19 September 2018
EMA/787363/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): gemcitabine

Procedure No.: PSUSA/00001519/201801

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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
Gembin 40 mg/ml concentrato per soluzione per infusione	PT/H/2035/001	040237012	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Gembin 40 mg/ml concentrato per soluzione per infusione	PT/H/2035/001	040237024	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Gembin 40 mg/ml concentrato per soluzione per infusione	PT/H/2035/001	040237036	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Gembin 40 mg/ml koncentratas infuziniam tirpalui	NL/H/1644/001	LT/1/10/2236/001-003	ACTAVIS GROUP PTC EHF.	LT
GEMBIN 40 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/1644/001	10-0583	ACTAVIS GROUP PTC EHF.	LV
Gembin, 40 mg/ml infusioonilahuse kontsentraat	NL/H/1644/001	708310	ACTAVIS GROUP PTC EHF.	EE
GEMCI-cell® 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	73750.00.00	STADAPHARM GMBH	DE
Gemcisela 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93427.00.00	ALCHEMIA LIMITED	DE
Gemcitabin "medac", koncentrat til infusionsvæske, opløsning	UK/H/5735/001	55640	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DK
Gemcitabin "SUN", infusionsvæske, opløsning	NL/H/3313/001	55023	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	DK
Gemcitabin "Accord"	UK/H/1124/003	46864	ACCORD HEALTHCARE LIMITED	DK
Gemcitabin "Accord", koncentrat til infusionsvæske, opløsning	NL/H/2136/001	48134	ACCORD HEALTHCARE LIMITED	DK
Gemcitabin "Accord", koncentrat til infusionsvæske, opløsning	NL/H/2136/001	48134	ACCORD HEALTHCARE LIMITED	DK
Gemcitabin "Accord", koncentrat til infusionsvæske, opløsning	NL/H/2136/001	48134	ACCORD HEALTHCARE LIMITED	DK
Gemcitabin "Accord", koncentrat til infusionsvæske, opløsning	NL/H/2136/001	48134	ACCORD HEALTHCARE LIMITED	DK
GEMCITABIN "ACTAVIS", KONCENTRAT TIL INFUSIONSVÆSKE, OPLØSNING	NL/H/1644/001	44586	ACTAVIS GROUP PTC EHF.	DK
Gemcitabin "Hospira", koncentrat til infusionsvæske, opløsning	UK/H/1862/001	44787	HOSPIRA UK LTD	DK
Gemcitabin "Hospira", koncentrat til	UK/H/1862/001	44787	HOSPIRA UK LTD	DK

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infusionsvæske, opløsning				
Gemcitabin "Hospira", koncentrat til infusionsvæske, opløsning	UK/H/1862/001	44787	HOSPIRA UK LTD	DK
Gemcitabin "Sandoz", koncentrat til infusionsvæske, opløsning	AT/H/0359/001	46179	SANDOZ A/S	DK
Gemcitabin Accord 100 mg/ml koncentrat za raztopino za infundiranje	UK/H/4287/001	H/12/00679/001	ACCORD HEALTHCARE LIMITED	SI
Gemcitabin Accord 100 mg/ml koncentrat za raztopino za infundiranje	UK/H/4287/001	H/12/00679/002	ACCORD HEALTHCARE LIMITED	SI
Gemcitabin Accord 100 mg/ml koncentrat za raztopino za infundiranje	UK/H/4287/001	H/12/00679/003	ACCORD HEALTHCARE LIMITED	SI
Gemcitabin Accord 100 mg/ml koncentrat za raztopino za infundiranje	UK/H/4287/001	H/12/00679/004	ACCORD HEALTHCARE LIMITED	SI
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	1-31286	ACCORD HEALTHCARE LIMITED	AT
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	1-31286	ACCORD HEALTHCARE LIMITED	AT
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	1-31286	ACCORD HEALTHCARE LIMITED	AT
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	1-31286	ACCORD HEALTHCARE LIMITED	AT
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	84099.00.00	ACCORD HEALTHCARE LIMITED	DE
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	84099.00.00	ACCORD HEALTHCARE LIMITED	DE
Gemcitabin Accord 100 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2136/001	84099.00.00	ACCORD HEALTHCARE LIMITED	DE
Gemcitabin Accord 100 mg/ml	NL/H/2136/001	84099.00.00	ACCORD HEALTHCARE LIMITED	DE

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Konzentrat zur Herstellung einer Infusionslösung				
Gemcitabin Accord 2 g Pulver zur Herstellung einer Infusionslösung	UK/H/1124/003	82092.00.00	ACCORD HEALTHCARE LIMITED	DE
Gemcitabin Actavis 40 mg/ml concentrate for solution for infusion	NL/H/1644/001	MA628/06301	ACTAVIS GROUP PTC EHF.	MT
Gemcitabin Actavis 40 mg/ml concentrate for solution for infusion	NL/H/1644/001	09-6488	ACTAVIS GROUP PTC EHF.	NO
Gemcitabin Actavis 40 mg/ml koncentrat till infusionsvätska, lösning	NL/H/1644/001	41996	ACTAVIS GROUP PTC EHF.	SE
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/004	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/002	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/003	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/005	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/006	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Actavis 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/3383/001	H/16/02154/001	ACTAVIS GROUP PTC EHF.	SI
Gemcitabin Aurobindo 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/1646/001/DC	77425.00.00	PUREN PHARMA GMBH & CO. KG	DE
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/004	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/005	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/006	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/007	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/008	EBEWE PHARMA	SI

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Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/009	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/001	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/002	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/003	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/010	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/011	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/012	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/013	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/014	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/015	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/016	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/017	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml koncentrat za raztopino za infundiranje	AT/H/0224/001	H/10/02011/018	EBEWE PHARMA	SI
Gemcitabin Ebewe 10 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0224/001	1-28015	EBEWE PHARMA	AT
Gemcitabin Ebewe 40 mg/ml koncentrát pro infuzní roztok	AT/H/0359/001	44/907/10-C	EBEWE PHARMA	CZ
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/004	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/005	EBEWE PHARMA	SI

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Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/006	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/007	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/008	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/009	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/010	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/001	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/002	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml koncentrat za raztopino za infundiranje	AT/H/0359/001	H/11/02012/003	EBEWE PHARMA	SI
Gemcitabin Ebewe 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0360/001	1-29789	EBEWE PHARMA	AT
Gemcitabin Fresenius Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	BE466755	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabin Fresenius Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	BE466764	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabin Fresenius Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	BE466773	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabin Fresenius Kabi 40 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2447/001	30208	FRESENIUS KABI ONCOLOGY PLC.	FI
Gemcitabin Fresenius Kabi 40 mg/ml innrennslisþykkni, lausn.	NL/H/2447/001	IS/1/12/129/01	FRESENIUS KABI ONCOLOGY PLC.	IS
Gemcitabin Fresenius Kabi 40 mg/ml koncentrat till infusionsvätska, lösning	NL/H/2447/001	30208	FRESENIUS KABI ONCOLOGY PLC.	FI
Gemcitabin Fresenius Kabi 40 mg/ml	NL/H/2447/001	46981	FRESENIUS KABI ONCOLOGY	SE

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koncentrat till infusionsvätska, lösning			PLC.	
Gemcitabin HEXAL® 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0360/001	80317.00.00	HEXAL AG	DE
Gemcitabin Hospira 38 mg/ml koncentrat till infusionsvätska, lösning	UK/H/1862/001	42265	HOSPIRA UK LTD	SE
Gemcitabin Kabi 38 mg/ml koncentrat za raztopino za infundiranje	NL/H/2447/002	H/14/00683/004	FRESENIUS KABI ONCOLOGY PLC.	SI
Gemcitabin Kabi 38 mg/ml koncentrat za raztopino za infundiranje	NL/H/2447/002	H/14/00683/005	FRESENIUS KABI ONCOLOGY PLC.	SI
Gemcitabin Kabi 38 mg/ml koncentrat za raztopino za infundiranje	NL/H/2447/002	H/14/00683/006	FRESENIUS KABI ONCOLOGY PLC.	SI
Gemcitabin Kabi 38 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2447/002	OGYI-T-21719/07	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 38 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2447/002	OGYI-T-21719/08	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 38 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2447/002	OGYI-T-21719/09	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	136344	FRESENIUS KABI ONCOLOGY PLC.	AT
Gemcitabin Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	90672.00.00	FRESENIUS KABI ONCOLOGY PLC.	DE
Gemcitabin Kabi 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/002	2017040097	FRESENIUS KABI DEUTSCHLAND GMBH	LU
Gemcitabin Kabi 40 mg/ml koncentrテ。tum oldatos infテコziテウhoz	NL/H/2447/001	OGYI-T-21719/04	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 40 mg/ml koncentrテ。tum oldatos infテコziテウhoz	NL/H/2447/001	OGYI-T-21719/05	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 40 mg/ml koncentrテ。tum oldatos infテコziテウhoz	NL/H/2447/001	OGYI-T-21719/06	FRESENIUS KABI ONCOLOGY PLC.	HU
Gemcitabin Kabi 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/001	1-31940	FRESENIUS KABI ONCOLOGY PLC.	AT
Gemcitabin Kabi 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/001	2013110396	FRESENIUS KABI ONCOLOGY PLC.	LU

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Gemcitabin Mylan 40 mg/ml infúzny koncentrát infúzny koncentrát	NL/H/1645/001	44/0774/10-S	MYLAN S.A.S	SK
Gemcitabin OMNICARE 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	94427.00.00	OMNICARE PHARMA GMBH	DE
Gemcitabin Pfizer 38 mg/ml infuusiokonsentraatti, liuosta varten	UK/H/1862/001	27346	PFIZER PFE FINLAND OY	FI
Gemcitabin Pfizer 38 mg/ml infuusiokonsentraatti, liuosta varten	UK/H/1862/001	27346	PFIZER PFE FINLAND OY	FI
Gemcitabin Pfizer 38 mg/ml infuusiokonsentraatti, liuosta varten	UK/H/1862/001	27346	PFIZER PFE FINLAND OY	FI
Gemcitabin Pfizer 38 mg/ml innrennslisþykkni, lausn	UK/H/1862/001	IS/1/10/001/01	PFIZER APS	IS
Gemcitabin Pfizer 38 mg/ml innrennslisþykkni, lausn	UK/H/1862/001	IS/1/10/001/01	PFIZER APS	IS
Gemcitabin Pfizer 38 mg/ml innrennslisþykkni, lausn	UK/H/1862/001	IS/1/10/001/01	PFIZER APS	IS
Gemcitabin Pfizer 38 mg/ml koncentrat till infusionsvätska, lösning	UK/H/1862/001	27346	PFIZER PFE FINLAND OY	FI
Gemcitabin Pfizer 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	1-30909	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Gemcitabin Pfizer 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	1-30909	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Gemcitabin Pfizer 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	1-30909	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Gemcitabin Profusio 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93425.00.00	PGD PROFUSIO LEIPZIG GESUNDHEITS GMBH DEUTSCHLAND	DE
Gemcitabin Ribosepharm 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93426.00.00	HIKMA PHARMA GMBH	DE
Gemcitabin Ribosepharm 40 mg/ml	not available	93426.00.00	HIKMA PHARMA GMBH	DE

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Konzentrat zur Herstellung einer Infusionslösung				
Gemcitabin Ribosepharm 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93426.00.00	HIKMA PHARMA GMBH	DE
Gemcitabin Sandoz 40 mg/ml concentraat voor oplossing voor infusie	AT/H/0359/001	BE379224	SANDOZ N.V.	BE
Gemcitabin Sandoz 40 mg/ml concentraat voor oplossing voor infusie	AT/H/0359/001	BE379233	SANDOZ N.V.	BE
Gemcitabin Sandoz 40 mg/ml concentraat voor oplossing voor infusie	AT/H/0359/001	BE379242	SANDOZ N.V.	BE
Gemcitabin Sandoz 40 mg/ml infuusiokonsentraatti, liuosta varten	AT/H/0359/001	28251	SANDOZ A/S	FI
Gemcitabin Sandoz 40 mg/ml koncentrat till infusionsvätska, lösning	AT/H/0359/001	43531	SANDOZ A/S	SE
Gemcitabin Sandoz 40 mg/ml konsentrat til infusjonsvæske, oppløsning	AT/H/0359/001	09-7261	SANDOZ A/S	NO
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	0782276	SANDOZ N.V.	LU
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	0782293	SANDOZ N.V.	LU
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	0782309	SANDOZ N.V.	LU
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	0782312	SANDOZ N.V.	LU
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	0782262	SANDOZ N.V.	LU
Gemcitabin Sandoz 40mg/ml solution à diluer pour perfusion	AT/H/0359/001	2015040072	SANDOZ N.V.	LU
Gemcitabin STADA 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	1-28435	STADA ARZNEIMITTEL GMBH	AT
Gemcitabin SUN 10 mg/ml Infusionslösung	NL/H/3313/001	93415.00.00	SUN PHARMACEUTICALS GERMANY GMBH	DE

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Gemcitabin SUN 10 mg/ml infusionsvätska, lösning	NL/H/3313/001	32841	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FI
Gemcitabin SUN 10 mg/ml infusionsvätska, lösning	NL/H/3313/001	52086	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	SE
Gemcitabin SUN 10 mg/ml infusjonsvæske, oppløsning	NL/H/3313/001	14-10403	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NO
Gemcitabin SUN 10 mg/ml infuusioneste, liuos	NL/H/3313/001	32841	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FI
Gemcitabin SUN 10 mg/ml infuzní roztok	NL/H/3313/001	44/587/15-C	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	CZ
Gemcitabín SUN 10 mg/ml infúzny roztok	NL/H/3313/001	44/0457/15-S	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	SK
Gemcitabin Teva 40 mg/ml koncentrát pro infuzní roztok	NL/H/1647/001	44/906/10-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Gemcitabin Teva 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/1647/001	H/11/00688/001	TEVA PHARMA B.V.	SI
Gemcitabin Teva 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/1647/001	H/11/00688/002	TEVA PHARMA B.V.	SI
Gemcitabin Teva 40 mg/ml koncentrat za raztopino za infundiranje	NL/H/1647/001	H/11/00688/003	TEVA PHARMA B.V.	SI
Gemcitabina Accord 1.000 mg concentrado para solución para perfusión	NL/H/2136/001	76166	ACCORD HEALTHCARE S.L.U.	ES
Gemcitabina Accord 1.500 mg concentrado para solución para perfusión	NL/H/2136/001	76157	ACCORD HEALTHCARE S.L.U.	ES
Gemcitabina Accord 100 mg l ml concentrat pentru solutie perfuzabila	NL/H/2136/001	6658/2014/01	ACCORD HEALTHCARE LIMITED	RO
Gemcitabina Accord 100 mg l ml concentrat pentru solutie perfuzabila	NL/H/2136/001	6658/2014/02	ACCORD HEALTHCARE LIMITED	RO
Gemcitabina Accord 100 mg l ml concentrat pentru solutie perfuzabila	NL/H/2136/001	6658/2014/03	ACCORD HEALTHCARE LIMITED	RO
Gemcitabina Accord 100 mg l ml concentrat pentru solutie perfuzabila	NL/H/2136/001	6658/2014/04	ACCORD HEALTHCARE LIMITED	RO
Gemcitabina Accord 100 mg/ml	NL/H/2136/001	5456843	ACCORD HEALTHCARE LIMITED	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
concentrado para solução para perfusão				
Gemcitabina Accord 100 mg/ml concentrado para solução para perfusão	NL/H/2136/001	5456850	ACCORD HEALTHCARE LIMITED	PT
Gemcitabina Accord 100 mg/ml concentrado para solução para perfusão	NL/H/2136/001	5456876	ACCORD HEALTHCARE LIMITED	PT
Gemcitabina Accord 100 mg/ml concentrado para solução para perfusão	NL/H/2136/001	5456868	ACCORD HEALTHCARE LIMITED	PT
Gemcitabina Accord 100 mg/ml concentrato per soluzione per infusione	NL/H/2136/001	040928044	ACCORD HEALTHCARE LIMITED	IT
Gemcitabina Accord 100 mg/ml concentrato per soluzione per infusione	NL/H/2136/001	040928018	ACCORD HEALTHCARE LIMITED	IT
Gemcitabina Accord 100 mg/ml concentrato per soluzione per infusione	NL/H/2136/001	040928020	ACCORD HEALTHCARE LIMITED	IT
Gemcitabina Accord 100 mg/ml concentrato per soluzione per infusione	NL/H/2136/001	040928032	ACCORD HEALTHCARE LIMITED	IT
Gemcitabina Accord 2.000 mg concentrado para solución para perfusión	NL/H/2136/001	76156	ACCORD HEALTHCARE S.L.U.	ES
Gemcitabina Accord 200 mg concentrado para solución para perfusión	NL/H/2136/001	76158	ACCORD HEALTHCARE S.L.U.	ES
Gemcitabina Accord 2000 mg Pó para solução para perfusão	UK/H/1124/003	5401625	ACCORD HEALTHCARE LIMITED	PT
Gemcitabina Accord 2000 mg polvo para solución para perfusión	UK/H/1124/003	74479	ACCORD HEALTHCARE S.L.U.	ES
Gemcitabina Accord Healthcare 2 g polvere per soluzione per infusione	UK/H/1124/003	039531037	ACCORD HEALTHCARE LIMITED	IT
Gemcitabina Aurobindo Pharma Italia 40 mg/ml concentrato per soluzione per infusione	NL/H/3390/001	044078018	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Gemcitabina Aurobindo Pharma Italia	NL/H/3390/001	044078020	AUROBINDO PHARMA (ITALIA)	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
40 mg/ml concentrato per soluzione per infusione			S.R.L.	
Gemcitabina Aurobindo Pharma Italia 40 mg/ml concentrato per soluzione per infusione	NL/H/3390/001	044078032	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Gemcitabina Aurovitas 1.000 mg concentrado para solución para perfusión	PT/H/2035/001	72.981	AUROVITAS SPAIN,S.A.U.	ES
Gemcitabina Aurovitas 2.000 mg concentrado para solución para perfusión	PT/H/2035/001	72.980	AUROVITAS SPAIN,S.A.U.	ES
Gemcitabina Aurovitas 200 mg concentrado para solución para perfusión	PT/H/2035/001	72.979	AUROVITAS SPAIN,S.A.U.	ES
Gemcitabina Aurovitas 40 mg/ml concentrado para solução para perfusão	NL/H/3390/001	5673421	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Aurovitas 40 mg/ml concentrado para solução para perfusão	NL/H/3390/001	5673439	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Aurovitas 40 mg/ml concentrado para solução para perfusão	NL/H/3390/001	5673447	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Aurovitas Spain 1.000 mg concentrado para solución para perfusión	NL/H/3390/001	80.846	AUROVITAS SPAIN,S.A.U.	ES
Gemcitabina Aurovitas Spain 2.000 mg concentrado para solución para perfusión	NL/H/3390/001	80.845	AUROVITAS SPAIN,S.A.U.	ES
Gemcitabina Fresenius 1.000 mg concentrado para solución para perfusión	NL/H/2447/001	78.459	FRESENIUS KABI ONCOLOGY PLC.	ES
Gemcitabina Fresenius 2.000 mg concentrado para solución para perfusión	NL/H/2447/001	78.463	FRESENIUS KABI ONCOLOGY PLC.	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
Gemcitabina Fresenius 200 mg concentrado para solución para perfusión	NL/H/2447/001	78.458	FRESENIUS KABI ONCOLOGY PLC.	ES
Gemcitabina Hospira 1000 mg concentrado para solución para perfusión	UK/H/1862/001	74082	HOSPIRA UK LTD	ES
Gemcitabina Hospira 200 mg concentrado para solución para perfusión	UK/H/1862/001	74083	HOSPIRA UK LTD	ES
Gemcitabina Hospira 2000 mg concentrado para solución para perfusión	UK/H/1862/001	74081	HOSPIRA UK LTD	ES
Gemcitabina Kabi 1000 mg concentrado para solución para perfusión	NL/H/2447/002	79.873	FRESENIUS KABI ONCOLOGY PLC.	ES
Gemcitabina Kabi 200 mg concentrado para solución para perfusión	NL/H/2447/002	79.872	FRESENIUS KABI ONCOLOGY PLC.	ES
Gemcitabina Kabi 2000 mg concentrado para solución para perfusión	NL/H/2447/002	79.874	FRESENIUS KABI ONCOLOGY PLC.	ES
Gemcitabina Kabi 38 mg/ml concentrado para solução para perfusão	NL/H/2447/002	5627963	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 38 mg/ml concentrado para solução para perfusão	NL/H/2447/002	5627971	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 38 mg/ml concentrado para solução para perfusão	NL/H/2447/002	5628003	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 38 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/002	7404/2015/01	FRESENIUS KABI ONCOLOGY PLC.	RO
Gemcitabina Kabi 38 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/002	7404/2015/02	FRESENIUS KABI ONCOLOGY PLC.	RO
Gemcitabina Kabi 38 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/002	7404/2015/03	FRESENIUS KABI ONCOLOGY PLC.	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
Gemcitabina Kabi 40 mg/ml concentrado para solução para perfusão	NL/H/2447/001	5553805	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 40 mg/ml concentrado para solução para perfusão	NL/H/2447/001	5553813	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 40 mg/ml concentrado para solução para perfusão	NL/H/2447/001	5553821	FRESENIUS KABI ONCOLOGY PLC.	PT
Gemcitabina Kabi 40 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/001	10586/2018/01	FRESENIUS KABI ONCOLOGY PLC.	RO
Gemcitabina Kabi 40 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/001	10586/2018/02	FRESENIUS KABI ONCOLOGY PLC.	RO
Gemcitabina Kabi 40 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/2447/001	10586/2018/03	FRESENIUS KABI ONCOLOGY PLC.	RO
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675152	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675178	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675210	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675202	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675160	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/ml concentrado para solução para perfusão	UK/H/5735/001	5675228	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	PT
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210019	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE	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
			MBH (WEDEL)	
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210021	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IT
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210033	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IT
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210045	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IT
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210058	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IT
Gemcitabina medac 38 mg/mL concentrato per soluzione per infusione	UK/H/5735/001	044210060	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	IT
Gemcitabina Mylan 40 mg/ml, concentrado para solução para perfusão	NL/H/1645/001	5346234	MYLAN, LDA	PT
Gemcitabina Mylan 40 mg/ml, concentrado para solução para perfusão	NL/H/1645/001	5346226	MYLAN, LDA	PT
Gemcitabina Mylan 40 mg/ml, concentrado para solução para perfusão	NL/H/1645/001	5346242	MYLAN, LDA	PT
Gemcitabina Pfizer 38 mg/ml Concentrado para solução para perfusão	UK/H/1862/001	5361019	LABORATÓRIOS PFIZER, LDA.	PT
Gemcitabina Pfizer 38 mg/ml Concentrado para solução para perfusão	UK/H/1862/001	5361001	LABORATÓRIOS PFIZER, LDA.	PT
Gemcitabina Pfizer 38 mg/ml Concentrado para solução para perfusão	UK/H/1862/001	5360979	LABORATÓRIOS PFIZER, LDA.	PT
Gemcitabina Pfizer 38 mg/ml	UK/H/1862/001	040638013	PFIZER 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
Concentrato per soluzione per infusione				
Gemcitabina Pfizer 38 mg/ml Concentrato per soluzione per infusione	UK/H/1862/001	040638037	PFIZER ITALIA S.R.L.	IT
Gemcitabina Pfizer 38 mg/ml Concentrato per soluzione per infusione	UK/H/1862/001	040638025	PFIZER ITALIA S.R.L.	IT
Gemcitabina Ritisca 40 mg/ml Concentrado para soluテテ」o para perfusテ」o	PT/H/2035/001	5330238	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Ritisca 40 mg/ml Concentrado para soluテテ」o para perfusテ」o	PT/H/2035/001	5330246	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Ritisca 40 mg/ml Concentrado para soluテテ」o para perfusテ」o	PT/H/2035/001	5330253	AUROVITAS UNIPESSOAL, LDA.	PT
Gemcitabina Stada 38 mg/ml Concentrado para solução para perfusão	DE/H/5530/001	5316401	STADA LDA.	PT
Gemcitabina Stada 38 mg/ml Concentrado para solução para perfusão	DE/H/5530/001	5316419	STADA LDA.	PT
Gemcitabina Stada 38 mg/ml Concentrado para solução para perfusão	DE/H/5530/001	5316435	STADA LDA.	PT
Gemcitabina Stada 38 mg/ml Concentrado para solução para perfusão	DE/H/5530/001	5316427	STADA LDA.	PT
Gemcitabină Stada 38 mg/ml concentrat pentru soluţie perfuzabilă	DE/H/5530/001	1883/2009/02	STADA ARZNEIMITTEL AG	RO
Gemcitabină Stada 38 mg/ml concentrat pentru soluţie perfuzabilă	DE/H/5530/001	1883/2009/04	STADA ARZNEIMITTEL AG	RO
Gemcitabină Stada 38 mg/ml concentrat pentru soluţie perfuzabilă	DE/H/5530/001	1883/2009/01	STADA ARZNEIMITTEL AG	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
Gemcitabină Stada 38 mg/ml concentrat pentru soluţie perfuzabilă	DE/H/5530/001	1883/2009/03	STADA ARZNEIMITTEL AG	RO
Gemcitabina SUN 10 mg/ml soluţie perfuzabilă	NL/H/3313/001	8514/2016/01-18	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	RO
Gemcitabina SUN 10 mg/ml, solución para perfusión	NL/H/3313/001	80814	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	ES
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167029	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167169	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167031	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167082	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167118	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167070	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167068	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167056	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167017	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167132	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167043	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167094	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167157	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167106	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml	NL/H/3313/001	044167183	SUN PHARMACEUTICAL	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			INDUSTRIES EUROPE B.V.	
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167144	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167120	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabina SUN Pharma 10 mg/ml soluzione per infusione	NL/H/3313/001	044167171	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	IT
Gemcitabină Teva 2000 mg pulbere pentru soluţie perfuzabilă	NL/H/1597/003	6171/2014/01-03	TEVA PHARMACEUTICALS S.R.L	RO
Gemcitabină Teva 40 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/1647/001	7041/2014/01-03	TEVA PHARMACEUTICALS S.R.L	RO
Gemcitabine 10 mg/ml, solution for infusion	NL/H/3313/001	PL 31750/0062	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	UK
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	PA1390/007/004	ACCORD HEALTHCARE LIMITED	IE
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	PA1390/007/004	ACCORD HEALTHCARE LIMITED	IE
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	PA1390/007/004	ACCORD HEALTHCARE LIMITED	IE
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	PA1390/007/004	ACCORD HEALTHCARE LIMITED	IE
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	MA 054/05403	ACCORD HEALTHCARE LIMITED	MT
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	MA 054/05402	ACCORD HEALTHCARE LIMITED	MT
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	MA 054/05401	ACCORD HEALTHCARE LIMITED	MT
Gemcitabine 100 mg/ml Concentrate for Solution for Infusion	NL/H/2136/001	MA 054/05404	ACCORD HEALTHCARE LIMITED	MT
Gemcitabine 2 g Powder for Solution for Infusion	UK/H/1124/003	PA1390/007/003	ACCORD HEALTHCARE LIMITED	IE
Gemcitabine 2 g Powder for Solution for Infusion	UK/H/1124/003	PL 20075/0262	ACCORD HEALTHCARE LIMITED	UK
Gemcitabine 2000 mg Pharmachemie, poeder voor oplossing voor infusie	NL/H/1597/003	RVG 104031	PHARMACHEMIE B.V	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gemcitabine 38 mg/ml powder for solution for infusion	not available	PL 40378/0096	APTIL PHARMA LIMITED	UK
Gemcitabine 38 mg/ml concentrate for solution for infusion	NL/H/2447/002	PA 1422/003/005	FRESENIUS KABI ONCOLOGY PLC.	IE
Gemcitabine 38 mg/ml Concentrate for Solution for Infusion	UK/H/1862/001	PA 437/63/1	HOSPIRA UK LTD	IE
Gemcitabine 38 mg/ml concentrate for solution for infusion	NL/H/2447/002	MA911/00602	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 38 mg/ml concentrate for solution for infusion	NL/H/2447/002	MA911/00603	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 38 mg/ml concentrate for solution for infusion	NL/H/2447/002	MA911/00604	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 38 mg/ml Concentrate for Solution for Infusion	UK/H/1862/001	MA 734/00501	HOSPIRA UK LTD	MT
Gemcitabine 38 mg/ml Concentrate for Solution for Infusion	UK/H/1862/001	MA 734/00502	HOSPIRA UK LTD	MT
Gemcitabine 38 mg/ml Concentrate for Solution for Infusion	UK/H/1862/001	MA 734/00503	HOSPIRA UK LTD	MT
Gemcitabine 38 mg/ml concentrate for solution for infusion	NL/H/2447/002	PL 18727/0038	FRESENIUS KABI ONCOLOGY PLC.	UK
Gemcitabine 38 mg/ml Concentrate for Solution for Infusion	UK/H/1862/001	PL 04515/0224	HOSPIRA UK LTD	UK
Gemcitabine 38 mg/ml concentrate for solution for infusion	UK/H/5735/001	PL 11587/0087	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	UK
Gemcitabine 40 mg/ml Concentrate for solution for infusion	NL/H/2447/001	PA 1422/003/004	FRESENIUS KABI ONCOLOGY PLC.	IE
Gemcitabine 40 mg/ml concentrate for solution for infusion	NL/H/2447/001	MA 911/00601	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 40 mg/ml concentrate for solution for infusion	NL/H/2447/001	MA 911/00605	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 40 mg/ml concentrate for solution for infusion	NL/H/2447/001	MA 911/00606	FRESENIUS KABI ONCOLOGY PLC.	MT
Gemcitabine 40 mg/ml Concentrate for solution for infusion	NL/H/2447/001	PL 18727/0028	FRESENIUS KABI ONCOLOGY PLC.	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gemcitabine 40mg/ml Concentrate for solution for Infusion	NL/H/1644/001	PA1380/015/003	ACTAVIS GROUP PTC EHF.	IE
Gemcitabine 40mg/ml Concentrate for Solution for Infusion	UH/H/6765/001	PL 30306/0654	ACTAVIS GROUP PTC EHF.	UK
Gemcitabine AB 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/3390/001	BE489786	AUROBINDO PHARMA B.V.	BE
Gemcitabine AB 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/3390/001	BE489795	AUROBINDO PHARMA B.V.	BE
Gemcitabine AB 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/3390/001	BE489777	AUROBINDO PHARMA B.V.	BE
Gemcitabine Accord 100 mg/ml Concentraat voor Oplossing voor Infusie	NL/H/2136/001	RVG 109019	ACCORD HEALTHCARE LIMITED	NL
Gemcitabine Accord 100 mg/ml Concentraat voor Oplossing voor Infusie	NL/H/2136/001	RVG 109019	ACCORD HEALTHCARE LIMITED	NL
Gemcitabine Accord 100 mg/ml Concentraat voor Oplossing voor Infusie	NL/H/2136/001	RVG 109019	ACCORD HEALTHCARE LIMITED	NL
Gemcitabine Accord 100 mg/ml Concentraat voor Oplossing voor Infusie	NL/H/2136/001	RVG 109019	ACCORD HEALTHCARE LIMITED	NL
Gemcitabine Accord 100 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2136/001	29406	ACCORD HEALTHCARE LIMITED	FI
Gemcitabine Accord 100 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2136/001	29406	ACCORD HEALTHCARE LIMITED	FI
Gemcitabine Accord 100 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2136/001	29406	ACCORD HEALTHCARE LIMITED	FI
Gemcitabine Accord 100 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2136/001	29406	ACCORD HEALTHCARE LIMITED	FI
Gemcitabine Accord 100 mg/ml infúzny koncentrát	NL/H/2136/001	44/0402/12-S	ACCORD HEALTHCARE LIMITED	SK
Gemcitabine Accord 100 mg/ml infúzny koncentrát	NL/H/2136/001	44/0402/12-S	ACCORD HEALTHCARE LIMITED	SK
Gemcitabine Accord 100 mg/ml infúzny	NL/H/2136/001	44/0402/12-S	ACCORD HEALTHCARE LIMITED	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
koncentrát				
Gemcitabine Accord 100 mg/ml infúzny koncentrát	NL/H/2136/001	44/0402/12-S	ACCORD HEALTHCARE LIMITED	SK
Gemcitabine Accord 100 mg/ml koncentrát pro infuzní roztok	NL/H/2136/001	44/607/12-C	ACCORD HEALTHCARE LIMITED	CZ
Gemcitabine Accord 100 mg/ml koncentrát pro infuzní roztok	NL/H/2136/001	44/607/12-C	ACCORD HEALTHCARE LIMITED	CZ
Gemcitabine Accord 100 mg/ml koncentrát pro infuzní roztok	NL/H/2136/001	44/607/12-C	ACCORD HEALTHCARE LIMITED	CZ
Gemcitabine Accord 100 mg/ml koncentrát pro infuzní roztok	NL/H/2136/001	44/607/12-C	ACCORD HEALTHCARE LIMITED	CZ
Gemcitabine Accord 100 mg/ml Koncentrat till infusionsvätska, lösning	NL/H/2136/001	45507	ACCORD HEALTHCARE LIMITED	SE
Gemcitabine Accord 100 mg/ml Koncentrat till infusionsvätska, lösning	NL/H/2136/001	45507	ACCORD HEALTHCARE LIMITED	SE
Gemcitabine Accord 100 mg/ml Koncentrat till infusionsvätska, lösning	NL/H/2136/001	45507	ACCORD HEALTHCARE LIMITED	SE
Gemcitabine Accord 100 mg/ml Koncentrat till infusionsvätska, lösning	NL/H/2136/001	45507	ACCORD HEALTHCARE LIMITED	SE
Gemcitabine Accord 100 mg/ml koncentratas infuziniam tirpalui	NL/H/2136/001	LT/1/12/2889/002	ACCORD HEALTHCARE LIMITED	LT
Gemcitabine Accord 100 mg/ml koncentratas infuziniam tirpalui	NL/H/2136/001	LT/1/12/2889/003	ACCORD HEALTHCARE LIMITED	LT
Gemcitabine Accord 100 mg/ml koncentratas infuziniam tirpalui	NL/H/2136/001	LT/1/12/2889/004	ACCORD HEALTHCARE LIMITED	LT
Gemcitabine Accord 100 mg/ml koncentratas infuziniam tirpalui	NL/H/2136/001	LT/1/12/2889/001	ACCORD HEALTHCARE LIMITED	LT
Gemcitabine Accord 100 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2136/001	12-0114	ACCORD HEALTHCARE LIMITED	LV
Gemcitabine Accord 100 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2136/001	12-0114	ACCORD HEALTHCARE LIMITED	LV
Gemcitabine Accord 100 mg/ml koncentrāts infūziju šķīduma	NL/H/2136/001	12-0114	ACCORD HEALTHCARE LIMITED	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pagatavošanai				
Gemcitabine Accord 100 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2136/001	12-0114	ACCORD HEALTHCARE LIMITED	LV
Gemcitabine Accord 100 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2136/001	OGYI-T-21772/03	ACCORD HEALTHCARE LIMITED	HU
Gemcitabine Accord 100 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2136/001	OGYI-T-21772/04	ACCORD HEALTHCARE LIMITED	HU
Gemcitabine Accord 100 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2136/001	OGYI-T-21772/05	ACCORD HEALTHCARE LIMITED	HU
Gemcitabine Accord 100 mg/ml koncentrátum oldatos infúzióhoz	NL/H/2136/001	OGYI-T-21772/06	ACCORD HEALTHCARE LIMITED	HU
Gemcitabine Accord 100 mg/ml konsentrat til infusjonsvæske	NL/H/2136/001	10-8088	ACCORD HEALTHCARE LIMITED	NO
Gemcitabine Accord 100 mg/ml konsentrat til infusjonsvæske	NL/H/2136/001	10-8088	ACCORD HEALTHCARE LIMITED	NO
Gemcitabine Accord 100 mg/ml konsentrat til infusjonsvæske	NL/H/2136/001	10-8088	ACCORD HEALTHCARE LIMITED	NO
Gemcitabine Accord 100 mg/ml konsentrat til infusjonsvæske	NL/H/2136/001	10-8088	ACCORD HEALTHCARE LIMITED	NO
Gemcitabine Accord 100 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος για έγχυση	NL/H/2136/001	21481	ACCORD HEALTHCARE LIMITED	CY
Gemcitabine Accord 100 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος για έγχυση	NL/H/2136/001	21481	ACCORD HEALTHCARE LIMITED	CY
Gemcitabine Accord 100 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος για έγχυση	NL/H/2136/001	21481	ACCORD HEALTHCARE LIMITED	CY
Gemcitabine Accord 100 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος για έγχυση	NL/H/2136/001	21481	ACCORD HEALTHCARE LIMITED	CY
Gemcitabine Accord 100 mg/ml Συμπύκνωμα για διάλυμα προς έγχυση	NL/H/2136/001	297230101/ 03-02-2015	ACCORD HEALTHCARE LIMITED	GR
Gemcitabine Accord 100 mg/ml	NL/H/2136/001	297230102/ 03-02-2015	ACCORD HEALTHCARE LIMITED	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
Συμπύκνωμα για διάλυμα προς έγχυση				
Gemcitabine Accord 100 mg/ml Συμπύκνωμα για διάλυμα προς έγχυση	NL/H/2136/001	297230103/ 03-02-2015	ACCORD HEALTHCARE LIMITED	GR
Gemcitabine Accord 100 mg/ml Συμπύκνωμα για διάλυμα προς έγχυση	NL/H/2136/001	297230104/ 03-02-2015	ACCORD HEALTHCARE LIMITED	GR
Gemcitabine Accord 100 mg/ml, infusioonilahuse kontsentraat	NL/H/2136/001	781312	ACCORD HEALTHCARE LIMITED	EE
Gemcitabine Accord 100 mg/ml, infusioonilahuse kontsentraat	NL/H/2136/001	781312	ACCORD HEALTHCARE LIMITED	EE
Gemcitabine Accord 100 mg/ml, infusioonilahuse kontsentraat	NL/H/2136/001	781312	ACCORD HEALTHCARE LIMITED	EE
Gemcitabine Accord 100 mg/ml, infusioonilahuse kontsentraat	NL/H/2136/001	781312	ACCORD HEALTHCARE LIMITED	EE
GEMCITABINE ACCORD 100 mg/ml, solution à diluer pour perfusion	not available	34009 218 993 5 1	ACCORD HEALTHCARE FRANCE SAS	FR
GEMCITABINE ACCORD 100 mg/ml, solution à diluer pour perfusion	not available	34009 218 994 1 2	ACCORD HEALTHCARE FRANCE SAS	FR
GEMCITABINE ACCORD 100 mg/ml, solution à diluer pour perfusion	not available	34009 218 995 8 0	ACCORD HEALTHCARE FRANCE SAS	FR
GEMCITABINE ACCORD 100 mg/ml, solution à diluer pour perfusion	not available	34009 218 992 9 0	ACCORD HEALTHCARE FRANCE SAS	FR
Gemcitabine Accord 2 g infuusiokuiva-aine liuosta varten	UK/H/1124/003	28683	ACCORD HEALTHCARE LIMITED	FI
Gemcitabine Accord 2 g por oldatos infúzióhoz	UK/H/1124/003	OGYI-T- 21772/07	ACCORD HEALTHCARE LIMITED	HU
Gemcitabine Accord 2 g prášok na infúzny roztok	UK/H/1124/003	44/0034/12-S	ACCORD HEALTHCARE LIMITED	SK
Gemcitabine Accord 2 g pulver til infusjonsvæske, oppløsning	UK/H/1124/003	10-7623	ACCORD HEALTHCARE LIMITED	NO
Gemcitabine Accord 2 g pulver till infusionsvätska, lösning	UK/H/1124/003	44339	ACCORD HEALTHCARE LIMITED	SE
Gemcitabine Accord 2000 mg infusioonilahuse pulber	UK/H/1124/003	757711	ACCORD HEALTHCARE LIMITED	EE
Gemcitabine Accord 2000 mg milteliai infuziniam tirpalui	UK/H/1124/003	LT/1/10/1940/003	ACCORD HEALTHCARE LIMITED	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
Gemcitabine Accord 2000 mg, poeder voor oplossing voor infusie	UK/H/1124/003	RVG 107507	ACCORD HEALTHCARE LIMITED	NL
Gemcitabine Accord Healthcare 100 mg/ml concentraat voor oplossing voor infusie	NL/H/2136/001	BE422922	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml concentraat voor oplossing voor infusie	NL/H/2136/001	BE422904	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml concentraat voor oplossing voor infusie	NL/H/2136/001	BE422931	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml concentraat voor oplossing voor infusie	NL/H/2136/001	BE422913	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml Solution à Diluer pour Perfusion	NL/H/2136/001	BE422922	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml Solution à Diluer pour Perfusion	NL/H/2136/001	BE422904	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml Solution à Diluer pour Perfusion	NL/H/2136/001	BE422931	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 100 mg/ml Solution à Diluer pour Perfusion	NL/H/2136/001	BE422913	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 2 g Poeder voor oplossing voor infusie	UK/H/1124/003	BE 426316	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord Healthcare 2 g Poudre pour solution pour perfusion	UK/H/1124/003	BE 426316	ACCORD HEALTHCARE LIMITED	BE
Gemcitabine Accord, 2 g, proszek do sporządzania roztworu do infuzji	UK/H/1124/003	18923	ACCORD HEALTHCARE LIMITED	PL
Gemcitabine Actavis 40 mg/ml concentrate for solution for infusion	NL/H/3383/001	PA 1380/182/001	ACTAVIS GROUP PTC EHF.	IE
Gemcitabine Actavis 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/1644/001	RVG 104283	ACTAVIS GROUP PTC EHF.	NL
Gemcitabine Actavis 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/1646/001/DC	RVG 104285	AUROBINDO PHARMA B.V.	NL
Gemcitabine Actavis PTC 40 mg/ml,	NL/H/3383/001	RVG 116660	ACTAVIS GROUP PTC EHF.	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
concentraat voor oplossing voor infusie				
Gemcitabine Actavis, 40 mg/ml, koncentrat do sporządzania roztworu do infuzji	DK/H/2783/001	19582	ACTAVIS GROUP PTC EHF.	PL
GEMCITABINE ARROW 40 mg/ml, solution à diluer pour perfusion	not available	37,669	ARROW GENERIQUES	FR
Gemcitabine Aurobindo 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/3390/001	RVG 116668	AUROBINDO PHARMA B.V.	NL
Gemcitabine CF 38 mg/ml, concentraat voor oplossing voor infusie	DE/H/5530/001	RVG 102616	CENTRAFARM B.V.	NL
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/07	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/09	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/13	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/11	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/06	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/08	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/05	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/12	EBEWE PHARMA	HU
Gemcitabine Ebewe 10 mg/ml koncentrátum oldatos infúzióhoz	AT/H/0224/001	OGYI-T-21167/10	EBEWE PHARMA	HU
Gemcitabine EG 38 mg/ml concentraat voor oplossing voor infusie	DE/H/5530/001	BE347191	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml concentraat voor oplossing voor infusie	DE/H/5530/001	BE347216	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml concentraat voor oplossing voor infusie	DE/H/5530/001	BE347207	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml concentraat	DE/H/5530/001	BE347182	EUROGENERICS 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
voor oplossing voor infusie				
Gemcitabine EG 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	BE347182	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	BE347207	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	BE347216	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	DE/H/5530/001	BE347191	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml solution à diluer pour perfusion	DE/H/5530/001	BE347182	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml solution à diluer pour perfusion	DE/H/5530/001	BE347207	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml solution à diluer pour perfusion	DE/H/5530/001	BE347191	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml solution à diluer pour perfusion	DE/H/5530/001	BE347216	EUROGENERICS N.V./S.A.	BE
Gemcitabine EG 38 mg/ml solution à diluer pour perfusion	DE/H/5530/001	0019/10010015	EUROGENERICS N.V./S.A.	LU
Gemcitabine Fresenius Kabi 38 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/002	BE466755	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 38 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/002	BE466764	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 38 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/002	BE466773	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 38 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/002	RVG 113984	FRESENIUS KABI ONCOLOGY PLC.	NL
Gemcitabine Fresenius Kabi 38 mg/ml, solution à diluer pour perfusion	NL/H/2447/002	BE466755	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 38 mg/ml, solution à diluer pour perfusion	NL/H/2447/002	BE466764	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 38 mg/ml, solution à diluer pour perfusion	NL/H/2447/002	BE466773	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/001	BE431453	FRESENIUS KABI ONCOLOGY PLC.	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
Gemcitabine Fresenius Kabi 40 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/001	BE431462	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml concentraat voor oplossing voor infusie	NL/H/2447/001	BE431471	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/001	BE431453	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/001	BE431462	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	NL/H/2447/001	BE431471	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml solution à diluer pour perfusion	NL/H/2447/001	BE431453	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml solution à diluer pour perfusion	NL/H/2447/001	BE431462	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40 mg/ml solution à diluer pour perfusion	NL/H/2447/001	BE431471	FRESENIUS KABI ONCOLOGY PLC.	BE
Gemcitabine Fresenius Kabi 40mg/ml, concentraat voor oplossing voor infusie	NL/H/2447/001	RVG 110700	FRESENIUS KABI ONCOLOGY PLC.	NL
Gemcitabine Hospira 38 mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	RVG 104695	HOSPIRA BENELUX BVBA	NL
Gemcitabine Hospira 38 mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	RVG 104695	HOSPIRA BENELUX BVBA	NL
Gemcitabine Hospira 38 mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	RVG 104695	HOSPIRA BENELUX BVBA	NL
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	BE390451	HOSPIRA BENELUX BVBA	BE
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	BE390467	HOSPIRA BENELUX BVBA	BE
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	BE390476	HOSPIRA BENELUX BVBA	BE
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	34009 579 213 5 4	HOSPIRA 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
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	34009 579 214 1 5	HOSPIRA FRANCE	FR
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	34009 579 215 8 3	HOSPIRA FRANCE	FR
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	2011060020 0612942	HOSPIRA BENELUX BVBA	LU
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	2011060020 0612956	HOSPIRA BENELUX BVBA	LU
GEMCITABINE HOSPIRA 38 mg/ml, solution à diluer pour perfusion	UK/H/1862/001	2011060020 0612973	HOSPIRA BENELUX BVBA	LU
Gemcitabine Hospira 38mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	BE390451	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	BE390467	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml concentraat voor oplossing voor infusie	UK/H/1862/001	BE390476	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	BE390451	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	BE390467	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	BE390476	HOSPIRA BENELUX BVBA	BE
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	2011060020 0612956	HOSPIRA BENELUX BVBA	LU
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	2011060020 0612973	HOSPIRA BENELUX BVBA	LU
Gemcitabine Hospira 38mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/1862/001	2011060020 0612942	HOSPIRA BENELUX BVBA	LU
Gemcitabine Hydrochloride Accord 2 g	UK/H/1124/003	11-0321	ACCORD HEALTHCARE LIMITED	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pulveris infuziju škiduma pagatavošanai				
Gemcitabine Kabi 38 mg/ml infúzny koncentrát	NL/H/2447/002	44/0186/15-S	FRESENIUS KABI ONCOLOGY PLC.	SK
Gemcitabine Kabi 38 mg/ml koncentrát pro infuzní roztok	NL/H/2447/002	44/077/15-C	FRESENIUS KABI ONCOLOGY PLC.	CZ
Gemcitabine Kabi 38 mg/ml koncentratas infuziniam tirpalui	NL/H/2447/002/DC	LT/1/13/3233/004	FRESENIUS KABI ONCOLOGY PLC.	LT
Gemcitabine Kabi 38 mg/ml koncentratas infuziniam tirpalui	NL/H/2447/002/DC	LT/1/13/3233/005	FRESENIUS KABI ONCOLOGY PLC.	LT
Gemcitabine Kabi 38 mg/ml koncentratas infuziniam tirpalui	NL/H/2447/002/DC	LT/1/13/3233/006	FRESENIUS KABI ONCOLOGY PLC.	LT
Gemcitabine Kabi 38 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2447/002	15-0028-01	FRESENIUS KABI ONCOLOGY PLC.	LV
Gemcitabine Kabi 38 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2447/002	15-0028-02	FRESENIUS KABI ONCOLOGY PLC.	LV
Gemcitabine Kabi 38 mg/ml koncentrāts infūziju šķīduma pagatavošanai	NL/H/2447/002	15-0028-03	FRESENIUS KABI ONCOLOGY PLC.	LV
Gemcitabine Kabi 38 mg/ml, infusioonilahuse kontsentraat	NL/H/2447/002	863014	FRESENIUS KABI ONCOLOGY PLC.	EE
GEMCITABINE KABI 38 mg/mL, solution à diluer pour perfusion	NL/H/2447/002	34009 550 013 6 2	FRESENIUS KABI ONCOLOGY PLC.	FR
GEMCITABINE KABI 38 mg/mL, solution à diluer pour perfusion	NL/H/2447/002	34009 550 013 7 9	FRESENIUS KABI ONCOLOGY PLC.	FR
GEMCITABINE KABI 38 mg/mL, solution à diluer pour perfusion	NL/H/2447/002	34009 550 013 9 3	FRESENIUS KABI ONCOLOGY PLC.	FR
Gemcitabine Kabi 40 mg/ml infúzny koncentrát	NL/H/2447/001	44/0167/13-S	FRESENIUS KABI ONCOLOGY PLC.	SK
Gemcitabine Kabi, 38 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/2447/002	22322	FRESENIUS KABI ONCOLOGY PLC.	PL
Gemcitabine medac 38 mg/ml	UK/H/5735/001	33196	MEDAC GESELLSCHAFT FÜR	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infuusiokonsentraatti, liuosta varten			KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	
Gemcitabine medac 38 mg/ml infuusiokonsentraatti, liuosta varten	UK/H/5735/001	33196	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FI
Gemcitabine medac 38 mg/ml infúzny koncentrát	UK/H/5735/001	44/0122/16-S	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SK
Gemcitabine medac 38 mg/ml koncentrát pro infuzní roztok	UK/H/5735/001	44/156/16-C	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	CZ
Gemcitabine medac 38 mg/ml koncentrat till infusionsvätska, lösning	UK/H/5735/001	52693	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	SE
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/001	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/002	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/003	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/004	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/005	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentratas infuziniam tirpalui	UK/H/5735/001	LT/1/16/3896/006	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LT
Gemcitabine medac 38 mg/ml koncentrāts infūziju šķīduma pagatavošanai	UK/H/5735/001	16-0051	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 3 1	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 6 2	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 8 6	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 5 5	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 7 9	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac 38 mg/mL solution à diluer pour perfusion	UK/H/5735/001	34009 550 419 9 3	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	FR
Gemcitabine medac, 38 mg/ml infusioonilahuse kontsentraat	UK/H/5735/001	905816	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	EE
Gemcitabine Mylan 40 mg/ml, concentrate for solution for infusion	NL/H/1645/001	RVG 104286	MYLAN S.P.A.	NL
Gemcitabine Mylan 40 mg/ml, solution à diluer pour perfusion	not available	NL 37 667	MYLAN S.A.S	FR
Gemcitabine Sandoz 40 mg/ml, concentraat voor oplossing voor infusie	AT/H/0359/001	RVG 106596	SANDOZ B.V.	NL
GEMCITABINE SANDOZ 40 mg/ml, solution à diluer pour perfusion	AT/H/0359/001	34009 578 730 6 6	SANDOZ	FR
GEMCITABINE SANDOZ 40 mg/ml, solution à diluer pour perfusion	AT/H/0359/001	34009 578 731 2 7	SANDOZ	FR
GEMCITABINE SANDOZ 40 mg/ml, solution à diluer pour perfusion	AT/H/0359/001	34009 578 732 9 5	SANDOZ	FR
GEMCITABINE SANDOZ 40 mg/ml, solution à diluer pour perfusion	AT/H/0359/001	34009 578 733 5 6	SANDOZ	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
GEMCITABINE SANDOZ 40 mg/ml, solution à diluer pour perfusion	AT/H/0359/001	34009 578 734 1 7	SANDOZ	FR
Gemcitabine STADA 38 mg/ml Concentrate for Solution for Infusion	DE/H/5530/001	PL 11204/0215	STADA ARZNEIMITTEL AG	UK
Gemcitabine STADA 38 mg/ml koncentratas infuziniam tirpalui	DE/H/5530/001	LT/1/10/1894/001	STADA ARZNEIMITTEL AG	LT
Gemcitabine STADA 38 mg/ml koncentratas infuziniam tirpalui	DE/H/5530/001	LT/1/10/1894/002	STADA ARZNEIMITTEL AG	LT
Gemcitabine STADA 38 mg/ml koncentratas infuziniam tirpalui	DE/H/5530/001	LT/1/10/1894/004	STADA ARZNEIMITTEL AG	LT
Gemcitabine STADA 38 mg/ml koncentratas infuziniam tirpalui	DE/H/5530/001	LT/1/10/1894/003	STADA ARZNEIMITTEL AG	LT
GEMCITABINE STADA, 38 mg/ml infusioonilahuse kontsentraat	DE/H/5530/001	643609	STADA ARZNEIMITTEL AG	EE
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 283 0 7	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 282 8 4	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 283 1 4	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 283 3 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 283 2 1	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
GEMCITABINE SUN 10 mg/mL, solution pour perfusion	NL/H/3313/001	34009 550 378 2 8	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	FR
Gemcitabine SUN 1200 mg oplossing voor infusie	NL/H/3313/001	RVG 116483	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN 1400 mg oplossing voor infusie	NL/H/3313/001	RVG 121182	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN 1600 mg oplossing voor infusie	NL/H/3313/001	RVG 117781	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN 1700 mg oplossing voor infusie	NL/H/3313/001	RVG 117782	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN 1800 mg oplossing	NL/H/3313/001	RVG 117784	SUN PHARMACEUTICAL	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
voor infusie			INDUSTRIES EUROPE B.V.	
Gemcitabine SUN 2000 mg oplossing voor infusie	NL/H/3313/001	RVG 117785	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN 2200 mg oplossing voor infusie	NL/H/3313/001	RVG 117786	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	NL
Gemcitabine SUN, 10 mg/ml, roztwór do infuzji	NL/H/3313/001	23323	SUN PHARMACEUTICAL INDUSTRIES EUROPE B.V.	PL
Gemcitabine Teva 40 mg/ml concentraat voor oplossing voor infusie	NL/H/1647/001	BE384675	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml concentraat voor oplossing voor infusie	NL/H/1647/001	BE384666	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml concentraat voor oplossing voor infusie	NL/H/1647/001	BE384684	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml Concentrate for Solution for Infusion	NL/H/1647/001	MA715/02201	TEVA PHARMA B.V.	MT
GEMCITABINE TEVA 40 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLÖSUNG	NL/H/1647/001	BE384684	TEVA PHARMA BELGIUM N.V./S.A	BE
GEMCITABINE TEVA 40 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLÖSUNG	NL/H/1647/001	BE384666	TEVA PHARMA BELGIUM N.V./S.A	BE
GEMCITABINE TEVA 40 MG/ML KONZENTRAT ZUR HERSTELLUNG EINER INFUSIONSLÖSUNG	NL/H/1647/001	BE384675	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml solution à diluer pour perfusion	NL/H/1647/001	BE384684	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml solution à diluer pour perfusion	NL/H/1647/001	BE384675	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml solution à diluer pour perfusion	NL/H/1647/001	BE384666	TEVA PHARMA BELGIUM N.V./S.A	BE
Gemcitabine Teva 40 mg/ml solution à diluer pour perfusion	NL/H/1647/001	20120300412012300041	TEVA PHARMA BELGIUM N.V./S.A	LU
Gemcitabine Teva 40 mg/ml, concentraat voor oplossing voor infusie	NL/H/1647/001	RVG 104284	TEVA NEDERLAND B.V.	NL
Gemcitabine/Actavis 40 mg/ml	DK/H/2783/001	2865701	ACTAVIS GROUP PTC EHF.	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
συμπύκνωμα για την παρασκευή διαλύματος προς έγχυση				
Gemcitabine/Hospira 38 mg/ml Πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	UK/H/1862/001	21784	HOSPIRA UK LTD	CY
Gemcitabine100 mg/ml Concentrate for Solution for Infusion	UK/H/4287/001	PL 20075/0235	ACCORD HEALTHCARE LIMITED	UK
Gemcitabine100 mg/ml Concentrate for Solution for Infusion	UK/H/4287/001	PL 20075/0235	ACCORD HEALTHCARE LIMITED	UK
Gemcitabine100 mg/ml Concentrate for Solution for Infusion	UK/H/4287/001	PL 20075/0235	ACCORD HEALTHCARE LIMITED	UK
Gemcitabine100 mg/ml Concentrate for Solution for Infusion	UK/H/4287/001	PL 20075/0235	ACCORD HEALTHCARE LIMITED	UK
Gemcitabin-Teva 40 mg/ml koncentrátum oldatos infúzióhoz	NL/H/1647/001	OGYI-T-21769/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Gemcitabin-Teva 40 mg/ml koncentrátum oldatos infúzióhoz	NL/H/1647/001	OGYI-T-21769/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Gemcitabin-Teva 40 mg/ml koncentrátum oldatos infúzióhoz	NL/H/1647/001	OGYI-T-21769/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Gemcitabinum Accord, 100 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/2136/001	20148	ACCORD HEALTHCARE LIMITED	PL
Gemcitabinum Accord, 100 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/2136/001	20148	ACCORD HEALTHCARE LIMITED	PL
Gemcitabinum Accord, 100 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/2136/001	20148	ACCORD HEALTHCARE LIMITED	PL
Gemcitabinum Accord, 100 mg/ml, koncentrat do sporządzania roztworu do infuzji	NL/H/2136/001	20148	ACCORD HEALTHCARE LIMITED	PL
Gemcitina 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93429.00.00	ALCHEMIA LIMITED	DE
Gemcitom 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	not available	93428.00.00	ALCHEMIA LIMITED	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gemedac 38 mg/ml Konzentrat zur Herstellung einer Infusionslösung	UK/H/5735/001	94878.00.00	MEDAC GESELLSCHAFT FÜR KLINISCHE SPEZIALPRÄPARATE MBH (WEDEL)	DE
Gemkabi 38 mg/ml infuusiokonsentraatti, liuosta varten	NL/H/2447/002	31697	FRESENIUS KABI ONCOLOGY PLC.	FI
Gemkabi 38 mg/ml innrennslisþykkni, lausn	NL/H/2447/002	IS/1/14/095/01	FRESENIUS KABI ONCOLOGY PLC.	IS
Gemkabi 38 mg/ml koncentrat till infusionsvätska, lösning	NL/H/2447/002	49702	FRESENIUS KABI ONCOLOGY PLC.	SE
Gemkabi 38 mg/ml konsentrat til infusjonsvæske, oppløsning	NL/H/2447/002	13-9656	FRESENIUS KABI ONCOLOGY PLC.	NO
Gemkabi 40 mg/ml konsentrat til infusjonsvæske, oppløsning	NL/H/2447/001	11-8655	FRESENIUS KABI ONCOLOGY PLC.	NO
Gemkabi, koncentrat til infusionsvæske, opløsning	NL/H/2447/001	49610	FRESENIUS KABI ONCOLOGY PLC.	DK
Gemkabi, koncentrat til infusionsvæske, opløsning	NL/H/2447/002	52930	FRESENIUS KABI ONCOLOGY PLC.	DK
Gemliquid 10 mg/ml infúzny koncentrát	AT/H/0224/001	44/0050/10-S	EBEWE PHARMA	SK
Gemliquid 40 mg/ml infúzny koncentrát	AT/H/0359/001	44/0840/10-S	EBEWE PHARMA	SK
Gemliquid 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0361/001	1-29790	EBEWE PHARMA	AT
GEMLIQUID 40 MG/ML, πυκνό διάλυμα για παρασκευή διαλύματος προς έγχυση	AT/H/0359/001	62147/16-09-2011	EBEWE PHARMA	GR
GEMLIQUID, 10 MG/ML, KONCENTRAT DO SPORZĄDZANIA ROZTWORU DO INFUZJI	AT/H/0224/001	16520	EBEWE PHARMA	PL
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/01	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/02	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/03	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
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/04	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/05	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/06	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/07	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/08	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/09	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	10441/2017/10	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/01	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/02	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/03	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/04	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/05	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/06	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/07	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/08	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/09	EBEWE PHARMA	RO
Gemsol 40 mg/ml concentrat pentru soluţie perfuzabilă	AT/H/0359/001	3294/2011/10	EBEWE PHARMA	RO
Gemsol 40 mg/ml koncentratas	AT/H/0359/001	LT/1/10/2265/002	EBEWE PHARMA	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
infuziniam tirpalui				
Gemsol 40 mg/ml koncentratas infuziniam tirpalui	AT/H/0359/001	LT/1/10/2265/003	EBEWE PHARMA	LT
Gemsol 40 mg/ml koncentratas infuziniam tirpalui	AT/H/0359/001	LT/1/10/2265/005	EBEWE PHARMA	LT
Gemsol 40 mg/ml koncentratas infuziniam tirpalui	AT/H/0359/001	LT/1/10/2265/004	EBEWE PHARMA	LT
Gemsol 40 mg/ml koncentratas infuziniam tirpalui	AT/H/0359/001	LT/1/10/2265/001	EBEWE PHARMA	LT
Gemsol 40 mg/ml koncentrāts infūziju šķīduma pagatavošanai	AT/H/0359/001	10-0549	EBEWE PHARMA	LV
Gemsol 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0359/001	1-29788	EBEWE PHARMA	AT
Gemsol, 40 mg/ml, koncentrat do sporzadzania roztworu do infuzji	AT/H/0359/001	18280	EBEWE PHARMA	PL
Gemstad 38 mg/ml koncentrátum oldatos infúzióhoz	DE/H/5530/001	OGYI-T-20935/04	STADA ARZNEIMITTEL AG	HU
Gemstad 38 mg/ml koncentrátum oldatos infúzióhoz	DE/H/5530/001	OGYI-T-20935/03	STADA ARZNEIMITTEL AG	HU
Gemstad 38 mg/ml koncentrátum oldatos infúzióhoz	DE/H/5530/001	OGYI-T-20935/05	STADA ARZNEIMITTEL AG	HU
Gemstad 38 mg/ml koncentrátum oldatos infúzióhoz	DE/H/5530/001	OGYI-T-20935/06	STADA ARZNEIMITTEL AG	HU
GEMSTAD 38 mg/ml, infúzny koncentrát	DE/H/5530/001	44/0725/09-S	STADA ARZNEIMITTEL AG	SK
Gemstada, koncentrat til infusionsvæske, opløsning	DE/H/5530/001	43338	STADA ARZNEIMITTEL AG	DK
Gemzar 1 g, prášek pro přípravu infuzního roztoku	SE/H/0261/002	44/153/96-B/C	ELI LILLY ČR, S.R.O.	CZ
Gemzar 1 g, pulver till infusionsvätska, lösning	SE/H/0261/002	12063	ELI LILLY SWEDEN AB	SE
Gemzar 1.000 mg innrennslisstofn, lausn	SE/H/0261/002	940119	ELI LILLY DANMARK A/S	IS
Gemzar 1.000 mg polvere per soluzione per infusione	SE/H/0261/002	029452012	ELI LILLY ITALIA 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
Gemzar 1000 mg milteliai infuziniam tirpalui	SE/H/0261/002	LT/1/97/1461/002	ELI LILLY HOLDINGS LIMITED (UK)	LT
Gemzar 1000 mg powder for solution for infusion	SE/H/0261/002	MA001/00301	ELI LILLY AND COMPANY LIMITED	MT
GEMZAR 1000 mg powder for solution for infusion	SE/H/0261/002	PL 00006/0302	ELI LILLY AND COMPANY LIMITED	UK
Gemzar 1000 mg pulver til infusjonsvæske, oppløsning	SE/H/0261/002	09/6924	ELI LILLY NORGE A.S.	NO
Gemzar 1000 mg, pó para solução para perfusão	SE/H/0261/002	2427185	LILLY PORTUGAL - PRODUTOS FARMACÊUTICOS, LDA	PT
GEMZAR 1000 mg, poudre pour solution pour perfusion	SE/H/0261/002	559 675-3 OU 3400955967538	LILLY FRANCE	FR
Gemzar 1000 mg, pulbere pentru soluţie perfuzabilă	SE/H/0261/002	1731/2009/01	LILLY FRANCE	RO
Gemzar 200 mg innrennslisstofn, lausn	SE/H/0261/001	940120	ELI LILLY DANMARK A/S	IS
Gemzar 200 mg milteliai infuziniam tirpalui	SE/H/0261/001	LT/1/97/1461/001	ELI LILLY HOLDINGS LIMITED (UK)	LT
Gemzar 200 mg polvere per soluzione per infusione	SE/H/0261/001	029452024	ELI LILLY ITALIA S.P.A.	IT
Gemzar 200 mg powder for solution for infusion	SE/H/0261/001	MA001/00302	ELI LILLY AND COMPANY LIMITED	MT
Gemzar 200 mg pulver til infusjonsvæske, oppløsning	SE/H/0261/001	94-1045	ELI LILLY NORGE A.S.	NO
Gemzar 200 mg, pó para solução para perfusão	SE/H/0261/001	2427086	LILLY PORTUGAL - PRODUTOS FARMACÊUTICOS, LDA	PT
GEMZAR 200 mg, poudre pour solution pour perfusion	SE/H/0261/001	559 674-7 OU 3400955967477	LILLY FRANCE	FR
Gemzar 200 mg, prášek pro přípravu infuzního roztoku	SE/H/0261/001	44/153/96-A/C	ELI LILLY ČR, S.R.O.	CZ
Gemzar 200 mg, pulbere pentru soluţie perfuzabilă	SE/H/0261/001	1730/2009/01	LILLY FRANCE	RO
Gemzar 200 mg, pulver till infusionsvätska, lösning	SE/H/0261/001	12062	ELI LILLY SWEDEN AB	SE
GEMZAR* 200 mg powder for solution for infusion	SE/H/0261/001	PL 00006/0301	ELI LILLY AND COMPANY LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Gemzar, 1000 mg infusioonilahuse pulber	SE/H/0261/002	209598	ELI LILLY HOLDINGS LIMITED (UK)	EE
Gemzar, 200 mg infusioonilahuse pulber	SE/H/0261/001	209498	ELI LILLY HOLDINGS LIMITED (UK)	EE
Gemzar, pulver til infusionsvæske, opløsning	SE/H/0261/002	16631	ELI LILLY DANMARK A/S	DK
Gemzar, pulver til infusionsvæske, opløsning	SE/H/0261/001	16630	ELI LILLY DANMARK A/S	DK
GEMZAR® 1g Pulver zur Herstellung einer Infusionslösung	SE/H/0261/002	1-21425	ELI LILLY GES. M.B.H	AT
GEMZAR® 1g Pulver zur Herstellung einer Infusionslösung	SE/H/0261/002	52222.01.00	LILLY DEUTSCHLAND GMBH	DE
GEMZAR® 200 mg Pulver zur Herstellung einer Infusionslösung	SE/H/0261/001	1-21426	ELI LILLY GES. M.B.H	AT
GEMZAR® 200 mg Pulver zur Herstellung einer Infusionslösung	SE/H/0261/001	52222.00.00	LILLY DEUTSCHLAND GMBH	DE
Getmisi, koncentrat til infusionsvæske, opløsning	NL/H/1647/001	44585	TEVA DENMARK A/S	DK
Gitrabin	NL/H/3383/001	14-10487	ACTAVIS GROUP PTC EHF.	NO
Gitrabin 40 mg/ml concentrat pentru soluţie perfuzabilă	NL/H/1644/001	8720/2016/01-03	ACTAVIS GROUP PTC EHF.	RO
Gitrabin 40 mg/ml innrennslisþykkni, lausn	NL/H/3383/001	IS/1/15/118/01	ACTAVIS GROUP PTC EHF.	IS
Gitrabin 40 mg/ml koncentrat till infusionsvätska, lösning	DK/H/2783/001	42031	ACTAVIS GROUP PTC EHF.	SE
Gitrabin 40 mg/ml koncentrátum oldatos infúzióhoz	NL/H/1644/001	OGYI-T-21026/05-07	ACTAVIS GROUP PTC EHF.	HU
Gitrabin, koncentrat til infusionsvæske, opløsning	DK/H/2783/001	44584	ACTAVIS GROUP PTC EHF.	DK
Ribozar 200 mg Pulver zur Herstellung einer Infusionslösung	DE/H/1275/001	70981.00.00	EBEWE PHARMA	DE
Ribozar 40 mg/ml Konzentrat zur Herstellung einer Infusionslösung	AT/H/0361/001	80318.00.00	EBEWE PHARMA	DE
ΓΚΕΜΖΑΡ 1000 mg κόνις για διάλυμα προς έγχυση	SE/H/0261/002	12140/3-2-2014	PHARMASERVE-LILLY SACI	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**
ΓΚΕΜΖΑΡ 200 mg κόνις για διάλυμα προς έγχυση	SE/H/0261/001	12139/3-2-2014	PHARMASERVE-LILLY SACI	GR
ГЕМСОЛ 40 MG/ML КОНЦЕНТРАТ ЗА ИНФУЗИОНЕН РАЗТВОР	AT/H/0359/001	20130102	EBEWE PHARMA	BG
Гемцитабин Акорд 100 mg/ml концентрат за инфузионен разтвор	NL/H/2136/001	20120379	ACCORD HEALTHCARE LIMITED	BG
Гемцитабин Акорд 100 mg/ml концентрат за инфузионен разтвор	NL/H/2136/001	20120379	ACCORD HEALTHCARE LIMITED	BG
Гемцитабин Акорд 100 mg/ml концентрат за инфузионен разтвор	NL/H/2136/001	20120379	ACCORD HEALTHCARE LIMITED	BG
Гемцитабин Акорд 100 mg/ml концентрат за инфузионен разтвор	NL/H/2136/001	20120379	ACCORD HEALTHCARE LIMITED	BG
Гемцитабин Актавис 40 mg/ml концентрат за инфузионен разтвор	NL/H/3383/001	20160089	ACTAVIS GROUP PTC EHF.	BG
Гемцитабин Хоспира 38 mg/ml концентрат за инфузионен разтвор	UK/H/1862/001	20110466	HOSPIRA UK LTD	BG