11 December 2025
EMADOC-1700519818-2864496
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): linezolid

EURD list No. PSUSA/00001888/202504

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
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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
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 100 mg/5 ml - Granulat zur Herstellung einer Suspension zum Einnehmen	IE/H/0644/003	1-24230	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid® 600 mg - Filmtabletten	IE/H/0644/002	1-24229	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Zyvoxid 2 mg/ml - Infusionslösung	IE/H/0644/001	1-24227	PFIZER CORPORATION AUSTRIA GES.M.B.H.	AT
Linezolid HCS 600 mg Filmtabletten	AT/H/0625/001	136603	HCS BVBA	AT
Linezolid Sandoz 2 mg/ml - Infusionslösung	HR/H/0263/001	136402	SANDOZ GMBH	AT
Linezolid 1A Pharma 600 mg - Filmtabletten	NL/H/2966/001	141447	1A PHARMA GMBH	AT
Ilenozyd 600 mg Filmtabletten	AT/H/0620/001	136602	KRKA, D.D., NOVO MESTO	AT
Linezolid STADA 600 mg Filmtabletten	DE/H/3490/001	135157	STADA ARZNEIMITTEL GMBH	AT
Linezolid Accord 600 mg Filmtabletten	NL/H/4545/001	136631	ACCORD HEALTHCARE B.V.	AT
Linezolid Hikma 2 mg/ml Infusionslösung	AT/H/1082/001	141619	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	135944	FRESENIUS KABI AUSTRIA GMBH	AT
Linezolid Pharmathen 600 mg Filmtabletten	AT/H/0491/001	135817	PHARMATHEN S.A.	AT
Linezolid Krka 2 mg/ml – Infusionslösung	AT/H/0639/001	137292	KRKA, D.D., NOVO MESTO	AT
Linezolid Sandoz 600 mg – Filmtabletten	NL/H/2965/001	135752	SANDOZ GMBH	AT
Linezolid Accord 2 mg/ml Infusionslösung	PT/H/1195/001	138023	ACCORD HEALTHCARE B.V.	AT
Linezolid ratiopharm 600 mg Filmtabletten	NL/H/2945/001	136042	TEVA B.V	AT
Linezolid ratiopharm 600 mg Filmtabletten	NL/H/2945/001	136042	TEVA B.V	AT
Zyvoxid 2 mg/ml solution pour perfusion	IE/H/0644/001	BE226651	PFIZER S.A.	BE
Zyvoxid 600 mg comprimés pelliculés	IE/H/0644/002	BE228313	PFIZER S.A.	BE
Zyvoxid 2 mg/ml Infusionslösung	IE/H/0644/001	BE397713	PFIZER S.A.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg filmomhulde tabletten	IE/H/0644/002	BE228304	PFIZER S.A.	BE
Zyvoxid 100 mg/5 ml granulaat voor orale suspensie	IE/H/0644/003	BE228322	PFIZER S.A.	BE
Linezolid Viatris 600 mg, Filmtabletten	NL/H/3011/001	BE471280	VIATRIS GX	BE
Linezolid Viatris 600 mg, comprimés pelliculés	NL/H/3011/001	BE471280	VIATRIS GX	BE
Linezolid Viatris 600 mg, filmomhulde tabletten	NL/H/3011/001	BE471280	VIATRIS GX	BE
Linezolid Fresenius Kabi 2 mg/ml, oplossing voor infusie	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2 mg/ml, solution pour perfusion	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	BE465973	FRESENIUS KABI NV/SA	BE
Linezolid Fresenius Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	BE483066	FRESENIUS KABI NV/SA	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 Sandoz 600 mg filmomhulde tabletten	NL/H/2965/001	BE460444	SANDOZ N.V.	BE
Linezolid Accord 2 mg/ml solution pour perfusion.	PT/H/1195/001	BE522062	ACCORD HEALTHCARE B.V.	BE
Linezolid Accord 2 mg/ml oplossing voor infusie.	PT/H/1195/001	BE522062	ACCORD HEALTHCARE B.V.	BE
Линезолид Крка 600 mg филмирани таблетки	AT/H/0620/001	20150383	KRKA, D.D., NOVO MESTO	BG
Linezolid Kabi 2 mg/ml solution for infusion	PT/H/1090/001	20170221	FRESENIUS KABI BULGARIA EOOD	BG
Линезолид Полфарма 2 mg/ml инфузионен разтвор	NL/H/2781/001	20150377	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	BG
Линезолид Крка 2 mg/ml инфузионен разтвор	AT/H/0639/001	20170003	KRKA, D.D., NOVO MESTO	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
Линестад 2 mg/ml инфузионен разтвор	not available	20150407	STADA ARZNEIMITTEL AG	BG
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	023036	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	023035	PFIZER HELLAS A.E.	CY
Linezid 2 mg/ml διάλυμα για ενδοφλέβια έγχυση	not available	023163	SAPIENS PHARMACEUTICALS LTD	CY
Linezolid Accord 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/4545/001	022370	ACCORD HEALTHCARE S.L.U.	CY
Linezolid Accord 2 mg/ml Διάλυμα για έγχυση.	PT/H/1195/001	022785	ACCORD HEALTHCARE S.L.U.	CY
ZYVOXID 600 mg potahované tablety	not available	15/068/02-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 600 mg potahované tablety	not available	15/068/02-C	PFIZER SPOL. S R.O.	CZ
ZYVOXID 600 mg potahované tablety	not available	15/068/02-C	PFIZER SPOL. S R.O.	CZ
ZYVOXID 2 mg/ml infuzní roztok	not available	15/069/02-C	PFIZER SPOL. S R.O.	CZ
ZYVOXID 2 mg/ml infuzní roztok	not available	15/069/02-C	PFIZER SPOL. S R.O.	CZ
ZYVOXID 600 mg potahované tablety	not available	15/068/02-C	PFIZER SPOL. S R.O.	CZ
Linezolid Olikla 600 mg potahované tablety	not available	15/864/16-C	OLIKLA S.R.O.	CZ
Linezolid Krka 600 mg potahované tablety	AT/H/0620/001	15/532/15-C	KRKA, D.D., NOVO MESTO	CZ
Linezolid Accord 600 mg potahované tablety	NL/H/4545/001	15/561/15-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Linezolid Kabi 2 mg/ml infuzní roztok	PT/H/1090/001	15/340/14-C	FRESENIUS KABI S.R.O.	CZ
Linezolid Krka 2 mg/ml infuzní roztok	AT/H/0639/001	15/708/15-C	KRKA, D.D., NOVO MESTO	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
Linezolid Sandoz 600 mg potahované tablety	NL/H/2965/001	15/300/14-C	SANDOZ S.R.O.	CZ
Linezolid Accord 2 mg/ml infuzní roztok	PT/H/1195/001	15/126/17-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Linezolid Olikla 2 mg/ml infuzní roztok	not available	15/739/16-C	OLIKLA S.R.O.	CZ
ZYVOXID® 2 mg/ml Infusionslösung	IE/H/0644/001	51712.00.00	PFIZER PHARMA GMBH	DE
ZYVOXID® 100 mg/5 ml Granulat zur Herstellung einer Suspension zum Einnehmen	IE/H/0644/003	51712.00.02	PFIZER PHARMA GMBH	DE
ZYVOXID® 600 mg Filmtabletten	IE/H/0644/002	51712.01.01	PFIZER PHARMA GMBH	DE
Linezolid TAD® 600 mg Filmtabletten	AT/H/0625/001	94723.00.00	TAD PHARMA GMBH	DE
Linezolid PUREN 600 mg Filmtabletten	AT/H/0491/001	89986.00.00	PUREN PHARMA GMBH & CO. KG	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	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 Zentiva® 600 mg Filmtabletten	DE/H/7727/001	89987.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid Glenmark 600 mg Filmtabletten	PT/H/1245/001	92186.00.00	GLENMARK ARZNEIMITTEL GMBH	DE
Linezolid STADA® 2 mg/ml Infusionslösung	DE/H/5027/001	92193.00.00	STADAPHARM GMBH	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid Denk 600 mg Filmtabletten	not available	2200044.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid Inresa 2 mg/ml Infusionslösung	DE/H/4765/001	94423.00.00	HELM PHARMACEUTICALS GMBH	DE
Linezolid AL 600 mg Filmtabletten	DE/H/3490/001	86422.00.00	ALIUD PHARMA GMBH	DE
Linezolid Dr. Eberth 2 mg/ml Infusionslösung	DE/H/7426/001/DC	7008401.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	DE
Linezolid PUREN 2 mg/ml Infusionslösung	PT/H/1629/001	97163.00.00	EUGIA PHARMA (MALTA) LTD	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 Accord 600 mg Filmtabletten	NL/H/4545/001	92264.00.00	ACCORD HEALTHCARE B.V.	DE
Linezolid Hikma 2 mg/ml Infusionslösung	AT/H/1082/001	2205922.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Linezolid Panpharma 2 mg/ml Infusionslösung Zur Anwendung bei Erwachsenen	DE/H/6119/001	91742.00.00	PANMEDICA	DE
Linezolid Panpharma 2 mg/ml Infusionslösung Zur Anwendung bei Erwachsenen	DE/H/6119/001	91742.00.00	PANMEDICA	DE
Linezolid Panpharma 2 mg/ml Infusionslösung Zur Anwendung bei Erwachsenen	DE/H/6119/001	91742.00.00	PANMEDICA	DE
Linezolid STADA® 600 mg Filmtabletten	DE/H/3489/001	86421.00.00	STADAPHARM GMBH	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	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 Amarox 600 mg Filmtabletten	DE/H/6978/001	93756.00.00	AMAROX PHARMA GMBH	DE
Linezolid Ascend 600 mg Filmtabletten	DE/H/6178/001	99427.00.00	ASCEND GMBH	DE
Linezolid Kabi 2 mg/ml Infusionslösung	PT/H/1090/001	90853.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Linezolid TAD® 2 mg/ml Infusionslösung	AT/H/0639/001	95668.00.00	TAD PHARMA GMBH	DE
Linezolid – 1 A Pharma 600 mg Filmtabletten	NL/H/2965/001	90203.00.00	1 A PHARMA GMBH	DE
Linezolid Accord 2 mg/ml Infusionslösung	PT/H/1195/001	2200088.00.00	ACCORD HEALTHCARE B.V.	DE
Linezolid Zentiva® 2 mg/ml Infusionslösung	DE/H/7979/001	86554.00.00	ZENTIVA PHARMA GMBH	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid Denk 2 mg/ml Infusionslösung	not available	2200045.00.00	DENK PHARMA GMBH & CO. KG	DE
Linezolid-ratiopharm® 600 mg Filmtabletten	NL/H/2945/001	90497.00.00	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
Linezolid-ratiopharm® 600 mg Filmtabletten	NL/H/2945/001	90497.00.00	RATIOPHARM GMBH	DE
Linezolid Demo 2 mg/ml Infusionslösung	PT/H/1143/001	90655.00.00	DEMO ABEE	DE
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, granulat til oral suspension	IE/H/0644/003	32327	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, granulat til oral suspension	IE/H/0644/003	32327	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK

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, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, granulat til oral suspension	IE/H/0644/003	32327	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK

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, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK

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, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, filmovertrukne tabletter	IE/H/0644/002	32326	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Zyvoxid, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK

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, infusionsvæske, opløsning	IE/H/0644/001	32324	PFIZER APS	DK
Linezolid ”Krka”, filmovertrukne tabletter	AT/H/0625/001	55654	KRKA, D.D., NOVO MESTO	DK
Linezolid ”Glenmark”, filmovertrukne tabletter	PT/H/1245/001	54065	GLENMARK ARZNEIMITTEL GMBH	DK
Linezolid "Accord", filmovertrukne tabletter	NL/H/4545/001	54073	ACCORD HEALTHCARE B.V.	DK
Linezolid "Accord", filmovertrukne tabletter	NL/H/4545/001	54073	ACCORD HEALTHCARE B.V.	DK
Linezolid Fresenius Kabi	PT/H/1090/001	53133	FRESENIUS KABI AB	DK
Linezolid ”Sandoz”, filmovertrukne tabletter	NL/H/2965/001	52754	SANDOZ A/S	DK
Linezolid "Accord", infusionsvæske, opløsning	PT/H/1195/001	59558	ACCORD HEALTHCARE B.V.	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK

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 B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK

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 B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK

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 B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Teva filmovertrukne tabletter	NL/H/2945/001	53077	TEVA B.V	DK
Linezolid Krka 600 mg õhukese polümeerikattega tabletid	AT/H/0620/001	893315	KRKA, D.D., NOVO MESTO	EE
Linezolid Accord 600 mg õhukese polümeerikattega tabletid	NL/H/4545/001	893215	ACCORD HEALTHCARE B.V.	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Krka 2 mg/ml infusioonilahus	AT/H/0639/001	924916	KRKA, D.D., NOVO MESTO	EE
Linezolid Accord 2 mg/ml infusioonilahus	PT/H/1195/001	958618	ACCORD HEALTHCARE B.V.	EE
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 600 mg comprimidos recubiertos con película	IE/H/0644/002	64.109	PFIZER S.L.	ES
Zyvoxid 100 mg/5 ml granulado para suspensión oral	IE/H/0644/003	64.107	PFIZER S.L.	ES
Zyvoxid 100 mg/5 ml granulado para suspensión oral	IE/H/0644/003	64.107	PFIZER S.L.	ES
Zyvoxid 100 mg/5 ml granulado para suspensión oral	IE/H/0644/003	64.107	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml solución para perfusión	IE/H/0644/001	64.106	PFIZER S.L.	ES
Linezolid Aurovitas Spain 600 mg comprimidos recubiertos con película EFG	AT/H/0491/001	79.231	AUROVITAS SPAIN,S.A.U.	ES
Linezolid Krka 600 mg comprimidos recubiertos con película EFG	AT/H/0620/001	80940	KRKA, D.D., NOVO MESTO	ES
Linezolid Kern Pharma 600 mg comprimidos recubiertos con película EFG	not available	84.474	KERN PHARMA, S.L.	ES
Linezolid Glenmark 600 mg comprimidos recubiertos con película EFG	PT/H/1245/001	81569	GLENMARK ARZNEIMITTEL GMBH	ES
Linezolid Normon 600 mg comprimidos recubiertos con película EFG	ES/H/0291/001	80368	LABORATORIOS NORMON, S.A.	ES
Linezolid Normon 2 mg/ml solución para perfusión EFG	PT/H/1220/001	80679	LABORATORIOS NORMON, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Aurovitas 2 mg/ml solución para perfusión EFG	PT/H/1629/001	83.053	EUGIA PHARMA (MALTA) LTD	ES
Linezolid Hikma 2 mg/ml solución para perfusión EFG	AT/H/1082/001	87990	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	ES
Linezolid Altan 2 mg/ml solución para perfusión EFG	DE/H/6119/001	80258	ALTAN PHARMACEUTICALS S.A.	ES
Linezolid Kabi 2 mg/ml solución para perfusión EFG	PT/H/1090/001	79295	FRESENIUS KABI ESPAÑA S.A.U.	ES
Linezolid B. Braun 2 mg/ml solución para perfusión EFG	ES/H/0348/001	81866	B.BRAUN MELSUNGEN AG	ES
Linezolid Farmaprojects 2 mg/ml solución para perfusión EFG	NL/H/2780/001	79.499	FARMAPROJECTS S.A.U.	ES
Linezolid Sandoz 600 mg comprimidos recubiertos con película EFG	NL/H/2965/001	79193	SANDOZ FARMACÉUTICA, S.A.	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Accordpharma 2 mg/ml solución para perfusión EFG	PT/H/1195/001	79964	ACCORD HEALTHCARE S.L.U.	ES
Linezolid Teva Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/2945/001	79489	TEVA PHARMA S.L.U.,	ES
Linezolid Teva Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/2945/001	79489	TEVA PHARMA S.L.U.,	ES
Linezolid Teva Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/2945/001	79489	TEVA PHARMA S.L.U.,	ES
Linezolid Teva Pharma 600 mg comprimidos recubiertos con película EFG	NL/H/2945/001	79489	TEVA PHARMA S.L.U.,	ES
Linezolid Demo 2 mg/ml solución para perfusión EFG	PT/H/1143/001	80350	DEMO ABEE	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg, filmdragerade tabletter	IE/H/0644/002	16423	PFIZER OY	FI
Zyvoxid 2 mg/ml infusionsvätska, lösning.	IE/H/0644/001	16421	PFIZER OY	FI
Zyvoxid 20 mg/ml rakeet oraalisuspensiota varten	IE/H/0644/003	16424	PFIZER OY	FI
Linezolid Krka 600 mg kalvopäällysteiset tabletit	AT/H/0625/001	33189	KRKA, D.D., NOVO MESTO	FI
Linezolid Sandoz 600 mg kalvopäällysteiset tabletit	NL/H/2965/001	31593	SANDOZ A/S	FI
Linezolid Accord 2 mg/ml infuusioneste, liuos	PT/H/1195/001	35197	ACCORD HEALTHCARE B.V.	FI
LINEZOLIDE EG 600 mg, comprimé pelliculé	not available	3400930086186	EG LABO LABORATOIRES EUROGENERICS	FR
ZYVOXID 600 mg, comprimé pelliculé	IE/H/0644/002	34009 563 139 5 2	PFIZER HOLDING FRANCE	FR
ZYVOXID 2 mg/ml, solution pour perfusion	IE/H/0644/001	34009 581 107 4 0	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 563 140 3 4	PFIZER HOLDING FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 565 126 8 3	PFIZER HOLDING FRANCE	FR
ZYVOXID 2 mg/ml, solution pour perfusion	IE/H/0644/001	34009 563 142 6 3	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 565 126 8 3	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 563 140 3 4	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 565 126 8 3	PFIZER HOLDING FRANCE	FR
ZYVOXID 100 mg/5 ml, granulés pour suspension buvable	IE/H/0644/003	34009 563 140 3 4	PFIZER HOLDING FRANCE	FR
LINEZOLIDE EG 2 mg/ml, solution pour perfusion	DE/H/5027/001	3400955019190	EG LABO LABORATOIRES EUROGENERICS	FR
LINEZOLIDE EG 2 mg/ml, solution pour perfusion	DE/H/5027/001	3400955019183	EG LABO LABORATOIRES EUROGENERICS	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
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 4 3	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 5 0	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 301 467 1 2	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 6 7	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 7 4	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 8 1	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 300 545 9 8	VIATRIS SANTE	FR
LINEZOLIDE VIATRIS 600 mg, comprimé pelliculé	NL/H/3011/001	34009 550 192 2 0	VIATRIS SANTE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 026 6 6	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 026 7 3	STRAGEN FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 026 8 0	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 026 9 7	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 0 3	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 1 0	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 2 7	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 3 4	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 4 1	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 5 8	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 6 5	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 7 2	STRAGEN FRANCE	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 027 8 9	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 028 0 2	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 028 1 9	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 028 2 6	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 028 3 3	STRAGEN FRANCE	FR
LINEZOLIDE STRAGEN 600 mg, comprimé pelliculé	AT/H/0491/001	34009 550 028 4 0	STRAGEN FRANCE	FR
LINEZOLIDE KALCEKS 2 mg/mL, solution pour perfusion	SE/H/2285/001	34009 551 077 3 6	KALCEKS	FR
LINEZOLIDE KALCEKS 2 mg/mL, solution pour perfusion	SE/H/2285/001	34009 551 077 4 3	KALCEKS	FR
LINEZOLIDE ARROW 2 mg/mL, solution pour perfusion	PT/H/1629/001	68316916	EUGIA PHARMA (MALTA) 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
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 558 5 4	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 558 6 1	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 558 8 5	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 558 9 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 559 0 8	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 559 1 5	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 550 199 9 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 559 2 2	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 300 559 4 6	ACCORD HEALTHCARE FRANCE SAS	FR
LINEZOLIDE ACCORD 600 mg, comprimé pelliculé	NL/H/4545/001	34009 550 200 0 4	ACCORD HEALTHCARE FRANCE SAS	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
LINEZOLIDE HIKMA 2 mg/mL, solution pour perfusion	AT/H/1082/001	34009 550 894 3 8	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	DE/H/6119/001	34009 550 065 2 7	PANMEDICA	FR
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	DE/H/6119/001	34009 550 065 3 4	PANMEDICA	FR
LINEZOLIDE PANPHARMA 2 mg/ml, solution pour perfusion	DE/H/6119/001	34009 550 065 4 1	PANMEDICA	FR
LINEZOLIDE ARROW 600 mg, comprimé pelliculé	not available	61722572	ARROW GENERIQUES	FR
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

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 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
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	IE/H/0644/002	41058/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Zyvoxid 2 mg/ml διάλυμα για έγχυση	IE/H/0644/001	41057/15-6-2015	PFIZER HELLAS A.E.	GR
Linesol 600 mg Επικαλυμμένα με λεπτό υμένιο δισκία	not available	92619/24/19-6-2025	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
Linezan 2 mg/ml Διάλυμα για Έγχυση	PT/H/1555/01/R/0	116063/10-11-2022	ANFARM HELLAS SA	GR
Linesol 2 mg/ml διάλυμα για ενδοφλέβια έγχυση.	not available	28977/23/02-04-2024	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	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
Antizolid 2 mg/ml διάλυμα για ενδοφλέβια έγχυση	not available	40816/7-5-2021	COOPER PHARMACEUTICALS S.A.	GR
Linezolid Kabi, 2 mg/ml διάλυμα για έγχυση	PT/H/1090/001	83407/24-11-2015	FRESENIUS KABI HELLAS SINGLE MEMBER S.A.	GR
Lizedia 600 mg επικαλυμμένα με λεπτό υμένιο δισκία	AT/H/0491/001	157069/17-12-2019	PHARMATHEN S.A.	GR
ZETALID 2 mg/ml Διάλυμα για έγχυση	PT/H/1143/001	3054401	DEMO ABEE	GR
Zyvoxid 600 mg filmom obložene tablete	not available	HR-H-202569569	PFIZER CROATIA D.O.O.	HR
Linezolid Sandoz 2 mg/ml otopina za infuziju	HR/H/0263/001	HR-H-249166135	SANDOZ D.O.O.	HR
Linezolid Krka 600 mg filmom obložene tablete	AT/H/0620/001	HR-H-094120244	KRKA-FARMA D.O.O.	HR
ZYVOXID 2 mg/ ml otopina za infuziju	not available	HR-H-711194389	PFIZER CROATIA D.O.O.	HR
Linezolid Kabi 2 mg/ml otopina za infuziju	PT/H/1090/001	HR-H-690522814	FRESENIUS KABI D.O.O.	HR
Linezolid Krka 2 mg/ml otopina za infuziju	AT/H/0639/001	HR-H-611386157	KRKA-FARMA 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
Lynz 600 mg filmom obložene tablete	NL/H/2965/001	HR-H-725335862	SANDOZ D.O.O.	HR
Linezolid Accord 2 mg/ml otopina za infuziju	PT/H/1195/001	HR-H-328955893	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA 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 Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA 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 Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA 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 Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Linezolid Pliva 600 mg filmom obložene tablete	NL/H/2945/001	HR-H-230450457	PLIVA HRVATSKA D.O.O.	HR
Anozilad 2 mg/ml oldatos infúzió	HU/H/0659/001	OGYI-T-22617/15	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/01	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/09	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/06	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/12	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/05	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/02	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/04	ACTAVIS GROUP PTC EHF.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/03	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/08	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/11	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/07	ACTAVIS GROUP PTC EHF.	HU
Anozilad 600 mg filmtabletta	HU/H/0834/001	OGYI-T-22617/10	ACTAVIS GROUP PTC EHF.	HU
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
Linesan 2 mg/ml oldatos infúzió	not available	0GYI-T-23736/01	ANFARM HELLAS SA	HU
Linesan 2 mg/ml oldatos infúzió	not available	0GYI-T-23736/02	ANFARM HELLAS SA	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linesan 2 mg/ml oldatos infúzió	not available	0GYI-T-23736/03	ANFARM HELLAS SA	HU
Linesan 2 mg/ml oldatos infúzió	not available	0GYI-T-23736/04	ANFARM HELLAS SA	HU
Linesan 2 mg/ml oldatos infúzió	not available	0GYI-T-23736/05	ANFARM HELLAS SA	HU
Linesan 2 mg/ml oldatos infiuzi0	not available	0GYI-T-23736/06	ANFARM HELLAS SA	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/01	FRESENIUS KABI HUNGARY KFT.	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/03	FRESENIUS KABI HUNGARY KFT.	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/05	FRESENIUS KABI HUNGARY KFT.	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/02	FRESENIUS KABI HUNGARY KFT.	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/04	FRESENIUS KABI HUNGARY KFT.	HU
Linezolid Fresenius Kabi 2 mg/ml oldatos infúzió	PT/H/1090/001/E001	OGYI-T-23774/06	FRESENIUS KABI HUNGARY KFT.	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PA 822/143/1	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PA 822/143/1	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PA 822/143/1	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PA0822/143/2	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PA 0822/143/004	PFIZER HEALTHCARE IRELAND UNLIMITED COMPANY	IE
Linezolid Krka 600 mg film-coated tablets	AT/H/0625/001	PA1347/057/001	KRKA, D.D., NOVO MESTO	IE
Linezolid 600 mg film-coated tablets	NL/H/4545/001	PA2315/104/001	ACCORD HEALTHCARE IRELAND LIMITED	IE
Linezolid Clonmel 600 mg film-coated tablets	DE/H/3489/001	PA0126/238/001	CLONMEL HEALTHCARE LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 2 mg/ml Solution for infusion	PT/H/1090/001	PA 2059/011/001	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Linezolid 2 mg/ml solution for infusion	AT/H/0639/001	PA1347/057/002	KRKA, D.D., NOVO MESTO	IE
Linezolid 600 mg Film-Coated Tablets	NL/H/2965/001	PA0711/230/001	ROWEX LTD	IE
Linezolid 2 mg/ml solution for infusion	PT/H/1195/001	PA2315/104/003	ACCORD HEALTHCARE IRELAND LIMITED	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Linezolid Teva 600 mg Film-coated Tablets	NL/H/2945/001	PA0749/204/001	TEVA PHARMA B.V.	IE
Zyvoxid 100 mg/5 ml mixtúrukyrni, dreifa.	IE/H/0644/003	IS/1/01/017/04	PFIZER APS	IS
Zyvoxid 100 mg/5 ml mixtúrukyrni, dreifa.	IE/H/0644/003	IS/1/01/017/04	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS

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 innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS

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 filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS

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 filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 600 mg filmuhúðaðar töflur.	IE/H/0644/002	IS/1/01/017/03	PFIZER APS	IS
Zyvoxid 100 mg/5 ml mixtúrukyrni, dreifa.	IE/H/0644/003	IS/1/01/017/04	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS

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 innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
Zyvoxid 2 mg/ml innrennslislyf, lausn.	IE/H/0644/001	IS/1/01/017/01	PFIZER APS	IS
ZYVOXID 100 mg/5 ml granulato per sospensione orale	IE/H/0644/003	035410075	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410024	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410024	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410226	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410315	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410253	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410366	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410380	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410289	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410291	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410240	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410238	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410366	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410226	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410051	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410226	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410265	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410289	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410416	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410291	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410380	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410430	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410428	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410378	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410339	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410404	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410392	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410277	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410303	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410327	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410238	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410366	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410341	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410354	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410315	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410024	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410253	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410240	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410048	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410036	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410063	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410327	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410339	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410265	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410341	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410303	PFIZER ITALIA S.R.L.	IT
ZYVOXID 600 mg compresse rivestite con film	IE/H/0644/002	035410226	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410012	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410366	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410380	PFIZER ITALIA S.R.L.	IT
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410380	PFIZER ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
ZYVOXID 2 mg/ml soluzione per infusione	IE/H/0644/001	035410024	PFIZER ITALIA S.R.L.	IT
ZYVOXID 100 mg/5 ml granulato per sospensione orale	IE/H/0644/003	035410075	PFIZER ITALIA S.R.L.	IT
ZYVOXID 100 mg/5 ml granulato per sospensione orale	IE/H/0644/003	035410075	PFIZER ITALIA S.R.L.	IT
ZYVOXID 100 mg/5 ml granulato per sospensione orale	IE/H/0644/003	035410075	PFIZER ITALIA S.R.L.	IT
ZYVOXID 100 mg/5 ml granulato per sospensione orale	IE/H/0644/003	035410075	PFIZER ITALIA S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948014	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948026	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948038	AUROBINDO PHARMA (ITALIA) S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948040	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948053	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948065	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948077	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948089	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948091	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948103	AUROBINDO PHARMA (ITALIA) S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948115	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948127	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948139	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948141	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948154	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948166	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948178	AUROBINDO PHARMA (ITALIA) S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Aurobindo Italia 600 mg compresse rivestite con film	AT/H/0491/001	044948180	AUROBINDO PHARMA (ITALIA) S.R.L.	IT
Linezolid Krka d.d. 600 mg compresse rivestite con film	AT/H/0620/001	AIC N. 044172017	KRKA, D.D., NOVO MESTO	IT
Linezolid Krka d.d. 600 mg compresse rivestite con film	AT/H/0620/001	044172029	KRKA, D.D., NOVO MESTO	IT
Linezolid Krka d.d. 600 mg compresse rivestite con film	AT/H/0620/001	AIC N. 044172031	KRKA, D.D., NOVO MESTO	IT
Linezolid S.A.L.F. 2 mg/ml soluzione per infusione	not available	045477015	S.A.L.F. SPA LABORATORIO FARMACOLOGICO	IT
Linezolid Aurobindo 2 mg/ml soluzione per infusione	PT/H/1629/001	045433012	EUGIA PHARMA (MALTA) LTD	IT
Linezolid Aurobindo 2 mg/ml soluzione per infusione	PT/H/1629/001	045433036	EUGIA PHARMA (MALTA) LTD	IT
Linezolid Aurobindo 2 mg/ml soluzione per infusione	PT/H/1629/001	045433048	EUGIA PHARMA (MALTA) 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 Aurobindo 2 mg/ml soluzione per infusione	PT/H/1629/001	045433024	EUGIA PHARMA (MALTA) LTD	IT
Linezolid Hikma 2 mg/ml soluzione per infusione.	AT/H/1082/001	049118019	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IT
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
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 Krka 2 mg/ml soluzione per infusione	AT/H/0639/001	044463014	KRKA, D.D., NOVO MESTO	IT
Linezolid Krka 2 mg/ml soluzione per infusione	AT/H/0639/001	044463026	KRKA, D.D., NOVO MESTO	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 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

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 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

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 600 mg compresse rivestite con film	NL/H/2945/001	043122124	TEVA ITALIA S.R.L.	IT
Linezolid Teva 600 mg compresse rivestite con film	NL/H/2945/001	043122050	TEVA ITALIA S.R.L.	IT
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

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 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	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	043122201	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 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
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
ZYVOXID 2 mg/ml infuzinis tirpalas	not available	LT/1/02/2448/001	PFIZER EUROPE MA EEIG	LT
Linezolid Kalceks 2 mg/ml infuzinis tirpalas	SE/H/2285/001	LT/1/25/5767/001	KALCEKS	LT
Linezolid Kalceks 2 mg/ml infuzinis tirpalas	SE/H/2285/001	LT/1/25/5767/002	KALCEKS	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/001	ACCORD HEALTHCARE B.V.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/002	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/003	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/004	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/005	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/006	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/007	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/008	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/009	ACCORD HEALTHCARE B.V.	LT
Linezolid Accord 600 mg plėvele dengtos tabletės	NL/H/4545/001	LT/1/15/3837/010	ACCORD HEALTHCARE B.V.	LT
Linezolid Krka 2 mg/ml infuzinis tirpalas	AT/H/0639/001	LT/1/16/4014/001	KRKA, D.D., NOVO MESTO	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Krka 2 mg/ml infuzinis tirpalas	AT/H/0639/001	LT/1/16/4014/002	KRKA, D.D., NOVO MESTO	LT
Linezolid Accord 2 mg/ml infuzinis tirpalas	PT/H/1195/001	LT/1/18/4185/001	ACCORD HEALTHCARE B.V.	LT
Zyvoxid 100 mg/5 ml granulés pour suspension buvable	IE/H/0644/003	2010020717	PFIZER S.A.	LU
Zyvoxid 2 mg/ml solution pour perfusion	IE/H/0644/001	2010020716	PFIZER S.A.	LU
Zyvoxid 600 mg comprimés pelliculés	IE/H/0644/002	2010020719	PFIZER S.A.	LU
Linezolid Fresenius Kabi 2 mg/1 ml Solution pour perfusion	PT/H/1090/001	2016040054	FRESENIUS KABI NV/SA	LU
Linezolid Krka 600 mg apvalkotās tabletes	AT/H/0620/001	15-0290	KRKA, D.D., NOVO MESTO	LV
Linezolid Kalceks 2 mg/ml šķīdums infūzijām	SE/H/2285/001	25-0114	KALCEKS	LV
Linezolid Kalceks 2 mg/ml šķīdums infūzijām	SE/H/2285/001	25-0114	KALCEKS	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 apvalkotās tabletes	NL/H/4545/001	15-0313	ACCORD HEALTHCARE B.V.	LV
ZYVOXID 2 mg/ml šķīdums infūzijām	not available	04-0288	PFIZER EUROPE MA EEIG	LV
Linezolid Krka 2 mg/ml šķīdums infūzijām	AT/H/0639/001	16-0225	KRKA, D.D., NOVO MESTO	LV
Linezolid Accord 2 mg/ml šķīdums infūzijām	PT/H/1195/001	18-0005	ACCORD HEALTHCARE B.V.	LV
Zyvox 2 mg/ml solution for infusion	not available	MA505/01803	PFIZER HELLAS A.E.	MT
Zyvox 600 mg film-coated tablets	not available	MA505/01802	PFIZER HELLAS A.E.	MT
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	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
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	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
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	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
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 2 mg/ml, oplossing voor infusie	IE/H/0644/001	RVG 26567	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Zyvoxid 600 mg, filmomhulde tabletten	IE/H/0644/002	RVG 26569	PFIZER B.V.	NL
Linezolid Sandoz 600 mg, filmomhulde tabletten	NL/H/2966/001	RVG 113824	SANDOZ 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
Linezolid Mylan 600 mg, filmomhulde tabletten	NL/H/3011/001	RVG 114163	MYLAN PHARMACEUTICALS LIMITED	NL
Dilizolen 2 mg/ml, oplossing voor infusie	NL/H/2505/001	RVG 111095	GLENMARK PHARMACEUTICALS S.R.O.	NL
Linezolid Accord 600 mg filmomhulde tabletten	NL/H/4545/001	RVG 115279	ACCORD HEALTHCARE B.V.	NL
Lynvox 2 mg/ml, oplossing voor infusie	NL/H/3443/001	RVG 115278	HIKMA PHARMA GMBH	NL
Lynvox 2 mg/ml, oplossing voor infusie	NL/H/3443/001	RVG 115278	HIKMA PHARMA GMBH	NL
Linezolid Fresenius Kabi 2 mg/ml, oplossing voor infusie	PT/H/1090/001	RVG 114296	FRESENIUS KABI NEDERLAND B.V.	NL
Linezolid Polpharma 2 mg/ml oplossing voor infusie	NL/H/2781/001	RVG 112520	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	NL
Linezolid Farmaprojects 2mg/ml, oplossing voor infusie	NL/H/2780/001/R/001	RVG 112519	FARMAPROJECTS S.A.U.	NL
Linezolid Polpharma 600 mg, filmomhulde tabletten	NL/H/3124/001	RVG 114804	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	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
Linezolid Sandoz 600 mg, filmomhulde tabletten	NL/H/2965/001	RVG 113822	SANDOZ B.V.	NL
Linezolid Teva 600 mg, filmomhulde tabletten	NL/H/2945/001	RVG 114205	TEVA NEDERLAND B.V.	NL
Linezolid Teva 600 mg, filmomhulde tabletten	NL/H/2945/001	RVG 114205	TEVA NEDERLAND B.V.	NL
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	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 tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	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 tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	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 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 2 mg/ml infusjonsvæske, oppløsning	IE/H/0644/001	01-2267	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	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 tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Zyvoxid 600 mg tabletter, filmdrasjerte.	IE/H/0644/002	01-2269	PFIZER AS	NO
Linezolid Krka 600 mg tabletter, filmdrasjerte	AT/H/0625/001	15-10604	KRKA, D.D., NOVO MESTO	NO
Linezolid Accord 600 mg filmdrasjerte tabletter	NL/H/4545/001	14-10050	ACCORD HEALTHCARE B.V.	NO
Linezolid Fresenius Kabi 2 mg/ml infusjonsvæske, oppløsning	PT/H/1090/001	13/9804	FRESENIUS KABI NORGE AS	NO
Linezolid Sandoz	NL/H/2965/001	13-9593	SANDOZ A/S	NO
Linezolid Accord 2 mg/ml infusjonsvæske, oppløsning.	PT/H/1195/001	17-11660	ACCORD HEALTHCARE B.V.	NO
Linezolid Krka, 600 mg, tabletki powlekane	AT/H/0620/001	22832	KRKA, D.D., NOVO MESTO	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
Dilizolen, 2 mg/ml, roztwór do infuzji	NL/H/2505/001	21192	GLENMARK PHARMACEUTICALS S.R.O.	PL
Linezolid Eugia, 2 mg/mL, roztwór do infuzji	PT/H/1629/001	24529	EUGIA PHARMA (MALTA) LTD	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 2 mg/ml, roztwór do infuzji	not available	9200	PFIZER EUROPE MA EEIG	PL
Zyvoxid, 600 mg, tabletki powlekane	not available	9198	PFIZER EUROPE MA EEIG	PL
Linezolid Kabi, 2 mg/ml, roztwór do infuzji	PT/H/1090/001	22370	FRESENIUS KABI POLSKA SP. Z O.O.	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
Linezolid Polpharma, 2 mg/ml, roztwór do infuzji	NL/H/2781/001	22018	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Linezolid Polpharma, 600 mg, tabletki powlekane	NL/H/3124/001	22745	ZAKLADY FARMACEUTYCZNE POLPHARMA S.A.	PL
Linezolid Krka, 2 mg/ml, roztwór do infuzji	AT/H/0639/001	23685	KRKA, D.D., NOVO MESTO	PL
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751880	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652286	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652682	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 2 mg/ml solução para perfusão	IE/H/0644/001	3650884	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652880	LABORATÓRIOS PFIZER 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
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3653185	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652484	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752284	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652583	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751989	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752680	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752789	LABORATÓRIOS PFIZER 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
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752383	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752581	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752284	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752482	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752086	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752581	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751781	LABORATÓRIOS PFIZER 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
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752185	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752482	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652385	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3653482	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652781	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3652989	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 2 mg/ml solução para perfusão	IE/H/0644/001	5413646	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752086	LABORATÓRIOS PFIZER 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
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752789	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3653284	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3653383	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3653086	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752383	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751880	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751781	LABORATÓRIOS PFIZER 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
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3751989	LABORATÓRIOS PFIZER LDA.	PT
Zyvoxid 600 mg comprimidos revestidos por película	IE/H/0644/002	3752680	LABORATÓRIOS PFIZER LDA.	PT
Linezolida Pharmakern, 600 mg, comprimidos revestidos por película	not available	5681937	PHARMAKERN PORTUGAL – PRODUTOS FARMACÊUTICOS, SOCIEDADE UNIPESSOAL, LDA.	PT
Linezolida Altan 2 mg/ml Solução para perfusão	not available	5712773	ALTAN PHARMACEUTICALS S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, 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 Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	5708128	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, 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 Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
Linezolida Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, 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 Aurovitas 600 mg comprimidos revestidos por película	AT/H/0491/001	AT/H/0491/001	GENERIS FARMACÊUTICA, S.A.	PT
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
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687470	GLENMARK ARZNEIMITTEL GMBH	PT
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687504	GLENMARK ARZNEIMITTEL GMBH	PT
Linezolida Glenmark 600 mg comprimidos revestidos por película	PT/H/1245/001	5687512	GLENMARK ARZNEIMITTEL GMBH	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 Normon 600 mg comprimidos revestidos por película.	ES/H/0291/001	5669056	LABORATÓRIOS NORMON, S.A.	PT
Linezolida Mylan 600 mg comprimidos revestidos por película	NL/H/3011/001	5723200	VIATRIS LIMITED	PT
Linezolida Normon 2 mg/ml solução para perfusão.	PT/H/1220/001	5674270	LABORATÓRIOS NORMON, S.A.	PT
Linezolida Anfarm 600 mg/300 ml Solução para Perfusão	PT/H/1555/001	PT/H/1555/01	ANFARM HELLAS SA	PT
Linezolida Aurobindo 2 mg/ml solução para perfusão	PT/H/1629/001	PT/H/1629/001	EUGIA PHARMA (MALTA) LTD	PT
Linezolida Aurobindo 2 mg/ml solução para perfusão	PT/H/1629/001	PT/H/1629/001	EUGIA PHARMA (MALTA) LTD	PT
Linezolida Aurobindo 2 mg/ml solução para perfusão	PT/H/1629/001	PT/H/1629/001	EUGIA PHARMA (MALTA) LTD	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 Aurobindo 2 mg/ml solução para perfusão	PT/H/1629/001	5793930	EUGIA PHARMA (MALTA) LTD	PT
Linezolida Accord 600 mg comprimidos revestidos por película	NL/H/4545/001	5684501	ACCORD HEALTHCARE S.L.U.	PT
Linezolida Accord 600 mg comprimidos revestidos por película	NL/H/4545/001	5684519	ACCORD HEALTHCARE S.L.U.	PT
Linezolida Farmoz 600 mg comprimidos revestidos por película	not available	5658778	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
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
Linezolida Hikma 2 mg/ml solução para perfusão	AT/H/1082/001	5846258	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Linezolida Kabi, 2 mg/ml solução para perfusão	PT/H/1090/001	5644414	FRESENIUS KABI PHARMA PORTUGAL, 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
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
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
Linezolida Accordpharma 2 mg/ml solução para perfusão	PT/H/1195/001	5669825	ACCORD HEALTHCARE S.L.U.	PT
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

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 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
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672720	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672738	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672746	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672753	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672761	DEMO ABEE	PT
Linezolida Demo 2 mg/ml Solução para perfusão	PT/H/1143/001	5672779	DEMO ABEE	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 Krka 600 mg comprimate filmate	AT/H/0620/001	13184/2020/01	KRKA, D.D., NOVO MESTO	RO
Linezolid Krka 600 mg comprimate filmate	AT/H/0620/001	13184/2020/02	KRKA, D.D., NOVO MESTO	RO
Linezolid Krka 600 mg comprimate filmate	AT/H/0620/001	13184/2020/03	KRKA, D.D., NOVO MESTO	RO
Zyvoxid 2 mg/ml soluţie perfuzabilă	not available	2047/2009/01	PFIZER EUROPE MA EEIG	RO
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 100 mg/5 ml granule pentru suspensie orală	not available	2048/2009/01	PFIZER EUROPE MA EEIG	RO
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	DE/H/6119/001	12826/2019/01	INFOMED FLUIDS SRL	RO
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	DE/H/6119/001	12826/2019/02	INFOMED FLUIDS SRL	RO
Linezolid Infomed 2 mg/ml soluţie perfuzabilă	DE/H/6119/001	12826/2019/03	INFOMED FLUIDS SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/01	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/02	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/03	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/04	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/05	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Kabi 2 mg/ml soluție perfuzabilă	PT/H/1090/001	15234/2023/06	FRESENIUS KABI ROMANIA SRL	RO
Linezolid Krka 2 mg/ml soluţie perfuzabilă	AT/H/0639/001	14216/2021/01	KRKA, D.D., NOVO MESTO	RO
Linezolid Krka 2 mg/ml soluţie perfuzabilă	AT/H/0639/001	14216/2021/02	KRKA, D.D., NOVO MESTO	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/01	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/02	SANDOZ PHARMACEUTICALS S.R.L.	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
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/03	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/04	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/05	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/06	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/07	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Sandoz 600 mg comprimate filmate	NL/H/2965/001	11582/2019/08	SANDOZ PHARMACEUTICALS S.R.L.	RO
Linezolid Accord 2 mg/ml soluţie perfuzabilă	PT/H/1195/001	12850/2019/01	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE

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 infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 20 mg/ml granulat till oral suspension	IE/H/0644/003	17169	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE

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 infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE

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 filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 600 mg filmdragerade tabletter	IE/H/0644/002	17168	PFIZER AB	SE

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 infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 20 mg/ml granulat till oral suspension	IE/H/0644/003	17169	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 2 mg/ml infusionsvätska, lösning	IE/H/0644/001	17166	PFIZER AB	SE
Zyvoxid 20 mg/ml granulat till oral suspension	IE/H/0644/003	17169	PFIZER AB	SE
Linezolid Krka 600 mg filmdragerade tabletter	AT/H/0625/001	52691	KRKA, D.D., NOVO MESTO	SE
Linezolid Glenmark 600 mg filmdragerade tabletter	PT/H/1245/001	51059	GLENMARK ARZNEIMITTEL GMBH	SE
Linezolid Kalceks 2 mg/ml infusionsvätska, lösning	SE/H/2285/001	64161	KALCEKS	SE

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 Kalceks 2 mg/ml infusionsvätska, lösning	SE/H/2285/001	64161	KALCEKS	SE
Linezolid Accord 600 mg filmdragerade tabletter	NL/H/4545/001	51043	ACCORD HEALTHCARE B.V.	SE
Linezolid Sandoz 600 mg filmdragerade tabletter	NL/H/2965/001	49543	SANDOZ A/S	SE
Linezolid Accord 2 mg/ml infusionsvätska, lösning.	PT/H/1195/001	56240	ACCORD HEALTHCARE B.V.	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE

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 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE

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 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE

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 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter.	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
Linezolid Teva 600 mg filmdragerade tabletter	NL/H/2945/001	49967	TEVA SWEDEN AB	SE
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
Zyvoxid 2 mg/ml raztopina za infundiranje	not available	H/04/01729/001	PFIZER LUXEMBOURG S.À.R.L.	SI
Zyvoxid 600 mg filmsko obložene tablete	not available	H/04/01729/002	PFIZER LUXEMBOURG S.À.R.L.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/001	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/002	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/003	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/004	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/005	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/006	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/007	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/008	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/009	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Accord 600 mg filmsko obložene tablete	NL/H/4545/001	H/16/02127/010	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/001	FRESENIUS KABI DEUTSCHLAND GMBH	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
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
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/004	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/005	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Linezolid Kabi 2 mg/ml raztopina za infundiranje	PT/H/1090/001	H/15/02023/006	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Linezolid Krka 2 mg/ml raztopina za infundiranje	AT/H/0639/001	H/17/02293/001	KRKA, D.D., NOVO MESTO	SI
Linezolid Krka 2 mg/ml raztopina za infundiranje	AT/H/0639/001	H/17/02293/002	KRKA, D.D., NOVO MESTO	SI
Linezolid Accord 2 mg/ml raztopina za infundiranje	PT/H/1195/001	H/18/02426/001	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SI
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

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 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
Linezolid Krka 600 mg filmom obalené tablety	AT/H/0620/001	15/0478/15-S	KRKA, D.D., NOVO MESTO	SK
Linezolid Accord 600 mg filmom obalené tablety	NL/H/4545/001	15/0188/16-S	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SK
Linezolid Kabi 2 mg/ml, infúzny roztok	PT/H/1090/001	15/0128/15-S	FRESENIUS KABI S.R.O.	SK
Linezolid Krka 2 mg/ml infúzny roztok	AT/H/0639/001	15/0482/16-S	KRKA, D.D., NOVO MESTO	SK
Linezolid Sandoz 600 mg filmom obalené tablety	NL/H/2965/001	15/0076/14-S	SANDOZ PHARMACEUTICALS D.D.	SK
Linezolid Accord 2 mg/ml infúzny roztok	PT/H/1195/001	15/0038/18-S	ACCORD HEALTHCARE POLSKA SP. Z O.O.	SK
Linezolid 600 mg Film-Coated Tablets	not available	PL 17907/0525	BRISTOL LABORATORIES LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PL 00057/1065	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PL 00057/1065	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI
Zyvox 100 mg/5 ml granules for oral suspension	IE/H/0644/003	PL 00057/1065	PFIZER LIMITED	XI
Zyvox 2 mg/ml solution for infusion	IE/H/0644/001	PL 00057/1066	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI
Zyvox 600 mg film-coated tablets	IE/H/0644/002	PL 00057/1068	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 2mg/ml solution for infusion	PT/H/1143/001	PL 24598/0046	NORIDEM ENTERPRISES LTD	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 00057/1418	PFIZER LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 00057/1419	PFIZER LIMITED	XI
Linezolid Krka 600 mg film-coated tablets	AT/H/0625/001	PL 01656/0209	KRKA, D.D., NOVO MESTO	XI
Linezolid 2 mg/ml Solution for Infusion	HR/H/0263/001	PL 04416/1428	SANDOZ LTD	XI
Linezolid 600mg Film-Coated Tablets	not available	PL 20416/0517	CRESCENT PHARMA LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 600 mg film-coated tablets	not available	PL 43805/0030	ASCEND LABORATORIES (UK) LIMITED	XI
Linezolid 600 mg film-coated tablets	UK/H/6727/001	PL 06831/0313	GENUS PHARMACEUTICALS LIMITED	XI
Linezolid 600 mg film-coated tablets	NL/H/3011/001	PL 04569/1640	GENERICS [UK] LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 16363/0491	MILPHARM LIMITED	XI
Linezolid 600 mg film-coated tablets	NL/H/4545/001	PL 20075/0394	ACCORD HEALTHCARE LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 15413/0161	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	XI
Linezolid 600 mg Film-coated Tablets	DE/H/6978/001	PL 49445/0092	AMAROX LIMITED	XI
Linezolid 600 mg film-coated tablets	not available	PL 17780/0902	ZENTIVA PHARMA UK LIMITED	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml solution for infusion	not available	PL 56284/0033	EUGIA (UK) LTD	XI
Linezolid 2 mg/ml Solution for infusion	PT/H/1090/001	PL 08828/0256	FRESENIUS KABI LIMITED	XI
Linezolid 2 mg/ml solution for infusion	ES/H/0348/001	PL 03551/0147	B.BRAUN MELSUNGEN AG	XI
Linezolid 2 mg/ml solution for infusion	AT/H/0639/001	PL 01656/0213	KRKA, D.D., NOVO MESTO	XI
Linezolid 600 mg Film-coated Tablets	NL/H/2965/001	PL 04416/1389	SANDOZ LTD	XI
Linezolid 2 mg/ml solution for infusion.	PT/H/1195/001	PL 20075/1159	ACCORD HEALTHCARE LIMITED	XI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Linezolid 2mg/ml solution for infusion	PT/H/1143/001	PL 24598/0046	NORIDEM ENTERPRISES LTD	XI