2 December 2021
EMA/719030/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: oxaliplatin

Procedure no.: PSUSA/00002229/202104

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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
Axiplatin 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88849.00.00	AXIONOVO GMBH	DE
Bendaplatin 5 mg/ml Pulver zur Herstellung einer Infusionslösung	DE/H/2980/001	68364.00.00	BENDALIS GMBH	DE
Ebeoxal 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0346/001	1-30058	EBEWE PHARMA	AT
Elatofen 5 mg / ml, σκόνη για διάλυμα προς έγχυση	EL/H/0162/002	79963/12-11-2012	DEMO ABEE	GR
Elatofen 5 mg / ml, σκόνη για διάλυμα προς έγχυση	EL/H/0162/002	31490/11/9-1-12	DEMO ABEE	GR
ELOXATIN 5 mg/ml concentraat voor oplossing voor infusie	FR/H/0144/002	BE281601	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml concentraat voor oplossing voor infusie	FR/H/0144/002	BE281592	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml concentraat voor oplossing voor infusie	FR/H/0144/002	BE288784	SANOFI BELGIUM	BE
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	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
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	ES
Eloxatin 5 mg/ml concentrado para solución para perfusión.	FR/H/0144/002	67390	SANOFI-AVENTIS, S.A.	ES
Eloxatin 5 mg/ml concentrate for solution for infusion	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Eloxatin 5 mg/ml concentrate for solution for infusion	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Eloxatin 5 mg/ml concentrate for solution for infusion	FR/H/0284/002	PA 540/148/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411037	SANOFI S.R.L.	IT
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411052	SANOFI S.R.L.	IT
ELOXATIN 5 mg/ml concentrato per soluzione per infusione	FR/H/0144/002	034411049	SANOFI S.R.L.	IT
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND 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
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ELOXATIN 5 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLOESUNG	FR/H/0144/002	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
ELOXATIN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0144/002	BE281592	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0144/002	BE281601	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	FR/H/0144/002	BE288784	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE281592	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE281601	SANOFI BELGIUM	BE
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	BE288784	SANOFI BELGIUM	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
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0422335	SANOFI BELGIUM	LU
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0436599	SANOFI BELGIUM	LU
ELOXATIN 5 mg/ml solution à diluer pour perfusion	FR/H/0144/002	0422321	SANOFI BELGIUM	LU
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-06	SANOFI-AVENTIS AEBE	GR
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-06	SANOFI-AVENTIS AEBE	GR
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-06	SANOFI-AVENTIS AEBE	GR
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5865886	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5710082	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Eloxatin, 5 mg/ml, concentrado para solução para perfusão	FR/H/0284/002	5710181	SANOFI - PRODUTOS FARMACEUTICOS LDA	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
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	XI
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	XI
ELOXATINE 5 MG/ML CONCENTRATE FOR SOLUTION FOR INFUSION	FR/H/0144/002	PL 04425/0296	AVENTIS PHARMA LTD	XI
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	34009 559 649 2 6	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	34009 559 648 6 5	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, POUDRE POUR SOLUTION POUR PERFUSION	FR/H/0144/001	34009 563 191 7 6	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 565 984 4 1	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 565 983 8 0	SANOFI-AVENTIS FRANCE	FR
ELOXATINE 5 MG/ML, SOLUTION A DILUER POUR PERFUSION	FR/H/0144/002	34009 569 560 4 3	SANOFI-AVENTIS FRANCE	FR

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GENEPLATIN® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	8750/9-2-2021	GENEPHARM S.A.	GR
medoxa 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88848.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Medoxa 5 mg/ml Pulver zur Herstellung einer Infusionslösung	FI/H/0584/001	66241.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/001	FRESENIUS KABI AUSTRIA GMBH	SI
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/002	FRESENIUS KABI AUSTRIA GMBH	SI
Oksaliplatin Kabi 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0893/001	H/11/01147/003	FRESENIUS KABI AUSTRIA GMBH	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/001	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/002	SANDOZ PHARMACEUTICALS D.D.	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
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/005	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/003	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/004	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/006	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/007	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin SUN 5 mg/ml konsentrat til infusjonsvæske, oppløsning	DE/H/5677/001	12-8909	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NO
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/003	TEVA PHARMA B.V.	SI
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/001	TEVA PHARMA B.V.	SI
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/002	TEVA PHARMA B.V.	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
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/004	TEVA PHARMA B.V.	SI
Oxali-Bendalis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200076.00.00	BENDALIS GMBH	DE
Oxali-Bendalis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200076.00.00	BENDALIS GMBH	DE
Oxaliplatin "Accord", koncentrat til infusionsvæske, opløsning	IE/H/0756/001	44183	ACCORD HEALTHCARE B.V.	DK
Oxaliplatin "Accord", koncentrat til infusionsvæske, opløsning	IE/H/0756/001	44183	ACCORD HEALTHCARE B.V.	DK
Oxaliplatin "Accord", koncentrat til infusionsvæske, opløsning	IE/H/0756/001	44183	ACCORD HEALTHCARE B.V.	DK
Oxaliplatin "Actavis", koncentrat til infusionsvæske, opløsning	DK/H/3039/001	46888	ACTAVIS GROUP PTC EHF.	DK
Oxaliplatin "Fresenius Kabi", koncentrat til infusionsvæske, opløsning	NL/H/4321/001	41945	FRESENIUS KABI AB	DK
Oxaliplatin "Sandoz", koncentrat til infusionsvæske, opløsning	AT/H/0341/001	45579	SANDOZ A/S	DK

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Oxaliplatin "Teva", koncentrat til infusionsvæske, opløsning	NL/H/0820/001	39379	TEVA B.V	DK
Oxaliplatin 5 mg/ml concentrate for solution for infusion	NL/H/4321/001	PA 2059/049/001	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Oxaliplatin 5 mg/ml concentrate for solution for infusion	DE/H/6302/001	PL 44570/0002	QILU PHARMA SPAIN S.L.U	XI
Oxaliplatin 5 mg/ml Concentrate for Solution for Infusion	AT/H/0341/001	PL 04416/1604	SANDOZ LTD	XI
Oxaliplatin 5 mg/ml concentrate for solution for infusion	NL/H/4321/001	PL 08828/0300	FRESENIUS KABI LIMITED	XI
Oxaliplatin 5 mg/ml Concentrate for Solution for Infusion	AT/H/0893/001	PL 08828/0304	FRESENIUS KABI LIMITED	XI
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	IE/H/0756/001	PA 2315/114/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	IE/H/0756/001	PA 2315/114/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Oxaliplatin 5mg/ml Concentrate for Solution for Infusion	IE/H/0756/001	PA 2315/114/001	ACCORD HEALTHCARE IRELAND LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	MA1269/02102	ACCORD HEALTHCARE IRELAND LIMITED	MT
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	MA1269/02103	ACCORD HEALTHCARE IRELAND LIMITED	MT
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	MA1269/02101	ACCORD HEALTHCARE IRELAND LIMITED	MT
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	PL 20075/0112	ACCORD HEALTHCARE LIMITED	XI
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	PL 20075/0112	ACCORD HEALTHCARE LIMITED	XI
Oxaliplatin 5mg/ml concentrate for Solution for Infusion	IE/H/0756/001	PL 20075/0112	ACCORD HEALTHCARE LIMITED	XI
Oxaliplatin 5mg/ml concentrate for solution for infusion	not available	PL 41013/0025	SEACROSS PHARMA (EUROPE) LTD	XI
Oxaliplatin 5mg/ml powder for solution for infusion	NL/H/2337/001	PL 42069/0001	CADIASUN PHARMA GMBH	XI
Oxaliplatin Accord 5 mg/ml concentrat pentru solutie perfuzabila	IE/H/0756/001	10763/2018/01	ACCORD HEALTHCARE POLSKA SP. Z O.O.	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
Oxaliplatin Accord 5 mg/ml concentrat pentru soluţie perfuzabilă	IE/H/0756/001	10763/2018/02	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Oxaliplatin Accord 5 mg/ml concentrat pentru soluţie perfuzabilă	IE/H/0756/001	10763/2018/03	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Oxaliplatin Accord 5 mg/ml koncentrát pro infuzní roztok	IE/H/0756/001	44/316/10-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Oxaliplatin Accord 5 mg/ml koncentrát pro prípravu infuzního roztoku	IE/H/0756/001	44/316/10-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Oxaliplatin Accord 5 mg/ml koncentrát pro prípravu infuzního roztoku	IE/H/0756/001	44/316/10-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, lösning	IE/H/0756/001/DC	41633	ACCORD HEALTHCARE B.V.	SE
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, lösning	IE/H/0756/001	41633	ACCORD HEALTHCARE B.V.	SE
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, lösning	IE/H/0756/001	41633	ACCORD HEALTHCARE B.V.	SE
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	IE/H/0756/001	LT/1/10/2144/001	ACCORD HEALTHCARE B.V.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	IE/H/0756/001	LT/1/10/2144/002	ACCORD HEALTHCARE B.V.	LT
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	IE/H/0756/001	LT/1/10/2144/003	ACCORD HEALTHCARE B.V.	LT
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	IE/H/0756/001	10-0151	ACCORD HEALTHCARE B.V.	LV
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	IE/H/0756/001	10-0151	ACCORD HEALTHCARE B.V.	LV
Oxaliplatin Accord 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	IE/H/0756/001	10-0151	ACCORD HEALTHCARE B.V.	LV
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	IE/H/0756/001	OGYI-T-21479/04	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HU
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	IE/H/0756/001	OGYI-T-21479/01	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HU
Oxaliplatin Accord 5 mg/ml koncentrátum oldatos infúzióhoz	IE/H/0756/001	OGYI-T-21479/02	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HU

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Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	1-29637	ACCORD HEALTHCARE B.V.	AT
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	1-29637	ACCORD HEALTHCARE B.V.	AT
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	1-29637	ACCORD HEALTHCARE B.V.	AT
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	76427.00.00	ACCORD HEALTHCARE B.V.	DE
Oxaliplatin Accord 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	76427.00.00	ACCORD HEALTHCARE B.V.	DE
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	IE/H/0756/001	674810	ACCORD HEALTHCARE B.V.	EE
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	IE/H/0756/001/DC	674810	ACCORD HEALTHCARE B.V.	EE
Oxaliplatin Accord 5 mg/ml, infusioonilahuse kontsentraat	IE/H/0756/001	674810	ACCORD HEALTHCARE B.V.	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
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI
Oxaliplatin Accord 5 mg/ml, Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	76427.00.00	ACCORD HEALTHCARE B.V.	DE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001	BE373992	ACCORD HEALTHCARE B.V.	BE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001	BE374001	ACCORD HEALTHCARE B.V.	BE
Oxaliplatin Accord Healthcare 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001/DC	BE418555	ACCORD HEALTHCARE 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
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	IE/H/0756/001	BE373992	ACCORD HEALTHCARE B.V.	BE
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	IE/H/0756/001	BE374001	ACCORD HEALTHCARE B.V.	BE
Oxaliplatin Accord Healthcare 5 mg/ml solution à diluer pour perfusion	IE/H/0756/001	BE418555	ACCORD HEALTHCARE B.V.	BE
Oxaliplatin Actavis 5 mg/ml innrennslisþykkni, lausn	DK/H/3039/001	IS/1/11/065/01	ACTAVIS GROUP PTC EHF.	IS
OXALIPLATIN ACTAVIS 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	LV/H/0156/001	11-0268	ACTAVIS GROUP PTC EHF.	LV
Oxaliplatin Actavis 5 mg/ml konsentrat til infusjonsvæske, oppløsning	DK/H/3039/001	10-7598	ACTAVIS GROUP PTC EHF.	NO
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88845.00.00	AQVIDA 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
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200078.00.00	AQVIDA GMBH	DE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200077.00.00	AQVIDA GMBH	DE
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200079.00.00	AQVIDA GMBH	DE
Oxaliplatin Aristo 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88714.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Oxaliplatin Aurobindo 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2068/001	82014.00.00	PUREN PHARMA GMBH & CO. KG	DE
Oxaliplatin Bendalis 5 mg/ml, Konzentrat zur Herstellung einer Infusionslösung	not available	81696.00.00	BENDALIS GMBH	DE
Oxaliplatin beta 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	86708.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/01	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/02	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/03	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/04	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/05	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/07	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/08	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/06	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/09	EBEWE PHARMA	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
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/10	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/11	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/13	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/14	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0341/001	12961/2020/12	EBEWE PHARMA	RO
Oxaliplatin Ebewe 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0341/001	1-30057	EBEWE PHARMA	AT
Oxaliplatin Ebewe 5 mg/ml Pulver zur Herstellung einer Infusionslösung	AT/H/0185/001	1-26435	EBEWE PHARMA	AT
Oxaliplatin Ebewe 5mg/ml concentrate for solution infusion	AT/H/0341/001	20120240	EBEWE PHARMA	BG
Oxaliplatin Fresenius Kabi 5 mg/ml konsentrat til infusjonsvæske,	NL/H/4321/001	07-5183	FRESENIUS KABI NORGE AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
oppløsning				
Oxaliplatin Fresenius Kabi 5 mg/ml, koncentrat till infusionsvätska, lösning	AT/H/0893/001	44409	FRESENIUS KABI AB	SE
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/01	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/02	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/05	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/03	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/04	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0893/001	10109/2017/06	FRESENIUS KABI ROMANIA SRL	RO
Oxaliplatin Kabi 5 mg/ml infúzny koncentrát	NL/H/4321/001	44/0590/09-S	FRESENIUS KABI S.R.O.	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
Oxaliplatin Kabi 5 mg/ml koncentrát pro infuzní roztok	NL/H/4321/001	44/624/09-C	FRESENIUS KABI S.R.O.	CZ
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/004	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/005	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Oxaliplatin Kabi 5 mg/ml koncentratas infuziniam tirpalui	AT/H/0893/001	LT/1/10/1980/003	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Oxaliplatin Kabi 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	AT/H/0893/001	11-0054	FRESENIUS KABI POLSKA SP. Z O.O.	LV
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/01	FRESENIUS KABI HUNGARY KFT.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/02	FRESENIUS KABI HUNGARY KFT.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/05	FRESENIUS KABI HUNGARY KFT.	HU
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	1-30553	FRESENIUS KABI AUSTRIA 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
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/4321/001	70865.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	2011110055	FRESENIUS KABI DEUTSCHLAND GMBH	LU
Oxaliplatin Kabi 5 mg/ml, infusioonilahuse kontsentraat	AT/H/0893/001	728111	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Oxaliplatin Kabi, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/4321/001	17086	FRESENIUS KABI POLSKA SP. Z O.O.	PL
Oxaliplatin medac 5 mg/ml concentrate for solution for infusion	DE/H/3915/001	PL 11587/0086	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	XI
Oxaliplatin medac 5 mg/ml powder for solution for infusion	FI/H/0584/001	PA0623/008/001	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IE
Oxaliplatin onkovis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88846.00.00	ONKOVIS 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
Oxaliplatin PhaRes 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90005.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin Pliva 5 mg/ml koncentrat za otopinu za infuziju	not available	HR-H-505929304	PLIVA HRVATSKA D.O.O.	HR
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/2769/001	81219.00.00	HIKMA PHARMA GMBH	DE
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/2769/001	81219.00.00	HIKMA PHARMA GMBH	DE
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/2769/001	81219.00.00	HIKMA PHARMA GMBH	DE
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90006.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90006.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	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
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90006.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Oxaliplatin Sandoz 5 mg/ml - Konzentrat zur Herstellung einer Infusionslösung	AT/H/0437/001	1-31680	SANDOZ GMBH	AT
Oxaliplatin Sandoz 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0341/001	BE386662	SANDOZ N.V.	BE
Oxaliplatin Sandoz 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0341/001	BE386671	SANDOZ N.V.	BE
Oxaliplatin Sandoz 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0341/001	BE386687	SANDOZ N.V.	BE
Oxaliplatin Sandoz 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0341/001	BE404476	SANDOZ N.V.	BE
Oxaliplatin Sandoz 5 mg/ml koncentrat till infusionsvätska, lösning.	AT/H/0341/001	43028	SANDOZ A/S	SE
Oxaliplatin Sandoz 5 mg/ml konsentrat til infusjonsvæske, oppløsning	AT/H/0341/001	09-6984	SANDOZ A/S	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
Oxaliplatin STADA® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88850.00.00	STADAPHARM GMBH	DE
Oxaliplatin SUN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5677/001	82817.00.00	SUN PHARMACEUTICALS GERMANY GMBH	DE
Oxaliplatin Teva 5 mg/ml, koncentrat till infusionsvätska, lösning	NL/H/0820/001	23844	TEVA SWEDEN AB	SE
Oxaliplatin Tillomed 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2202103.00.00	TILLOMED PHARMA GMBH	DE
Oxaliplatin Vianex 5 mg/ml pulver till infusionsvätska, lösning	SE/H/0960/001	23806	VIANEX S.A.	SE
Oxaliplatin Vipharm® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	81697.00.00	VIPHARM GMBH	DE
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	ES/H/0609/001	24611/12/26-07-2018	PFIZER HELLAS, A.E.	GR
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος	ES/H/0609/001	24611/12/26-07-2018	PFIZER HELLAS, A.E.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
προς έγχυση.				
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	ES/H/0609/001	24611/12/26-07-2018	PFIZER HELLAS, A.E.	GR
OXALIPLATIN/MEDICUS 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	65258/23-06-2020	MEDICUS A.E	GR
Oxaliplatin/Teva 5 mg/ml, πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	NL/H/0820/001	55775/21-8-2015	TEVA PHARMA B.V.	GR
Oxaliplatin® HEXAL 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0346/001	80373.00.00	HEXAL AG	DE
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	IE/H/0756/001	5273735	ACCORD HEALTHCARE S.L.U.	PT
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	IE/H/0756/001	5273743	ACCORD HEALTHCARE S.L.U.	PT
Oxaliplatina Accord 5 mg/ml Concentrado para solução para perfusão	IE/H/0756/001	5422431	ACCORD HEALTHCARE S.L.U.	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
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	PT/H/2068/001	5388160	GENERIS FARMACÊUTICA, S.A.	PT
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	PT/H/2068/001	5388178	GENERIS FARMACÊUTICA, S.A.	PT
Oxaliplatina Aurovitas 5 mg/ml concentrado para solução para perfusão	PT/H/2068/001	5388202	GENERIS FARMACÊUTICA, S.A.	PT
Oxaliplatina Aurovitas 5 mg/ml pó para solução para perfusão	PT/H/1702/001	5087655	GENERIS FARMACÊUTICA, S.A.	PT
Oxaliplatina Aurovitas 5 mg/ml pó para solução para perfusão	PT/H/1702/001	5087663	GENERIS FARMACÊUTICA, S.A.	PT
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5201132	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5396171	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Oxaliplatina Kabi 5 mg/ml concentrado para solução para perfusão	NL/H/4321/001	5201124	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Oxaliplatina Mylan 5 mg/ml prášek pro infuzní roztok	FR/H/0324/001	44/646/07-C	MYLAN S.A.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
Oxaliplatina Mylan 5 mg/ml prášok na infúzny roztok	FR/H/0324/001	44/0345/07-S	MYLAN S.A.S	SK
Oxaliplatine Accord 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001	RVG 103779	ACCORD HEALTHCARE B.V.	NL
Oxaliplatine Accord 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001	RVG 103779	ACCORD HEALTHCARE B.V.	NL
Oxaliplatine Accord 5 mg/ml concentraat voor oplossing voor infusie	IE/H/0756/001	RVG 103779	ACCORD HEALTHCARE B.V.	NL
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	IE/H/0756/001	34009 576 842 1 1	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	IE/H/0756/001	34009 579 546 4 2	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ACCORD 5 mg/ml, solution à diluer pour perfusion	IE/H/0756/001	34009 576 841 5 0	ACCORD HEALTHCARE FRANCE SAS	FR
OXALIPLATINE ARROW 5 mg/ml, solution à diluer pour perfusion	not available	NL 39965	ARROW GENERIQUES	FR
Oxaliplatine Cadiasun 5 mg/ml poeder voor oplossing voor infusie	NL/H/2337/001	RVG 110045	CADIASUN PHARMA GMBH	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
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE384632	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE 428391	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	AT/H/0893/001	BE384623	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	NL/H/4321/001	RVG 100834	FRESENIUS KABI NEDERLAND B.V.	NL
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE384632	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE428391	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	BE384623	FRESENIUS KABI NV/SA	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
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE384632	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE428391	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/ml solution à diluer pour perfusion	AT/H/0893/001	BE384623	FRESENIUS KABI NV/SA	BE
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 572 480 8 6	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 572 482 0 8	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 574 402 4 4	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 572 696 0 9	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 572 697 7 7	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	ES/H/0609/001	34009 574 403 0 5	PFIZER HOLDING FRANCE	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
OXALIPLATINE MEDAC 5 mg/ml, solution à diluer pour perfusion	DE/H/3915/001	34009 586 857 1 2	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
OXALIPLATINE MEDAC 5 mg/ml, solution à diluer pour perfusion	DE/H/3915/001	34009 586 858 8 0	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
OXALIPLATINE MEDAC 5 mg/ml, solution à diluer pour perfusion	DE/H/3915/001	34009 586 859 4 1	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Oxaliplatine Mylan 5 mg/ml, concentraat voor oplossing voor infusie	NL/H/1977/001	RVG 107390	MYLAN B.V.	NL
OXALIPLATINE MYLAN 5 mg/ml, poudre pour solution pour perfusion	FR/H/0324/001	NL 32863	MYLAN S.A.S	FR
OXALIPLATINE MYLAN 5 mg/ml, solution a diluer pour perfusion	NL/H/1977/001	NL 39768	MYLAN S.A.S	FR
Oxaliplatine SUN 5 mg/ml, concentraat voor oplossing voor infusie	DE/H/5677/001	RVG 108042	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 161 2 1	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	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
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 158 1 0	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 156 9 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 159 8 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 160 6 0	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
OXALIPLATINE SUN 5 mg/ml, solution à diluer pour perfusion	DE/H/5677/001	34009 582 157 5 9	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/001	BE303834	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/001	BE303825	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/001	BE318832	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml concentraat voor oplossing voor infusie	NL/H/0820/001	BE303816	TEVA PHARMA BELGIUM N.V./S.A	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
Oxaliplatine Teva 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	BE303834	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	BE318832	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	BE303816	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	BE303825	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion	NL/H/0820/001	BE303816	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion	NL/H/0820/001	BE303834	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion	NL/H/0820/001	BE318832	TEVA PHARMA BELGIUM N.V./S.A	BE
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion	NL/H/0820/001	BE303825	TEVA PHARMA BELGIUM N.V./S.A	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
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion.	NL/H/0820/001	2007110014	TEVA PHARMA BELGIUM N.V./S.A	LU
OXALIPLATINE TEVA 5 mg/ml, solution à diluer pour perfusion	NL/H/0820/001	NL32938	TEVA SANTÉ	FR
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	34009 567 117 6 5	SANOFI-AVENTIS FRANCE	FR
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	34009 569 815 2 6	SANOFI-AVENTIS FRANCE	FR
OXALIPLATINE WINTHROP 5 MG/ML, SOLUTION À DILUER POUR PERFUSION	FR/H/0284/002	34009 567 115 3 6	SANOFI-AVENTIS FRANCE	FR
OXALIPLATIN-EBEWE, 5 MG/ML, KONCENTRAT DO SPORZĄDZANIA ROZTWORU DO INFUZJI	AT/H/0341/001	18330	EBEWE PHARMA	PL
Oxaliplatin-GRY® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	66264.00.00	TEVA GMBH	DE
Oxaliplatino 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107013	TEVA ITALIA S.R.L.	IT
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión	IE/H/0756/001	72386	ACCORD HEALTHCARE S.L.U.	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
EFG				
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión EFG	IE/H/0756/001	72386	ACCORD HEALTHCARE S.L.U.	ES
Oxaliplatino Accord 5 mg/ml concentrado para solución para perfusión EFG	IE/H/0756/001	72386	ACCORD HEALTHCARE S.L.U.	ES
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	IE/H/0756/001	041274010	ACCORD HEALTHCARE S.L.U.	IT
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	IE/H/0756/001	041274022	ACCORD HEALTHCARE S.L.U.	IT
Oxaliplatino Accord 5 mg/ml concentrato per soluzione per infusione	IE/H/0756/001	041274034	ACCORD HEALTHCARE S.L.U.	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione	PT/H/2068/001/DC	039999014	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione	PT/H/2068/001/DC	039999038	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Oxaliplatino Aurobindo 5 mg/ml concentrato per soluzione per infusione.	PT/H/2068/001/DC	039999026	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
Oxaliplatino Aurovitas 5 mg/ml concentrado para solución para perfusión EFG	PT/H/2068/001	74.655	AUROVITAS SPAIN,S.A.U.	ES
Oxaliplatino Cadiasun 5 mg/ml polvo para solución para perfusión EFG	NL/H/2337/001	77233	CADIASUN PHARMA GMBH	ES
Oxaliplatino GP-Pharm 5 mg/ml, polvo para solución para perfusión EFG	not available	72666	GP-PHARM S.A.	ES
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	ES/H/0609/001	69518	PFIZER, S.L.	ES
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	ES/H/0609/001	69518	PFIZER, S.L.	ES
Oxaliplatino Hospira 5 mg/ml concentrado para solución para perfusión EFG	ES/H/0609/001	69518	PFIZER, S.L.	ES
Oxaliplatino Kabi 5 mg/ml concentrado para solución para perfusión EFG	NL/H/4321/001	71346	FRESENIUS KABI ESPAÑA S.A.U.	ES
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170028	FRESENIUS KABI 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
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170030	FRESENIUS KABI ITALIA S.R.L.	IT
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170016	FRESENIUS KABI ITALIA S.R.L.	IT
Oxaliplatino Qilu 5 mg/ml concentrado para solución para perfusión EFG	DE/H/6302/001	81823	QILU PHARMA SPAIN S.L.U	ES
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654055	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654028	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654067	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654016	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654042	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz 5 mg/ml concentrato per soluzione per infusione	AT/H/0341/001	040654030	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
Oxaliplatino SUN 5 mg/ml concentrado para solución para perfusión EFG	DE/H/5677/001	75278	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761038	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761053	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761026	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761040	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761014	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761065	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107025	TEVA ITALIA S.R.L.	IT
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107076	TEVA 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
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusione.	NL/H/0820/001	038107037	TEVA ITALIA S.R.L.	IT
Oxaliplatino TEVA 5 mg/ml concentrado para solución para perfusión EFG	NL/H/0820/001	69288	TEVA PHARMA S.L.U.,	ES
Oxaliplatin-Tchaikapharma 100 mg powder for solution for infusion	not available	20110481	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Oxaliplatin-Tchaikapharma 50 mg powder for solution for infusion	not available	20110480	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/004	TEVA B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/005	TEVA B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/006	TEVA B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentratas infuziniam tirpalui	NL/H/0820/001	LT/1/07/0921/007	TEVA B.V	LT
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/02	TEVA GYÓGYSZERGYÁR ZRT	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
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/0820/001	OGYI-T-20505/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Oxaliplatin-Teva 5 mg/ml, koncentrát pro infuzní roztok	NL/H/0820/001	44/591/07-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji.	IE/H/0756/001	17070	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji.	IE/H/0756/001	17070	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporządzania roztworu do infuzji.	IE/H/0756/001	17070	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
OXALIPROL® 5mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	91329/11/ 23-01-2014	PHARMAZAC SA	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxalisan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	86711.00.00	MEDICOPHARM AG	DE
Oxalisin 5 mg/ml, concentraat voor oplossing voor infusie.	NL/H/0820/001	RVG 34033	PHARMACHEMIE BV	NL
OXALIZOR 5mg/mL πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	not available	104908/05-11-2020	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
OXAVIATIN 5mg/mL πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	29264/16/23-03-2017	VIANEX S.A.	GR
OXAVIATIN® 5 mg/ml concentrate for solution for infusion	not available	AA721/00601	VIANEX S.A.	MT
OXAVIATIN® 5 mg/ml κόνις για διάλυμα προς έγχυση	SE/H/0960/001	21057	VIANEX S.A.	CY
OXAVIATIN® 5 mg/ml κόνις για διάλυμα προς έγχυση	SE/H/0960/001	26163	VIANEX S.A.	GR
RECTOXAL® 5 mg/ml κόνις για διάλυμα προς έγχυση	EL/H/0123/001	57125/14-11-2012	ARITI SA	GR
Sinoxal 100 mg prašak za otopinu za infuziju	not available	HR-H-640588353-02	ACTAVIS GROUP PTC EHF.	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
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	DK/H/3039/001	H/11/01425/002	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	DK/H/3039/001	H/11/01425/003	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 5 mg/ml koncentrat za raztopino za infundiranje	DK/H/3039/001	H/11/01425/001	ACTAVIS GROUP PTC EHF.	SI
Sinoxal 50 mg prašak za otopinu za infuziju	not available	HR-H-640588353-01	ACTAVIS GROUP PTC EHF.	HR
VELMINOX®	not available	3197	VIOFAR LTD	GR
Оксалиплатин Акорд 5 mg/ml концентрат за инфузионен разтвор.	IE/H/0756/001	20120332	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Оксалиплатин Акорд 5 mg/ml концентрат за инфузионен разтвор.	IE/H/0756/001	20120332	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Оксалиплатин Акорд 5 mg/ml концентрат за инфузионен разтвор.	IE/H/0756/001	20120332	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Оксалиплатин Актавис 5 mg/ml концентрат за инфузионен разтвор	DK/H/3039/001	20110406	ACTAVIS GROUP PTC EHF.	BG