERIQUES	FR
OXYCODONE ARROW 5 mg, gélule	not available	NL 42518	ARROW GENERIQUES	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927784651	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785542	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785252	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927784712	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927784880	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785481	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927784941	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération	FR/H/0621/002/DC	3400927785191	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 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785023	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785771	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785313	BIOGARAN	FR
OXYCODONE BIOGARAN LP 10 mg, comprimé pelliculé à libération prolongée	FR/H/0621/002/DC	3400927785603	BIOGARAN	FR
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
OXYCODONE BIOGARAN LP 15 mg, comprimé pelliculé à libération	FR/H/0621/003/DC	3400927787263	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 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 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927787324	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération	FR/H/0621/004/DC	3400927787843	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 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927788093	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927788154	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927787492	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927787782	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927787553	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927788383	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927787904	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927788215	BIOGARAN	FR
OXYCODONE BIOGARAN LP 20 mg, comprimé pelliculé à libération	FR/H/0621/004/DC	3400927787614	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 20 mg, comprimé pelliculé à libération prolongée	FR/H/0621/004/DC	3400927788444	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927788505	BIOGARAN	FR
OXYCODONE BIOGARAN LP 30 mg, comprimé pelliculé à libération prolongée	FR/H/0621/005/DC	3400927789564	BIOGARAN	FR
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	FR/H/0621/005/DC	3400927789625	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 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 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/DC	3400927789793	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927789915	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790164	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790225	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790515	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération	FR/H/0621/006/DC	3400927791116	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 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790393	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790744	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927791055	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790973	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790454	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790805	BIOGARAN	FR
OXYCODONE BIOGARAN LP 40 mg, comprimé pelliculé à libération prolongée	FR/H/0621/006/DC	3400927790683	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927783241	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération	FR/H/0621/001/DC	3400927783821	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 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927784132	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927783999	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927783302	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927784422	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927783760	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927784361	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927784590	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927783470	BIOGARAN	FR
OXYCODONE BIOGARAN LP 5 mg, comprimé pelliculé à libération	FR/H/0621/001/DC	3400927783531	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 5 mg, comprimé pelliculé à libération prolongée	FR/H/0621/001/DC	3400927784071	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 BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927792694	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793127	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793417	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927792755	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération	FR/H/0621/008/DC	3400927793356	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 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793646	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793295	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793585	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793707	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927793066	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927792816	BIOGARAN	FR
OXYCODONE BIOGARAN LP 80 mg, comprimé pelliculé à libération prolongée	FR/H/0621/008/DC	3400927792984	BIOGARAN	FR
Oxycodone Depot ”Sandoz”, depottabletter 10 mg	DE/H/1154/002	41836	SANDOZ A/S	DK
Oxycodone Depot ”Sandoz”, depottabletter 20 mg	DE/H/1154/003	41837	SANDOZ A/S	DK
Oxycodone Depot	DE/H/1154/004	48908	SANDOZ 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
”Sandoz”, depottabletter 40 mg				
Oxycodone Depot ”Sandoz”, depottabletter 5 mg	DE/H/1154/001	41835	SANDOZ A/S	DK
Oxycodone Depot ”Sandoz”, depottabletter 60mg	DE/H/1154/005	48909	SANDOZ A/S	DK
Oxycodone Depot ”Sandoz”, depottabletter 80 mg	DE/H/1154/006	48910	SANDOZ A/S	DK
Oxycodone Depot 1A Farma 10 mg depottablett	DE/H/1154/002	25823	1A FARMA A/S	SE
Oxycodone Depot 1A Farma 20 mg depottablett	DE/H/1154/003	25824	1A FARMA A/S	SE
Oxycodone Depot 1A Farma 40 mg depottabletter	DE/H/1154/004	46151	1A FARMA A/S	SE
Oxycodone Depot 1A Farma 5 mg depottablett	DE/H/1154/001	25822	1A FARMA A/S	SE
Oxycodone Depot 1A Farma 80 mg depottabletter	DE/H/1154/006	46152	1A FARMA A/S	SE
Oxycodone Depot Orion 10 mg depottabletter	SE/H/1493/02	48494	ORION CORPORATION	SE
Oxycodone Depot Orion 15 mg depottabletter	SE/H/1493/03	48495	ORION CORPORATION	SE
Oxycodone Depot Orion 20 mg depottabletter	SE/H/1493/04	48496	ORION CORPORATION	SE
Oxycodone Depot Orion 30 mg depottabletter	SE/H/1493/05	48497	ORION CORPORATION	SE
Oxycodone Depot Orion 40 mg depottabletter	SE/H/1493/06	48498	ORION CORPORATION	SE
Oxycodone Depot Orion	SE/H/1493/01	48493	ORION CORPORATION	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
5 mg depottabletter				
Oxycodone Depot Orion 60 mg depottabletter	SE/H/1493/07	48499	ORION CORPORATION	SE
Oxycodone Depot Orion 80 mg depottabletter	SE/H/1493/08	48500	ORION CORPORATION	SE
Oxycodone Depot Teva Sweden 10 mg prolonged-release tablets	SE/H/1549/002	53283	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 15 mg prolonged-release tablets	SE-H-1549-001-008-DC	53284	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 20 mg prolonged-release tablets	SE-H-1549-001-008-DC	53285	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 30 mg prolonged-release tablets	SE-H-1549-001-008-DC	53286	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 40 mg prolonged-release tablets	SE-H-1549-001-008-DC	53287	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 5 mg depottablett	SE/H/1549/001	53282	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 60 mg depottablett	SE/H/1549/007	53288	TEVA SWEDEN AB	SE
Oxycodone Depot Teva Sweden 80 mg prolonged-release tablets	SE/H/1549/008	53289	TEVA SWEDEN AB	SE
OXYCODONE EG LP 10 mg, comprimé pelliculé à libération prolongée	DE/H/3641/002	NL42843	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 15 mg, comprimé pelliculé à libération prolongée	DE/H/3641/003	NL42844	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 20	DE/H/3641/004	NL42845	EG LABO LABORATOIRES	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, comprimé pelliculé à libération prolongée			EUROGENERICS	
OXYCODONE EG LP 30 mg, comprimé pelliculé à libération prolongée	DE/H/3641/005	NL42846	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 40 mg, comprimé pelliculé à libération prolongée	DE/H/3641/006	NL42847	EG LABO LABORATOIRES EUROGENERICS	FR
OXYCODONE EG LP 5 mg, comprimé pelliculé à libération prolongée	DE/H/3641/001	NL42842	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 EG LP 80 mg, comprimé pelliculé à libération prolongée	DE/H/3641/008	NL42849	EG LABO LABORATOIRES EUROGENERICS	FR
Oxycodone Ethypharm 10 mg forðahylki, hörð	not available	IS/1/09/157/01	ETHYPHARM	IS
Oxycodone Ethypharm 20 mg forðahylki, hörð	not available	IS/1/09/157/02	ETHYPHARM	IS
Oxycodone Ethypharm 40 mg forðahylki, hörð	not available	IS/1/09/157/03	ETHYPHARM	IS
Oxycodone Ethypharm 80 mg forðahylki, hörð	not available	IS/1/09/157/04	ETHYPHARM	IS
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 372 7 0	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 372 8 7	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération	DE/H/4081/003	34009 300 372 9 4	ETHYPHARM	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 0 0	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 1 7	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 2 4	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 3 1	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 4 8	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 373 5 5	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 371 5 7	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 371 6 4	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération	DE/H/4081/002	34009 300 371 7 1	ETHYPHARM	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 371 8 8	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 371 9 5	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 372 1 8	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 372 2 5	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 372 4 9	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 10 mg, comprimé à libération prolongée	DE/H/4081/002	34009 300 372 5 6	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 20 mg, comprimé à libération prolongée	DE/H/4081/003	34009 300 372 6 3	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 373 6 2	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération	DE/H/4081/004	34009 300 373 8 6	ETHYPHARM	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 373 9 3	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 0 9	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 1 6	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 2 3	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 4 7	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 5 4	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 6 1	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 40 mg, comprimé à libération prolongée	DE/H/4081/004	34009 300 374 7 8	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération	DE/H/4081/001	34009 300 370 3 4	ETHYPHARM	FR

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 370 4 1	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 370 5 8	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 370 6 5	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 370 7 2	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 370 9 6	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 371 0 2	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 371 1 9	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération prolongée	DE/H/4081/001	34009 300 371 2 6	ETHYPHARM	FR
OXYCODONE ETHYPHARM LP, 5 mg, comprimé à libération	DE/H/4081/001	34009 300 371 3 3	ETHYPHARM	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolongée				
Oxycodone Ethypharm, 10 mg, tabletki o przedłużonym uwalnianiu	DE/H/3738/002	22158	ETHYPHARM	PL
Oxycodone Ethypharm, 20 mg, tabletki o przedluzonym uwalnianiu	DE/H/3738/003	22159	ETHYPHARM	PL
Oxycodone Ethypharm, 40 mg, tabletki o przedluzonym uwalnianiu	DE/H/3738/004	22160	ETHYPHARM	PL
Oxycodone Ethypharm, 5 mg, tabletki o przedluzonym uwalnianiu	DE/H/3738/001	22157	ETHYPHARM	PL
Oxycodone G.L. 10 mg filmdragerade tabletter	DE/H/2669/002	43552	G.L. PHARMA GMBH	SE
Oxycodone G.L. 20 mg filmdragerade tabletter	DE/H/2669/004	53510	G.L. PHARMA GMBH	SE
Oxycodone G.L. 5 mg filmdragerade tabletter	DE/H/2669/001	43551	G.L. PHARMA GMBH	SE
Oxycodone G.L. Pharma 10 mg prolonged-release tablets	not available	PL 21597/0022	G.L. PHARMA GMBH	UK
Oxycodone G.L. Pharma 20 mg prolonged-release tablets	not available	PL 21597/0023	G.L. PHARMA GMBH	UK
Oxycodone G.L. Pharma 40 mg prolonged-release tablets	not available	PL 21597/0024	G.L. PHARMA GMBH	UK
Oxycodone G.L. Pharma 5 mg prolonged-release tablets	not available	PL 21597/0021	G.L. PHARMA GMBH	UK
Oxycodone G.L. Pharma 80 mg prolonged-release tablets	not available	PL 21597/0025	G.L. PHARMA GMBH	UK
Oxycodone Glenmark 10 mg prolonged-release	DE/H/3645/002	PL 25258/0218	GLENMARK PHARMACEUTICALS EUROPE	UK

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
tablets			LIMITED	
Oxycodone Glenmark 15 mg prolonged-release tablets	DE/H/3645/003	PL 25258/0219	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 20 mg prolonged-release tablets	DE/H/3645/004	PL 25258/0220	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 30 mg prolonged-release tablets	DE/H/3645/005	PL 25258/0221	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 40 mg prolonged-release tablets	DE/H/3645/006	PL 25258/0222	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 5 mg prolonged-release tablets	DE/H/3645/001	PL 25258/0217	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 60 mg prolonged-release tablets	DE/H/3645/007	PL 25258/0223	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone Glenmark 80 mg prolonged-release tablets	DE/H/3645/008	PL 25258/0224	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Oxycodone HCl Lannacher 10 mg ilgstošās darbības tabletes	DE/H/2182/002	10-0258	G.L. PHARMA GMBH	LV
Oxycodone HCl Lannacher 20 mg ilgstošās darbības tabletes	DE/H/2182/003	10-0259	G.L. PHARMA GMBH	LV
Oxycodone HCl Lannacher 40 mg ilgstošās darbības tabletes	DE/H/2182/004	10-0260	G.L. PHARMA GMBH	LV
Oxycodone HCl Lannacher 5 mg ilgstošās	DE/H/2182/001	10-0257	G.L. PHARMA GMBH	LV

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
darbības tabletes				
Oxycodone HCl Lannacher 80 mg ilgstošās darbības tabletes	DE/H/2182/005	10-0261	G.L. PHARMA GMBH	LV
Oxycodone Hydrochloride 10 mg/ml Solution for Injection or Infusion	UK/H/2915/001	PA 1339/025/001	WOCKHARDT UK LTD	IE
Oxycodone Hydrochloride 10 mg/ml Solution for Injection or Infusion	UK/H/2915/001	MA154/05101	WOCKHARDT UK LTD	MT
Oxycodone Hydrochloride 10mg/ml Oral Solution	UK/H/2780/002	MA154/06902	WOCKHARDT UK LTD	MT
Oxycodone Hydrochloride 10mg/ml Oral Solution	UK/H/2780/02/DC	29831/0459	WOCKHARDT UK LTD	UK
Oxycodone Hydrochloride 10mg/ml Solution for Injection or Infusion	UK/H/2915/01/DC	29831/0359	WOCKHARDT UK LTD	UK
Oxycodone Hydrochloride 50mg/ml Solution for Injection or Infusion	UK/H/5246/01/DC	PA 1339/025/002	WOCKHARDT UK LTD	IE
Oxycodone Hydrochloride 50mg/ml Solution for Injection or Infusion	UK/H/5246/01/DC	29831/0367	WOCKHARDT UK LTD	UK
Oxycodone Hydrochloride 5mg/5ml Oral Solution	UK/H/2780/001	MA 154/06901	WOCKHARDT UK LTD	MT
Oxycodone Hydrochloride 5mg/5ml	UK/H/2780/01/DC	29831/0458	WOCKHARDT UK LTD	UK

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
Oral Solution				
Oxycodone hydrochloride Rowex 10 mg Hard capsules	DE/H/4502/002	PA0711/286/002	ROWEX LTD	IE
Oxycodone hydrochloride Rowex 20 mg Hard capsules	DE/H/4502/003	PA0711/286/003	ROWEX LTD	IE
Oxycodone hydrochloride Rowex 5 mg Hard capsules	DE/H/4502/001	PA0711/286/001	ROWEX LTD	IE
Oxycodone Hydrochloride Wockhardt, 10 mg/ml, roztwór do wstrzykiwań lub infuzji	UK/H/2915/001	22711	WOCKHARDT UK LTD	PL
Oxycodone Lannacher 10 mg depottabletter	DE/H/2182/002	41899	LANNACHER HEILMITTEL GES.M.B.H.,	SE
Oxycodone Lannacher 10 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/2182/002	699510	G.L. PHARMA GMBH	EE
Oxycodone Lannacher 20 mg depottabletter	DE/H/2182/003	41900	LANNACHER HEILMITTEL GES.M.B.H.,	SE
Oxycodone Lannacher 20 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/2182/003	699610	G.L. PHARMA GMBH	EE
Oxycodone Lannacher 40 mg depottabletter	DE/H/2182/004	41901	LANNACHER HEILMITTEL GES.M.B.H.,	SE
Oxycodone Lannacher 40 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/2182/004	699710	G.L. PHARMA GMBH	EE
Oxycodone Lannacher 5 mg depottabletter	DE/H/2182/001	41898	LANNACHER HEILMITTEL GES.M.B.H.,	SE
Oxycodone Lannacher 5	DE/H/2182/001	699810	G.L. PHARMA GMBH	EE

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 toimeainet prolongeeritult vabastavad tabletid				
Oxycodone Lannacher 80 mg depottabletter	DE/H/2182/005	41902	LANNACHER HEILMITTEL GES.M.B.H.,	SE
Oxycodone Lannacher 80 mg toimeainet prolongeeritult vabastavad tabletid	DE/H/2182/005	699910	G.L. PHARMA GMBH	EE
Oxycodone Lannacher SR 10 mg prolonged-release tablet	DE/H/2183/002	PA0947/003/002	LANNACHER HEILMITTEL GES.M.B.H.,	IE
Oxycodone Lannacher SR 20 mg prolonged-release tablet	DE/H/2183/003	PA0947/003/003	LANNACHER HEILMITTEL GES.M.B.H.,	IE
Oxycodone Lannacher SR 40 mg prolonged-release tablet	DE/H/2183/004	PA0947/003/004	LANNACHER HEILMITTEL GES.M.B.H.,	IE
Oxycodone Lannacher SR 5 mg prolonged-release tablet	DE/H/2183/001	PA0947/003/001	LANNACHER HEILMITTEL GES.M.B.H.,	IE
Oxycodone Lannacher SR 80 mg prolonged-release tablet	DE/H/2183/005	PA0947/003/005	LANNACHER HEILMITTEL GES.M.B.H.,	IE
OXYCODONE MEDAC 50 mg/ml, solution Injectable/pour perfusion	UK/H/5881/002	34009 300 923 0 9	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	FR
Oxycodone Molteni, 10 mg/ml, rozt solution for injection or infusion	UK/H/5881/001/DC	PL 46373/0001	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	UK
Oxycodone Molteni, 10 mg/ml, rozt solution for injection or infusion	UK/H/5881/001/DC	PL 46373/0002	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	UK
Oxycodone Molteni, 10 mg/ml, roztwór do wstrzykiwań / do infuzji	UK/H/5881/001	23408	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodone Molteni, 50 mg/ml, roztwór do wstrzykiwań / do infuzji	UK/H/5881/002	23409	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
OXYCODONE MYLAN LP 10 mg, comprimé pelliculé à libération prolongée	DE/H/2827/002	NL 39918	MYLAN S.A.S	FR
OXYCODONE MYLAN LP 20 mg, comprimé pelliculé à libération prolongée	DE/H/2827/003	NL 39919	MYLAN S.A.S	FR
OXYCODONE MYLAN LP 40 mg, comprimé pelliculé à libération prolongée	DE/H/2827/004	NL 39920	MYLAN S.A.S	FR
OXYCODONE MYLAN LP 5 mg, comprimé pelliculé à libération prolongée	DE/H/2827/001	NL 39917	MYLAN S.A.S	FR
OXYCODONE MYLAN LP 80 mg, comprimé pelliculé à libération prolongée	DE/H/2827/005	NL 39921	MYLAN S.A.S	FR
Oxycodone Orion 10 mg depottabletter	SE/H/1493/002	31050	ORION CORPORATION	FI
Oxycodone Orion 10 mg depottabletti	SE/H/1493/02	31050	ORION CORPORATION	FI
Oxycodone Orion 10 mg/ml injektio-/infuusioneste, liuos	SE/H/1097/001	29117	ORION CORPORATION	FI
Oxycodone Orion 10 mg/ml injektions-/infusionsvätska, lösning	SE/H/1097/001	29117	ORION CORPORATION	FI
Oxycodone Orion 10 mg/ml injektions-/infusionsvätska, lösning	SE/H/1097/001	44897	ORION CORPORATION	SE
Oxycodone Orion 15 mg	SE/H/1493/003	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
depottabletter				
Oxycodone Orion 15 mg depottabletti	SE/H/1493/03	31051	ORION CORPORATION	FI
Oxycodone Orion 20 mg depottabletter	SE/H/1493/004	31052	ORION CORPORATION	FI
Oxycodone Orion 20 mg depottabletti	SE/H/1493/04	31052	ORION CORPORATION	FI
Oxycodone Orion 30 mg depottabletter	SE/H/1493/005	31053	ORION CORPORATION	FI
Oxycodone Orion 30 mg depottabletti	SE/H/1493/05	31053	ORION CORPORATION	FI
Oxycodone Orion 40 mg depottabletter	SE/H/1493/006	31054	ORION CORPORATION	FI
Oxycodone Orion 40 mg depottabletti	SE/H/1493/06	31054	ORION CORPORATION	FI
Oxycodone Orion 5 mg depottabletter	SE/H/1493/001	31049	ORION CORPORATION	FI
Oxycodone Orion 5 mg depottabletti	SE/H/1493/01	31049	ORION CORPORATION	FI
Oxycodone Orion 60 mg depottabletter	SE/H/1493/007	31055	ORION CORPORATION	FI
Oxycodone Orion 60 mg depottabletti	SE/H/1493/07	31055	ORION CORPORATION	FI
Oxycodone Orion 80 mg depottabletter	SE/H/1493/008	31056	ORION CORPORATION	FI
Oxycodone Orion 80 mg depottabletti	SE/H/1493/08	31056	ORION CORPORATION	FI
Oxycodone ratiopharm 10 mg depottabletti	DE/H/1395/001	24289	RATIOPHARM GMBH	FI
Oxycodone ratiopharm 10 mg forðatöflur	DE/H/1395/002	IS/1/13/014/01	RATIOPHARM GMBH	IS
Oxycodone ratiopharm 20 mg depottabletti	DE/H/0790/001	22390	RATIOPHARM GMBH	FI
Oxycodone ratiopharm 20 mg forðatöflur	DE/H/0790/001	IS/1/12/114/01	RATIOPHARM GMBH	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
Oxycodone ratiopharm 40 mg depottabletti	DE/H/0790/002	22391	RATIOPHARM GMBH	FI
Oxycodone ratiopharm 40 mg forðatöflur	DE/H/0790/002	IS/1/12/114/02	RATIOPHARM GMBH	IS
Oxycodone ratiopharm 5 mg depottabletti	DE/H/2947/001	29116	RATIOPHARM GMBH	FI
Oxycodone ratiopharm 5 mg forðatöflur.	DE/H/2947/001	IS/1/13/015/01	RATIOPHARM GMBH	IS
Oxycodone ratiopharm 80 mg depottabletti	DE/H/0790/003	22392	RATIOPHARM GMBH	FI
Oxycodone ratiopharm 80 mg forðatöflur	DE/H/0790/003	IS/1/12/114/03	RATIOPHARM GMBH	IS
OXYCODONE SANDOZ 10 mg kapseli, kova	DE/H/4502/002	35354	SANDOZ A/S	FI
Oxycodone Sandoz 10 mg kemény kapszula	DE/H/4502/002	OGYI-T-23294/04	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 10 mg kemény kapszula	DE/H/4502/002	OGYI-T-23294/05	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 10 mg kemény kapszula	DE/H/4502/002	OGYI-T-23294/06	SANDOZ HUNGÁRIA KFT	HU
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 688 9 2	SANDOZ	FR
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 689 0 8	SANDOZ	FR
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 689 1 5	SANDOZ	FR
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 689 2 2	SANDOZ	FR
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 689 3 9	SANDOZ	FR
OXYCODONE SANDOZ 10 mg, gélule	not available	34009 300 689 5 3	SANDOZ	FR
Oxycodone Sandoz 20 mg depottabletti	DE/H/1154/003	23943	SANDOZ A/S	FI
OXYCODONE SANDOZ 20	DE/H/4502/003	35355	SANDOZ A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg kapseli, kova				
Oxycodone Sandoz 20 mg kemény kapszula	DE/H/4502/003	OGYI-T-23294/07	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 20 mg kemény kapszula	DE/H/4502/003	OGYI-T-23294/08	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 20 mg kemény kapszula	DE/H/4502/003	OGYI-T-23294/09	SANDOZ HUNGÁRIA KFT	HU
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 689 6 0	SANDOZ	FR
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 689 7 7	SANDOZ	FR
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 689 8 4	SANDOZ	FR
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 689 9 1	SANDOZ	FR
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 690 0 4	SANDOZ	FR
OXYCODONE SANDOZ 20 mg, gélule	not available	34009 300 690 1 1	SANDOZ	FR
Oxycodone Sandoz 40 mg depottabletti	DE/H/1154/004	29779	SANDOZ A/S	FI
OXYCODONE SANDOZ 5 mg kapseli, kova	DE/H/4502/001	35353	SANDOZ A/S	FI
Oxycodone Sandoz 5 mg kemény kapszula	DE/H/4502/001	OGYI-T-23294/01	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 5 mg kemény kapszula	DE/H/4502/001	OGYI-T-23294/02	SANDOZ HUNGÁRIA KFT	HU
Oxycodone Sandoz 5 mg kemény kapszula	DE/H/4502/001	OGYI-T-23294/03	SANDOZ HUNGÁRIA KFT	HU
OXYCODONE SANDOZ 5 mg, gélule	not available	34009 300 688 3 0	SANDOZ	FR
OXYCODONE SANDOZ 5 mg, gélule	not available	34009 300 688 4 7	SANDOZ	FR
OXYCODONE SANDOZ 5 mg, gélule	not available	34009 300 688 7 8	SANDOZ	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 SANDOZ 5 mg, gélule	not available	34009 300 688 6 1	SANDOZ	FR
OXYCODONE SANDOZ 5 mg, gélule	not available	34009 300 688 5 4	SANDOZ	FR
OXYCODONE SANDOZ 5 mg, gélule	not available	34009 300 688 2 3	SANDOZ	FR
Oxycodone Sandoz 80 mg depottabletti	DE/H/1154/006	29781	SANDOZ A/S	FI
Oxycodone STADA 10 mg kapseli, kova	DE/H/4774/002	34403	STADA ARZNEIMITTEL AG	FI
Oxycodone STADA 20 mg kapseli, kova	DE/H/4774/003	34404	STADA ARZNEIMITTEL AG	FI
Oxycodone STADA 5 mg kapseli, kova	DE/H/4774/001	34402	STADA ARZNEIMITTEL AG	FI
Oxycodone Teva 10 mg hårda kapslar	SE/H/1423/002	50981	TEVA SWEDEN AB	SE
Oxycodone Teva 10 mg, tabletten met verlengde afgifte	SE/H/1549/002	BE499031	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 10 mg, tabletten met verlengde afgifte	SE/H/1549/002	BE499022	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 20 mg hårda kapslar	SE/H/1423/003	50982	TEVA SWEDEN AB	SE
Oxycodone Teva 20 mg, tabletten met verlengde afgifte	SE/H/1549/004	BE499057	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 20 mg, tabletten met verlengde afgifte	SE/H/1549/004	BE499040	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 40 mg, tabletten met verlengde afgifte	SE/H/1549/006	BE499075	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 40 mg, tabletten met verlengde afgifte	SE/H/1549/006	BE499066	TEVA PHARMA BELGIUM N.V./S.A	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
Oxycodone Teva 5 mg hårda kapslar	SE/H/1423/001	50980	TEVA SWEDEN AB	SE
Oxycodone Teva 5 mg, tabletten met verlengde afgifte	SE/H/1549/001	BE499013	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 5 mg, tabletten met verlengde afgifte	SE/H/1549/001	BE499004	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 80 mg tabletten met verlengde afgifte	SE/H/1549/008	BE499093	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Teva 80 mg, tabletten met verlengde afgifte	SE/H/1549/008	BE499084	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxycodone Torrent 10 mg gélule	not available	34009300683-684	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 10 mg, gélule	not available	34009 300 683 9 7	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 10 mg, gélule	not available	34009 300 684 0 3	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 10 mg, gélule	not available	34009 300 6841 0	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 10 mg, gélule	not available	34009 300 684 3 4	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 10 mg, gélule	not available	34009 300 684 41	TORRENT PHARMA GMBH	FR
Oxycodone Torrent 20 mg gélule	not available	3400930068458-502	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 20 mg, gélule	not available	34009 300 684 6 5	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 20 mg, gélule	not available	34009 300 684 7 2	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 20 mg, gélule	not available	34009 300 684 8 9	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT	not available	34009 300 684 9 6	TORRENT PHARMA GMBH	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
20 mg, gélule				
OXYCODONE TORRENT 20 mg, gélule	not available	34009 300 685 0 2	TORRENT PHARMA GMBH	FR
Oxycodone Torrent 5 mg gélule	not available	3400930068311-7	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 5 mg, gélule	not available	34009 300 683 2 8	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 5 mg, gélule	not available	34009 300 683 3 5	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 5 mg, gélule	not available	34009 300 683 4 2	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 5 mg, gélule	not available	34009 300 683 5 9	TORRENT PHARMA GMBH	FR
OXYCODONE TORRENT 5 mg, gélule	not available	34009 300 683 7 3	TORRENT PHARMA GMBH	FR
Oxycodone Vitabalans 10 mg apvalkotās tabletes	SE/H/1119/002	12-0183	VITABALANS OY	LV
Oxycodone Vitabalans 10 mg filmdragerade tabletter	SE/H/1119/002	46198	VITABALANS OY	SE
Oxycodone Vitabalans 10 mg filmtabletta	SE/H/1119/002	OGYI-T-22287/03	VITABALANS OY	HU
Oxycodone Vitabalans 10 mg filmtabletta	SE/H/1119/002	OGYI-T-22287/04	VITABALANS OY	HU
Oxycodone Vitabalans 10 mg kalvopäällysteiset tabletit	SE/H/1119/002	29839	VITABALANS OY	FI
Oxycodone Vitabalans 10 mg plėvele dengtos tabletės	SE/H/1119/002	LT/1/13/3201/003	VITABALANS OY	LT
Oxycodone Vitabalans 10 mg plėvele dengtos tabletės	SE/H/1119/002	LT/1/13/3201/004	VITABALANS OY	LT
Oxycodone Vitabalans 10 mg potahované tablety	SE/H/1119/002	65/566/12-C	VITABALANS OY	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
Oxycodone Vitabalans 10 mg tabletter, filmdrasjerte	SE/H/1119/002	11-8362	VITABALANS OY	NO
Oxycodone Vitabalans 5 mg apvalkotās tabletes	SE/H/1119/001	12-0182	VITABALANS OY	LV
Oxycodone Vitabalans 5 mg filmdragerade tabletter	SE/H/1119/001	46197	VITABALANS OY	SE
Oxycodone Vitabalans 5 mg filmtabletta	SE/H/1119/001	OGYI-T-22287/01	VITABALANS OY	HU
Oxycodone Vitabalans 5 mg filmtabletta	SE/H/1119/001	OGYI-T-22287/02	VITABALANS OY	HU
Oxycodone Vitabalans 5 mg kalvopäällysteiset tabletit	SE/H/1119/001	29838	VITABALANS OY	FI
Oxycodone Vitabalans 5 mg plėvele dengtos tabletės	SE/H/1119/001	LT/1/13/3201/001	VITABALANS OY	LT
Oxycodone Vitabalans 5 mg plėvele dengtos tabletės	SE/H/1119/001	LT/1/13/3201/002	VITABALANS OY	LT
Oxycodone Vitabalans 5 mg potahované tablety	SE/H/1119/001	65/565/12-C	VITABALANS OY	CZ
Oxycodone Vitabalans 5 mg tabletter, filmdrasjerte	SE/H/1119/001	11-8361	VITABALANS OY	NO
Oxycodone Vitabalans, 10 mg õhukese polümeerikattega tabletid	SE/H/1119/002	788412	VITABALANS OY	EE
Oxycodone Vitabalans, 10 mg, tabletki powlekane	SE/H/1119/002	20450	VITABALANS OY	PL
Oxycodone Vitabalans, 5 mg õhukese polümeerikattega	SE/H/1119/001	788512	VITABALANS OY	EE

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
tabletid				
Oxycodone Vitabalans, 5 mg, tabletki powlekane	SE/H/1119/001	20449	VITABALANS OY	PL
Oxycodone Wockhardt 10 mg/ml ενέσιμο διάλυμα ή διάλυμα για έγχυση	UK/H/2915/001	ML 21405	WOCKHARDT UK LTD	CY
Oxycodon-HCl HEXAL 10 mg Retardtabletten	DE/H/1084/002	67704.00.00	HEXAL AG	DE
Oxycodon-HCl - 1 A Pharma 10 mg Hartkapseln	not available	99688.00.00	1 A PHARMA GMBH	DE
Oxycodon-HCl - 1 A Pharma 20 mg Hartkapseln	not available	99689.00.00	1 A PHARMA GMBH	DE
Oxycodon-HCl - 1 A Pharma 5 mg Hartkapseln	not available	99687.00.00	1 A PHARMA GMBH	DE
Oxycodon-HCl AbZ 10 mg Retardtabletten	not available	71327.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 20 mg Retardtabletten	not available	66596.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 30 mg Retardtabletten	DE/H/3495/001	86471.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 40 mg Retardtabletten	not available	66597.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 5 mg Retardtabletten	DE/H/2947/001	82776.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl AbZ 60 mg Retardtabletten	DE/H/3495/002	86472.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl Accord 10 mg tabletten met verlengde afgifte	NL/H/3192/002/E/001	RVG 109542	ACCORD HEALTHCARE LIMITED	NL
Oxycodon-HCl Accord 20 mg tabletten met verlengde afgifte	NL/H/3192/003/E/001	RVG 109543	ACCORD HEALTHCARE LIMITED	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodon-HCl Accord 40 mg tabletten met verlengde afgifte	NL/H/3192/004/E/001	RVG 109544	ACCORD HEALTHCARE LIMITED	NL
Oxycodon-HCl Accord 5 mg tabletten met verlengde afgifte	NL/H/3192/001/E/001	RVG 109541	ACCORD HEALTHCARE LIMITED	NL
Oxycodon-HCl Accord 80 mg tabletten met verlengde afgifte	NL/H/3192/005/E/001	RVG 109545	ACCORD HEALTHCARE LIMITED	NL
Oxycodon-HCl AL 10 mg Retardtabletten	DE/H/3860/002	89917.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 10 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/1091/002	69173.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 15 mg Retardtabletten	DE/H/3860/003	89918.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 20 mg Retardtabletten	DE/H/3860/004	89919.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 20 mg Retardtabletten	DE/H/0669/001	65363.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 30 mg Retardtabletten	DE/H/3860/005	89920.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 40 mg Retardtabletten	DE/H/3860/006	89921.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 40 mg Retardtabletten	DE/H/0669/002	65364.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 5 mg Retardtabletten	DE/H/3860/001	89916.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 60 mg Retardtabletten	DE/H/3860/007	89922.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 80 mg Retardtabletten	DE/H/3860/008	89923.00.00	ALIUD PHARMA GMBH	DE
Oxycodon-HCl AL 80 mg Retardtabletten	DE/H/0669/003	65365.00.00	ALIUD 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 Amneal 10 mg Retardtabletten	DE/H/3639/002	87928.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 15 mg Retardtabletten	DE/H/3639/003	87929.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 20 mg Retardtabletten	DE/H/3639/004	87930.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 30 mg Retardtabletten	DE/H/3639/005	87931.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 40 mg Retardtabletten	DE/H/3639/006	87932.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 5 mg Retardtabletten	DE/H/3639/001	87927.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 60 mg Retardtabletten	DE/H/3639/007	87933.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCL Amneal 80 mg Retardtabletten	DE/H/3639/008	87934.00.00	MORNINGSIDE HEALTHCARE LTD	DE
Oxycodon-HCl Aristo akut 20 mg Filmtabletten	DE/H/4047/001	91325.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon-HCl Aristo® 10 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	not available	83015.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon-HCl Aristo® 20 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	not available	83016.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon-HCl Aristo® 40 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	not available	83017.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon-HCl Aristo® 5 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	not available	83014.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxycodon-HCl Aristo® 80 mg Retardtabletten	not available	83018.00.00	ARISTO PHARMA GMBH (ART 57)	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
Wirkstoff: Oxycodonhydrochlorid				
Oxycodon-HCl beta 1 x 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 10 mg Retardtabletten	DE/H/1085/002	67705.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 20 mg Retardtabletten	DE/H/0667/001	65357.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 30 mg Retardtabletten	DE/H/0667/004	69806.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 40 mg Retardtabletten	DE/H/0667/002	65358.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 5 mg Retardtabletten	not available	83009.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 60 mg Retardtabletten	DE/H/0667/005	69807.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta 80 mg Retardtabletten	DE/H/0667/003	65359.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta akut 10 mg Hartkapseln	not available	99658.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta akut 20 mg Hartkapseln	not available	99659.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl beta akut 5 mg Hartkapseln	not available	99657.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Oxycodon-HCl dura 10	DE/H/2439/0012	78322.00.00	MYLAN DURA 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
mg Retardtabletten				
Oxycodon-HCl dura 20 mg Retardtabletten	DE/H/2439/003	78323.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl dura 40 mg Retardtabletten	DE/H/2439/004	78324.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl dura 5 mg Retardtabletten	DE/H/2439/001	78321.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl dura 80 mg Retardtabletten	DE/H/2439/005	78325.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl HEXAL 10 mg Retardtabletten	DE/H/1155/002	69184.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL 20 mg Retardtabletten	DE/H/1155/003	69185.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL 5 mg Retardtabletten	DE/H/1155/001	69183.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL akut 10 mg Hartkapseln	DE/H/4502/002	92111.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL akut 20 mg Hartkapseln	DE/H/4502/003	92112.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL akut 5 mg Hartkapseln	DE/H/4502/001	92110.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL® 40 mg Retardtabletten	DE/H/1155/004	84874.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL® 60 mg Retardtabletten	DE/H/1155/005	84875.00.00	HEXAL AG	DE
Oxycodon-HCl HEXAL® 80 mg Retardtabletten	DE/H/1155/006	84876.00.00	HEXAL AG	DE
Oxycodon-HCl Krugmann 1 mg/ml Lösung zum Einnehmen	DE/H/1026/007	83866.00.00	KRUGMAN GMBH	DE
Oxycodon-HCl Krugmann 10 mg Retardtabletten	DE/H/0366/002	53004.00.00	KRUGMAN GMBH	DE
Oxycodon-HCl Krugmann 20 mg Retardtabletten	DE/H/1026/003	53004.01.00	KRUGMAN GMBH	DE
Oxycodon-HCl Krugmann	DE/H/1026/004	53004.02.00	MUNDIPHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
40 mg Retardtabletten				
Oxycodon-HCl Krugmann 5 mg Retardtabletten	DE/H/1026/001	53004.04.00	MUNDIPHARMA GMBH	DE
Oxycodon-HCl Krugmann 80 mg Retardtabletten	DE/H/1026/005	53004.03.00	MUNDIPHARMA GMBH	DE
Oxycodon-HCl Mylan 10 mg Retardtabletten	DE/H/4246/002	93682.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 20 mg Retardtabletten	DE/H/4246/003	93683.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 30 mg Retardtabletten	DE/H/4246/004	93684.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 40 mg Retardtabletten	DE/H/4246/005	93685.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 5 mg Retardtabletten	DE/H/4246/001	93681.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 60 mg Retardtabletten	DE/H/4246/006	93686.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl Mylan 80 mg Retardtabletten	DE/H/4246/007	93687.00.00	MYLAN DURA GMBH	DE
Oxycodon-HCl ratiopharm 20 mg Retardtabletten	DE/H/0668/001	1-27779	TEVA B.V	AT
Oxycodon-HCl ratiopharm 40 mg Retardtabletten	DE/H/0668/002	1-27780	TEVA B.V	AT
Oxycodon-HCl ratiopharm 80 mg Retardtabletten	DE/H/0668/003	1-27781	TEVA B.V	AT
Oxycodon-HCL Sandoz 10 mg Retardtabletten	DE/H/1154/002	69180.00.00	HEXAL AG	DE
OXYCODON-HCL SANDOZ 20 MG RETARDTABLETTEN	DE/H/1154/003	69181.00.00	HEXAL AG	DE
Oxycodon-HCL Sandoz 5 mg Retardtabletten	DE/H/1154/001	69179.00.00	HEXAL AG	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 Sandoz® 10 mg, Retardtabletten	DE/H/1419/002	71451.00.00	HEXAL AG	DE
Oxycodon-HCL Sandoz® 40 mg Retardtabletten	DE/H/0789/002	66539.00.00	HEXAL AG	DE
Oxycodon-HCL Sandoz® 80 mg Retardtabletten	DE/H/0789/003	66540.00.00	HEXAL AG	DE
Oxycodon-HCl STADA® 10 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/1083/002	67703.00.00	STADAPHARM GMBH	DE
Oxycodon-HCl STADA® 20 mg Retardtabletten	DE/H/0665/001	65351.00.00	STADAPHARM GMBH	DE
Oxycodon-HCl STADA® 40 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/0665/002	65352.00.00	STADAPHARM GMBH	DE
Oxycodon-HCl STADA® 80 mg Retardtabletten	DE/H/0665/003	65353.00.00	STADAPHARM GMBH	DE
Oxycodon-HCl Winthrop 10 mg Retardtabletten	DE/H/2438/002	78317.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Winthrop 20 mg Retardtabletten	DE/H/2438/003	78318.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Winthrop 40 mg Retardtabletten	DE/H/2438/004	78319.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Winthrop 5 mg Retardtabletten	DE/H/2438/001	78316.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Winthrop 80 mg Retardtabletten	DE/H/2438/005	78320.00.00	WINTHROP ARZNEIMITTEL GMBH	DE
Oxycodon-HCl Zentiva 10 mg Hartkapseln	not available	92114.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl Zentiva 20 mg Hartkapseln	not available	92115.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl Zentiva 30 mg Retardtabletten	DE/H/3476/001	86290.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl Zentiva 5	not available	92113.00.00	ZENTIVA 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
mg Hartkapseln				
Oxycodon-HCl Zentiva 60 mg Retardtabletten	DE/H/3476/002	86291.00.00	ZENTIVA PHARMA GMBH	DE
Oxycodon-HCl-Acino 10 mg Retardtabletten	DE/H/1153/002	69200.00.00	ACINO AG	DE
Oxycodon-HCl-Acino 20 mg Retardtabletten	DE/H/1153/003	69201.00.00	ACINO AG	DE
Oxycodon-HCl-Acino 40 mg Retardtabletten	DE/H/1153/004	69202.00.00	ACINO AG	DE
Oxycodon-HCl-Acino 80 mg Retardtabletten	DE/H/1153/005	69203.00.00	ACINO AG	DE
Oxycodon-HCl-CT 10 mg Retardtabletten	not available	71325.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-CT 20 mg Retardtabletten	not available	66593.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-CT 30 mg Retardtabletten	DE/H/3494/001	86469.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-CT 40 mg Retardtabletten	not available	66594.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-CT 60 mg Retardtabletten	DE/H/3494/002	86470.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-CT 80 mg Retardtabletten	not available	66595.00.00	ABZ-PHARMA GMBH	DE
Oxycodon-HCl-ratiopharm® 10 mg Retardtabletten	DE/H/1086/002	67706.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 20 mg Retardtabletten	DE/H/0668/001	65360.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 30 mg Retardtabletten	DE/H/3475/001	86288.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 40 mg Retardtabletten	DE/H/0668/002	65361.00.00	RATIOPHARM 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-ratiopharm® 5 mg Retardtabletten	not available	70022.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 60 mg Retardtabletten	DE/H/3475/002	86289.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® 80 mg Retardtabletten	DE/H/0668/003	65362.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® akut 10 mg Hartkapseln	SE/H/1423/002	92061.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® akut 20 mg Hartkapseln	SE/H/1423/003	92062.00.00	RATIOPHARM GMBH	DE
Oxycodon-HCl-ratiopharm® akut 5 mg Hartkapseln	SE/H/1423/001	92060.00.00	RATIOPHARM GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 10 mg Retardtabletten	DE/H/1156/002	69141.00.00	1 A PHARMA GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 20 mg Retardtabletten	DE/H/1156/003	69142.00.00	1 A PHARMA GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 40 mg Retardtabletten	DE/H/1154/004	84845.00.00	1 A PHARMA GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 5 mg Retardtabletten	DE/H/1156/001	69140.00.00	1 A PHARMA GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 60 mg Retardtabletten	DE/H/1154/005	84846.00.00	1 A PHARMA GMBH	DE
Oxycodonhydrochlorid - 1 A Pharma® 80 mg Retardtabletten	DE/H/1154/006	84847.00.00	1 A 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
Oxycodonhydrochlorid "Actavis", hårde kapsler	SE/H/1226/001	50043	ACTAVIS GROUP PTC EHF.	DK
Oxycodonhydrochlorid "Actavis", hårde kapsler	SE/H/1226/002	50044	ACTAVIS GROUP PTC EHF.	DK
Oxycodonhydrochlorid "Actavis", hårde kapsler	SE/H/1226/003	50045	ACTAVIS GROUP PTC EHF.	DK
Oxycodonhydrochlorid "G.L."	DE/H/2669/004	56759	G.L. PHARMA GMBH	DK
Oxycodonhydrochlorid ”Acino”, depottabletter	DE/H/1153/005	41757	ACINO AG	DK
Oxycodonhydrochlorid ”Acino”, depottabletter	DE/H/1153/003	41755	ACINO AG	DK
Oxycodonhydrochlorid ”Acino”, depottabletter	DE/H/1153/004	41756	ACINO AG	DK
Oxycodonhydrochlorid ”Acino”, depottabletter	DE/H/1153/002	41754	ACINO AG	DK
Oxycodonhydrochlorid ”G.L.”, filmovertrukne tabletter	DE/H/2669/001	46271	G.L. PHARMA GMBH	DK
Oxycodonhydrochlorid ”G.L.”, filmovertrukne tabletter	DE/H/2669/002	46272	G.L. PHARMA GMBH	DK
Oxycodonhydrochlorid ”Lannacher”, depottabletter	DE/H/2182/001	44426	LANNACHER HEILMITTEL GES.M.B.H.,	DK
Oxycodonhydrochlorid ”Lannacher”, depottabletter	DE/H/2182/002	44427	LANNACHER HEILMITTEL GES.M.B.H.,	DK
Oxycodonhydrochlorid ”Lannacher”, depottabletter	DE/H/2182/003	44428	LANNACHER HEILMITTEL GES.M.B.H.,	DK
Oxycodonhydrochlorid ”Lannacher”, depottabletter	DE/H/2182/004	44429	LANNACHER HEILMITTEL GES.M.B.H.,	DK
Oxycodonhydrochlorid ”Lannacher”,	DE/H/2182/005	44430	LANNACHER HEILMITTEL GES.M.B.H.,	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				
Oxycodonhydrochlorid "Orion", injektions- og infusionsvæske, opløsning	SE/H/1097/001	47477	ORION CORPORATION	DK
Oxycodonhydrochlorid axcount 10 mg Retardtabletten	DE/H/4245/002	93675.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 20 mg Retardtabletten	DE/H/4245/003	93676.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 30 mg Retardtabletten	DE/H/4245/004	93677.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 40 mg Retardtabletten	DE/H/4245/005	93678.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 5 mg Retardtabletten	DE/H/4245/001	93674.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 60 mg Retardtabletten	DE/H/4245/006	93679.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid axcount 80 mg Retardtabletten	DE/H/4245/007	93680.00.00	AXCOUNT GENERIKA GMBH	DE
Oxycodonhydrochlorid Ethypharm 10 mg Retardtabletten	DE/H/4081/002	91780.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid Ethypharm 10 mg Retardtabletten	DE/H/4080/002	91750.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid Ethypharm 20 mg Retardtabletten	DE/H/4081/003	91781.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid	DE/H/4080/003	91751.00.00	ETHYPHARM	DE

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ethypharm 20 mg Retardtabletten				
Oxycodonhydrochlorid Ethypharm 40 mg Retardtabletten	DE/H/4081/004	91782.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid Ethypharm 40 mg Retardtabletten	DE/H/4080/004	91752.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid Ethypharm 5 mg Retardtabletten	DE/H/4081/001	91779.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid Ethypharm 5 mg Retardtabletten	DE/H/4080/001	91749.00.00	ETHYPHARM	DE
Oxycodonhydrochlorid G.L. 10 mg Filmtabletten	DE/H/2669/002	80364.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 10 mg Retardtabletten	DE/H/2827/002	81682.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 20 mg Filmtabletten	DE/H/2669/004	96062.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 20 mg Retardtabletten	DE/H/2827/003	81683.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 40 mg Retardtabletten	DE/H/2827/004	81684.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 5 mg Filmtabletten	DE/H/2669/001	80363.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 5 mg Retardtabletten	DE/H/2827/001	81681.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid G.L. 80 mg Retardtabletten	DE/H/2827/005	81685.00.00	G.L. PHARMA GMBH	DE
Oxycodonhydrochlorid	not available	78001.00.00	HENNIG ARZNEIMITTEL	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
Hennig 10 mg Retardtabletten			GMBH & CO. KG	
Oxycodonhydrochlorid Hennig 20 mg Retardtabletten	not available	78002.00.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Oxycodonhydrochlorid Hennig 40 mg Retardtabletten	not available	78003.00.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Oxycodonhydrochlorid Hennig 5 mg Retardtabletten	not available	78000.00.00	HENNIG ARZNEIMITTEL GMBH & CO. KG	DE
Oxycodonhydrochlorid Heumann 10 mg Hartkapseln	not available	92108.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 10 mg Retardtabletten	SE/H/1381/002/DC	91465.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 15 mg Retardtabletten	SE/H/1381/003/DC	91466.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 20 mg Hartkapseln	not available	92109.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 20 mg Retardtabletten	SE/H/1381/004/DC	91467.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 30 mg Retardtabletten	SE/H/1381/005/DC	91468.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 40 mg Retardtabletten	SE/H/1381/006/DC	91469.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 5 mg Hartkapseln	not available	92107.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid	SE/H/1381/001/DC	91464.00.00	HEUMANN 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
Heumann 5 mg Retardtabletten			CO. GENERICA KG	
Oxycodonhydrochlorid Heumann 60 mg Retardtabletten	SE/H/1381/007/DC	91470.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid Heumann 80 mg Retardtabletten	SE/H/1381/008/DC	91471.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Oxycodonhydrochlorid HEXAL 5 mg Retardtabletten	DE/H/1419/001	71450.00.00	HEXAL AG	DE
Oxycodonhydrochlorid PUREN 10 mg Hartkapseln	NL/H/4429/002	86977.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 10 mg Retardtabletten	DE/H/5356/002	89095.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 15 mg Retardtabletten	DE/H/5356/004	89096.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid PUREN 20 mg Hartkapseln	NL/H/4429/003	86978.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 20 mg Retardtabletten	DE/H/5356/003	89097.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 30 mg Retardtabletten	DE/H/5356/005	89098.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 40 mg Retardtabletten	DE/H/5356/006	89099.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid PUREN 5 mg Hartkapseln	NL/H/4429/001	86976.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 5 mg	DE/H/5356/001	89094.00.00	PUREN PHARMA GMBH & CO. KG	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
Retardtabletten				
Oxycodonhydrochlorid Puren 60 mg Retardtabletten	DE/H/5356/007	89100.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid Puren 80 mg Retardtabletten	DE/H/5356/008	89101.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxycodonhydrochlorid STADA® 10 mg Retardtabletten	DE/H/3641/002	87944.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 15 mg Retardtabletten	DE/H/3641/003	87945.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 20 mg Retardtabletten	DE/H/3641/004	87946.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 30 mg Retardtabletten	DE/H/3641/005	87947.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 40 mg Retardtabletten	DE/H/3641/006	87948.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 5 mg Retardtabletten	DE/H/3641/001	87943.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 60 mg Retardtabletten	DE/H/3641/007	87949.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid STADA® 80 mg Retardtabletten	DE/H/3641/008	87950.00.00	STADAPHARM GMBH	DE
Oxycodonhydrochlorid-CT 5 mg Retardtabletten	not available	82794.00.00	ABZ-PHARMA GMBH	DE
Oxycodonhydrochlorid-ratiopharm® 10 mg Retardtabletten	DE/H/1395/002	71323.00.00	RATIOPHARM 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
Oxycodonhydrochlorid-ratiopharm® 20 mg Retardtabletten	DE/H/0790/001	66590.00.00	RATIOPHARM GMBH	DE
Oxycodonhydrochlorid-ratiopharm® 40 mg Retardtabletten	DE/H/0790/002	66591.00.00	RATIOPHARM GMBH	DE
Oxycodonhydrochlorid-ratiopharm® 80 mg Retardtabletten	DE/H/0790/003	66592.00.00	RATIOPHARM GMBH	DE
Oxycon 40 mg Retardtabletten	DE/H/3293/001	84871.00.00	HEXAL AG	DE
Oxycon 60 mg Retardtabletten	DE/H/3293/002	84872.00.00	HEXAL AG	DE
Oxycon 80 mg Retardtabletten	DE/H/3293/003	84873.00.00	HEXAL AG	DE
Oxyconica 10 mg Retardtabletten	DE/H/3644/002	87968.00.00	ACINO AG	DE
Oxyconica 15 mg Retardtabletten	DE/H/3644/003	87969.00.00	ACINO AG	DE
Oxyconica 20 mg Retardtabletten	DE/H/3644/004	87970.00.00	ACINO AG	DE
Oxyconica 30 mg Retardtabletten	DE/H/3644/005	87971.00.00	ACINO AG	DE
Oxyconica 40 mg Retardtabletten	DE/H/3644/006	87972.00.00	ACINO AG	DE
Oxyconica 5 mg Retardtabletten	DE/H/3644/001	87967.00.00	ACINO AG	DE
Oxyconica 60 mg Retardtabletten	DE/H/3644/007	87973.00.00	ACINO AG	DE
Oxyconica 80 mg Retardtabletten	DE/H/3644/008	87974.00.00	ACINO AG	DE
Oxyconicur 10 mg Retardtabletten	DE/H/3643/002	87960.00.00	ACINO AG	DE
Oxyconicur 15 mg Retardtabletten	DE/H/3643/003	87961.00.00	ACINO AG	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
Oxyconicur 20 mg Retardtabletten	DE/H/3643/004	87962.00.00	ACINO AG	DE
Oxyconicur 30 mg Retardtabletten	DE/H/3643/005	87963.00.00	ACINO AG	DE
Oxyconicur 40 mg Retardtabletten	DE/H/3643/006	87964.00.00	ACINO AG	DE
Oxyconicur 5 mg Retardtabletten	DE/H/3643/001	87959.00.00	ACINO AG	DE
Oxyconicur 60 mg Retardtabletten	DE/H/3643/007	87965.00.00	ACINO AG	DE
Oxyconicur 80 mg Retardtabletten	DE/H/3643/008	87966.00.00	ACINO AG	DE
Oxyconoica 10 mg Retardtabletten	DE/H/3645/002	87976.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 15 mg Retardtabletten	DE/H/3645/003	87977.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 20 mg Retardtabletten	DE/H/3645/004	87978.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 30 mg Retardtabletten	DE/H/3645/005	87979.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 40 mg Retardtabletten	DE/H/3645/006	87980.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 5 mg Retardtabletten	DE/H/3645/001	87975.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 60 mg Retardtabletten	DE/H/3645/007	87981.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Oxyconoica 80 mg Retardtabletten	DE/H/3645/008	87982.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
OxyContin ®160 mg	not available	PA 1688/5/10	MUNDIPHARMA	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
prolonged release tablets			PHARMACEUTICALS LIMITED	
OxyContin 1 mg/ml Lösung zum Einnehmen	UK/H/2780/001	83866.00.00	MUNDIPHARMA GMBH	DE
OxyContin 10 mg comprimidos de liberación prolongada	IE/H/0112/001	63.446	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 10 mg depottabletter	not available	94-2768	MUNDIPHARMA AS.	NO
OxyContin 10 mg depottabletter	not available	13830	MUNDIPHARMA AB	SE
OxyContin 10 mg prolonged release tablets	not available	18608	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 10 mg prolonged release tablets	not available	PL 16950/0097	NAPP PHARMACEUTICALS LTD	UK
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/01/01201/002	MEDIS, D.O.O.	SI
OxyContin 10 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/001	MEDIS, D.O.O.	SI
OxyContin 10 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/003	MEDIS, D.O.O.	SI
OXYCONTIN 10 mg, comprimés à libération prolongée	DE/H/0366/001	BE253197	MUNDIPHARMA COMM VA	BE
OXYCONTIN 10 mg, comprimés à libération prolongée	DE/H/0366/001	1495/03/10/0017	MUNDIPHARMA COMM VA	LU
OXYCONTIN 10 mg, forðatöflur	DE/H/0366/001	IS/1/02/047/01	NORPHARMA A/S	IS
OxyContin 10 mg, tabletten met verlengde afgifte	DE/H/0366/001	BE253197	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
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 tablety s prodlouženým uvolňováním	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 prolonged release tablets	not available	PL 16950/0150	NAPP PHARMACEUTICALS LTD	UK
OXYCONTIN 120 mg, comprimés à libération prolongée	DE/H/0366/012	BE348993	MUNDIPHARMA COMM VA	BE
OXYCONTIN 120 mg, comprimés à libération prolongée	DE/H/0366/012	1495/10030087	MUNDIPHARMA COMM VA	LU
OxyContin 120 mg, tabletten met verlengde afgifte	DE/H/0366/012	BE348993	MUNDIPHARMA COMM VA	BE
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 15 mg prolonged release tablets	not available	PL 16950/ 0139	NAPP PHARMACEUTICALS LTD	UK
OXYCONTIN 15 mg, comprimés à libération prolongée	DE/H/0366/009	BE348966	MUNDIPHARMA COMM VA	BE
OXYCONTIN 15 mg, comprimés à libération prolongée	DE/H/0366/009	1495/10030084	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
OxyContin 15 mg, tabletten met verlengde afgifte	DE/H/0366/009	BE348966	MUNDIPHARMA COMM VA	BE
OxyContin 15 mg, tabletten met verlengde afgifte	not available	RVG 100819	MUNDIPHARMA PHARMACEUTICALS BV	NL
OXYCONTIN 160 mg, comprimés à libération prolongée	DE/H/0366/013	BE349002	MUNDIPHARMA COMM VA	BE
OXYCONTIN 160 mg, comprimés à libération prolongée	DE/H/0366/013	1495/10030088	MUNDIPHARMA COMM VA	LU
OxyContin 160 mg, tabletten met verlengde afgifte	DE/H/0366/013	BE349002	MUNDIPHARMA COMM VA	BE
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	63.447	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	not available	PL 16950/0098	NAPP PHARMACEUTICALS LTD	UK
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/01/01201/005	MEDIS, D.O.O.	SI
OxyContin 20 mg tablete s podaljšanim	not available	H/01/01201/004	MEDIS, D.O.O.	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
sproščanjem				
OxyContin 20 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/006	MEDIS, D.O.O.	SI
OxyContin 20 mg tablete s produljenim oslobadanjem	DE/H/3862/003	HR-H-188823708	MUNDIPHARMA GESELLSCHAFT M.B.H.	HR
OXYCONTIN 20 mg, comprimés à libération prolongée	DE/H/0366/002	BE253242	MUNDIPHARMA COMM VA	BE
OXYCONTIN 20 mg, comprimés à libération prolongée	DE/H/0366/002	1495/03/10/0018	MUNDIPHARMA COMM VA	LU
OXYCONTIN 20 mg, forðatöflur	DE/H/0366/002	IS/1/02/047/02	NORPHARMA A/S	IS
OxyContin 20 mg, tabletten met verlengde afgifte	DE/H/0366/002	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 tablety s prodlouženým uvolňováním	not available	65/258/00-C	MUNDIPHARMA GES.M.B.H	CZ
OxyContin 30 mg depottabletter	not available	07-4848	MUNDIPHARMA AS.	NO
OxyContin 30 mg prolonged release tablets	not available	PL 16950/0140	NAPP PHARMACEUTICALS LTD	UK
OXYCONTIN 30 mg, comprimés à libération prolongée	DE/H/0366/010	BE348975	MUNDIPHARMA COMM VA	BE
OXYCONTIN 30 mg,	DE/H/0366/010	1495/10030085	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
comprimés à libération prolongée				
OxyContin 30 mg, tabletten met verlengde afgifte	DE/H/0366/010	BE348975	MUNDIPHARMA COMM VA	BE
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	63.448	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 prolonged release tablets	not available	18610	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 40 mg prolonged release tablets	not available	PL 16950/0099	NAPP PHARMACEUTICALS LTD	UK
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/01/01201/008	MEDIS, D.O.O.	SI
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/007	MEDIS, D.O.O.	SI
OxyContin 40 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/009	MEDIS, D.O.O.	SI
OxyContin 40 mg tablete s produljenim oslobadanjem	DE/H/1026/004	HR-H-849997645	MUNDIPHARMA GESELLSCHAFT M.B.H.	HR
OXYCONTIN 40 mg, comprimés à libération prolongée	DE/H/0366/003	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
OXYCONTIN 40 mg, comprimés à libération prolongée	DE/H/0366/003	1495/03/10/0019	MUNDIPHARMA COMM VA	LU
OXYCONTIN 40 mg, forðatöflur	DE/H/0366/003	IS/1/02/047/03	NORPHARMA A/S	IS
OxyContin 40 mg, tabletten met verlengde afgifte	DE/H/0366/003	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 tablety s prodlouženým uvolňováním	not available	65/259/00-C	MUNDIPHARMA GES.M.B.H	CZ
OxyContin 5 mg comprimidos de liberación prolongada	IE/H/0112/005	68.605	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 film-coated, prolonged release tablets	not available	20346	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyContin 5 mg prolonged release tablets	not available	PL 16950/0123	NAPP PHARMACEUTICALS LTD	UK
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

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, forðatöflur	DE/H/0366/005	IS/1/05/021/01	NORPHARMA A/S	IS
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 60 mg prolonged release tablets	not available	PL 16950/0141	NAPP PHARMACEUTICALS LTD	UK
OXYCONTIN 60 mg, comprimés à libération prolongée	DE/H/0366/011	BE348984	MUNDIPHARMA COMM VA	BE
OXYCONTIN 60 mg, comprimés à libération prolongée	DE/H/0366/011	1495/10030086	MUNDIPHARMA COMM VA	LU
OxyContin 60 mg, tabletten met verlengde afgifte	DE/H/0366/011	BE348984	MUNDIPHARMA COMM VA	BE
OxyContin 60 mg, tabletten met verlengde afgifte	not available	RVG 100821	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyContin 80 mg comprimidos de liberación prolongada	IE/H/0112/004	63.449	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyContin 80 mg depottabletter	not available	01-2425	MUNDIPHARMA AS.	NO
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	not available	PL 16950/0100	NAPP PHARMACEUTICALS LTD	UK
OxyContin 80 mg retard	not available	OGYI-T-7166/06	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
filmtabletta				
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/011	MEDIS, D.O.O.	SI
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/010	MEDIS, D.O.O.	SI
OxyContin 80 mg tablete s podaljšanim sproščanjem	not available	H/01/01201/012	MEDIS, D.O.O.	SI
OxyContin 80 mg tablete s produljenim oslobadanjem	DE/H/1026/005	HR-H-76344 1426	MUNDIPHARMA GESELLSCHAFT M.B.H.	HR
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	1495/03/10/0020	MUNDIPHARMA COMM VA	LU
OXYCONTIN 80 mg, forðatöflur	DE/H/0366/004	IS/1/02/047/04	NORPHARMA A/S	IS
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 80mg tablety s prodlouženým uvolňováním	not available	65/260/00-C	MUNDIPHARMA GES.M.B.H	CZ
OxyContin I 0 mg tablete s produljenim	DE/H/1026/002	HR-H- 11 8192252	MUNDIPHARMA GESELLSCHAFT 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
oslobadanjem				
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 210 9 8	MUNDIPHARMA	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 209 0 9	MUNDIPHARMA	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 208 4 8	MUNDIPHARMA	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 207 8 7	MUNDIPHARMA	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 211 5 9	MUNDIPHARMA	FR
OXYCONTIN LP 10 mg, comprimé pelliculé à libération prolongée	not available	34009 354 206 1 9	MUNDIPHARMA	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 604 5 2	MUNDIPHARMA	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 603 9 1	MUNDIPHARMA	FR
OXYCONTIN LP 120 mg, comprimé pelliculé à libération prolongée	not available	34009 384 602 2 3	MUNDIPHARMA	FR
OXYCONTIN LP 15 mg, comprimé pelliculé à libération prolongée	not available	34009 384 586 7 1	MUNDIPHARMA	FR
OXYCONTIN LP 15 mg, comprimé pelliculé à libération prolongée	not available	34009 384 585 0 3	MUNDIPHARMA	FR
OXYCONTIN LP 15 mg, comprimé pelliculé à	not available	34009 384 584 4 2	MUNDIPHARMA	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 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 215 0 0	MUNDIPHARMA	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 216 7 8	MUNDIPHARMA	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 214 4 9	MUNDIPHARMA	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 213 8 8	MUNDIPHARMA	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 217 3 9	MUNDIPHARMA	FR
OXYCONTIN LP 20 mg, comprimé pelliculé à libération prolongée	not available	34009 354 212 1 0	MUNDIPHARMA	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à libération prolongée	not available	34009 384 587 3 2	MUNDIPHARMA	FR
OXYCONTIN LP 30 mg, comprimé pelliculé à libération prolongée	not available	34009 384 589 6 1	MUNDIPHARMA	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 223 3 3	MUNDIPHARMA	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 220 4 0	MUNDIPHARMA	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 221 0 1	MUNDIPHARMA	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à	not available	34009 354 222-7 0	MUNDIPHARMA	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 354 225 6 9	MUNDIPHARMA	FR
OXYCONTIN LP 40 mg, comprimé pelliculé à libération prolongée	not available	34009 354 219 6 8	MUNDIPHARMA	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 905 7 8	MUNDIPHARMA	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 904 0 9	MUNDIPHARMA	FR
OXYCONTIN LP 5 mg, comprimé pelliculé à libération prolongée	not available	34009 366 903 4 9	MUNDIPHARMA	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 599 1 3	MUNDIPHARMA	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 601 6 2	MUNDIPHARMA	FR
OXYCONTIN LP 60 mg, comprimé pelliculé à libération prolongée	not available	34009 384 598 5 2	MUNDIPHARMA	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 296 0 5	MUNDIPHARMA	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 295 4 4	MUNDIPHARMA	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 228 5 9	MUNDIPHARMA	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à	not available	34009 354 227 9 8	MUNDIPHARMA	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 354 294 8 3	MUNDIPHARMA	FR
OXYCONTIN LP 80 mg, comprimé pelliculé à libération prolongée	not available	34009 354 226 2 0	MUNDIPHARMA	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, tabletki o przedłu??onym uwalnianiu	DE/H/1026/002	14525	NORPHARMA A/S	PL
OxyContin, 20 mg, tabletki o przedłu??onym uwalnianiu	DE/H/1026/003	14526	NORPHARMA A/S	PL
OxyContin, 40 mg, tabletki o przedłu??onym uwalnianiu	DE/H/1026/004	14527	NORPHARMA A/S	PL
OxyContin, 5 mg, tabletki o przedłu??onym uwalnianiu	DE/H/1026/001	14524	NORPHARMA A/S	PL
OxyContin, 80 mg, tabletki o przedluzonym uwalnianiu	DE/H/1026/005	14528	NORPHARMA A/S	PL
OxyContin, depottabletter	not available	16893	MUNDIPHARMA A/S	DK
OxyContin, depottabletter	not available	41088	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, depottabletter	not available	41089	MUNDIPHARMA A/S	DK
OxyContin, depottabletter	not available	42029	MUNDIPHARMA A/S	DK
OxyContin, depottabletter	not available	16894	MUNDIPHARMA A/S	DK
OxyContin, depottabletter	not available	30177	MUNDIPHARMA A/S	DK
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435038	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435053	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435040	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435026/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435014	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg compresse a rilascio prolungato	IE/H/0112/001	034435065	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 10 mg depottabletit	not available	11969	MUNDIPHARMA OY	FI
OxyContin® 10 mg prolonged release tablets	IE/H/0112/001	PA 1688/5/2	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 120 mg prolonged release tablets	not available	PA 1688/5/9	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 15 mg prolonged release tablets	not available	PA 1688/5/6	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 20 mg compresse a rilascio	IE/H/0112/002	034435077	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
prolungato				
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/002	034435089	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/002	034435091	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/002	034435115	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/002	034435127	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg compresse a rilascio prolungato	IE/H/0112/002	034435103	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 20 mg prolonged release tablets	IE/H/0112/002	PA 1688/5/3	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 20 mg depottabletit	not available	11970	MUNDIPHARMA OY	FI
OxyContin® 30 mg prolonged release tablets	not available	PA 1688/5/7	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/003	034435141	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/003	034435178	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/003	034435180	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/003	034435154	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg compresse a rilascio	IE/H/0112/003	034435139	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
prolungato				
OxyContin® 40 mg compresse a rilascio prolungato	IE/H/0112/003	034435166	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 40 mg prolonged release tablets	IE/H/0112/003	PA 1688/5/4	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 40 mg depottabletit	not available	11971	MUNDIPHARMA OY	FI
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	034435255	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	034435279	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	034435267	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	034435293/M	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg compresse a rilascio prolungato	IE/H/0112/005	034435281	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 5 mg depottabletit	not available	16771	MUNDIPHARMA OY	FI
OxyContin® 5 mg prolonged release tablets	IE/H/0112/005	PA 1688/5/1	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 60 mg prolonged release tablets	not available	PA 1688/5/8	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/004	034435192	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/004	034435230	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg	IE/H/0112/004	034435242	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	IE/H/0112/004	034435216	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/004	034435228	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg compresse a rilascio prolungato	IE/H/0112/004	034435204	MUNDIPHARMA PHARMACEUTICALS SRL	IT
OxyContin® 80 mg depottabletit	not available	13673	MUNDIPHARMA OY	FI
OxyContin® 80 mg prolonged release tablets	IE/H/0112/004	PA 1688/5/5	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
Oxydol 10 mg hörð forðahylki	IS/H/0196/001	IS/1/13/034/01	ETHYPHARM	IS
Oxydol 20 mg hörð forðahylki	IS/H/0196/002	IS/1/13/034/02	ETHYPHARM	IS
Oxydol 40 mg hörð forðahylki	IS/H/0196/003	IS/1/13/034/03	ETHYPHARM	IS
Oxydol 80 mg hörð forðahylki	IS/H/0196/004	IS/1/13/034/04	ETHYPHARM	IS
Oxydol, hårde depotkapsler	IS/H/0196/001	49886	ETHYPHARM	DK
Oxydol, hårde depotkapsler	IS/H/0196/002	49887	ETHYPHARM	DK
Oxydol, hårde depotkapsler	IS/H/0196/003	49888	ETHYPHARM	DK
Oxydol, hårde depotkapsler	IS/H/0196/004	49889	ETHYPHARM	DK
Oxydolor, 10 mg, tabletki o przedłużonym uwalnianiu	DE/H/2182/002	17818	G.L. PHARMA GMBH	PL
Oxydolor, 20 mg,	DE/H/2182/003	17817	G.L. PHARMA GMBH	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletki o przedłużonym uwalnianiu				
Oxydolor, 40 mg, tabletki o przedłużonym uwalnianiu	DE/H/2182/004	17816	G.L. PHARMA GMBH	PL
Oxydolor, 5 mg, tabletki o przedłużonym uwalnianiu	DE/H/2182/001	17819	G.L. PHARMA GMBH	PL
Oxydolor, 80 mg, tabletki o przedłużonym uwalnianiu	DE/H/2182/005	17815	G.L. PHARMA GMBH	PL
Oxydon 10 mg Prolonged-Release Tablets	DE/H/1419/002	PA 711/115/5	ROWEX LTD	IE
Oxydon 10mg Prolonged-Release Tablets	DE/H/1419/002	PA 0711/115/005	ROWEX LTD	IE
Oxydon 40mg Prolonged-Release Tablets	DE/H/0789/002	PA0711/115/002	ROWEX LTD	IE
Oxydon 40mg Prolonged-Release Tablets	DE/H/0789/002	PA 711/115/2	ROWEX LTD	IE
Oxydon 5 mg Prolonged-release tablets	DE/H/1419/001	PA 0711/115/004	ROWEX LTD	IE
Oxydon 80mg Prolonged-Release Tablets	DE/H/0789/003	PA0711/115/003	ROWEX LTD	IE
Oxygerolan 10 mg-Filmtabletten	DE/H/2669/002	1-30178	G.L. PHARMA GMBH	AT
Oxygerolan 10 mg-Retardtabletten	DE/H/2182/002	1-29620	G.L. PHARMA GMBH	AT
Oxygerolan 20 mg-Filmtabletten	DE/H/2669/004	137551	G.L. PHARMA GMBH	AT
Oxygerolan 20 mg-Retardtabletten	DE/H/2182/003	1-29621	G.L. PHARMA GMBH	AT
Oxygerolan 40 mg-Retardtabletten	DE/H/2182/004	1-29622	G.L. PHARMA GMBH	AT
Oxygerolan 5 mg-	DE/H/2669/001	1-30177	G.L. PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Filmtabletten				
Oxygerolan 5 mg-Retardtabletten	DE/H/2182/001	1-29619	G.L. PHARMA GMBH	AT
Oxygerolan 80 mg-Retardtabletten	DE/H/2182/005	1-29624	G.L. PHARMA GMBH	AT
Oxygesic 10 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/0366/001	41153.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 10 mg/ml Lösung zum Einnehmen	UK/H/2780/002	83865.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 120 mg Retardtabletten	DE/H/0366/012	67600.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 15 mg Retardtabletten	DE/H/0366/009	67597.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 160 mg Retardtabletten	DE/H/0366/013	67601.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 20 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/0366/002	41153.01.00	MUNDIPHARMA GMBH	DE
Oxygesic 30 mg Retardtabletten	DE/H/0366/010	67598.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 40 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/0366/003	41153.02.00	MUNDIPHARMA GMBH	DE
Oxygesic 5 mg Retardtabletten Wirkstoff: Oxycodonhydrochlorid	DE/H/0366/005	56594.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 60 mg Retardtabletten	DE/H/0366/011	67599.00.00	MUNDIPHARMA GMBH	DE
Oxygesic 80 mg Retardtabletten Wirkstoff:	DE/H/0366/004	41153.03.00	MUNDIPHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxycodonhydrochlorid				
Oxygesic infusio 50 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/1026/014	72147.00.00	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/0366/003	68388.00.00	MUNDIPHARMA GMBH	DE
Oxygesic® Dispersa 20 mg Schmelztabletten	DE/H/0366/002	68389.00.00	MUNDIPHARMA GMBH	DE
Oxygesic® Dispersa 5 mg Schmelztabletten	DE/H/0366/001	68387.00.00	MUNDIPHARMA GMBH	DE
Oxygesic® injekt 10 mg/1 ml Injektionslösung	DE/H/0366/015	57886.00.00	MUNDIPHARMA GMBH	DE
Oxygesic® injekt 20 mg/2 ml Injektionslösung	not available	57887.00.00	MUNDIPHARMA GMBH	DE
Oxykodon Mylan 10 mg tablety s prodlouženým uvolňováním	DE/H/4246/002	65/541/15-C	GENERICS [UK] LIMITED	CZ
Oxykodon Mylan 20 mg tablety s prodlouženým uvolňováním	DE/H/4246/003	65/542/15-C	GENERICS [UK] LIMITED	CZ
Oxykodon Mylan 40 mg tablety s prodlouženým uvolňováním	DE/H/4246/005	65/544/15-C	GENERICS [UK] LIMITED	CZ
Oxykodon Mylan 80 mg tablety s prodlouženým uvolňováním	DE/H/4246/007	65/546/15-C	GENERICS [UK] LIMITED	CZ
Oxykodón Sandoz 10 mg	DE/H/4502/002	65/0337/17-S	SANDOZ PHARMACEUTICALS	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tvrdé kapsuly			D.D.	
Oxykodón Sandoz 20 mg tvrdé kapsuly	DE/H/4502/003	65/0338/17-S	SANDOZ PHARMACEUTICALS D.D.	SK
Oxykodón Sandoz 5 mg tvrdé kapsuly	DE/H/4502/001	65/0336/17-S	SANDOZ PHARMACEUTICALS D.D.	SK
Oxykodon Stada 10 mg tvrdé tobolky	DE/H/4774/002	65/776/16-C	STADA ARZNEIMITTEL AG	CZ
Oxykodon Stada 20 mg tvrdé tobolky	DE/H/4774/003	65/777/16-C	STADA ARZNEIMITTEL AG	CZ
Oxykodon Stada 5 mg tvrdé tobolky	DE/H/4774/001	65/775/16-C	STADA ARZNEIMITTEL AG	CZ
Oxykodon Teva 10 mg tablety s prodlouženým uvolňováním	SE/H/1313/002	65/196/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxykodon Teva 20 mg tablety s prodlouženým uvolňováním	SE/H/1313/004	65/198/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxykodon Teva 40 mg tablety s prodlouženým uvolňováním	SE/H/1313/006	65/200/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxykodon Teva 5 mg tablety s prodlouženým uvolňováním	SE/H/1313/001	65/195/14-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxylan 10 mg prolonged-release tablets	DE/H/2183/002	PL 17049/00010 - 0001	LANNACHER HEILMITTEL GES.M.B.H.,	UK
Oxylan 20 mg prolonged-release tablets	DE/H/2183/002	PL 17049/00011 - 0001	LANNACHER HEILMITTEL GES.M.B.H.,	UK
Oxylan 40 mg prolonged-release tablets	DE/H/2183/004	PL 17049/00012 - 0001	LANNACHER HEILMITTEL GES.M.B.H.,	UK
Oxylan 5 mg prolonged-release tablets	DE/H/2183/001	PL 17049/0009 - 0001		UK
Oxylan 80 mg prolonged-release tablets	DE/H/2183/005	PL 17049/00013 - 0001	LANNACHER HEILMITTEL GES.M.B.H.,	UK
Oxylor Depot 10 mg depotkapslar, hårda	IS/H/0196/001	47193	ETHYPHARM	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
Oxylor Depot 20 mg depotkapslar, hårda	IS/H/0196/002	47194	ETHYPHARM	SE
Oxylor Depot 40 mg depotkapslar, hårda	IS/H/0196/003	47195	ETHYPHARM	SE
Oxylor Depot 80 mg depotkapslar, hårda	IS/H/0196/004	47196	ETHYPHARM	SE
OxyNorm 1 mg/ml drank	DE/H/0366/016	BE422037	MUNDIPHARMA COMM VA	BE
OxyNorm 1 mg/ml mikstur, oppløsning	not available	01-9979	MUNDIPHARMA AS.	NO
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	not available	PL 16950/0107	NAPP PHARMACEUTICALS LTD	UK
OxyNorm 10 mg cápsulas	IE/H/0139/004	66575	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 10 mg capsules	IE/H/0139/004	PA 1688/6/5	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 Hartkapseln	not available	58373.00.00	MUNDIPHARMA GMBH	DE
OxyNorm 10 mg Kapseln	IE/H/0139/004	1-25660	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyNorm 10 mg smeltetabletter	not available	07-4737	MUNDIPHARMA AS.	NO
OXYNORM 10 mg, gélule	not available	34009 362 899 9 7	MUNDIPHARMA	FR
OXYNORM 10 mg, gélule	not available	34009 361 898 2 9	MUNDIPHARMA	FR
OXYNORM 10 mg, gélule	not available	34009 361 897 6 8	MUNDIPHARMA	FR
OXYNORM 10 mg, gélule	not available	34009 362 900 7 8	MUNDIPHARMA	FR
OXYNORM 10 mg, gélule	not available	34009 362 421 5 9	MUNDIPHARMA	FR
OXYNORM 10 mg, gélule	not available	34009 362 420 9 8	MUNDIPHARMA	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 10 mg, harde capsules	not available	RVG 27510	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm 10 mg/ml injeksjonsvæske/infusjonsvæske, oppløsning	not available	02-216	MUNDIPHARMA AS.	NO
OxyNorm 10 mg/ml Injektions-/Infusionslösung	IE/H/0139/006	1-28001	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
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 oral lösning	not available	15793	MUNDIPHARMA AB	SE
OxyNorm 10 mg/ml solución inyectable y para perfusión	IE/H/0139/006	71.243	MUNDIPHARMA PHARMACEUTICALS SL	ES
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 stungulyf, lausn, stungulyfs-/innrennslisþykkni, lausn	DE/H/0366/015	IS/1/09/016/01	NORPHARMA A/S	IS
OxyNorm 10 mg/ml, oplessing voer injectie/concentraat veer eplossing veor injectie of infusie	DE/H/0366/015	BE339814	MUNDIPHARMA COMM VA	BE
OxyNorm 10 mg/ml, oplossing voor injectie/concentraat voor oplossing voor injectie of infusie	DE/H/0366/015	BE339823	MUNDIPHARMA COMM VA	BE
OXYNORM 10 mg/ml, solution buvable	not available	34009 368 458 8 6	MUNDIPHARMA	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 10 mg/ml, solution buvable	not available	34009 366 912 3 0	MUNDIPHARMA	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	not available	PL 16950/0128	NAPP PHARMACEUTICALS LTD	UK
OXYNORM 10 mg/ml, solution injectable	not available	34009 392 317 1 6	MUNDIPHARMA	FR
OXYNORM 10 mg/ml, solution injectable	not available	34009 366 915 2 0	MUNDIPHARMA	FR
OXYNORM 10 mg/ml, solution injectable	not available	34009 366 914 6 9	MUNDIPHARMA	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 10 mg/ml, solution injectable/solution à diluer injectable ou perfusion	DE/H/0366/015	2009080027	MUNDIPHARMA COMM VA	LU
OxyNorm 20 mg	not available	20305	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm 20 mg	not available	PL 16950/0108	NAPP PHARMACEUTICALS LTD	UK
OxyNorm 20 mg cápsulas	IE/H/0139/005	66576	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 20 mg	IE/H/0139/005	PA 1688/6/6	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
capsules			PHARMACEUTICALS LIMITED	
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 Hartkapseln	not available	58374.00.00	MUNDIPHARMA GMBH	DE
OxyNorm 20 mg Kapseln	IE/H/0139/005	1-25661	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT
OxyNorm 20 mg smeltetabletter	not available	07-4738	MUNDIPHARMA AS.	NO
OXYNORM 20 mg, gélule	not available	34009 362 423 8 8	MUNDIPHARMA	FR
OXYNORM 20 mg, gélule	not available	34009 361 905 9 7	MUNDIPHARMA	FR
OXYNORM 20 mg, gélule	not available	34009 361 904 2 9	MUNDIPHARMA	FR
OXYNORM 20 mg, gélule	not available	34009 361 903 6 8	MUNDIPHARMA	FR
OXYNORM 20 mg, gélule	not available	34009 361 901 3 9	MUNDIPHARMA	FR
OXYNORM 20 mg, gélule	not available	34009 362 422 1 0	MUNDIPHARMA	FR
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	not available	PL 16950/0106	NAPP PHARMACEUTICALS LTD	UK
OxyNorm 5 mg cápsulas	IE/H/0139/003	66574	MUNDIPHARMA PHARMACEUTICALS SL	ES
OxyNorm 5 mg capsules	IE/H/0139/003	PA 1688/6/4	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 Hartkapseln	not available	58372.00.00	MUNDIPHARMA GMBH	DE
OxyNorm 5 mg Kapseln	IE/H/0139/003	1-25659	MUNDIPHARMA GESELLSCHAFT M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OxyNorm 5 mg smeltetabletter	not available	07-4736	MUNDIPHARMA AS.	NO
OXYNORM 5 mg, gélule	not available	34009 361 892 4 0	MUNDIPHARMA	FR
OXYNORM 5 mg, gélule	not available	34009 361 895 3 9	MUNDIPHARMA	FR
OXYNORM 5 mg, gélule	not available	34009 361 893 0 0	MUNDIPHARMA	FR
OXYNORM 5 mg, gélule	not available	34009 362 419 0 9	MUNDIPHARMA	FR
OXYNORM 5 mg, gélule	not available	34009 361 894 7 8	MUNDIPHARMA	FR
OXYNORM 5 mg, gélule	not available	34009 362 418 4 8	MUNDIPHARMA	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 injektionsvätska, lösning	not available	26691	MUNDIPHARMA AB	SE
OxyNorm 50 mg/ml, concentraat voor oplossing voor infusie	DE/H/0366/014	BE346701	MUNDIPHARMA COMM VA	BE
OXYNORM 50 mg/ml, solution à diluer pour perfusion	DE/H/0366/014	BE346701	MUNDIPHARMA GMBH	BE
OXYNORM 50 mg/ml, solution à diluer pour perfusion	DE/H/0366/014	1495/09120027	MUNDIPHARMA COMM VA	LU
OXYNORM 50 mg/ml, solution injectable	not available	34009 387 625 3 2	MUNDIPHARMA	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	not available	PL 16950/0004	NAPP PHARMACEUTICALS LTD	UK
OxyNorm concentrate 10mg/ml oral solution	IE/H/0139/002	PA 1688/6/3	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 10	not available	PA 1688/6/8	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
mg orodispersible tablets			PHARMACEUTICALS LIMITED	
OxyNorm Dispersa 10 mg orodispersible tablets	not available	PL06934/0220	ETHYPHARM	UK
OxyNorm Dispersa 10 mg, munndreifitöflur	DE/H/0366/007	IS/1/07/015/02	NORPHARMA A/S	IS
OxyNorm Dispersa 20 mg orodispersible tablets	not available	PA 1688/6/9	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 20 mg orodispersible tablets	not available	PL06934/0219	ETHYPHARM	UK
OxyNorm Dispersa 20 mg, munndreifitöflur	DE/H/0366/008	IS/1/07/015/03	NORPHARMA A/S	IS
OxyNorm Dispersa 5 mg orodispersible tablets	not available	PA 1688/6/7	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm Dispersa 5 mg orodispersible tablets	not available	PL06934/0219	ETHYPHARM	UK
OxyNorm Dispersa 5 mg, munndreifitöflur	DE/H/0366/006	IS/1/07/015/01	NORPHARMA A/S	IS
OxyNorm Dispersa, smeltetabletter	not available	40537	NORPHARMA A/S	DK
OxyNorm Dispersa, smeltetabletter	not available	40538	NORPHARMA A/S	DK
OxyNorm Dispersa, smeltetabletter	not available	40539	NORPHARMA A/S	DK
OxyNorm drank 10 mg/ml	DE/H/0366/017	BE422046	MUNDIPHARMA COMM VA	BE
OxyNorm drank 10 mg/ml	not available	RVG 27940	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm drank 5 mg I 5 ml	not available	RVG 27939	MUNDIPHARMA PHARMACEUTICALS BV	NL
Oxynorm hårde kapsler	not available	31296	NORPHARMA 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	not available	RVG 101605	MUNDIPHARMA PHARMACEUTICALS BV	NL

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
injectie of infusie				
OxyNorm Instant 10 mg, comprimés orodispersibles	DE/H/0366/007	1495/08110045	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 10 mg, orodispergeerbare tabletten	DE/H/0366/007	BE319076	MUNDIPHARMA COMM VA	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	BE 319076	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 20 mg, comprimés orodispersibles	DE/H/0366/008	1495/08110043	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 20 mg, orodispergeerbare tabletten	DE/H/0366/008	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/0366/008	BE 319085	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 5 mg, comprimés orodispersibles	DE/H/0366/006	1495/08110044	MUNDIPHARMA COMM VA	LU
OxyNorm Instant 5 mg, orodispergeerbare tabletten	DE/H/0366/006	BE319067	MUNDIPHARMA COMM VA	BE
OxyNorm Instant 5 mg, orodispergeerbare tabletten	not available	RVG 34775	MUNDIPHARMA PHARMACEUTICALS BV	NL
OxyNorm Instant 5 mg, comprimés	DE/H/0366/006	BE 319067	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
orodispersibles				
OxyNorm liquid 1mg/ml oral solution	IE/H/0139/001	PA 1688/6/2	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm liquid 5mg/5ml	not available	20972	MUNDIPHARMA PHARMACEUTICALS LTD	CY
OxyNorm liquid 5mg/5ml oral solution	not available	PL 16950/0003	NAPP PHARMACEUTICALS LTD	UK
OxyNorm, 1 mg/ml, roztwór doustny	DE/H/1026/007	20461	NORPHARMA A/S	PL
OxyNorm, 10 mg/ml, roztwór do wstrzykiwan/koncentral do sporz~dzan.ia roztworu do infuzji	PL/H/0355/006	15220	NORPHARMA A/S	PL
Oxynorm, hårde kapsler	not available	31294	NORPHARMA A/S	DK
Oxynorm, hårde kapsler	not available	31295	NORPHARMA A/S	DK
Oxynorm, injektions- og infusionsvæske, opløsning	not available	33361	MUNDIPHARMA A/S	DK
Oxynorm, oral opløsning	not available	31158	NORPHARMA A/S	DK
Oxynorm, oral opløsning	not available	31157	NORPHARMA A/S	DK
OxyNorm® 1 mg/ml oraaliliuos	not available	14807	MUNDIPHARMA OY	FI
OxyNorm® 10 mg kapseli, kova	not available	15321	MUNDIPHARMA OY	FI
OxyNorm® 10 mg/ml oraaliliuos	not available	14808	MUNDIPHARMA OY	FI
OxyNorm® 10 mg/ml, injektio-/infuusioneste, liuos	not available	17150	MUNDIPHARMA OY	FI
OxyNorm® 20 mg kapseli, kova	not available	15322	MUNDIPHARMA OY	FI
OxyNorm® 5 mg, kapseli, kova	not available	15609	MUNDIPHARMA OY	FI
OxyNorm® 50 mg/ml,	not available	24824	MUNDIPHARMA OY	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
injektio-/infuusioneste, liuos				
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 1688/6/10	MUNDIPHARMA PHARMACEUTICALS LIMITED	IE
OxyNorm® 50 mg/ml, solution for injection or infusion	not available	PL 16950/0155	NAPP PHARMACEUTICALS LTD	UK
OxyNorm1 mg/ml solution buvable	DE/H/0366/016	BE422037	MUNDIPHARMA COMM VA	BE
OxyNorm10 mg/ml solution buvable	DE/H/0366/017	BE422046	MUNDIPHARMA COMM VA	BE
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 426 5 8	MUNDIPHARMA	FR
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 427 1 9	MUNDIPHARMA	FR
OXYNORMORO 10 mg, comprimé orodispersible	not available	34009 380 425 9 7	MUNDIPHARMA	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 429 4 8	MUNDIPHARMA	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 430 2 0	MUNDIPHARMA	FR
OXYNORMORO 20 mg, comprimé orodispersible	not available	34009 380 428 8 7	MUNDIPHARMA	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 424 2 9	MUNDIPHARMA	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 423 6 8	MUNDIPHARMA	FR
OXYNORMORO 5 mg, comprimé orodispersible	not available	34009 380 421 3 9	MUNDIPHARMA	FR
Oxypro 10 mg Retardtabletten	DE/H/2183/002	76282.00.00	G.L. 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
Oxypro 10 mg tablety s predĺženým uvoľňovaním	DE/H/2182/002	65/0321/10-S	G.L. PHARMA GMBH	SK
Oxypro 20 mg Retardtabletten	DE/H/2183/003	76283.00.00	G.L. PHARMA GMBH	DE
Oxypro 20 mg tablety s predĺženým uvoľňovaním	DE/H/2182/003	65/0322/10-S	G.L. PHARMA GMBH	SK
Oxypro 40 mg Retardtabletten	DE/H/2183/004	76284.00.00	G.L. PHARMA GMBH	DE
Oxypro 40 mg tablety s predĺženým uvoľňovaním	DE/H/2182/004	65/0323/10-S	G.L. PHARMA GMBH	SK
Oxypro 5 mg Retardtabletten	DE/H/2183/001	76281.00.00	G.L. PHARMA GMBH	DE
Oxypro 5 mg tablety s predĺženým uvoľňovaním	DE/H/2182/001	65/0320/10-S	G.L. PHARMA GMBH	SK
Oxypro 80 mg Retardtabletten	DE/H/2183/005	76285.00.00	G.L. PHARMA GMBH	DE
Oxypro 80 mg tablety s predĺženým uvoľňovaním	DE/H/2182/005	65/0324/10-S	G.L. PHARMA GMBH	SK
Oxyratio 10 mg kapseli, kova	SE/H/1423/002	32188	TEVA GMBH; ULM, GERMANY	FI
Oxyratio 20 mg kapseli, kova	SE/H/1423/003	32189	RATIOPHARM GMBH	FI
Oxyratio 5 mg kapseli, kova	SE/H/1423/001	32187	RATIOPHARM GMBH	FI
Oxytina 10 mg Retardtabletten	not available	93431.00.00	ALCHEMIA LIMITED	DE
Oxytina 15 mg Retardtabletten	not available	93432.00.00	ALCHEMIA LIMITED	DE
Oxytina 20 mg Retardtabletten	not available	93433.00.00	ALCHEMIA LIMITED	DE
Oxytina 30 mg Retardtabletten	not available	93434.00.00	ALCHEMIA LIMITED	DE
Oxytina 40 mg Retardtabletten	not available	93435.00.00	ALCHEMIA LIMITED	DE
Oxytina 5 mg	not available	93430.00.00	ALCHEMIA LIMITED	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
Retardtabletten				
Oxytina 60 mg Retardtabletten	not available	93436.00.00	ALCHEMIA LIMITED	DE
Oxytina 80 mg Retardtabletten	not available	93437.00.00	ALCHEMIA LIMITED	DE
Reltebon 10mg Prolongedrelease|Tablets	UK/H/6837/002	PL 30306/0465	ACTAVIS GROUP PTC EHF.	UK
Reltebon 15 mg Prolongedrelease Tablets	UK/H/6837/003	PL 30306/0466	ACTAVIS GROUP PTC EHF.	UK
Reltebon 20 mg Prolongedrelease Tablets	UK/H/6837/004	PL 30306/0467	ACTAVIS GROUP PTC EHF.	UK
Reltebon 30mg Prolongedrelease Tablets	UK/H/6837/005	PL 30306/0468	ACTAVIS GROUP PTC EHF.	UK
Reltebon 40mg Prolongedrelease Tablets	UK/H/6837/006	PL 30306/0469	ACTAVIS GROUP PTC EHF.	UK
Reltebon 5mg Prolongedrelease Tablets	UK/H/6837/001	PL 30306/0464	ACTAVIS GROUP PTC EHF.	UK
Reltebon 60mg Prolongedrelease Tablets	UK/H/6837/007	PL 30306/0470	ACTAVIS GROUP PTC EHF.	UK
Reltebon 80 mg prolongedrelease tablets	UK/H/6837/008	PL 30306/0471	ACTAVIS GROUP PTC EHF.	UK
Reltebon Depot, depottabletter	SE/H/1313/006	51828	ACTAVIS GROUP PTC EHF.	DK
Reltebon Depot, depottabletter	SE/H/1313/001	51823	ACTAVIS GROUP PTC EHF.	DK
Reltebon Depot, depottabletter	SE/H/1313/002	51824	ACTAVIS GROUP PTC EHF.	DK
Reltebon Depot, depottabletter	SE/H/1313/004	51826	ACTAVIS GROUP PTC EHF.	DK
Renocontin 10 mg prolonged-release tablets	DE/H/3645/002	PL 25258/0218	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 15 mg prolonged-release tablets	DE/H/3645/003	PL 25258/0219	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK

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
Renocontin 20 mg prolonged-release tablets	DE/H/3645/004	PL 25258/0220	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 30 mg prolonged-release tablets	DE/H/3645/005	PL 25258/0221	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 40 mg prolonged-release tablets	DE/H/3645/006	PL 25258/0222	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 5 mg prolonged-release tablets	DE/H/3645/001	PL 25258/0217	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 60 mg prolonged-release tablets	DE/H/3645/007	PL 25258/0223	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Renocontin 80 mg prolonged-release tablets	DE/H/3645/008	PL 25258/0224	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Shortec 10 mg capsules, hard	not available	PL 40431/0011	QDEM PHARMACEUTICALS LIMITED	UK
Shortec 10 mg/ml, solution for injection or infusion	not available	PL 40431/0015	QDEM PHARMACEUTICALS LIMITED	UK
Shortec 20 mg capsules, hard	not available	PL 40431/0012	QDEM PHARMACEUTICALS LIMITED	UK
Shortec 5 mg capsules, hard	not available	PL 40431/0010	QDEM PHARMACEUTICALS LIMITED	UK
Shortec 50 mg/ml, solution for injection or infusion	not available	PL 40431/0016	QDEM PHARMACEUTICALS LIMITED	UK
Shortec concentrate 10 mg/ml oral solution	not available	PL 40431/0014	QDEM PHARMACEUTICALS LIMITED	UK
Shortec liquid 5mg/5ml oral solution	not available	PL 40431/0013	QDEM PHARMACEUTICALS LIMITED	UK
Taioma 10 mg comprimidos de	DE/H/2827/002	77030	ARISTO PHARMA IBERIA, S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
liberación prolongada EFG				
Taioma 20 mg comprimidos de liberación prolongada EFG	DE/H/2827/003	77031	ARISTO PHARMA IBERIA, S.L.	ES
Taioma 40 mg comprimidos de liberación prolongada EFG	DE/H/2827/004	77032	ARISTO PHARMA IBERIA, S.L.	ES
Taioma 80 mg comprimidos de liberación prolongada EFG	DE/H/2827/005	77033	ARISTO PHARMA IBERIA, S.L.	ES
Taioma 5 mg comprimidos de liberación prolongada EFG	DE/H/2827/001	77029	ARISTO PHARMA IBERIA, S.L.	ES
Xancodal , 20 mg, kapsulki, twarde	DE/H/4502/003	24546	SANDOZ GMBH	PL
Xancodal , 5 mg, kapsulki, twarde	DE/H/4502/001	24544	SANDOZ GMBH	PL
Xancodal, 10 mg, kapsulki, twarde	DE/H/4502/002	24545	SANDOZ GMBH	PL
XANCODAL, 40 MG, TABLETKI O PRZEDŁUŻONYM UWALNIANIU	DE/H/3293/001	20885	SANDOZ GMBH	PL
XANCODAL, 60 MG, TABLETKI O PRZEDŁUŻONYM UWALNIANIU	DE/H/3293/002	20886	SANDOZ GMBH	PL
XANCODAL, 80 MG, TABLETKI O PRZEDŁUŻONYM UWALNIANIU	DE/H/3293/003	20887	SANDOZ GMBH	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zarenoxin 10 mg Retardtabletten	DE/H/3642/002	87952.00.00	ACINO AG	DE
Zarenoxin 15 mg Retardtabletten	DE/H/3642/003	87953.00.00	ACINO AG	DE
Zarenoxin 20 mg Retardtabletten	DE/H/3642/004	87954.00.00	ACINO AG	DE
Zarenoxin 30 mg Retardtabletten	DE/H/3642/005	87955.00.00	ACINO AG	DE
Zarenoxin 40 mg Retardtabletten	DE/H/3642/006	87956.00.00	ACINO AG	DE
Zarenoxin 5 mg Retardtabletten	DE/H/3642/001	87951.00.00	ACINO AG	DE
Zarenoxin 60 mg Retardtabletten	DE/H/3642/007	87957.00.00	ACINO AG	DE
Zarenoxin 80 mg Retardtabletten	DE/H/3642/008	87958.00.00	ACINO AG	DE
Zomestine 10 mg Prolonged-release Tablets	NL/H/3192/002	PA1390/40/002	ACCORD HEALTHCARE LIMITED	IE
Zomestine 10 mg prolonged-release tablets	NL/H/3192/002	PL 20075/0327	ACCORD HEALTHCARE LIMITED	UK
Zomestine 20 mg Prolonged-release Tablets	NL/H/3192/003	PA1390/40/003	ACCORD HEALTHCARE LIMITED	IE
Zomestine 20 mg prolonged-release tablets	NL/H/3192/003	PL 20075/0328	ACCORD HEALTHCARE LIMITED	UK
Zomestine 40 mg Prolonged-release Tablets	NL/H/3192/004	PA1390/40/004	ACCORD HEALTHCARE LIMITED	IE
Zomestine 40 mg prolonged-release tablets	NL/H/3192/004	PL 20075/0329	ACCORD HEALTHCARE LIMITED	UK
Zomestine 5 mg Prolonged-release Tablets	NL/H/3192/001	PA1390/40/001	ACCORD HEALTHCARE LIMITED	IE
Zomestine 5 mg	NL/H/3192/001	PL 20075/0326	ACCORD HEALTHCARE	UK

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			LIMITED	
Zomestine 80 mg Prolonged-release Tablets	NL/H/3192/005	PA1390/40/005	ACCORD HEALTHCARE LIMITED	IE
Zomestine 80 mg prolonged-release tablets	NL/H/3192/005	PL 20075/0330	ACCORD HEALTHCARE LIMITED	UK
Оксикодон Актавис 10 mg таблетки с удължено освобождаване	SE/H/1313/002	20140166	ACTAVIS GROUP PTC EHF.	BG
Оксикодон Актавис 10 mg твърди капсули	SE/H/1226/002	20130045	ACTAVIS GROUP PTC EHF.	BG
Оксикодон Актавис 20 mg таблетки с удължено освобождаване	SE/H/1313/004	20140168	ACTAVIS GROUP PTC EHF.	BG
Оксикодон Актавис 20 mg твърди капсули	SE/H/1226/003	20130046	ACTAVIS GROUP PTC EHF.	BG
Оксикодон Актавис 40 mg таблетки с удължено освобождаване	SE/H/1313/006	20140170	ACTAVIS GROUP PTC EHF.	BG
ОКСИКОНТИН 10 mg таблетки с изменено освобождаване	not available	20010056	MUNDIPHARMA GES.M.B.H	BG
ОКСИКОНТИН 20 mg таблетки с изменено освобождаване	not available	20010057	MUNDIPHARMA GES.M.B.H	BG
ОКСИКОНТИН 40 mg таблетки с изменено освобождаване	not available	20010058	MUNDIPHARMA GES.M.B.H	BG
ОКСИКОНТИН 80 mg таблетки с изменено освобождаване	not available	20010059	MUNDIPHARMA GES.M.B.H	BG
Оксилан 10 mg таблетки с удължено	DE/H/2182/002	20100387	G.L. PHARMA GMBH	BG

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
освобождаване				
Оксилан 20 mg таблетки с удължено освобождаване	DE/H/2182/003	20100388	G.L. PHARMA GMBH	BG
Оксилан 40 mg таблетки с удължено освобождаване	DE/H/2182/004	20100389	G.L. PHARMA GMBH	BG
Оксилан 5 mg таблетки с удължено освобождаване	DE/H/2182/001	20100386	G.L. PHARMA GMBH	BG
Оксилан 80 mg таблетки с удължено освобождаване	DE/H/2182/005	20100390	G.L. PHARMA GMBH	BG