30 November 2017
EMA/811742/2017
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): linezolid

Procedure No.: PSUSA/00001888/201704

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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
Linezolid Sandoz 600 mg/300 ml infúzny roztok	NL/H/2873/001	15/0075/14-S	SANDOZ PHARMACEUTICALS D.D.	SK
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/01	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/02	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/04	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml solu?ie perfuzabila	NL/H/2873/001	6328/2014/03	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/08	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/09	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/05	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/06	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/07	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/10	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/11	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 2 mg/ml soluție perfuzabilă	NL/H/2873/001	6328/2014/12	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg/ 300 ml, oplossing voor infusie	NL/H/2873/001	RVG 113204	SANDOZ B.V.	NL
LINEZOLID SANDOZ 2 MG/ML INFUZNÍ ROZTOK	NL/H/2873/001	15/174/14-C	SANDOZ S.R.O.	CZ
ЛИНЕЗОЛИД САНДОЗ 2MG/ML ИНФУЗИОНЕН РАЗТВОР	NL/H/2873/001	20140118	SANDOZ PHARMACEUTICALS D.D.	BG
Linezolid Sandoz 600 mg filmomhulde tabletten	NL/H/2965/001	BE460444	SANDOZ N.V.	BE
Linezolid Sandoz 2 mg/ml otopina za infuziju	NL/H/2873/001	UP/I-530-09/13-01/32	SANDOZ D.O.O.	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Sandoz 600 mg filmom obalené tablety	NL/H/2965/001	15/0076/14-S	SANDOZ PHARMACEUTICALS D.D.	SK
Linezolid Sandoz 600 mg potahované tablety	NL/H/2965/001	15/300/14-C	SANDOZ S.R.O.	CZ
Linezolid Sandoz, 600 mg, tabletki powlekane	NL/H/2965/001	22143	SANDOZ GMBH	PL
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/01	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/02	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/03	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/04	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/05	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/07	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/06	S.C. SANDOZ S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	6812/2014/08	S.C. SANDOZ S.R.L.	RO
Linezolid "Sandoz", filmovertrukne tabletter	NL/H/2965/001	52754	SANDOZ A/S	DK
Linezolid Sandoz	NL/H/2966/001	043491024	SANDOZ S.P.A.	IT
Linezolid Sandoz	NL/H/2966/001	043491012	SANDOZ S.P.A.	IT
Linezolid Sandoz	NL/H/2966/001	043491036	SANDOZ S.P.A.	IT
Linezolid – 1 A Pharma 600 mg Filmtabletten	NL/H/2965/001	90203.00.00	1 A PHARMA GMBH	DE
Linezolid Sandoz	NL/H/2965/001	852414	SANDOZ PHARMACEUTICALS D.D.	EE
Linezolid Sandoz 600 mg – Filmtabletten	NL/H/2965/001	135752	SANDOZ GMBH	AT
Linezolid HEXAL 600 mg Filmtabletten	NL/H/2966/001	90221.00.00	HEXAL AG	DE
Linezolid Sandoz 600 mg,	NL/H/2965/001	RVG 113822	SANDOZ B.V.	NL

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filmomhulde tabletten				
Linezolid Sandoz 600 mg, filmomhulde tabletten	NL/H/2966/001	RVG 113824	SANDOZ B.V.	NL
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/001	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/002	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/004	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/005	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/006	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/007	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/008	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg plėvele dengtos tabletės	NL/H/2965/001	LT/1/14/3604/003	SANDOZ PHARMACEUTICALS D.D.	LT
Linezolid Sandoz 600 mg comprimidos recubiertos con película EFG	NL/H/2965/001	79193	SANDOZ FARMACÉUTICA, S.A.	ES
Linezolida Sandoz 600 mg comprimidos revestidos por película	NL/H/2965/001	5639539	SANDOZ FARMACÊUTICA LDA.	PT
Linezolida Sandoz 600 mg comprimidos revestidos por película	NL/H/2965/001	5639521	SANDOZ FARMACÊUTICA LDA.	PT
Linezolida Sandoz 600 mg comprimidos revestidos por película	NL/H/2965/001	5639547	SANDOZ FARMACÊUTICA LDA.	PT
Linezolid Sandoz 600 mg tabletti, kalvopäällysteinen	NL/H/2965/001	31593	SANDOZ A/S	FI
Linezolid Sandoz	NL/H/2965/001	13-9593	SANDOZ A/S	NO
Linezolid Sandoz 600 mg filmdragerade tabletter	NL/H/2965/001	49543	SANDOZ A/S	SE

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Linezolid 600 mg Film-coated Tablets	NL/H/2965/001	PL 04416/1389	SANDOZ LTD	UK
Linezolid 600 mg Film-Coated Tablets	NL/H/2965/001	PA0711/230/001	ROWEX LTD	IE
Linezolid Sandoz, 2 mg/ml, roztwór do infuzji	NL/H/2873/001	22046	SANDOZ GMBH	PL
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 402 7	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 403 3	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 406 2	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 405 6	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 409 1	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 408 5	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	279 407 9	SANDOZ	FR
LINEZOLIDE SANDOZ 600 mg, comprimé pelliculé	NL/H/2965/001	586 989 5	SANDOZ	FR
Lynz 600 mg filmom obložene tablete	NL/H/2965/001	HR-H-725335862	SANDOZ D.O.O.	HR
Linezolid HEXAL 2 mg/ml Infusionslösung	NL/H/2873/001	92697.00.00	HEXAL AG	DE
Linezolid Sandoz 2 mg/ml - Infusionslösung	NL/H/2873/001	136402	SANDOZ GMBH	AT
Linezolid Sandoz 2mg/ml solución para perfusión EFG	NL/H/2873/001	80027	SANDOZ FARMACÉUTICA, S.A.	ES
Linezolid 2 mg/ml Solution for Infusion	NL/H/2873/001	PL 04416/1428	SANDOZ LTD	UK
Linezolid Rowex 2 mg/ml solution for infusion	NL/H/2873/001	PA 711/230/2	ROWEX LTD	IE
Linezolid Sandoz GmbH 2	NL/H/2873/001	044079010	SANDOZ GMBH	IT

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mg/ml soluzione per infusione				
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079022	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079034	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079046	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079059	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079061	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079073	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079085	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079097	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079109	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079111	SANDOZ GMBH	IT
Linezolid Sandoz GmbH 2 mg/ml soluzione per infusione	NL/H/2873/001	044079123	SANDOZ GMBH	IT
Linezolid "Sandoz"	NL/H/2965/001	52754	SANDOZ A/S	DK
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/003	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/004	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/005	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/002	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/009	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml	NL/H/2873/001	H/16/02126/008	SANDOZ PHARMACEUTICALS D.D.	SI

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raztopina za infundiranje				
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/001	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/010	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/012	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/006	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/011	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/014	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/013	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/020	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/016	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/007	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/017	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/015	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/019	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/018	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/024	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/022	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/023	SANDOZ PHARMACEUTICALS D.D.	SI

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Linezolid Sandoz 2 mg/ml raztopina za infundiranje	NL/H/2873/001	H/16/02126/021	SANDOZ PHARMACEUTICALS D.D.	SI
Linezolid Mylan 2 mg/ml solution pour perfusion	DK/H/2666/001	BE459120	MYLAN BVBA/SPRL	BE
Linezolid Mylan 2 mg/ml oplossing voor infusie	DK/H/2666/001	BE459120	MYLAN BVBA/SPRL	BE
Linezolid Mylan 2 mg/ml Infusionslösung	DK/H/2666/001	BE459120	MYLAN BVBA/SPRL	BE
Linezolid 2 mg/ml solution for infusion	DK/H/2666/001	PL 04569/1630	GENERICS [UK] LIMITED	UK
Linezolid "Mylan", infusionsvæske, opløsning	DK/H/2666/001	56097	MYLAN HOSPITAL AS	DK
Linezolid 2mg/ml solution for infusion	DK/H/2666/001	PA0405/094/001	GENERICS [UK] LIMITED	IE
Linezolid Mylan 2 mg/ml Infusionslösung	DK/H/2666/001	95119.00.00	MYLAN DURA GMBH	DE
Linezolid Mylan 2 mg/ml infúzny roztok	DK/H/2666/001	15/0422/15-S	GENERICS [UK] LIMITED	SK
Linezolida Mylan 2 mg/ml solução para perfusão	DK/H/2666/001	5668272	MYLAN, LDA	PT
Linezolida Mylan 2 mg/ml solução para perfusão	DK/H/2666/001	5669841	MYLAN, LDA	PT
Linezolida Mylan 2 mg/ml solução para perfusão	DK/H/2666/001	5668264	MYLAN, LDA	PT
Linezolid Mylan 2 mg/ml infusjonsvæske, oppløsning	DK/H/2666/001	15-10734	MYLAN HOSPITAL AS	NO
Linezolid Mylan 2 mg/ml infuusioneste, liuos	DK/H/2666/001	33442	MYLAN HOSPITAL AS	FI
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499010	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499022	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499034	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml	DK/H/2666/001	044499046	MYLAN S.P.A.	IT

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soluzione per infusione				
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499059	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499061	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499073	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499085	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499097	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499109	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499111	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml soluzione per infusione	DK/H/2666/001	044499123	MYLAN S.P.A.	IT
Linezolid Mylan 2 mg/ml infusionsvätska, lösning	DK/H/2666/001	53032	MYLAN HOSPITAL AS	SE
Linézolide Mylan 2 mg/ml, solution pour perfusion	DK/H/2666/001	NL 46105	MYLAN S.A.S	FR
Linezolid Mylan 2 mg/ml solución para perfusión EFG	DK/H/2666/001	80425	MYLAN PHARMACEUTICALS S.L.	ES
Linezolid Mylan 2 mg/mi infuZnl roztok	DK/H/2666/001	15/187/14-C	MYLAN S.A.S	CZ
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/01	ALVOGEN IPCO S.AR.L	RO
Pneumolid 2mg/ml otopina za infuziju	NL/H/2869/001	UP/I-530-09/13-01/31	ALVOGEN IPCO S.AR.L	HR
Pneumolid, 2 mg/ml, roztwór do infuzji	NL/H/2869/001	21937	ALVOGEN IPCO S.AR.L	PL
Pneumolid 2 mg/ml, oplossing voor infusie	NL/H/2869/001	RVG 113200	ALVOGEN IPCO S.AR.L	NL
Пневмолид 2 mg/ml инфузионен разтвор	NL/H/2869/001	II-25490	ALVOGEN IPCO S.AR.L	BG

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/01	ALVOGEN IPCO S.AR.L	HU
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/02	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/03	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/04	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/05	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/06	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/07	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/08	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/09	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/10	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/11	ALVOGEN IPCO S.AR.L	RO
PNEUMOLID 2 mg/ml soluţie perfuzabilă	NL/H/2869/001	6301/2014/12	ALVOGEN IPCO S.AR.L	RO
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/02	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/03	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/04	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/05	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/06	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos	NL/H/2869/001	OGYI-T-22637/07	ALVOGEN IPCO S.AR.L	HU

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infúzió				
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/08	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/09	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/10	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/11	ALVOGEN IPCO S.AR.L	HU
Ziplemol 2 mg/ml oldatos infúzió	NL/H/2869/001	OGYI-T-22637/12	ALVOGEN IPCO S.AR.L	HU
Lineurlub 600 mg filmuhúðaðar töflur	IS/H/0234/001	IS/1/15/033/01	SIGILLATA LIMITED	IS
Lineurlub 600 mg Filmtabletten	IS/H/0234/001	92721.00.00	SIGILLATA LIMITED	DE
Zyvox 2 mg/ml Solution for Infusion	UK/H/0439/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND	IE
Zyvox 100 mg/5 ml granules for oral suspension	UK/H/0439/004	PA 822/143/1	PFIZER HEALTHCARE IRELAND	IE
Zyvoxid 2 mg/ml Infusionslösung	UK/H/0439/001	BE397713	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 600 mg Filmtabletten zur Anwendung bei Erwachsenen	UK/H/0439/003	BE228304	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 2 mg/ml oplossing voor infusie	UK/H/0439/001	BE397713	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 2 mg/ml solution pour perfusion	UK/H/0439/001	BE397713	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 2 mg/ml Infusionslösung	UK/H/0439/001	BE226651	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 600 mg Filmtabletten zur Anwendung bei Erwachsenen	UK/H/0439/003	BE228313	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 100mg/5ml Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/0439/004	BE228322	PFIZER S.A. (BELGIUM)	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg Filmtabletten zur Anwendung bei Erwachsenen	UK/H/0439/003	0194/10020719	PFIZER S.A. (BELGIUM)	LU
Zyvoxid 100mg/5ml Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/0439/004	0194/10020717	PFIZER S.A. (BELGIUM)	LU
Zyvoxid 2 mg/ml Infusionslösung	UK/H/0439/001	2010020716	PFIZER S.A. (BELGIUM)	LU
Zyvox 2 mg/ml Solution for Infusion	UK/H/0439/001	PL 00057/1066	PFIZER LIMITED	UK
Zyvoxid 2 mg/ml διάλυμα για ενδοφλέβια έγχυση	UK/H/0439/001	41057/15-6-2015	PFIZER HELLAS, A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/0439/003	41058/15-6-2015	PFIZER HELLAS, A.E.	GR
Zyvoxid 600 mg - Filmtabletten	UK/H/0439/003	1-24229	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410404	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410416	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410380	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410430	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410392	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410366	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410378	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410428	PFIZER LIMITED	IT
Zyvoxid 600 mg comprimidos recubiertos con película	UK/H/0439/003	64.109	PFIZER, S.L.	ES
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410238	PFIZER LIMITED	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410289	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410240	PFIZER LIMITED	IT
Zyvoxid 600 mg tabletti, kalvopäällysteinen	UK/H/0439/003	16423	PFIZER OY	FI
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410341	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410291	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410277	PFIZER LIMITED	IT
Zyvoxid 2 mg/ml solución para perfusión	UK/H/0439/001	64.106	PFIZER, S.L.	ES
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410354	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410265	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410327	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410253	PFIZER LIMITED	IT
Zyvoxid 2 mg/ml infuusioneste, liuos	UK/H/0439/001	16421	PFIZER OY	FI
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410303	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410339	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410315	PFIZER LIMITED	IT
ZYVOXID® 2 mg/ml Infusionslösung	UK/H/0439/001	51712.00.00	PFIZER PHARMA PFE GMBH	DE
ZYVOXID® 100 mg/5 ml Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/0439/004	51712.00.02	PFIZER PHARMA PFE GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID® 600 mg Filmtabletten	UK/H/0439/003	51712.01.01	PFIZER PHARMA PFE GMBH	DE
Zyvoxid 2 mg/ml solution pour perfusion	UK/H/0439/001	2010020716	PFIZER S.A. (BELGIUM)	LU
Zyvoxid 2 mg/ml oplossing voor infusie	UK/H/0439/001	BE226651	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652989	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752383	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652583	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652781	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3653383	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752284	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752185	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 2 mg/ml solução para perfusão	UK/H/0439/001	3650884	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752581	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3653185	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752482	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652286	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 2 mg/ml solução para perfusão	UK/H/0439/001	5413646	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3751989	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	UK/H/0439/001	01-2267	PFIZER AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3751781	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652682	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652385	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3653284	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3653086	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg filmuhúðaðar töflur	UK/H/0439/003	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn	UK/H/0439/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3653482	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652484	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3652880	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752086	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752789	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3752680	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg, filmdragerade tabletter	UK/H/0439/003	17168	PFIZER AB	SE
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410051	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410024	PFIZER LIMITED	IT
ZYVOXID 600 mg compresse rivestite con film	UK/H/0439/003	035410226	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per	UK/H/0439/001	035410048	PFIZER LIMITED	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
infusione				
Zyvoxid, filmovertrukne tabletter	UK/H/0439/003	32326	PFIZER APS	DK
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410036	PFIZER LIMITED	IT
Zyvoxid, infusionsvæske, opløsning	UK/H/0439/001	32324	PFIZER APS	DK
Zyvoxid 2 mg/ml infusionsvätska, lösning	UK/H/0439/001	171660	PFIZER AB	SE
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410063	PFIZER LIMITED	IT
ZYVOXID 2 mg/ml soluzione per infusione	UK/H/0439/001	035410012	PFIZER LIMITED	IT
Zyvoxid 100 mg/5 ml granulado para suspensão oral	UK/H/0439/004	3752987	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 100 mg/5 ml granulés pour suspension buvable	UK/H/0439/004	0194/10020717	PFIZER S.A. (BELGIUM)	LU
ZYVOXID 2 mg/ml, solution pour perfusion	UK/H/0439/001	34009 581 107 4 0	PFIZER HOLDING FRANCE	FR
Zyvoxid 600 mg tabletter, filmdrasjerte	UK/H/0439/003	01-2269	PFIZER AS	NO
Zyvoxid 100 mg/5 ml granulaat voor orale suspensie	UK/H/0439/004	BE228322	PFIZER S.A. (BELGIUM)	BE
Zyvoxid, granulat til oral suspension	UK/H/0439/004	32327	PFIZER APS	DK
Zyvoxid 20 mg/ml, granulat till oral suspension	UK/H/0439/004	17169	PFIZER AB	SE
ZYVOXID 2 mg/ml, solution pour perfusion	UK/H/0439/001	34009 563 142 6 3	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml - Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/0439/004	1-24230	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Zyvoxid 100 mg/5 ml granulado para suspensão oral	UK/H/0439/004	3653581	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg, filmomhulde	UK/H/0439/003	RVG 26569	PFIZER 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
tabletten				
Zyvoxid 20 mg/ml rakeet oraalisuspensiota varten	UK/H/0439/004	16424	PFIZER OY	FI
Zyvoxid 100 mg/5 ml mixtúrukyrni, dreifa	UK/H/0439/004	IS/1/01/017/04	PFIZER APS	IS
Zyvoxid 600 mg comprimidos revestidos por película	UK/H/0439/003	3751880	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 600 mg filmomhulde tabletten	UK/H/0439/003	BE228313	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 600 mg filmomhulde tabletten	UK/H/0439/003	BE228304	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 100 mg/5 ml granulado para suspensão oral	UK/H/0439/004	3752888	LABORATÓRIOS PFIZER, LDA.	PT
Zyvoxid 100 mg/5 ml granulado para suspensión oral	UK/H/0439/004	64.107	PFIZER, S.L.	ES
Zyvoxid 600 mg comprimés pelliculés	UK/H/0439/003	0194/10020719	PFIZER S.A. (BELGIUM)	LU
ZYVOXID 2 mg/ml - Infusionslösung	UK/H/0439/001	1-24227	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	UK/H/0439/004	34009 565 126 8 3	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	UK/H/0439/004	34009 563 140 3 4	PFIZER HOLDING FRANCE (S.C.A.)	FR
ZYVOXID 600 mg, comprimé pelliculé	UK/H/0439/003	34009 563 139 5 2	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml granulato per sospensione orale	UK/H/0439/004	035410075	PFIZER LIMITED	IT
Zyvoxid 2 mg/ml, oplossing voor infusie	UK/H/0439/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 600 mg comprimés pelliculés	UK/H/0439/003	BE228313	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 2 mg/ml solution pour perfusion	UK/H/0439/001	BE226651	PFIZER S.A. (BELGIUM)	BE
Zyvoxid 100 mg/5 ml granulés	UK/H/0439/004	BE228322	PFIZER S.A. (BELGIUM)	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pour suspension buvable				
Zyvoxid 600 mg comprimés pelliculés	UK/H/0439/003	BE228304	PFIZER S.A. (BELGIUM)	BE
Zyvox 600 mg film-coated tablets	UK/H/0439/003	PA 0822/143/004	PFIZER HEALTHCARE IRELAND	IE
ZYVOXID 20 mg/ml granulat för oral suspension	UK/H/0439/004	16424	PFIZER OY	FI
ZYVOXID 600 mg filmdragerade tabletter	UK/H/0439/003	16423	PFIZER OY	FI
ZYVOXID 2 mg/ml infusionsvätska, lösning	UK/H/0439/001	16421	PFIZER OY	FI
Linezolid Normon 600 mg comprimidos recubiertos con película EFG	ES/H/0291/001	80368	LABORATORIOS NORMON, S.A.	ES
Linezolid Normon 600 mgFilmtabletten	ES/H/0291/001	92996.00.00	LABORATORIOS NORMON, S.A.	DE
Linezolida Normon 600 mg comprimidos revestidos por película	ES/H/0291/001	5669056	LABORATÓRIOS NORMON, S.A.	PT
Linezolid Normon 2 mg/ml solución para perfusión EFG	PT/H/1220/001	80679	LABORATORIOS NORMON, S.A.	ES
Linezolida Normon 2 mg/ml solução para perfusão	PT/H/1220/001	5674270	LABORATÓRIOS NORMON, S.A.	PT
Linezolid Normon 2 mg/ml Infusionslösung	PT/H/1220/001	91574.00.00	LABORATORIOS NORMON, S.A.	DE
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 1 4	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 2 1	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 3 8	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 5 2	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 6 9	PHARMAKI GENERICS LTD	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
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 7 6	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 8 3	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 9 0	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 1 3	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 2 0	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 3 7	PHARMAKI GENERICS LTD	FR
BAGRIZIDINE 600 mg, comprimé pelliculé	not available	34009 300 857 4 4	PHARMAKI GENERICS LTD	FR
Linezolid Denk 600 mg Filmtabletten	NL/H/3125/001	91563.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid 600 mg film-coated tablets	ES/H/0384/001	PL 36390/0188	CIPLA (EU) LIMITED	UK
Linezolid 600 mg film-coated tablets	ES/H/0384/001	PA1809/023/001	CIPLA (EU) LIMITED	IE
Linezolid Cipla 600 mg Filmtabletten	ES/H/0384/001	96632.00.00	CIPLA (EU) LIMITED	DE
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617013	CIPLA (EU) LIMITED	IT
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617025	CIPLA (EU) LIMITED	IT
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617037	CIPLA (EU) LIMITED	IT
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617049	CIPLA (EU) LIMITED	IT
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617052	CIPLA (EU) LIMITED	IT
Linezolid Cipla 600 mg compresse rivestite con film	ES/H/0384/001	044617064	CIPLA (EU) LIMITED	IT
ZYVOXID 2 mg/ml infuzní roztok	not available	15/069/02-C	PFIZER, SPOL. S R.O.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml soluţie perfuzabilă	not available	2047/2009/01	PFIZER EUROPE MA EEIG	RO
ZYVOXID 2 mg/ml infúzny roztok	not available	15/0361/00-S	PFIZER EUROPE MA EEIG	SK
ZYVOXID 2 mg/ml infúzny roztok	not available	15/0361/00-S	PFIZER EUROPE MA EEIG	SK
ZYVOXID 2 mg/ml infuzinis tirpalas	not available	LT/1/02/2448/001	PFIZER EUROPE MA EEIG	LT
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
ZYVOXID 2 mg/ml infuzní roztok	not available	15/069/02-C	PFIZER, SPOL. S R.O.	CZ
ZYVOXID 600 mg plėvele dengtos tabletės	not available	LT/1/02/2448/003	PFIZER EUROPE MA EEIG	LT
ZYVOXID 600 mg apvalkotās tabletes	not available	04-0286	PFIZER EUROPE MA EEIG	LV
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid 2 mg/ml raztopina za infundiranje	not available	H/04/01729/001	PFIZER LUXEMBOURG SARL	SI
Zyvoxid 600 mg filmsko obložene tablete	not available	H/04/01729/002	PFIZER LUXEMBOURG SARL	SI
Zyvoxid, 2 mg/ml infusioonilahus	not available	393802	PFIZER EUROPE MA EEIG	EE
Zyvoxid, 600 mg õhukese polümeerikattega tabletid	not available	394002	PFIZER EUROPE MA EEIG	EE
Zyvoxid 400 mg comprimate filmate	not available	2045/2009/02	PFIZER EUROPE MA EEIG	RO
Zyvoxid 400 mg comprimate filmate	not available	2045/2009/01	PFIZER EUROPE MA EEIG	RO
ZYVOXID 600 mg potahované tablety	not available	15/068/02-C	PFIZER, SPOL. S R.O.	CZ
Zyvoxid, 600 mg, tabletki powlekane	not available	9198	PFIZER EUROPE MA EEIG	PL
ЗИВОКСИД 2 mg/ml инфузионен разтвор	not available	20020808	PFIZER ENTERPRISES SARL	BG

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg comprimate filmate	not available	2046/2009/01	PFIZER EUROPE MA EEIG	RO
Zyvoxid 600 mg comprimate filmate	not available	2046/2009/02	PFIZER EUROPE MA EEIG	RO
ZYVOXID 2 mg/ml šķīdums infūzijām	not available	04-0288	PFIZER EUROPE MA EEIG	LV
Zyvoxid 100 mg/5 ml granule pentru suspensie orală	not available	2048/2009/01	PFIZER EUROPE MA EEIG	RO
ZYVOXID 100 mg/5 ml granulės geriamajai suspensijai	not available	LT/1/02/2448/002	PFIZER EUROPE MA EEIG	LT
Zyvox 600 mg film-coated tablets	not available	MA505/01802	PFIZER HELLAS, A.E.	MT
Zyvox 100 mg/5 ml granules for oral suspension	not available	MA505/01801	PFIZER HELLAS, A.E.	MT
ZYVOXID 2 mg/ ml otopina za infuziju	not available	UP/I-530-09/12-02/47	PFIZER CROATIA D.O.O.	HR
ZYVOXID 600 mg filmom obložene tablete	not available	UP/I-530-09/12-02/45	PFIZER CROATIA D.O.O.	HR
Linezolid Accordpharma 2 mg/ml solución para perfusión EFG	PT/H/1195/001	79964	ACCORD HEALTHCARE S.L.U.	ES
Linezolid Eberth 2 mg/ml, Infusionslösung	not available	92197.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	DE
Linezolid Amneal 600 mg filmomhulde tabletten	NL/H/3292/001	RVG: 115933	AMNEAL PHARMA EUROPE LIMITED	NL
Linezolid Amneal 600 mg comprimidos recubiertos con película EFG	NL/H/3292/001/DC	80446	AMNEAL PHARMA EUROPE LIMITED	ES
Linezolid Amneal 600 mg Filmtabletten	NL/H/3292/001/DC	93313.00.00	AMNEAL PHARMA EUROPE LIMITED	DE
Linezolid Dr. Reddy's 600 mg film-coated tablets	DE/H/4853/001	PL 08553/0572	DR. REDDY'S LABORATORIES (UK) LTD.	UK
Linezolid beta 600 mg Filmtabletten	DE/H/4853/001	95751.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
ANTIZOLID, sol.iv.inf., 2 mg/ml	not available	27538/14	COOPER S.A. PHARMACEUTICALS	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
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 026 6 6	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 026 7 3	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 026 8 0	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 026 9 7	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 0 3	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 1 0	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 2 7	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 3 4	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 4 1	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 5 8	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 6 5	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 7 2	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 027 8 9	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 028 0 2	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 028 1 9	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 028 2 6	OHRE PHARMA	FR
Linezolide Ohre Pharma 600 mg, comprimé pelliculé	AT/H/0491/001/DC	34009 550 028 3 3	OHRE PHARMA	FR
Linezolide Ohre Pharma 600	AT/H/0491/001/DC	34009 550 028 4 0	OHRE PHARMA	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
mg, comprimé pelliculé				
Linezolid Farmaprojects 600 mg, filmomhulde tabletten	NL/H/3125/001	RVG: 114805	FARMAPROJECTS S.A.U.	NL
Linezolid Zentiva® 2 mg/ml Infusionslösung	PT/H/1669/001	86554.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 2 mg/ml Infusionslösung	PT/H/1669/001	86554.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva 2 mg/ml infuzní roztok	PT/H/1669/001	15/108/15-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 2 mg/ml infuzní roztok	PT/H/1669/001	15/108/15-C	ZENTIVA, K.S.	CZ
Linezolida Zentiva 600 mg/300 ml solução para perfusão	PT/H/1669/001	5639661	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Linezolida Zentiva 600 mg/300 ml solução para perfusão	PT/H/1669/001	5639653	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Linezolid Zentiva, 2 mg/ml, roztwór do infuzji	PT/H/1669/001	22475	ZENTIVA, K.S.	PL
Linezolid Zentiva, 2 mg/ml, roztwór do infuzji	PT/H/1669/001	22475	ZENTIVA, K.S.	PL
Linezolid Vancombex 2 mg/ml Solución para perfusión	PT/H/1508/001/DC	81770	MEDICHEM S.A.	ES
Linezolida Vancombex 600mg/300ml solução para perfusão	PT/H/1508/001	PT/H/1508/001	MEDICHEM S.A.	PT
Linezolid Aurovitas Spain 2 mg/ml solución para perfusión EFG	NL/H/2780/001	79.499	AUROVITAS SPAIN,S.A.U.	ES
Linezolida Zentiva 600 mg comprimidos revestidos por película	PT/H/1595/001	5621750	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Linezolida Zentiva 600 mg comprimidos revestidos por película	PT/H/1595/001	5621768	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
potahované tablety				
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva 600 mg potahované tablety	PT/H/1595/001	15/339/14-C	ZENTIVA, K.S.	CZ
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva® 600 mg Filmtabletten	PT/H/1595/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg,	PT/H/1595/001	22312	ZENTIVA, K.S.	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletki powlekane				
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolid Zentiva, 600 mg, tabletki powlekane	PT/H/1595/001	22312	ZENTIVA, K.S.	PL
Linezolida Zentiva 600 mg comprimidos revestidos por	PT/H/1595/001	5645148	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
película				
Linezolid Teva 600 mg filmdragerade tabletter	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/2945/001	79489	TEVA PHARMA S.L.U	ES
Linezolid-ratiopharm® 600 mg Filmtabletten	NL/H/2945/001	90497.00.00	RATIOPHARM GMBH	DE
Linezolid 600 mg Film-coated Tablets	NL/H/2945/001	PL 00289/1853	TEVA UK LIMITED	UK
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid ratiopharm 600 mg Filmtabletten	NL/H/2945/001	136042	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Linezolid Teva 600 mg, filmomhulde tabletten	NL/H/2945/001	RVG 114205	TEVA NEDERLAND B.V.	NL
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5627039	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5626973	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5627013	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5626965	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5627021	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolida Teva 600 mg Comprimidos Revestidos por Película	NL/H/2945/001	5627005	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid "Teva", filmovertrukne tabletter	NL/H/2945/001	53077	TEVA DENMARK A/S	DK
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	MA1060/01601	TEVA B.V	MT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122151	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122264	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122237	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122112	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122302	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122187	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122175	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122163	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122249	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122225	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122276	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122252	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122100	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122148	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122124	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg	NL/H/2945/001	043122050	TEVA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
compresse rivestite con film				
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122074	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122047	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122023	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122290	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122086	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122035	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122199	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122136	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122213	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122136	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122062	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122098	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122252	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122201	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122098	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122288	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122011	TEVA ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 600 mg Film-coated Tablets	PT/H/1362/001	PL 37222/0036	HETERO EUROPE S.L.	UK
Linezolida Hetero Europe, 600 mg, Comprimido revestido por película	PT/H/1362/001	PT/H/1362/001	HETERO EUROPE S.L.	PT
Linezolid Hetero Europe 600 mg Filmtabletten	PT/H/1362/001	93756.00.00	HETERO EUROPE S.L.	DE
Linezolid 2 mg/ml Solution for Infusion	UK/H/5156/001	PL 00057/1418	PFIZER LIMITED	UK
Linezolid 600 mg film-coated tablets	UK/H/5156/002	PL 00057/1419	PFIZER LIMITED	UK
Linezolid 100 mg/5 ml Granules for Oral Suspension	UK/H/5156/003	PL 00057/1420	PFIZER LIMITED	UK
Linezolid Pfizer 600 mg tabletti, kalvopäällysteinen	UK/H/5156/002	30456	PFIZER OY	FI
Linezolid Pfizer 2 mg/ml infuusioneste, liuos	UK/H/5156/001	30455	PFIZER OY	FI
Linezolid Pfizer 2 mg/ml infusionsvätska, lösning	UK/H/5156/001	30455	PFIZER OY	FI
Linezolid Pfizer 600 mg filmdragerade tabletter	UK/H/5156/002	30456	PFIZER OY	FI
Linezolid Pfizer 20 mg/ml granulat för oral suspension	UK/H/5156/003	30457	PFIZER OY	FI
Linezolid Pfizer 20 mg/ml rakeet oraalisuspensiota varten	UK/H/5156/003	30457	PFIZER OY	FI
Linezolid Kern Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/3125/001	80.126	KERN PHARMA, S.L.	ES
Linczolid 2 mg/ml solution for infusion	PT/H/1404/001	PL 42357/0173	AMNEAL PHARMA EUROPE LIMITED	UK
Linezolid Amneal 2 mg/ml Infusionslosung	PT/H/1404/001	94439.00.00	AMNEAL PHARMA EUROPE LIMITED	DE
Linezolid "Amneal", infusionsvæske, opløsning	PT/H/1404/001	55353	AMNEAL PHARMA EUROPE LIMITED	DK
Linezolid Amneal 2 mg/ml infuusioneste, liuos	PT/H/1404/001	32995	AMNEAL PHARMA EUROPE LIMITED	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
Linezolid Amneal 2 mg/ml infusionsvätska, lösning	PT/H/1404/001	52397	AMNEAL PHARMA EUROPE LIMITED	SE
Linezolid STADA® 600 mg Filmtabletten	DE/H/3489/001	86421.00.00	STADAPHARM GMBH	DE
Linezolid Clonmel 600 mg film-coated tablets	DE/H/3489/001	PA0126/238/001	CLONMEL HEALTHCARE LTD.	IE
Linezolid AL 600 mg Filmtabletten	DE/H/3490/001	86422.00.00	ALIUD PHARMA GMBH	DE
Linezolid STADA 600 mg Filmtabletten	DE/H/3490/001	135157	STADA ARZNEIMITTEL GMBH	AT
Tanturb 600 mg filmuhúðaðar töflur	IS/H/0235/001	IS/1/15/034/01	SIGILLATA LIMITED	IS
Anozilad 600 mg filmtabletta	IS/H/0222/001	OGYI-T-22617/01-12	ACTAVIS GROUP PTC EHF.	HU
Linezolid Actavis 600 mg filmuhúðaðar töflur	IS/H/0222/001	IS/1/14/012/01	ACTAVIS GROUP PTC EHF.	IS
Anozilad, 600 mg, tabletki powlekane	IS/H/0222/001	21893	ACTAVIS GROUP PTC EHF.	PL
Linezolid 600mg Film-coated Tablets	IS/H/0222/001/E01	PL 30306/0579	ACTAVIS GROUP PTC EHF.	UK
Linezolid Actavis 600 mg filmdragerade tabletter	IS/H/0222/001/E01	517850	ACTAVIS GROUP PTC EHF.	SE
Linezolid 600 mg film-coated tablets	IS/H/0222/001/E01	PA 1380/178/1	ACTAVIS GROUP PTC EHF.	IE
Linezolid Actavis 600 mg Filmtabletten	IS/H/0222/001	136107	ACTAVIS GROUP PTC EHF.	AT
Linezolid Actavis 600 mg potahované tablety	IS/H/0222/001	15/218/15-C	ACTAVIS GROUP PTC EHF.	CZ
Linezolid Actavis 600 mg filmsko obložene tablete	IS/H/0222/001	H/15/02056/001	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 600 mg filmsko obložene tablete	IS/H/0222/001/E01	H/15/02056/002-012	ACTAVIS GROUP PTC EHF.	SI
Linezolid Fresenius 600 mg, filmomhulde tabletten	NL/H/2777/001	RVG112420	FRESENIUS KABI NEDERLAND B.V.	NL
Linezolid Fresenius 600 mg compresse rivestite con film	NL/H/2777/001	042359012	FRESENIUS KABI ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 048 7 5	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 049 3 6	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 049 1 2	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 048 9 9	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 048 8 2	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 600 mg, comprimé pelliculé.	not available	34009 550 049 0 5	FRESENIUS KABI FRANCE S.A.S.	FR
Linezolid Krka 600 mg film-coated tablets	AT/H/0625/001	PA1347/057/001	KRKA, D.D., NOVO MESTO	IE
Linezolid "Krka", filmovertrukne tabletter	AT/H/0625/001	55654	KRKA, D.D., NOVO MESTO	DK
Linezolid Krka 600 mg tabletti, kalvopäällysteinen	AT/H/0625/001	33189	KRKA, D.D., NOVO MESTO	FI
Linezolid Krka 600 mg film-coated tablets	AT/H/0625/001	PL 01656/0209	KRKA, D.D., NOVO MESTO	UK
Linezolid HCS 600 mg Filmtabletten	AT/H/0625/001	136603	HCS BVBA	AT
Linezolid Krka 600 mg tabletter, filmdrasjerte	AT/H/0625/001	15-10604	KRKA, D.D., NOVO MESTO	NO
Linezolid Krka 600 mg filmdragerade tabletter	AT/H/0625/001	52691	KRKA, D.D., NOVO MESTO	SE
Linezolid TAD® 600 mg Filmtabletten	AT/H/0625/001	94723.00.00	TAD PHARMA GMBH	DE
Linezolid 2mg/ml solution for infusion	PT/H/1143/001	PL 24598/0046	NORIDEM ENTERPRISES LTD	UK
Linezolid Demo 2 mg/ml Infusionslösung	PT/H/1143/001	90655.00.00	DEMO ABEE	DE
Linezolid DEMO 2 mg/ml solución para perfusión EFG	PT/H/1143/001	80350	DEMO ABEE	ES
ZETALID 2 mg/ml Διάλυμα για	PT/H/1143/001	87463/10-12-2015	DEMO ABEE	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
έγχυση				
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672720	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672738	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672746	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672753	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672761	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Soluɬɬɹo para perfusɬɹo	PT/H/1143/001	5672779	DEMO ABEE	PT
Linezan 2 mg/ml Solution for Infusion	PT/H/1555/01/DC	2446/13-1-2017	ANFARM HELLAS SA	GR
Linezolida Anfarm 600mg/300ml Solucao para perfusao	PT/H/1555/01/DC	PT/H/1555/01/DC	ANFARM HELLAS SA	PT
Linezolid Accordpharma 600 mg comprimidos recubiertos con película EFG	NL/H/2776/001	78296	ACCORD HEALTHCARE S.L.U.	ES
Linezolid Accord 600 mg tabletti, kalvopäällysteinen	UK/H/5732/001	32219	ACCORD HEALTHCARE LIMITED	FI
Linezolid 600 mg film-coated tablets	UK/H/5732/001	PA1390/098/001	ACCORD HEALTHCARE LIMITED	IE
Linezolid Accord 600 mg Filmtabletten	UK/H/5732/001	136631	ACCORD HEALTHCARE LIMITED	AT
Linezolid Accord 600 mg filmdrasjerte tabletter	UK/H/5732/001	14-10050	ACCORD HEALTHCARE LIMITED	NO
Linezolid Accord 600 mg Filmtabletten	UK/H/5732/001	92264	ACCORD HEALTHCARE LIMITED	DE
Linezolid Accord 600 mg filmomhulde tabletten	UK/H/5732/001	RVG 115279	ACCORD HEALTHCARE LIMITED	NL
Linezolid Accord 600 mg apvalkotās tabletes	UK/H/5732/001	15-0313	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
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/001	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/002	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/003	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/004	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/005	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/006	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/007	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/008	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/009	ACCORD HEALTHCARE LIMITED	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	UK/H/5732/001	LT/1/15/3837/010	ACCORD HEALTHCARE LIMITED	LT
Linezolid "Accord", filmovertrukne tabletter	UK/H/5732/001	54073	ACCORD HEALTHCARE LIMITED	DK
Linezolid Accord 600 mg filmdragerade tabletter	UK/H/5732/001	51043	ACCORD HEALTHCARE LIMITED	SE
Linezolid Accord 600 mg comprimidos recubiertos con película EFG	UK/H/5732/001	80619	ACCORD HEALTHCARE S.L.U.	ES
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456019	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456021	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456033	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456045	ACCORD HEALTHCARE LIMITED	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456058	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456060	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456072	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456084	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456096	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg compresse rivestite con film	UK/H/5732/001	043456108	ACCORD HEALTHCARE LIMITED	IT
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/001	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/002	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/003	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/004	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/005	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/006	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/007	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/008	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/009	ACCORD HEALTHCARE LIMITED	SI
Linezolid Accord 600 mg filmsko obložene tablete	UK/H/5732/001	H/16/02127/010	ACCORD HEALTHCARE LIMITED	SI
Linezolid 600 mg film-coated tablets	UK/H/5732/001	MA 054/08901	ACCORD HEALTHCARE LIMITED	MT
Linezolida Accord 600 mg	UK/H/5732/001	5684501	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
comprimidos revestidos por película				
Linezolid Accord 600 mg filmom obalené tablety	UK/H/5732/001	15/0188/16-S	ACCORD HEALTHCARE LIMITED	SK
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 558 5 4	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 558 6 1	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 558 8 5	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 558 9 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 559 0 8	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 559 1 5	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 550 199 9 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 559 2 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 300 559 4 6	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	UK/H/5732/001	34009 550 200 0 4	ACCORD HEALTHCARE FRANCE SAS	FR
Linezolid Accord 600 mg potahované tablety	UK/H/5732/001	15/561/15-C	ACCORD HEALTHCARE LIMITED	CZ
Linezolid 600 mg film-coated tablets	UK/H/5732/001	PL 20075/0394	ACCORD HEALTHCARE LIMITED	UK
Linezolida Accord 600 mg comprimidos revestidos por película	UK/H/5732/001	5684519	ACCORD HEALTHCARE LIMITED	PT
Linezolid Accord 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5732/001	022370	ACCORD HEALTHCARE LIMITED	CY
Lizedia 2 mg/ml solution for infusion	AT/H/0563/001	PL 17277/0342	PHARMATHEN S.A.	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
Lizedia 2 mg/ml διάλυμα για έγχυση	AT/H/0563/001	57290/18-07-2016	PHARMATHEN S.A.	GR
Linezolid Pharmathen 2 mg/ml Infusionslösung	AT/H/0563/001	137316	PHARMATHEN S.A.	AT
Linezolid Pharmathen 2 mg/ml Infusionslösung	AT/H/0563/001	93336.00.00	PHARMATHEN S.A.	DE
Linezolid Hetero 600 mg Filmtabletten	NL/H/2776/001	88580.00.00	HETERO EUROPE S.L.	DE
Linezolid Accordpharma 600 mg comprimidos recubiertos con película EFG	NL/H/2776/001	78296	ACCORD HEALTHCARE S.L.U.	ES
Linezolid 600 mg Film-Coated Tablets	NL/H/2776/001	PL 37222/0020	HETERO EUROPE S.L.	UK
Linezolid Hetero 600 mg filmomhulde tabletten	NL/H/2776/001	RVG 112419	HETERO EUROPE S.L.	NL
Linezolid Polpharma 2 mg/ml oplossing voor infusie	NL/H/2781/001	RVG 112520	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	NL
Linezolid Polpharma, 2 mg/ml, roztwór do infuzji	NL/H/2781/001	22018	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Linezolid Medana 2 mg/ml infuzinis tirpalas	NL/H/2781/001	LT/1/15/3782/001	MEDANA PHARMA SPOLKA AKCYJNA	LT
Linezolid Medana 2 mg/ml infuzinis tirpalas	NL/H/2781/001	LT/1/15/3782/002	MEDANA PHARMA SPOLKA AKCYJNA	LT
LINEZA 2 mg/ml infuzní roztok	NL/H/2781/001	15/372/15-C	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	CZ
LINEZA 2 mg/ml infúzny roztok	NL/H/2781/001	15/0383/15-S	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	SK
Линезолид Полфарма 2 mg/ml инфузионен разтвор	NL/H/2781/001	20150377	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	BG
Linezolid Medana 2 mg/ml šķīdums infūzijām	NL/H/2781/001	15-0301	MEDANA PHARMA SPOLKA AKCYJNA	LV
Linezolida Aurovitas 2 mg/ml solução para perfusão	not available	5695143	AUROVITAS UNIPESSOAL, LDA.	PT
Linezolid 600 mg film-coated tablets	NL/H/3012/001	PL 25174/0023	LABORATORIO REIG JOFRE, S.A.	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
Linezolid Sala 600 mg comprimidos recubiertos con película E F G	NL/H/3012/001	79472	LABORATORIO RAMON SALA, S.L	ES
Synzolid 600 mg, filmomhulde tabletten	NL/H/3012/001	RVG 114164	SYNTHON BV	NL
Zolic 600 mg kalvopäällysteiset tabletit	NL/H/3012/001	31756	AVANSOR PHARMA OY	FI
Linezolid Reig Jofre 600 mg tabletter, filmdrasjerte	NL/H/3012/001/DC	13-9723	LABORATORIO REIG JOFRE, S.A.	NO
Linezolid Reig Jofre 600 mg, filmdragerade tabletter	NL/H/3012/001	49885	LABORATORIO REIG JOFRE, S.A.	SE
Synzolid 600 mg compresse rivestite con film	NL/H/3012/001	043560	SYNTHON BV	IT
Linezolid Adamed, 600 mg, tabletki powlekane	NL/H/3012/001	22833	ADAMED	PL
Linezolid Mylan, 2 mg/ml, roztwór do infuzji	NL/H/3530/001	23431	MYLAN S.A.S	PL
Lorezogram 2 mg/ml, oplossing voor infusie	NL/H/3530/001	RVG 117017	MYLAN B.V.	NL
Linezolid EG 2 mg/ml Infusionslösung	DE/H/5027/001	BE478311	EUROGENERICS SA	BE
Linezolid EG 2 mg/ml oplossing voor infusie	DE/H/5027/001	BE478311	EUROGENERICS SA	BE
Linezolid EG 2 mg/ml solution pour perfusion	DE/H/5027/001	BE478311	EUROGENERICS SA	BE
Linezolid Clonmel 2 mg/ml solution for infusion	DE/H/5027/001	PA0126/238/002	CLONMEL HEALTHCARE LTD.	IE
Linezolid STADA 2 mg/ml Infusionslösung	DE/H/5027/001	92193.00.00	STADAPHARM GMBH	DE
Linezolid Hikma 2 mg/ml, Infusionslösung	NL/H/3443/001	94483.00.00	HIKMA PHARMA GMBH	DE
Lynvox 2 mg/ml oplossing voor infusie	NL/H/3443/001	RVG 115278	HIKMA PHARMA GMBH	NL
Линестад 2 mg/ml инфузионен разтвор	not available	20150407	STADA ARZNEIMITTEL AG	BG

Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Inresa 2 mg/ml Infusionslösung	NL/H/3417/001	94423.00.00	INRESA ARZNEIMITTEL GMBH	DE
Lizox 2 mg/ml, oplossing voor infusie	NL/H/3417/001	RVG 115277	INRESA ARZNEIMITTEL GMBH	NL
Dilizolen 2 mg/ml infuzní roztok	NL/H/2505/001	15/315/13-C	GLENMARK PHARMACEUTICALS S.R.O.	CZ
Dilizolen 2 mg/ml infúzny roztok	NL/H/2505/001	15/0181/13-S	GLENMARK PHARMACEUTICALS S.R.O.	SK
Dilizolen 2 mg/ml, oplossing voor infusie	NL/H/2505/001	RVG 111095	GLENMARK PHARMACEUTICALS S.R.O.	NL
Dilizolen, 2 mg/ml, roztwór do infuzji	NL/H/2505/001	21192	GLENMARK PHARMACEUTICALS S.R.O.	PL
Linezolid 600mg Film-coated Tablet	UK/H/6566/001/DC	29831/0653	WOCKHARDT UK LTD	UK
Linezolid 600mg Film-coated Tablet	UK/H/6566/001/DC	PA1339/62/1	WOCKHARDT UK LTD	IE
Grampolid 600 mg, filmomhulde tabletten	NL/H/3013/001	RVG 114165	SYNTHON BV	NL
Grampolid 600 mg Filmtabletten	NL/H/3013/001	2016040046	SYNTHON BV	LU
Linezolida Teva 2 mg/ml solução para perfusão	DK/H/2079/001	5476841	TEVA PHARMA – PRODUTOS FARMACÊUTICOS LDA	PT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879049	TEVA ITALIA S.R.L.	IT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879013	TEVA ITALIA S.R.L.	IT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879052	TEVA ITALIA S.R.L.	IT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879025	TEVA ITALIA S.R.L.	IT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879037	TEVA ITALIA S.R.L.	IT
Linezolid Teva Italia 2 mg/ml soluzione per infusione	DK/H/2079/001	040879064	TEVA ITALIA S.R.L.	IT
Linezolid-ratiopharm® 2 mg/ml	DK/H/2079/001	84513.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Infusionslösung				
Linezolid "Teva", infusionsvæske, opløsning	DK/H/2079/001	48366	TEVA DENMARK A/S	DK
Linezolid 2 mg/ml solution for infusion	DK/H/2079/001	PL 00289/1638	TEVA UK LIMITED	UK
Linezolid ratiopharm 2 mg/ml Infusionslösung	DK/H/2079/001	1-31376	RATIOPHARM ARZNEIMITTEL VERTRIEBS-GMBH	AT
Linezolid Krka 600 mg potahované tablety	AT/H/0620/001	15/532/15-C	KRKA, D.D., NOVO MESTO	CZ
Linezolid Krka 600 mg filmtabletta	AT/H/0620/001	OGYI-T-22928/01	KRKA, D.D., NOVO MESTO	HU
Linezolid Krka 600 mg filmtabletta	AT/H/0620/001	OGYI-T-22928/02	KRKA, D.D., NOVO MESTO	HU
Linezolid Krka 600 mg filmtabletta	AT/H/0620/001	OGYI-T-22928/03	KRKA, D.D., NOVO MESTO	HU
Linezolid Krka 600 mg filmom obložene tablete	AT/H/0620/001	HR-H-094120244	KRKA, D.D., NOVO MESTO	HR
Ilenozyd 600 mg Filmtabletten	AT/H/0620/001	136602	KRKA, D.D., NOVO MESTO	AT
Linezolid Krka, 600 mg õhukese polümeerikattega tabletid	AT/H/0620/001	893315	KRKA, D.D., NOVO MESTO	EE
Линезолид Крка 600 mg филмирани таблетки	AT/H/0620/001	20150383	KRKA, D.D., NOVO MESTO	BG
Linezolid Krka 600 mg plėvele dengtos tabletės	AT/H/0620/001	LT/1/15/3838/001	KRKA, D.D., NOVO MESTO	LT
Linezolid Krka 600 mg plėvele dengtos tabletės	AT/H/0620/001	LT/1/15/3838/002	KRKA, D.D., NOVO MESTO	LT
Linezolid Krka 600 mg plėvele dengtos tabletės	AT/H/0620/001	LT/1/15/3838/003	KRKA, D.D., NOVO MESTO	LT
Linezolid Krka 600 mg apvalkotās tabletes	AT/H/0620/001	15-0290	KRKA, D.D., NOVO MESTO	LV
Linezolid Krka 600 mg filmom obalené tablety	AT/H/0620/001	15/0478/15-S	KRKA, D.D., NOVO MESTO	SK
Linezolid Krka, 600 mg, tabletki powlekane	AT/H/0620/001	22832	KRKA, D.D., NOVO MESTO	PL
Linezolid Krka 600 mg	AT/H/0620/001	8462/2015/01	KRKA, D.D., NOVO MESTO	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
comprimate filmate				
Linezolid Krka 600 mg comprimate filmate	AT/H/0620/001	8462/2015/02	KRKA, D.D., NOVO MESTO	RO
Linezolid Krka 600 mg comprimate filmate	AT/H/0620/001	8462/2015/03	KRKA, D.D., NOVO MESTO	RO
Linezolida Krka 600 mg comprimidos revestidos por película	AT/H/0620/001	5665211	KRKA, D.D., NOVO MESTO	PT
Linezolida Krka 600 mg comprimidos revestidos por película	AT/H/0620/001	5665229	KRKA, D.D., NOVO MESTO	PT
Linezolida Krka 600 mg comprimidos revestidos por película	AT/H/0620/001	5665237	KRKA, D.D., NOVO MESTO	PT
Linezolid Krka 600 mg filmsko obložene tablete	AT/H/0620/001	H/16/02098/001	KRKA, D.D., NOVO MESTO	SI
Linezolid Krka 600 mg filmsko obložene tablete	AT/H/0620/001	H/16/02098/002	KRKA, D.D., NOVO MESTO	SI
Linezolid Krka 600 mg filmsko obložene tablete	AT/H/0620/001	H/16/02098/003	KRKA, D.D., NOVO MESTO	SI
Linezolid Krka 600 mg comprimidos recubiertos con película EFG	AT/H/0620/001	80940	KRKA, D.D., NOVO MESTO	ES
Linezolid TAD 600 mg compresse rivestite con film	AT/H/0620/001	044172017	TAD PHARMA GMBH	IT
Linezolid TAD 600 mg compresse rivestite con film	AT/H/0620/001	044172029	TAD PHARMA GMBH	IT
Linezolid TAD 600 mg compresse rivestite con film	AT/H/0620/001	044172031	TAD PHARMA GMBH	IT
LINEZOLIDE ARROW 600 mg, comprimé pelliculé	not available	NL 46215	ARROW GENERIQUES	FR
Linezolid Sala 2 mg/ml solución para perfusión EFG	not available	80637	LABORATORIO RAMON SALA, S.L	ES
Linezolida Farmoz 600 mg comprimidos revestidos por película	not available	658778	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	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
Linezolida Farmoz 600 mg comprimidos revestidos por película	not available	5658802	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Linezolida Farmoz 600 mg comprimidos revestidos por película	not available	5658810	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Linezolid Mylan 600 mg Filmtabletten	NL/H/3011/001	90681.00.00	MYLAN DURA GMBH	DE
Linezolid 600 mg film-coated tablets	NL/H/3011/001	PL 04569/1640	GENERICS [UK] LIMITED	UK
Linezolid Mylan 600 mg, Filmtabletten	NL/H/3011/001	BE471280	MYLAN BVBA/SPRL	BE
Linezolid Mylan 600 mg, filmomhulde tabletten	NL/H/3011/001	RVG 114163	MYLAN B.V.	NL
Linezolid Mylan 600 mg comprimidos recubiertos con película EFG	NL/H/3011/001	79834	MYLAN DURA GMBH	ES
Linezolid Mylan Pharma 600 mg compresse rivestite con film	NL/H/3011/001	043621022	MYLAN S.P.A.	IT
Linézolide Mylan Pharma 600 mg, comprimé pelliculé	NL/H/3011/001	NL43789	MYLAN S.A.S	FR
Linezolid 2mg/ml Solution for Infusion	AT/H/0564/001	PL 30306/0640	ACTAVIS GROUP PTC EHF.	UK
Linezolid Actavis 2 mg/ml Infusionslösung	AT/H/0564/001	136711	ACTAVIS GROUP PTC EHF.	AT
Anozilad 2 mg/ml oldatos infúzió	AT/H/0564/001	OGYI-T-22617/13	ACTAVIS GROUP PTC EHF.	HU
Linezolid Actavis 2 mg/ml soluţie perfuzabilă	AT/H/0564/001	8569/2016/01-08	ACTAVIS GROUP PTC EHF.	RO
Линезолид Актавис 2 mg/ml инфузионен разтвор	AT/H/0564/001	20160055	ACTAVIS GROUP PTC EHF.	BG
Linezolid Actavis 2 mg/ml infuzní roztok	AT/H/0564/001	15/013/16-C	ACTAVIS GROUP PTC EHF.	CZ
Linezolid Actavis 2 mg/ml otopina za infuziju	AT/H/0564/001	HR-H-898922487	ACTAVIS GROUP PTC EHF.	HR
Linezolid Actavis 2 mg/ml	AT/H/0564/001	H/16/02262/008	ACTAVIS GROUP PTC EHF.	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
raztopina za infundiranje				
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/005	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/002	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/007	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/003	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/004	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/006	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis 2 mg/ml raztopina za infundiranje	AT/H/0564/001	H/16/02262/001	ACTAVIS GROUP PTC EHF.	SI
Linezolid Actavis	AT/H/0564/001/DC	54645	ACTAVIS GROUP PTC EHF.	DK
ANOZILAD	AT/H/0564/001/DC	23237	ACTAVIS GROUP PTC EHF.	PL
Anozilad 2 mg/ml oldatos infúzió	AT/H/0564/001/DC	OGYI-T-22617/14	ACTAVIS GROUP PTC EHF.	HU
Linezolide Ohre Pharma 2 mg/ml, solution pour perfusion	AT/H/0563/001/DC	34009 550 201 8 9	OHRE PHARMA	FR
Linezolide Ohre Pharma 2 mg/ml, solution pour perfusion	AT/H/0563/001/DC	34009 550 202 0 2	OHRE PHARMA	FR
Linezolide Ohre Pharma 2 mg/ml, solution pour perfusion	AT/H/0563/001/DC	34009 550 202 2 6	OHRE PHARMA	FR
Linezolid 600 mg Film-Coated Tablets	UK/H/5978/001	PL 17907/0525	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Linezolid axcount 600 mg Filmtabletten	UK/H/5978/001	94156.00.00	AXCOUNT GENERIKA GMBH	DE
Linezolid "Orion", filmovertrukne tabletter	FI/H/0909/001	57207	ORION CORPORATION	DK
Linezolid Orion 600 mg, filmdragerade tabletter	FI/H/0909/001	54250	ORION CORPORATION	SE
Linezolid Orion 600 mg tabletter, filmdrasjerte	FI/H/0909/001	15-11019	ORION CORPORATION	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Orion 600 mg kalvopäällysteiset tabletit	FI/H/0909/001	33912	ORION CORPORATION	FI
Linezolid Orion 600 mg filmdragerade tabletter	FI/H/0909/001	33912	ORION CORPORATION	FI
Linezolid Hetero 2 mg/ml Infusionslösung	NL/H/2776/002	95874.00.00	HETERO EUROPE S.L.	DE
Linezolid Hetero 2 mg/ml, oplossing voor infusie	NL/H/2776/002	RVG 117819	HETERO EUROPE S.L.	NL
Linezolid Krka 2 mg/ml otopina za infuziju	AT/H/0639/001	HR-H-611386157	KRKA-FARMA D.O.O.	HR
Linezolid Krka, 2 mg/ml infusioonilahus	AT/H/0639/001	924916	KRKA, D.D., NOVO MESTO	EE
Linezolid Krka 2 mg/ml infúzny roztok	AT/H/0639/001	15/0482/16-S	KRKA, D.D., NOVO MESTO	SK
Linezolid Krka 2 mg/ml – Infusionslösung	AT/H/0639/001	137292	KRKA, D.D., NOVO MESTO	AT
Linezolid Krka 2 mg/ml šķīdums infūzijām	AT/H/0639/001	16-0225	KRKA, D.D., NOVO MESTO	LV
Linezolid Krka 2 mg/ml infuzinis tirpalas	AT/H/0639/001	LT/1/16/4014/001	KRKA, D.D., NOVO MESTO	LT
Linezolid Krka 2 mg/ml infuzinis tirpalas	AT/H/0639/001	LT/1/16/4014/002	KRKA, D.D., NOVO MESTO	LT
Линезолид Крка 2 mg/ml инфузионен разтвор	AT/H/0639/001	20170003	KRKA, D.D., NOVO MESTO	BG
Linezolid Krka 2 mg/ml oldatos infúzió	AT/H/0639/001	OGYI-T-22928/04	KRKA, D.D., NOVO MESTO	HU
Linezolid Krka 2 mg/ml oldatos infúzió	AT/H/0639/001	OGYI-T-22928/05	KRKA, D.D., NOVO MESTO	HU
Linezolid Krka 2 mg/ml solución para perfusión EFG	AT/H/0639/001	81634	KRKA, D.D., NOVO MESTO	ES
Linezolid TAD® 2 mg/ml Infusionslösung	AT/H/0639/001	95668.00.00	TAD PHARMA GMBH	DE
Linezolid Krka 2 mg/ml infuzní roztok	AT/H/0639/001	15/708/15-C	KRKA, D.D., NOVO MESTO	CZ
Linezolid Krka 2 mg/ml	AT/H/0639/001	H/17/02293/001	KRKA, D.D., NOVO MESTO	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
raztopina za infundiranje				
Linezolid Krka 2 mg/ml raztopina za infundiranje	AT/H/0639/001	H/17/02293/002	KRKA, D.D., NOVO MESTO	SI
Linezolid Krka 2 mg/ml soluţie perfuzabilă	AT/H/0639/001	9707/2017/01	KRKA, D.D., NOVO MESTO	RO
Linezolid Krka 2 mg/ml soluţie perfuzabilă	AT/H/0639/001	9707/2017/02	KRKA, D.D., NOVO MESTO	RO
Linezolida Krka 2 mg/ml solução para perfusão	AT/H/0639/001	5713631	KRKA, D.D., NOVO MESTO	PT
Linezolid Krka 2 mg/ml roztwór do infuzji	AT/H/0639/001	23685	KRKA, D.D., NOVO MESTO	PL
Linezolid 2 mg/ml solution for infusion	AT/H/0639/001	PA1347/057/002	KRKA, D.D., NOVO MESTO	IE
Linezolid Krka 2 mg/ml soluzione per infusione	AT/H/0639/001	AIC N. 044463014	KRKA, D.D., NOVO MESTO	IT
Linezolid Krka 2 mg/ml soluzione per infusione	AT/H/0639/001	AIC N. 044463026	KRKA, D.D., NOVO MESTO	IT
Linezolid Krka 2 mg/ml Solution for Infusion	AT/H/0639/001	PL 01656/0213 - 0001	KRKA, D.D., NOVO MESTO	UK
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 855 8 5	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 855 9 2	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 0 8	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 1 5	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 2 2	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé	not available	34009 300 856 3 9	PHARMAKI GENERICS LTD	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
pelliculé				
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 4 6	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 5 3	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 6 0	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 7 7	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 856 9 1	PHARMAKI GENERICS LTD	FR
LINEZOLIDE PHARMAKI GENERICS 600 mg, comprimé pelliculé	not available	34009 300 857 0 7	PHARMAKI GENERICS LTD	FR
Linezolid 600 mg Film-Coated Tablets	UK/H/5978/001	PL 17907/0525	BRISTOL LABORATORIES LTD (BERKHAMSTED)	UK
Linezolid axcount 600 mg Filmtabletten	UK/H/5978/001	94156.00.00	AXCOUNT GENERIKA GMBH	DE
Linezolida Pharmakern, 600 mg, comprimidos revestidos por película	not available	5681937	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Linezolid Betapharm 2 mg/ml, oplossing voor infusie	NL/H/3529/001	RVG 116997	BETAPHARM ARZNEIMITTEL GMBH	NL
Linezolid beta 2 mg/ml Infusionslösung	NL/H/3529/001	95108.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Linezolid Dr. Reddys 2 mg/ml solución para perfusión EFG	NL/H/3529/001	80583	DR. REDDY'S LABORATORIES (UK) LTD.	ES
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504013	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504025	DR. REDDY'S LABORATORIES (UK) LTD.	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
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504037	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504052	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504049	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504064	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504076	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504088	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504090	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504102	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504126	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504114	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504138	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504140	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504165	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504177	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504189	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504191	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504227	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml	NL/H/3529/001	044504215	DR. REDDY'S LABORATORIES (UK)	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			LTD.	
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504203	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504239	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	044504241	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolid Dr. Reddy's 2 mg/ml soluzione per infusione	NL/H/3529/001	04450415 3	DR. REDDY'S LABORATORIES (UK) LTD.	IT
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 205 9 2	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 1 5	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 2 2	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 3 9	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 4 6	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 5 3	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 6 0	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 7 7	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 9 1	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 207 0 7	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 205 8 5	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolide Dr. Reddy´s 2 mg/ml, Solution pour perfusion	NL/H/3529/001	34009 550 206 8 4	DR. REDDY'S LABORATORIES (UK) LTD.	FR
Linezolid Denk 2mg/ml Infusionslösung	NL/H/2780/001	94451.00.00	DENK PHARMA GMBH & CO. KG	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid "Glenmark", filmovertrukne tabletter	PT/H/1245/001	54065	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DK
Linezolid 600mg Film-coated Tablets	PT/H/1245/001	PL 25258/0155	GLENMARK PHARMACEUTICALS EUROPE LIMITED	UK
Linezolid Glenmark 600 mg Filmtabletten	PT/H/1245/001	92186.00.00	GLENMARK PHARMACEUTICALS EUROPE LIMITED	DE
Linezolid Glenmark 600 mg filmdragerade tabletter	PT/H/1245/001	51059	GLENMARK PHARMACEUTICALS EUROPE LIMITED	SE
Dilizolen, 600 mg, tabletki powlekane	PT/H/1245/001	22858	GLENMARK PHARMACEUTICALS S.R.O.	PL
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687470	GLENMARK PHARMACEUTICALS EUROPE LIMITED	PT
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687504	GLENMARK PHARMACEUTICALS EUROPE LIMITED	PT
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687512	GLENMARK PHARMACEUTICALS EUROPE LIMITED	PT
Linezolid Glenmark 600 mg comprimidos recubiertos con película EFG	PT/H/1245/001	81569	GLENMARK PHARMACEUTICALS EUROPE LIMITED	ES
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	UK/H/5511/001	7669/2015/01	INFOMED FLUIDS SRL	RO
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	UK/H/5511/001	7669/2015/02	INFOMED FLUIDS SRL	RO
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	UK/H/5511/001	7669/2015/03	INFOMED FLUIDS SRL	RO
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	UK/H/5511/001	34009 550 065 2 7	PANMEDICA	FR
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	UK/H/5511/001	34009 550 065 3 4	PANMEDICA	FR
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	UK/H/5511/001	34009 550 065 4 1	PANMEDICA	FR
Linezolid 2 mg/ml Solution for Infusion	UK/H/5511/001	PL 34328/0012	PANMEDICA	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
Linezolid 2 mg/ml Solution for Infusion	UK/H/5511/001	PL 34328/0012	PANMEDICA	UK
Linezolid 2 mg/ml Solution for Infusion	UK/H/5511/001	PL 34328/0012	PANMEDICA	UK
LINEZOLID ROTEXMEDICA 2 mg/ml Infusionslösung	UK/H/5511/001	91742.00.00	PANMEDICA	DE
LINEZOLID ROTEXMEDICA 2 mg/ml Infusionslösung	UK/H/5511/001	91742.00.00	PANMEDICA	DE
LINEZOLID ROTEXMEDICA 2 mg/ml Infusionslösung	UK/H/5511/001	91742.00.00	PANMEDICA	DE
LINEZOLID PANPHARMA 2 mg/ml Infusionslösung	UK/H/5511/001	136408	PANMEDICA	AT
LINEZOLID PANPHARMA 2 mg/ml Infusionslösung	UK/H/5511/001	136408	PANMEDICA	AT
LINEZOLID PANPHARMA 2 mg/ml Infusionslösung	UK/H/5511/001	136408	PANMEDICA	AT
LINEZOLID G.E.S. 2 mg/ml Solución para perfusión EFG	UK/H/5511/DC	80258	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	ES
Linezolid Infomed 2 mg/ml Soluzione per Infusione	UK/H/5511/001	043331014	INFOMED FLUIDS SRL	IT
Linezolid Infomed 2 mg/ml Soluzione per Infusione	UK/H/5511/001	043331026	INFOMED FLUIDS SRL	IT
Linezolid Infomed 2 mg/ml Soluzione per Infusione	UK/H/5511/001	043331038	INFOMED FLUIDS SRL	IT
Linezolid Infomed, 2 mg/ml, roztwór do infuzji	UK/H/5511/001	22918	INFOMED FLUIDS SRL	PL
LINEZOLIDA G.E.S. 2 mg/ml Soluçao para perfusao	not available	5712773	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Линезолид-Чайкафарма 2 mg/ml инфузионен разтвор	not available	20160325	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG
Natlinez 2 mg/ml, oplossing voor infusie	UK/H/5167/001	RVG 114369	HOSPIRA BENELUX BVBA	NL
Linezolid Fresenius Kabi 2 mg/ml, oplossing voor infusie	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE

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mg/ml, solution pour perfusion				
Linezolid Fresenius Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE
Linezolid Kabi 2 mg/ml solución para perfusión EFG	PT/H/1090/001	79295	FRESENIUS KABI ESPAÑA S.A.U.	ES
Linezolid Fresenius Kabi 2 mg/ml infusjonsvæske, oppløsning	PT/H/1090/001	13-9804	FRESENIUS KABI NORGE AS	NO
Linezolid Fresenius Kabi 2 mg/ml, oplossing voor infusie	PT/H/1090/001	RVG 114296	FRESENIUS KABI NEDERLAND B.V.	NL
Linezolid Kabi 2 mg/ml otopina za infuziju	PT/H/1090/001	HR-H-690522814	FRESENIUS KABI D.O.O.	HR
Linezolid Kabi 2 mg/ml soluţie perfuzabilă	PT/H/1090/001	7392/2015/01	S.C. FRESENIUS KABI ROMÂNIA S.R.L.	RO
Linezolid Kabi 2 mg/ml solutie perfuzabila	PT/H/1090/001	7392/2015/02	S.C. FRESENIUS KABI ROMÂNIA S.R.L.	RO
Linezolid Kabi 2 mg/ml solutie perfuzabila	PT/H/1090/001	7392/2015/03	S.C. FRESENIUS KABI ROMÂNIA S.R.L.	RO
Linezolid 2 mg/ml Solution for infusion	PT/H/1090/001	PL 08828/0256	FRESENIUS KABI LIMITED	UK
Linezolid 2 mg/ml Solution for infusion	PT/H/1090/001	PA 566/069/001	FRESENIUS KABI LIMITED	IE
Linezolid Kabi 2 mg/ml infuzní roztok	PT/H/1090/001	15/340/14-C	FRESENIUS KABI S.R.O	CZ
Linezolid Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	135944	FRESENIUS KABI AUSTRIA GMBH	AT
Linezolid "Fresenius Kabi", infusionsvæske, opløsning	PT/H/1090/001	53133	FRESENIUS KABI AB	DK
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5644414	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5644422	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5644430	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolid Kabi, 2 mg/ml,	PT/H/1090/001	22370	FRESENIUS KABI POLSKA SP. Z O.O.	PL

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roztwór do infuzji				
Linezolid Kabi 2 mg/ml infúzny roztok	PT/H/1090/001	15/0128/15-S	FRESENIUS KABI S.R.O	SK
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5657838	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5657846	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5657853	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Linezolid Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	90853.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/001	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/002	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/003	FRESENIUS KABI DEUTSCHLAND GMBH	SI
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 036 5 6	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 036 6 3	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 090 9 2	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 091 0 8	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 091 1 5	FRESENIUS KABI FRANCE S.A.S.	FR
LINEZOLIDE KABI 2 mg/ml, solution pour perfusion	PT/H/1090/001	34009 550 036 4 9	FRESENIUS KABI FRANCE S.A.S.	FR
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113012	FRESENIUS KABI ITALIA S.R.L.	IT
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113024	FRESENIUS KABI ITALIA S.R.L.	IT
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113036	FRESENIUS KABI ITALIA S.R.L.	IT

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Linezolid Kabi, 2 mg/ml διάλυμα για έγχυση	PT/H/1090/001	83407/24-11-2015	FRESENIUS KABI HELLAS A.E.	GR
Linezolid Fresenius Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	BE483066	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2 mg/ml, oplossing voor infusie	PT/H/1090/001	BE483066	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2 mg/ml, solution pour perfusion	PT/H/1090/001	BE483066	FRESENIUS KABI NV/SA	BE
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113048	FRESENIUS KABI ITALIA S.R.L.	IT
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113051	FRESENIUS KABI ITALIA S.R.L.	IT
Linezolid Kabi 2 mg/ml soluzione per infusione	PT/H/1090/001	043113063	FRESENIUS KABI ITALIA S.R.L.	IT
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	7392/2015/04	FRESENIUS KABI ROMANIA S.R.L.	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	7392/2015/05	FRESENIUS KABI ROMANIA S.R.L.	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	7392/2015/06	FRESENIUS KABI ROMANIA S.R.L.	RO
Линезолид Каби 2 mg/ml инфузионен разтвор	PT/H/1090/001	20170221	FRESENIUS KABI BULGARIA EOOD	BG
Linezolid Adamed, 2 mg/ml, roztwór do infuzji	NL/H/2870/001	22061	ADAMED	PL
Ziloxon 2 mg/ml innrennslislyf, lausn	NL/H/2870/001	IS/1/14/006/01	WILLIAMS & HALLS EHF	IS
Gramposimide 2 mg/ml, oplossing voor infusie	NL/H/2870/001	RVG 113203	SYNTHON BV	NL
Linezolid Polpharma 600 mg, filmomhulde tabletten	NL/H/3124/001	RVG 114804	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	NL
LINEZA 600 mg potahované tablety	NL/H/3124/001	15/373/15-C	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	CZ
Linezolid Polpharma, 600 mg, tabletki powlekane	NL/H/3124/001	22745	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	PL
Linezolid Polpharma 600 mg	NL/H/3124/001	15-0281	ZAKLADY FARMACEUTYCZNE	LV

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apvalkotās tabletes			"POLPHARMA" SPOLKA AKCYJNA	
Линезолид Полфарма 600 mg филмирани таблетки	NL/H/3124/001	20150406	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	BG
LINEZA 600 mg filmom obalené tablety	NL/H/3124/001	15/0531/15-S	ZAKLADY FARMACEUTYCZNE "POLPHARMA" SPOLKA AKCYJNA	SK
Linezolid Pharmathen 600 mg Filmtabletten	AT/H/0491/001	135817	PHARMATHEN S.A.	AT
Lizedia 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	AT/H/0491/001	67953/9-10-2015	PHARMATHEN S.A.	GR
Livegramide 2 mg/ml, oplossing voor infusie	not available	RVG 117013	SYNTHON BV	NL
Linezolid Amneal 2 mg/ml solución para perfusión EFG	NL/H/3532/001/MR	81.328	AMNEAL PHARMA EUROPE LIMITED	ES
Linezolid 2 mg/ml solution for infusion	ES/H/0348/001	PL 03551/0147	B.BRAUN MELSUNGEN AG	UK
Linezolida B. Braun 2 mg/ml solução para perfusão	ES/H/0348/001	5711239	B.BRAUN MELSUNGEN AG	PT
Linezolida B. Braun 2 mg/ml solução para perfusão	ES/H/0348/001	5711221	B.BRAUN MELSUNGEN AG	PT
Linezolid B. Braun 2 mg/ml solución para perfusión EFG	ES/H/0348/001	81866	B.BRAUN MELSUNGEN AG	ES
LINEZOLIDA G.E.S. 2 mg/ml Soluçao para perfusao	not available	5712773	G.E.S. GENÉRICOS ESPAÑOLES LABORATORIO, S.A.	PT
Linezolid Teva 2 mg/ml solución para perfusión EFG	DK/H/1781/001	75868	TEVA PHARMA S.L.U	ES
ZOLINID 2 mg/ml solution for infusion	DK/H/1781/001	20110241	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Linezolid Teva 2 mg/ml infuzní roztok	DK/H/1781/001	15/457/11-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Linozid, infusionsvæske, opløsning	DK/H/1781/001	45970	TEVA DENMARK A/S	DK
Linezolid 2 mg/ml Solution for Infusion	DK/H/1781/001	PA0749/113/001	TEVA PHARMA B.V.	IE
Linezolid Teva 2 mg/ml soluţie perfuzabilă	DK/H/1781/001	8559/2016/01-06	TEVA PHARMACEUTICALS S.R.L	RO

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ЗОЛИНИД 2 mg/ml инфузионен разтвор	DK/H/1781/001	20110241	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Linezolid Pfizer 100 mg/5 ml Granules for Oral Suspension	UK/H/5515/003	PL 00057/1503	PFIZER LIMITED	UK
Linezolid Pfizer 600 mg film-coated tablets	UK/H/5515/002	PL 00057/1502	PFIZER LIMITED	UK
Linezolid Pfizer 2 mg/ml solution for infusion	UK/H/5515/001	PL 00057/1501	PFIZER LIMITED	UK
Linezolid Pfizer 600 mg Filmtabletten	UK/H/5515/002	136048	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Linezolid Pfizer 2 mg/ml Infusionslösung	UK/H/5515/001	136007	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Linezolid Pfizer100 mg/5 ml Granulat zur Herstellung einer Suspension zum Einnehmen	UK/H/5515/003	136009	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Linezolid Pfizer 600 mg filmomhulde tabletten	UK/H/5515/002	RVG 114541	PFIZER B.V.	NL
Linezolid Pfizer 100 mg/5 ml granulaat voor orale suspensie	UK/H/5515/003	RVG 114543	PFIZER B.V.	NL
Linezolid Pfizer 2 mg/ml oplossing voor infusie	UK/H/5515/001	RVG 114540	PFIZER B.V.	NL
Linezolid Pfizer 600 mg film-coated tablets	UK/H/5515/002	PA 0822/181/02	PFIZER HEALTHCARE IRELAND	IE
LINEZOLIDE PFIZER 100 mg/5 ml, granulés pour suspension buvable	UK/H/5515/003	34009 550 140 5 8	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 52	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 140 3 4	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 139 9 0	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 7 6	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 3 8	PFIZER HOLDING FRANCE	FR

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LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 140 1 0	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 6 9	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 2 1	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 140 2 7	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 100 mg/5 ml, granulés pour suspension buvable	UK/H/5515/003	34009 550 140 4 1	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 140 0 3	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 600 mg, comprimé pelliculé	UK/H/5515/002	34009 550 139 4 5	PFIZER HOLDING FRANCE	FR
LINEZOLIDE PFIZER 2 mg/ml, solution pour perfusion	UK/H/5515/001	34009 550 139 8 3	PFIZER HOLDING FRANCE	FR
Linezolid Pfizer 100 mg/5 ml granules for oral suspension	UK/H/5515/003	PA 0822/181/03	PFIZER HEALTHCARE IRELAND	IE
Linezolid Pfizer 2 mg/ml solution for infusion	UK/H/5515/001	PA 0822/181/01	PFIZER HEALTHCARE IRELAND	IE
Linezolid/Pfizer 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	UK/H/5515/002	32870/13-04-2016	PFIZER HELLAS, A.E.	GR
Linezolid/Pfizer 2 mg/ml διάλυμα για ενδοφλέβια έγχυση	UK/H/5515/001	32869/13-04-2016	PFIZER HELLAS, A.E.	GR
Linezolid Farmaprojects 2mg/ml, oplossing voor infusie	NL/H/2780/001	RVG 112519	FARMAPROJECTS S.A.U.	NL