14 June 2019
EMA/348220/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ceftazidime

Procedure no.: PSUSA/00000608/201810

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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
Ceftazidim "Stragen", pulver til injektions-/infusionsvæske, opløsning	DK/H/0925/004	38965	STRAGEN NORDIC A/S	DK
Ceftazidim "Stragen", pulver til injektions-/infusionsvæske, opløsning	DK/H/0925/005	38966	STRAGEN NORDIC A/S	DK
Ceftazidim "Stragen", pulver til injektions-/infusionsvæske, opløsning	DK/H/0925/006	38967	STRAGEN NORDIC A/S	DK
Ceftazidim Kabi 2000 mg prašek za raztopino za injiciranje ali infundiranje	PT/H/0186/003	H/10/00357/003	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Ceftazidim Kabi 2000 mg prašek za raztopino za injiciranje ali infundiranje	PT/H/0186/003	H/10/00357/004	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Ceftazidim Sandoz 1 g – Pulver zur Herstellung einer Injektions-/Infusionslösung	AT/H/0880/003	1-25964	SANDOZ GMBH	AT
Ceftazidim Sandoz 1 g injektiokuiva-aine, liuosta varten	SE/H/0366/003	18824	SANDOZ GMBH	FI
Ceftazidim Sandoz 2 g – Pulver zur Herstellung einer Injektions-/Infusionslösung	AT/H/0880/004	1-25965	SANDOZ GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ceftazidim Sandoz 250 mg – Pulver zur Herstellung einer Injektionslösung	AT/H/0880/001	1-25962	SANDOZ GMBH	AT
Ceftazidim Sandoz 500 mg – Pulver zur Herstellung einer Injektionslösung	AT/H/0880/002	1-25963	SANDOZ GMBH	AT
Ceftazidime 1 g Powder for solution for injection or infusion	DK/H/0925/004	PL 21844/0006	STRAGEN UK LTD.	UK
Ceftazidime 1.0 g Powder for Solution for Injection or Infusion	not available	PL 28395/0112	PHARMADREAMS LTD	UK
Ceftazidime 2 g Powder for solution for injection or infusion	DK/H/0925/005	PL 21844/0007	STRAGEN UK LTD.	UK
Ceftazidime 2.0 g Powder for Solution for Injection or Infusion	not available	PL 28395/0113	PHARMADREAMS LTD	UK
Ceftazidime 3 g Powder for solution for injection or infusion	DK/H/0925/006	PL 21844/0008	STRAGEN UK LTD.	UK
Ceftazidime 500 mg Powder for Solution for Injection	not available	PL 28395/0111	PHARMADREAMS LTD	UK
Ceftazidime Fresenius Kabi 2000 mg pulveris injekciju vai infūziju	PT/H/0186/003	08-0389	FRESENIUS KABI POLSKA SP. Z O.O.	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
šķīduma pagatavošanai				
Ceftazidime Fresenius Kabi 2000 mg, süste- või infusioonilahuse pulber	PT/H/0186/003	610708	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Ceftazidime Fresenius Kabi 500 mg pulveris injekciju šķīduma pagatavošanai	PT/H/0186/001	08-0387	FRESENIUS KABI POLSKA SP. Z O.O.	LV
Ceftazidime Fresenius Kabi 500 mg, süstelahuse pulber	PT/H/0186/001	610808	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Fortam 1 g polvo para solución para perfusión	IE/H/0249/001	63.670	GLAXOSMITHKLINE S.A.	ES
Fortam 2 g polvo para solución para perfusión	UK/H/4985/05	56. 613	GLAXOSMITHKLINE S.A.	ES
Fortum 1 g por oldatos injekcióhoz vagy infúzióhoz	IE/H/0249/001	OGYI-T-1265/02	GLAXOSMITHKLINE KFT.	HU
Fortum 1 g powder for solution - for infusion, monovial	IE/H/0249/001	PA 1077/7/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Fortum 1 g powder for solution - for injection or infusion, vial - for infusion, monovial	IE/H/0249/01/MR	PA 1077/7/2	GLAXOSMITHKLINE (IRELAND) LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Fortum 1 g powder for solution for injection or infusion	UK/H/4985/03	PL 00004/0293	GLAXO OPERATIONS UK LTD	UK
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/001	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/002	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/003	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/004	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/005	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/006	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/007	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/008	GLAXOSMITHKLINE D.O.O.	SI

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Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/009	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/010	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/011	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/012	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/013	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/014	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/015	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/016	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/017	GLAXOSMITHKLINE D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/018	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/019	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prašek za raztopino za injiciranje ali infundiranje	IE/H/0249/001	H/93/00650/020	GLAXOSMITHKLINE D.O.O.	SI
Fortum 1 g prášok na injekčný alebo infúzny roztok	IE/H/0249/01	15/0013/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Fortum 1 g pulver til injeksjonsvæske eller infusjonsvæske, oppløsning	IE/H/0249/01/MR	11-8525	GLAXOSMITHKLINE AS	NO
Fortum 1 g pulver till injektionsvätska, lösning	UK/H/4985/03	48221	GLAXOSMITHKLINE AB	SE
Fortum 1 g Pulver zur Herstellung einer Injektionslösung	UK/H/4985/03	1-18139	GLAXOSMITHKLINE PHARMA GMBH.	AT
Fortum 1 g stungulyfsstofn, lausn	IE/H/0249/01/MR	843303	GLAXOSMITHKLINE PHARMA A/S	IS
Fortum 1 g κόνις για ενέσιμο διάλυμα	UK/H/4985/03	19519	GLAXO GROUP LIMITED	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
FORTUM 1 g, poudre pour solution injectable (IM, IV) ou pour perfusion	IE/H/0249/01	NL13649-1	LABORATOIRE GLAXOSMITHKLINE	FR
Fortum 1 g, prášek pro injekční nebo infuzní roztok	SE/H/1875/003	15/207/85-B/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Fortum 1000 mg, poeder voor oplossing voor injectie	UK/H/4985/03	RVG 10540	GLAXOSMITHKLINE B.V.	NL
Fortum 2 g milteliai injekciniam ar infuziniam tirpalui	UK/H/4985/05	LT/1/94/0727/007	GLAXOSMITHKLINE LIETUVA UAB	LT
Fortum 2 g por oldatos injekcióhoz vagy infúzióhoz	UK/H/4985/005	OGYI-T-1265/03	GLAXOSMITHKLINE KFT.	HU
Fortum 2 g powder for solution - for infusion, monovial	UK/H/4985/005	PA 1077/7/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Fortum 2 g powder for solution - for injection or infusion, vial	UK/H/4985/05	PA 1077/7/3	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Fortum 2 g powder for solution for infusion (Monovial)	UK/H/4985/005	PL 00004/0424	GLAXO OPERATIONS UK LTD	UK
Fortum 2 g powder for solution for injection or infusion	UK/H/4985/005	PL 00004/0424	GLAXO OPERATIONS UK LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Fortum 2 g prášok na injekčný alebo infúzny roztok	UK/H/4985/05	15/0014/13-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Fortum 2 g pulver til injeksjonsvæske eller infusjonsvæske, oppløsning	UK/H/4985/05	11-8526	GLAXOSMITHKLINE AS	NO
Fortum 2 g pulver till injektions-/infusionsvätska, lösning	UK/H/4985/05	20655	GLAXOSMITHKLINE AB	SE
Fortum 2 g Pulver zur Herstellung einer Injektions- oder Infusionslösung	UK/H/4985/05	1-18140	GLAXOSMITHKLINE PHARMA GMBH.	AT
Fortum 2 g stungulyfs-/innrennslisstofn, lausn	UK/H/4985/05	843304	GLAXOSMITHKLINE PHARMA A/S	IS
FORTUM 2 g, poudre pour solution injectable (IV) ou pour perfusion	UK/H/4985/05	NL17978	LABORATOIRE GLAXOSMITHKLINE	FR
Fortum 2 g, prášek pro injekční nebo infuzní roztok	SE/H/1875/004	15/207/85-C/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Fortum 2000 mg, poeder voor oplossing voor injectie of infusie	UK/H/4985/005	RVG 12848	GLAXOSMITHKLINE B.V.	NL
FORTUM 250 mg ENFANTS ET NOURRISSONS, poudre	UK/H/4985/01	NL13653-1	LABORATOIRE GLAXOSMITHKLINE	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
pour solution injectable (IM, IV)				
Fortum 250 mg powder for solution for injection	UK/H/4985/01	PL 00004/0304	GLAXO OPERATIONS UK LTD	UK
Fortum 250 mg pulver till injektionsvätska, lösning	UK/H/4985/01	49655	GLAXOSMITHKLINE AB	SE
Fortum 3 g milteliai injekciniam ar infuziniam tirpalui	UK/H/4985/04	LT/1/94/0727/008	GLAXOSMITHKLINE LIETUVA UAB	LT
Fortum 3 g powder for solution for injection or infusion	UK/H/4985/04	PL 00004/0294	GLAXO OPERATIONS UK LTD	UK
FORTUM 500 mg ENFANTS ET NOURRISSONS, poudre pour solution injectable (IM, IV)	UK/H/4985/02	NL13652-1	LABORATOIRE GLAXOSMITHKLINE	FR
Fortum 500 mg milteliai ir tirpiklis injekciniam tirpalui	LT/H/0109/001	LT/1/94/0727/005	GLAXOSMITHKLINE LIETUVA UAB	LT
Fortum 500 mg powder for solution for injection	UK/H/4985/02	PA 1077/7/1	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Fortum 500 mg powder for solution for injection	UK/H/4985/02	PL 00004/0292	GLAXO OPERATIONS UK LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Fortum 500 mg prášok na injekcný roztok	UK/H/4985/02	15/0208/85-CS	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
Fortum 500 mg pulver til injeksjonsvæske, oppløsning	UK/H/4985/02	7140	GLAXOSMITHKLINE AS	NO
Fortum 500 mg pulver till injektionsvätska, lösning	UK/H/4985/02	48220	GLAXOSMITHKLINE AB	SE
Fortum 500 mg, poeder voor oplossing voor injectie	UK/H/4985/02	RVG 12847	GLAXOSMITHKLINE B.V.	NL
Fortum 500 mg, prášek pro injekční roztok	SE/H/1875/002	15/207/85-A/C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Fortum, 1 g süste-/infusioonilahuse pulber	IE/H/0249/001	214898	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Fortum, 1 g, proszek do sporzadzania roztworu do wstrzykiwan / do infuzji	IE/H/0249/001	R/0695	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Fortum, 2 g, proszek do sporzadzania roztworu do wstrzykiwan / do infuzji	UK/H/4985/005	R/0696	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Fortum, 2 g, proszek do sporządzania roztworu do wstrzykiwań / do infuzji	UK/H/4985/005	R/0696	GLAXOSMITHKLINE (IRELAND) LIMITED	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
Fortum, 250 mg, proszek do sporządzania roztworu do wstrzykiwań	SE/H/1875/001	R/0693	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Fortum, 500 mg, proszek do sporządzania roztworu do wstrzykiwań	SE/H/1875/002	R/0694	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Fortum, pulver til injektionsvæske, opløsning	UK/H/4985/05	11449	GLAXOSMITHKLINE PHARMA A/S	DK
Fortum® 0,5 g Pulver zur Herstellung einer Injektionslösung	DE/H/3529/001	4374.00.00	RATIOPHARM GMBH	DE
Fortum® 1,0 g Pulver zur Herstellung einer Injektions- oder Infusionslösung	DE/H/3529/002	4374.01.00	RATIOPHARM GMBH	DE
Fortum® 2,0 g Pulver zur Herstellung einer Infusionslösung	DE/H/3529/003	4374.02.00	RATIOPHARM GMBH	DE
Fortum® for Injection 1 g	UK/H/4985/03	MA 169/00101	GLAXO OPERATIONS UK LTD	MT
FORTUMSET 1 g, poudre pour solution pour perfusion (IV)	IE/H/0249/001	NL21549	LABORATOIRE GLAXOSMITHKLINE	FR
FORTUMSET 2 g, poudre pour solution pour perfusion (IV)	UK/H/4985/005	NL21548	LABORATOIRE GLAXOSMITHKLINE	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
Glazidim 1 g injektiokuiva-aine, liuosta varten	UK/H/4985/003	30689	GLAXO OPERATIONS UK LTD	FI
Glazidim 1 g poeder voor oplossing voor infusie	IE/H/0249/001	BE127321	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g poeder voor oplossing voor injectie of infusie	IE/H/0249/001	BE168122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g polvere per soluzione iniettabile	IT/H/0361/003	025212034	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 1 g polvere per soluzione iniettabile	IT/H/0361/004	025212046	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 1 g polvere per soluzione iniettabile o per infusione	IE/H/0249/001	025212123	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 1 g polvere per soluzione iniettabile o per infusione	IE/H/0249/001	025212111	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 1 g polvere per soluzione per infusione (Monovial)	IE/H/0249/001	025212073	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 1 g polvere per soluzione per infusione (Monovial)	IT/H/0361/005	025212097	GLAXOSMITHKLINE 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
Glazidim 1 g poudre pour solution injectable ou pour solution pour perfusion	IE/H/0249/001	BE168122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g poudre pour solution injectable ou pour solution pour perfusion	IE/H/0249/001	260/08 02 9693	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Glazidim 1 g poudre pour solution pour perfusion (Monovial)	IE/H/0249/001	BE127321	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g poudre pour solution pour perfusion (Monovial)	IE/H/0249/001	260/08 02 9695	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Glazidim 1 g pulver till injektionsvätska, lösning	UK/H/4985/03	30689	GLAXO OPERATIONS UK LTD	FI
Glazidim 1 g Pulver zur Herstellung einer Infusionslösung	IE/H/0249/001	BE127321	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g Pulver zur Herstellung einer Infusionslösung	IE/H/0249/001	260/08 02 9695	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Glazidim 1 g Pulver zur Herstellung einer Injektions-oder Infusionslösung	IE/H/0249/001	BE168122	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 1 g Pulver zur Herstellung einer Injektions-oder Infusionslösung	IE/H/0249/001	260/08 02 9693	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU

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Glazidim 2 g injektio- tai infuusiokuiva-aine, liuosta varten	UK/H/4985/005	9245	GLAXOSMITHKLINE OY	FI
Glazidim 2 g poeder voor oplossing voor injectie of infusie	UK/H/4985/05	BE168113	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 2 g polvere per soluzione iniettabile o per infusione	UK/H/4985/05	025212059	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 2 g polvere per soluzione per infusione (Monovial)	UK/H/4985/005	025212085	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 2 g polvere per soluzione per infusione (Monovial)	IT/H/0361/006	025212109	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 2 g poudre pour solution injectable ou pour solution pour perfusion	UK/H/4985/005	BE168113	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 2 g poudre pour solution pour perfusion (Monovial)	UK/H/4985/05	260/08 02 9696	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Glazidim 2 g pulver till injektions- eller infusionsvätska, lösning	UK/H/4985/05	9245	GLAXOSMITHKLINE OY	FI
Glazidim 2 g Pulver zur Herstellung einer Infusionslösung	UK/H/4985/05	260/08 02 9696	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glazidim 2 g Pulver zur Herstellung einer Injektions-oder Infusionslösung	UK/H/4985/005	BE168113	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 250 mg polvere per soluzione iniettabile	IT/H/0361/001	025212010	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 3 g injektio- tai infuusiokuiva-aine, liuosta varten	UK/H/4985/004	9245	GLAXOSMITHKLINE OY	FI
Glazidim 3 g pulver till injektions- eller infusionsvätska, lösning	UK/H/4985/004	9245	GLAXOSMITHKLINE OY	FI
Glazidim 500 mg injektiokuiva-aine, liuosta varten	UK/H/4985/02	9244	GLAXO OPERATIONS UK LTD	FI
Glazidim 500 mg poeder voor oplossing voor injectie	UK/H/4985/002	BE127337	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 500 mg polvere per soluzione iniettabile	IT/H/0361/002	025212022	GLAXOSMITHKLINE S.P.A.	IT
Glazidim 500 mg poudre pour solution injectable	UK/H/4985/02	BE127337	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 500 mg poudre pour solution injectable	UK/H/4985/02	260/08 02 9692	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glazidim 500 mg pulver till injektionsvätska, lösning	UK/H/4985/02	9244	GLAXO OPERATIONS UK LTD	FI
Glazidim 500 mg Pulver zur Herstellung einer Injektionslösung	UK/H/4985/002	BE127337	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Glazidim 500 mg Pulver zur Herstellung einer Injektionslösung	UK/H/4985/002	260/08 02 9692	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Kefadim 1 g, poeder voor oplossing voor injectie of infusie	not available	BE145801	EUROCEPT B.V.	BE
Kefadim 1 g, poudre pour solution injectable ou pour perfusion	not available	BE145801	EUROCEPT B.V.	BE
Kefadim 1 g, poudre pour solution injectable ou pour perfusion	not available	2005028683	EUROCEPT B.V.	LU
Kefadim 1 g, Pulver zur Herstellung einer Injektions- bzw. Infusionslösung	not available	BE145801	EUROCEPT B.V.	BE
Kefadim 1 g, Pulver zur Herstellung einer Injektions- bzw. Infusionslösung	not available	2005028683	EUROCEPT B.V.	LU
Kefadim 2 g, poeder voor oplossing voor injectie of infusie	not available	BE145792	EUROCEPT B.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Kefadim 2 g, poudre pour solution injectable ou pour perfusion	not available	BE145792	EUROCEPT B.V.	BE
Kefadim 2 g, poudre pour solution injectable ou pour perfusion	not available	2005028684	EUROCEPT B.V.	LU
Kefadim 2 g, Pulver zur Herstellung einer Injektions- bzw. Infusionslösung	not available	BE145792	EUROCEPT B.V.	BE
Kefadim 2 g, Pulver zur Herstellung einer Injektions- bzw. Infusionslösung	not available	2005028684	EUROCEPT B.V.	LU
Kefazim 1 g Trockenstechampulle	not available	1-18437	ASTRO-PHARMA GMBH	AT
Kefazim 2 g Trockenstechampulle	not available	1-18435	ASTRO-PHARMA GMBH	AT
Kefazim 500 mg Trockenstechampulle	not available	1-18436	ASTRO-PHARMA GMBH	AT
Solvetan 1 g κόνις για ενέσιμο διάλυμα ή διάλυμα προς έγχυση	UK/H/4985/003	1884202	GLAXOSMITHKLINE AEBE	GR
Solvetan 2 g κόνις για ενέσιμο διάλυμα ή διάλυμα προς έγχυση	UK/H/4985/05	1884203	GLAXOSMITHKLINE AEBE	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
Starcef 1 g/3 ml polvere e solvente per soluzione iniettabile	not available	025859 036	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
Starcef 250 mg/1 ml polvere e solvente per soluzione iniettabile	not available	025859 012	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT
Starcef 500 mg/1,5 ml polvere e solvente per soluzione iniettabile	not available	025859 024	FABBRICA ITALIANA RITROVATI MEDICINALI ED AFFINI F.I.R.M.A. - S.P.A.	IT