12 December 2024
EMA/360417/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): oxaliplatin

Procedure No. PSUSA/00002229/202404

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 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 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	BE418555	ACCORD HEALTHCARE B.V.	BE

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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 Konzentrat zur Herstellung einer Infusionslösung	IE/H/0756/001	76427.00.00	ACCORD HEALTHCARE B.V.	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 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
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI
Oxaliplatin Accord 5 mg/ml,	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI

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infuusiokonsentraatti, liuosta varten.				
Oxaliplatin Accord 5 mg/ml, infuusiokonsentraatti, liuosta varten.	IE/H/0756/001	26943	ACCORD HEALTHCARE B.V.	FI
Oxaliplatin Accord 5 mg/ml koncentratas infuziniam tirpalui	IE/H/0756/001	LT/1/10/2144/001	ACCORD HEALTHCARE B.V.	LT
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
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
Oxaliplatin Accord 5 mg/ml koncentrat till infusionsvätska, lösning	IE/H/0756/001/DC	41633	ACCORD HEALTHCARE B.V.	SE

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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
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
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
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
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

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Оксалиплатин Акорд 5 mg/ml концентрат за инфузионен разтвор.	IE/H/0756/001	20120332	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
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 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 infuzní roztok	IE/H/0756/001	44/316/10-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
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
Oxaliplatinum Accord, 5 mg/ml, koncentrat do sporzadzania 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
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

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Oxaliplatin Accord 5 mg/ml concentrat pentru soluţie perfuzabilă	IE/H/0756/001	10763/2018/01	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
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 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
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
Oxaliplatin "Actavis", koncentrat til infusionsvæske, opløsning	DK/H/3039/001	46888	ACTAVIS GROUP PTC EHF.	DK
Sinoxal 50 mg prašak za otopinu za infuziju	not available	HR-H-640588353-01	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 100 mg prašak za otopinu za infuziju	not available	HR-H-640588353-02	ACTAVIS GROUP PTC EHF.	HR
Oxaliplatin Actavis 5 mg/ml innrennslisþykkni, lausn	DK/H/3039/001	IS/1/11/065/01	ACTAVIS GROUP PTC EHF.	IS
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
Oxaliplatin AqVida 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88845.00.00	AQVIDA GMBH	DE
Oxaliplatin Bendalis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	81696.00.00	BENDALIS GMBH	DE
Oxali-Bendalis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	2200076.00.00	BENDALIS GMBH	DE
Оксалиплатин Булгермед 5 mg/ml концентрат за инфузионен разтвор	not available	20170376	BULGERMED VE	BG
Elatofen 5 mg / ml, σκόνη για διάλυμα προς έγχυση	EL/H/0162/002	31490/11/9-1-12	DEMO ABEE	GR
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

Product 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 Konzentrat zur Herstellung einer Infusionslösung	AT/H/0341/001	1-30057	EBEWE PHARMA	AT
Ebeoxal 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0346/001	1-30058	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-EBEWE, 5 MG/ML, KONCENTRAT DO SPORZĄDZANIA ROZTWORU DO INFUZJI	AT/H/0341/001	18330	EBEWE PHARMA	PL
Oxaliplatin 5 mg/ml Concentrate for Solution for infusion	not available	PL 56284/0009	EUGIA (UK) LTD	XI
Oxaliplatin Eugia 5 mg/ml concentraat voor oplossing voor infusie	SE/H/2172/001	BE661053	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml solution à diluer pour perfusion	SE/H/2172/001	BE661053	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	SE/H/2172/001	BE661053	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml concentraat voor oplossing voor infusie	SE/H/2172/001	BE661054	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml solution à diluer pour perfusion	SE/H/2172/001	BE661054	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5	SE/H/2172/001	BE661054	EUGIA PHARMA (MALTA) LTD	BE

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mg/ml Konzentrat zur Herstellung einer Infusionslösung				
Oxaliplatin Eugia 5 mg/ml concentraat voor oplossing voor infusie	SE/H/2172/001	BE661055	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml solution à diluer pour perfusion	SE/H/2172/001	BE661055	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin Eugia 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	SE/H/2172/001	BE661055	EUGIA PHARMA (MALTA) LTD	BE
Oxaliplatin PUREN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	SE/H/2172/001	7006423.00.00	EUGIA PHARMA (MALTA) LTD	DE
Oxaliplatino Aurovit 5 mg/ml concentrado para solución para perfusión EFG	SE/H/2172/001	88.574	EUGIA PHARMA (MALTA) LTD	ES
OXALIPLATINE ARROW LAB 5 mg/mL, solution à diluer pour perfusion	SE/H/2172/001	NL53685	EUGIA PHARMA (MALTA) LTD	FR
Oxaliplatino Aurobindo Italia 5 mg/ml concentrato per soluzione per infusione	SE/H/2172/001	050405012	EUGIA PHARMA (MALTA) LTD	IT
Oxaliplatino Aurobindo Italia 5 mg/ml concentrato per soluzione per infusione	SE/H/2172/001	050405024	EUGIA PHARMA (MALTA) LTD	IT
Oxaliplatino Aurobindo Italia 5 mg/ml concentrato per soluzione per infusione	SE/H/2172/001	050405036	EUGIA PHARMA (MALTA) LTD	IT
Oxaliplatin Eugia, 5 mg/mL, koncentrat do sporządzania roztworu	SE/H/2172/001	27730	EUGIA PHARMA (MALTA) LTD	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
do infuzji				
Oxaliplatina Generis 5 mg/ml concentrado para solução para perfusão	SE/H/2172/001	5858030	EUGIA PHARMA (MALTA) LTD	PT
Oxaliplatina Generis 5 mg/ml concentrado para solução para perfusão	SE/H/2172/001	5858048	EUGIA PHARMA (MALTA) LTD	PT
Oxaliplatina Generis 5 mg/ml concentrado para solução para perfusão	SE/H/2172/001	5858055	EUGIA PHARMA (MALTA) LTD	PT
Oxaliplatin Eugia 5 mg/ml, koncentrat till infusionsvätska, lösning	SE/H/2172/001	62413	EUGIA PHARMA (MALTA) LTD	SE
Oxaliplatin Fresenius Kabi	NL/H/4321/001	41945	FRESENIUS KABI AB	DK
Oxaliplatin Fresenius Kabi 5 mg/ml, koncentrat till infusionsvätska, lösning	AT/H/0893/001	44409	FRESENIUS KABI AB	SE
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0893/001	1-30553	FRESENIUS KABI AUSTRIA GMBH	AT
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
Oxaliplatin Kabi 5 mg/ml koncentrat za otopinu za infuziju	not available	HR-H-771037007	FRESENIUS KABI D.O.O.	HR
Oxaliplatin Kabi 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/4321/001	70865.00.00	FRESENIUS KABI 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
Oxaliplatin 5 mg/ml concentrate for solution for infusion	NL/H/4321/001	PA 2059/049/001	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Oxaliplatino Kabi 5 mg/ml concentrado para solución para perfusión EFG	NL/H/4321/001	71346	FRESENIUS KABI ESPAÑA S.A.U.	ES
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/06	FRESENIUS KABI HUNGARY KFT.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/07	FRESENIUS KABI HUNGARY KFT.	HU
Oxaliplatin Kabi 5 mg/ml koncentrátum oldatos infúzióhoz	NL/H/4321/001	OGYI-T-21085/08	FRESENIUS KABI HUNGARY KFT.	HU
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
Oxaliplatino Kabi 5 mg/ml concentrato per soluzione per infusione	NL/H/4321/001	039170028	FRESENIUS KABI ITALIA S.R.L.	IT
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
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	AT/H/0893/001	PL 08828/0304	FRESENIUS KABI LIMITED	XI

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for Infusion				
Oxaliplatine Fresenius Kabi 5 mg/ml concentraat voor oplossing voor infusie	NL/H/4321/001	RVG 100834	FRESENIUS KABI NEDERLAND B.V.	NL
Oxaliplatin Fresenius Kabi 5 mg/ml konsentrat til infusjonsvæske, oppløsning	NL/H/4321/001	07-5183	FRESENIUS KABI NORGE AS OSLO	NO
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	BE428391	FRESENIUS KABI NV/SA	BE
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 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 concentraat voor oplossing voor infusie	AT/H/0893/001	BE384623	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	BE384623	FRESENIUS KABI NV/SA	BE
Oxaliplatine Fresenius Kabi 5 mg/1 ml Solution à diluer pour perfusion	AT/H/0893/001	2011110055	FRESENIUS KABI NV/SA	LU
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
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 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, koncentrat do sporządzania roztworu do infuzji	NL/H/4321/001	17086	FRESENIUS KABI POLSKA SP. Z O.O.	PL
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	AT/H/0893/001	10109/2017/02	FRESENIUS KABI ROMANIA SRL	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
perfuzabilă				
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 koncentrát pro infuzní roztok	NL/H/4321/001	44/624/09-C	FRESENIUS KABI S.R.O.	CZ
Oxaliplatin Kabi 5 mg/ml, koncentrát na infúzny roztok	NL/H/4321/001	44/0590/09-S	FRESENIUS KABI S.R.O.	SK
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 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
Oxaliplatin HEXAL 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0346/001	80373.00.00	HEXAL AG	DE
Oxaliplatin Hikma 5	PT/H/2612/001	141043	HIKMA FARMACÊUTICA	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
mg/ml Konzentrat zur Herstellung einer Infusionslösung			(PORTUGAL), S.A.	
Oxaliplatin Hikma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2612/001	141043	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Oxaliplatin Hikma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2612/001	141043	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
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
Oxaliplatin Ribosepharm 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90006.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Oxaliplatin Hikma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2612/001	7000285.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Oxaliplatin Hikma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2612/001	7000285.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Oxaliplatin Hikma 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	PT/H/2612/001	7000285.00.00	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 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
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
Oxaliplatino Hikma 5 mg/ml concentrado para solución para perfusión EFG	PT/H/2612/001	86380	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	ES
Oxaliplatino Hikma 5 mg/ml concentrado para solución para perfusión EFG	PT/H/2612/001	86380	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	ES
Oxaliplatino Hikma 5 mg/ml concentrado para solución para perfusión EFG	PT/H/2612/001	86380	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	ES
OXALIPLATINE HIKMA 5 mg/mL, solution à diluer pour perfusion	PT/H/2612/001	34009 550 870 5 2	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
OXALIPLATINE HIKMA 5 mg/mL, solution à diluer pour perfusion	PT/H/2612/001	34009 550 870 6 9	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
OXALIPLATINE HIKMA 5	PT/H/2612/001	34009 550 870 7 6	HIKMA FARMACÊUTICA	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg/mL, solution à diluer pour perfusion			(PORTUGAL), S.A.	
Oxaliplatino Hikma 5 mg/ml, concentrato per soluzione per infusione	PT/H/2612/001	049146018	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IT
Oxaliplatino Hikma 5 mg/ml, concentrato per soluzione per infusione	PT/H/2612/001	049146020	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IT
Oxaliplatino Hikma 5 mg/ml, concentrato per soluzione per infusione	PT/H/2612/001	049146032	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IT
Oxaliplatin Hikma 5 mg/ml concentraat voor oplossing voor infusie	PT/H/2612/001	RVG 126655	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Oxaliplatin Hikma 5 mg/ml concentraat voor oplossing voor infusie	PT/H/2612/001	RVG 126655	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Oxaliplatin Hikma 5 mg/ml concentraat voor oplossing voor infusie	PT/H/2612/001	RVG 126655	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Oxaliplatina Hikma, 5 mg/ml, concentrado para soluテアテ」o para perfusテ」o	PT/H/2612/001	5833041	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Oxaliplatina Hikma, 5 mg/ml, concentrado para soluテアテ」o para perfusテ」o	PT/H/2612/001	5833058	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Oxaliplatina Hikma, 5 mg/ml, concentrado para soluテアテ」o para perfusテ」o	PT/H/2612/001	5833066	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Oxaliplatin 5 mg/ml concentrate for solution for infusion	PT/H/2612/001	PL 15413/0107	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	XI
Oxaliplatin 5 mg/ml concentrate for solution for infusion	PT/H/2612/001	PL 15413/0107	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Oxaliplatin 5 mg/ml concentrate for solution for infusion	PT/H/2612/001	PL 15413/0107	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	XI
Oxaliplatin Hospira 5 mg/ml concentrate for solution for infusion	EL/H/0317/001	PL 04515/0215	HOSPIRA UK LIMITED,WALTON OAKS	XI
Oxaliplatin Hospira 5 mg/ml concentrate for solution for infusion	EL/H/0317/001	PL 04515/0215	HOSPIRA UK LIMITED,WALTON OAKS	XI
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662340	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662340	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks	LV/H/0244/001	BE662340	KALCEKS	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
5 mg/ml Konzentrat zur Herstellung einer Infusionslösung				
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml concentraat voor oplossing voor infusie	LV/H/0244/001	BE662340	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml solution à diluer pour perfusion	LV/H/0244/001	BE662340	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	BE662338	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	BE662339	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	BE662340	KALCEKS	BE
Oxaliplatin Kalceks 5 mg/ml koncentrát pro infuzní roztok	LV/H/0244/001	44/228/22-C	KALCEKS	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
Oxaliplatin Kalceks 5 mg/ml koncentrát pro infuzní roztok	LV/H/0244/001	44/228/22-C	KALCEKS	CZ
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	7010017.00.00	KALCEKS	DE
Oxaliplatin Kalceks 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	LV/H/0244/001	7010017.00.00	KALCEKS	DE
Oxaliplatino Kalceks 5 mg/ml concentrado para solución para perfusión EFG	LV/H/0244/001	89.397	KALCEKS	ES
Oxaliplatino Kalceks 5 mg/ml concentrado para solución para perfusión EFG	LV/H/0244/001	89.397	KALCEKS	ES
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 1 5	KALCEKS	FR
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 2 2	KALCEKS	FR
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 3 9	KALCEKS	FR
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 1 5	KALCEKS	FR
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 2 2	KALCEKS	FR
OXALIPLATINE KALCEKS 5 mg/mL, solution à diluer pour perfusion	LV/H/0244/001	34009 550 992 3 9	KALCEKS	FR
Oksaliplatin Kalceks 5 mg/ml koncentrat za	LV/H/0244/001	HR-H-434230309	KALCEKS	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
otopinu za infuziju				
Oksaliplatin Kalceks 5 mg/ml koncentrat za otopinu za infuziju	LV/H/0244/001	HR-H-434230309	KALCEKS	HR
Oxaliplatin Kalceks 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	LV/H/0244/001	23-0221	KALCEKS	LV
Oxaliplatin Kalceks 5 mg/ml koncentrāts infūziju šķīduma pagatavošanai	LV/H/0244/001	23-0221	KALCEKS	LV
Oxaliplatin Kalceks 5 mg/ml konsentrat til infusjonsvæske, oppløsning	LV/H/0244/001	22-14844	KALCEKS	NO
Oxaliplatin Kalceks 5 mg/ml konsentrat til infusjonsvæske, oppløsning	LV/H/0244/001	22-14844	KALCEKS	NO
Oxaliplatin Kalceks, 5 mg/mL, koncentrat do sporządzania roztworu do infuzji	LV/H/0244/001	28394	KALCEKS	PL
Oxaliplatin Kalceks, 5 mg/mL, koncentrat do sporządzania roztworu do infuzji	LV/H/0244/001	28394	KALCEKS	PL
Oxaliplatin Kelix bio 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7010816.00.00	KELIX BIO (MALTA) LIMITED	DE
Medoxa 5 mg/ml Pulver zur Herstellung einer Infusionslösung	DE/H/6920/001	66241.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
medoxa 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88848.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	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
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
Oxaliplatin medac 5 mg/ml powder for solution for infusion	DE/H/6920/001	PA0623/008/001	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IE
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
Oxalisan 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	86711.00.00	MEDICOPHARM AG	DE
OXALIPLATIN/MEDICUS 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	65258/23-06-2020	MEDICUS A.E	GR
Oxaliplatin onkovis 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	88846.00.00	ONKOVIS GMBH	DE
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	EL/H/0317/001	24611/12/26-07-2018	PFIZER HELLAS A.E.	GR
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	EL/H/0317/001	24611/12/26-07-2018	PFIZER HELLAS A.E.	GR
OXALIPLATIN/HOSPIRA 5 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	EL/H/0317/001	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
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 572 480 8 6	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 572 482 0 8	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 574 402 4 4	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 572 696 0 9	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 572 697 7 7	PFIZER HOLDING FRANCE	FR
OXALIPLATINE HOSPIRA 5 mg/ml, solution à diluer pour perfusion	EL/H/0317/001	34009 574 403 0 5	PFIZER HOLDING FRANCE	FR
Oxaliplatin PhaRes 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7002825.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin PhaRes 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	90005.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin PhaRes 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7002826.00.00	PHARMA RESOURCES GMBH	DE
Oxaliplatin PhaRes 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7002827.00.00	PHARMA RESOURCES GMBH	DE
Oxalisin 5 mg/ml, concentraat voor oplossing voor infusie.	NL/H/0820/001	RVG 34033	PHARMACHEMIE BV	NL
OXALIPROL® 5mg/ml πυκνό διάλυμα για	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
παρασκευή διαλύματος προς έγχυση				
Oxaliplatin Pliva 5 mg/ml koncentrat za otopinu za infuziju	not available	HR-H-505929304	PLIVA HRVATSKA D.O.O.	HR
Oxaliplatino Qilu 5 mg/ml concentrado para solución para perfusión EFG	DE/H/6302/001	81823	QILU PHARMA SPAIN S.L.	ES
Oxaliplatin 5 mg/ml concentrate for solution for infusion	DE/H/6302/001	PL 44570/0002	QILU PHARMA SPAIN S.L.	XI
Oxaliplatin "Sandoz", koncentrat til infusionsvæske, opløsning	AT/H/0341/001	45579	SANDOZ A/S	DK
Oxaliplatin Sandoz 5 mg/ml konsentrat til infusjonsvæske, oppløsning	AT/H/0341/001	09-6984	SANDOZ A/S	NO
Oxaliplatin Sandoz 5 mg/ml koncentrat till infusionsvätska, lösning.	AT/H/0341/001	43028	SANDOZ A/S	SE
Oxaliplatin Sandoz 5 mg/ml - Konzentrat zur Herstellung einer Infusionslösung	AT/H/0437/001	1-31680	SANDOZ GMBH	AT
Oxaliplatin 5 mg/ml Concentrate for Solution for Infusion	AT/H/0341/001	PL 04416/1604	SANDOZ LTD	XI
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
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/005	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/11/01146/003	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 Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/014	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/008	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/009	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/010	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/011	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/012	SANDOZ PHARMACEUTICALS D.D.	SI
Oksaliplatin Sandoz 5 mg/ml koncentrat za raztopino za infundiranje	AT/H/0341/001	H/11/01146/013	SANDOZ PHARMACEUTICALS D.D.	SI
Oxaliplatino Sandoz	AT/H/0341/001	040654055	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz	AT/H/0341/001	040654028	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz	AT/H/0341/001	040654067	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz	AT/H/0341/001	040654016	SANDOZ S.P.A.	IT
Oxaliplatino Sandoz	AT/H/0341/001	040654042	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 Sandoz	AT/H/0341/001	040654030	SANDOZ S.P.A.	IT
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 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 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 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	0422321	SANOFI BELGIUM	LU
ELOXATINE 5 mg/ml, solution à diluer pour perfusion	FR/H/0144/002	34009 565 984 4 1	SANOFI WINTHROP INDUSTRIE	FR
ELOXATINE 5 mg/ml, solution à diluer pour perfusion	FR/H/0144/002	34009 565 983 8 0	SANOFI WINTHROP INDUSTRIE	FR
ELOXATINE 5 mg/ml, poudre pour solution pour perfusion	FR/H/0144/001	34009 559 649 2 6	SANOFI WINTHROP INDUSTRIE	FR
ELOXATINE 5 mg/ml, poudre pour solution pour perfusion	FR/H/0144/001	34009 559 648 6 5	SANOFI WINTHROP INDUSTRIE	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
ELOXATINE 5 mg/ml, poudre pour solution pour perfusion	FR/H/0144/001	34009 563 191 7 6	SANOFI WINTHROP INDUSTRIE	FR
OXALIPLATINE WINTHROP 5 mg/ml, solution à diluer pour perfusion	FR/H/0284/002	34009 567 117 6 5	SANOFI WINTHROP INDUSTRIE	FR
OXALIPLATINE WINTHROP 5 mg/ml, solution à diluer pour perfusion	FR/H/0284/002	34009 567 115 3 6	SANOFI WINTHROP INDUSTRIE	FR
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-2006	SANOFI-AVENTIS AEBE	GR
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-2006	SANOFI-AVENTIS AEBE	GR
ELOXATIN 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	FR/H/0284/002	54178/31-08-2006	SANOFI-AVENTIS AEBE	GR
ELOXATIN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	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	63264.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Oxaliplatin 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7000049.00.00	SEACROSS PHARMA (EUROPE) LTD	DE
Oxaliplatin 5mg/ml concentrate for solution for infusion	not available	PL 41013/0025	SEACROSS PHARMACEUTICALS LIMITED	XI
Oxaliplatin STADA 5	not available	88850.00.00	STADAPHARM 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/ml Konzentrat zur Herstellung einer Infusionslösung				
Oxaliplatino SUN 5 mg/ml concentrado para solución para perfusión EFG	DE/H/5677/001	75278	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
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
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
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	DE/H/5677/001	041761014	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
soluzione per infusione				
Oxaliplatino SUN 5 mg/ml concentrato per soluzione per infusione	DE/H/5677/001	041761065	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Oxaliplatine SUN 5 mg/ml, concentraat voor oplossing voor infusie	DE/H/5677/001	RVG 108042	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Oksaliplatin SUN 5 mg/ml konsentrat til infusjonsvæske, oppløsning	DE/H/5677/001	12-8909	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NO
Oxaliplatin SUN 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5677/001	82817.00.00	SUN PHARMACEUTICALS GERMANY GMBH	DE
Оксалиплатин-Чайкафарма 50 mg прах за инфузионен разтвор	not available	20110480	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Оксалиплатин-Чайкафарма 100 mg прах за инфузионен разтвор	not available	20110481	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Oxaliplatin Teva 5 mg/ml innrennslisþykkni, lausn	NL/H/0820/001	IS/1/23/017/01	TEVA B.V	IS
Oxaliplatin-GRY® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/0820/001	66264.00.00	TEVA GMBH	DE
Oxaliplatino Teva 5 mg / ml, concentrato per soluzione per infusion.	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
Oxaliplatino 5 mg / ml, concentrato per soluzione per infusione	NL/H/0820/001	038107013	TEVA ITALIA S.R.L.	IT
Oxaliplatino Teva 5 mg /	NL/H/0820/001	038107037	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
ml, concentrato per soluzione per infusione.				
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
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
Oksaliplatin Teva 5 mg/ml koncentrat za raztopino za infundiranje	NL/H/0820/001	H/07/01150/004	TEVA PHARMA B.V.	SI
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 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 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

Product 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 concentraat voor oplossing voor infusie	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	BE303825	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
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
Oxaliplatine Teva 5 mg/ml solution à diluer pour perfusion.	NL/H/0820/001	2007110014	TEVA PHARMA BELGIUM N.V./S.A	LU
Oxaliplatino TEVA 5 mg/ml concentrado para solución para perfusión EFG	NL/H/0820/001	69288	TEVA PHARMA S.L.U.,	ES
Oxaliplatin-Teva 5 mg/ml, koncentrát pro infuzní roztok	NL/H/0820/001	44/591/07-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
OXALIPLATINE TEVA 5 mg/ml, solution à diluer pour perfusion	NL/H/0820/001	NL32938	TEVA SANTÉ	FR
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

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
OXAVIATIN® 5 mg/ml κόνις για διάλυμα προς έγχυση	SE/H/0960/001	26163	VIANEX S.A.	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
Oxaliplatin Vianex 5 mg/ml pulver till infusionsvätska, lösning	SE/H/0960/001	23806	VIANEX S.A.	SE
Oxaliplatina Mylan 5 mg/ml prášok na infúzny roztok	FR/H/0324/001	44/0345/07-S	VIATRIS LIMITED	SK
OXALIPLATINE VIATRIS 5 mg/ml, poudre pour solution pour perfusion	FR/H/0324/001	NL32863	VIATRIS SANTE	FR
VELMINOX 5 mg/ml πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	not available	24330/17/01/01-07-2019	VIOFAR LTD	GR
Oxaliplatin Vipharm® 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	81697.00.00	VIPHARM GMBH	DE
Oxaliplatin Vitane 5 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	7010817.00.00	VITANE PHARMA GMBH	DE
OXALIZOR 5mg/mL πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση.	not available	67193/01-09-2022	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR