15 October 2020
EMA/647607/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: vancomycin

Procedure no.: PSUSA/00003097/202001

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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
Bactocin, pulver til infusionsvæske, opløsning	FI/H/0882/002	46553	MIP PHARMA GMBH	DK
Bactocin, pulver til infusionsvæske, opløsning	FI/H/0882/001	46552	MIP PHARMA GMBH	DK
Edicin 0,5 g prašak za otopinu za infuziju	not available	UP/I-530-09/13-02/237	SANDOZ D.O.O.	HR
EDICIN 0,5 g prášek pro koncentrát pro infuzní roztok	not available	15/165/97-C	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
EDICIN 0,5 g prášok na infúzny roztok	NL/H/4445/001	15/0869/09-S	SANDOZ PHARMACEUTICALS D.D.	SK
Edicin 1 g prašak za otopinu za infuziju	not available	UP/I-530-09/13-02/238	SANDOZ D.O.O.	HR
EDICIN 1 g prášek pro koncentrát pro infuzní roztok	not available	15/164/97-C	LEK PHARMACEUTICALS D.D. LJUBLJANA	CZ
EDICIN 1 g prášok na infúzny roztok	NL/H/4445/002	15/0870/09-S	SANDOZ PHARMACEUTICALS D.D.	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
Edicin 1000 mg prašek za koncentrat za raztopino za infundiranje	not available	H/93/00525/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Edicin 500 mg prašek za koncentrat za raztopino za infundiranje	not available	H/93/00525/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
EDICIN, 1 G, PROSZEK DO SPORZĄDZANIA ROZTWORU DO INFUZJI	not available	R/7011	SANDOZ GMBH	PL
EDICIN, 500 MG, PROSZEK DO SPORZĄDZANIA ROZTWORU DO INFUZJI	not available	R/7010	SANDOZ GMBH	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
LEVOVANOX 1 g polvere per soluzione per infusione endovenosa e per soluzione orale	not available	035003021	GENETIC SPA	IT
LEVOVANOX 250 mg Capsule rigide	not available	035003033	GENETIC SPA	IT
LEVOVANOX 500 mg polvere per soluzione per infusione endovenosa e per soluzione orale	not available	035003019	GENETIC SPA	IT
MAXIVANIL 1 g polvere per soluzione per infusione endovenosa e per soluzione orale	not available	034984029	GENETIC SPA	IT
MAXIVANIL 250 mg Capsule rigide	not available	034984031	GENETIC SPA	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
MAXIVANIL 500 mg polvere per soluzione per infusione endovenosa e per soluzione orale	not available	034984017	GENETIC SPA	IT
Selamat 1000 mg por oldatos infúzióhoz való koncentrátumhoz	HU/H/0542/002	OGYI-T-22781/02	ACTAVIS GROUP PTC EHF.	HU
Selamat 500 mg por oldatos infúzióhoz való koncentrátumhoz	HU/H/0542/001	OGYI-T-22781/01	ACTAVIS GROUP PTC EHF.	HU
Vamysin 1000 mg poeder voor concentraat voor oplossing voor infusie.	DE/H/5737/002	BE405291	TEVA PHARMA BELGIUM N.V./S.A	BE
Vamysin 1000 mg, poudre pour solution à diluer pour solution pour perfusion	DE/H/5737/002	2014090289	TEVA PHARMA BELGIUM N.V./S.A	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vamysin 500 mg poeder voor concentraat voor oplossing voor infusie.	DE/H/5737/001	BE405282	TEVA PHARMA BELGIUM N.V./S.A	BE
Vamysin 500 mg, poudre pour solution à diluer pour solution pour perfusion	DE/H/5737/001	2014090288	TEVA PHARMA BELGIUM N.V./S.A	LU
Vanco Sapiens 1g Σκόνη για διάλυμα προς έγχυση	not available	22213	SAPIENS PHARMACEUTICALS LTD	CY
Vanco Sapiens 1g Σκόνη για διάλυμα προς έγχυση	not available	22213	SAPIENS PHARMACEUTICALS LTD	CY
Vanco Sapiens 1g Σκόνη για διάλυμα προς έγχυση	not available	22213	SAPIENS PHARMACEUTICALS LTD	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
Vanco Sapiens 500mg κόνις για διάλυμα προς έγχυση	not available	022837	SAPIENS PHARMACEUTICALS LTD	CY
Vanco Sapiens 500mg κόνις για διάλυμα προς έγχυση	not available	022837	SAPIENS PHARMACEUTICALS LTD	CY
Vanco Sapiens 500mg κόνις για διάλυμα προς έγχυση	not available	022837	SAPIENS PHARMACEUTICALS LTD	CY
Vancocaps 250 mg harde capsules	NL/H/3359/001	RVG116655	EUROCEPT INTERNATIONAL BV	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
Vancocaps 250 mg, capsules, hard	NL/H/3359/001	BE503253	EUROCEPT INTERNATIONAL BV	BE
Vancocaps 250 mg, gélules	NL/H/3359/001	BE503253	EUROCEPT INTERNATIONAL BV	BE
Vancocaps 250 mg, gélules	NL/H/3359/001	2017040148	EUROCEPT B.V.	LU
Vancocaps 250 mg, Hartkapseln	NL/H/3359/001	BE503253	EUROCEPT INTERNATIONAL BV	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
Vancocaps 250 mg, Hartkapseln	NL/H/3359/001	2017040148	EUROCEPT B.V.	LU
Vancocin 1 g Pulver für ein Konzentrat zur Herstellung einer Infusionslösung und einer Lösung zum Einnehmen	not available	1-19487	ASTRO-PHARMA GMBH	AT
Vancocin 1 g powder for concentrate for solution for infusion and powder for oral solution.	not available	PA1226/005/004	FLYNN PHARMA LTD	IE
Vancocin 1000 mg powder for solution for infusion and oral solution.	not available	PL 13621/0033	FLYNN PHARMA LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancocin 125 mg kapsel, hård	not available	10514	STADA NORDIC APS	SE
Vancocin 500 mg - Pulver für ein Konzentrat zur Herstellung einer Infusionslösung und einer Lösung zum Einnehmen	not available	17.126	ASTRO-PHARMA GMBH	AT
Vancocin 500 mg powder for concentrate for solution for infusion and powder for oral solution.	not available	PA1226/005/003	FLYNN PHARMA LTD	IE
Vancocin 500 mg powder for solution for infusion and oral solution.	not available	PL 13621/0033	FLYNN PHARMA LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancocin CP 250 mg, capsules	not available	RVG 11984	EUROCEPT INTERNATIONAL BV	NL
Vancocin Matrigel 125mg Hard Capsules	not available	PA 1226/005/001	FLYNN PHARMA LTD	IE
Vancocin Matrigel 125mg Hard capsules	not available	PL 13621/0030	FLYNN PHARMA LTD	UK
Vancocin, kapsler, hårde	not available	30087	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancocin, kapsler, hårde	not available	12076	STRIDES PHARMA (CYPRUS) LIMITED	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
VANCOCINA A.P. 500 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	016334029	VANCOCIN ITALIA S.R.L	IT
VANCOCINA A.P. 500 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	016334029	VANCOCIN ITALIA S.R.L	IT
VANCOMICINA ACCORD 1 g IV EFG	not available	62292	ACCORD HEALTHCARE S.L.U.	ES
VANCOMICINA ACCORD 500 mg IV EFG	not available	62293	ACCORD HEALTHCARE S.L.U.	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
Vancomicina Azevedos, 1000 mg, Pó para solução para perfusão	not available	5623723	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 1000 mg, Pó para solução para perfusão	not available	5623731	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 1000 mg, Pó para solução para perfusão	not available	5623749	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 1000 mg, Pó para solução para perfusão	not available	5623756	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 500 mg, Pó para solução para perfusão	not available	5623665	LABORATÓRIOS AZEVEDOS - INDÚSTRIA 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
Vancomicina Azevedos, 500 mg, Pó para solução para perfusão	not available	5623673	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 500 mg, Pó para solução para perfusão	not available	5623707	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Azevedos, 500 mg, Pó para solução para perfusão	not available	5623715	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
VANCOMICINA COMBINO 1000 mg, pó para solução para perfusão	not available	5214465	ACCORD HEALTHCARE LIMITED	PT
VANCOMICINA COMBINO 500 mg, pó para solução para perfusão	not available	5214457	ACCORD HEALTHCARE LIMITED	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomicina Generis 1000 mg pó para concentrado para solução para perfusão	not available	2679280	GENERIS FARMACÊUTICA, S.A.	PT
Vancomicina Generis 1000 mg pó para concentrado para solução para perfusão	not available	2679389	GENERIS FARMACÊUTICA, S.A.	PT
Vancomicina Generis 500 mg pó para concentrado para solução para perfusão	not available	2679082	GENERIS FARMACÊUTICA, S.A.	PT
Vancomicina Generis 500 mg pó para concentrado para solução para perfusão	not available	2679181	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
VANCOMICINA HIKMA 1000 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	035004050	HIKMA ITALIA S.P.A.	IT
VANCOMICINA HIKMA 1000 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	035004023	HIKMA ITALIA S.P.A.	IT
VANCOMICINA HIKMA 500 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	035004047	HIKMA ITALIA S.P.A.	IT
VANCOMICINA HIKMA 500 mg polvere per concentrato per soluzione per infusione e per soluzione orale	not available	035004011	HIKMA ITALIA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomicina Hikma, 1000 mg, pó para concentrado para solução para perfusão	PT/H/0771/002	5570429	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 1000 mg, pó para concentrado para solução para perfusão	PT/H/0771/002	5570437	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 1000 mg, Pó para solução para perfusão	PT/H/0339/002	5182951	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 1000 mg, Pó para solução para perfusão	PT/H/0339/002	5182944	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 500 mg, pó para concentrado para solução para perfusão	PT/H/0771/001	5569876	HIKMA FARMACÊUTICA (PORTUGAL), 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
Vancomicina Hikma, 500 mg, pó para concentrado para solução para perfusão	PT/H/0771/001	5569900	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 500 mg, Pó para solução para perfusão	PT/H/0339/001	5182928	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Hikma, 500 mg, Pó para solução para perfusão	PT/H/0339/001	5182936	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	PT
Vancomicina Kabi 1000 mg pó para concentrado para solução para perfusão.	PT/H/2179/002	5571849	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Vancomicina Kabi 1000 mg pó para concentrado para solução para perfusão.	PT/H/2179/002	5379854	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
Vancomicina Kabi 1000 mg pulbere pentru concentrat pentru soluţie perfuzabilă	PT/H/2179/002	8513/2016/01	FRESENIUS KABI ROMANIA SRL	RO
Vancomicina Kabi 1000 mg pulbere pentru concentrat pentru soluţie perfuzabilă	PT/H/2179/002	8513/2016/02	FRESENIUS KABI ROMANIA SRL	RO
Vancomicina Kabi 500 mg pó para concentrado para solução para perfusão.	PT/H/2179/001	5571831	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Vancomicina Kabi 500 mg pó para concentrado para solução para perfusão.	PT/H/2179/001	5378039	FRESENIUS KABI PHARMA PORTUGAL, LDA.	PT
Vancomicina Kabi 500 mg pulbere pentru concentrat pentru soluţie perfuzabilă	PT/H/2179/001	8512/2016/01	FRESENIUS KABI ROMANIA SRL	RO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomicina Kabi 500 mg pulbere pentru concentrat pentru soluţie perfuzabilă	PT/H/2179/001	8512/2016/02	FRESENIUS KABI ROMANIA SRL	RO
Vancomicina Labesfal 1000 mg Pó para concentrado para solução para perfusão.	not available	3211398	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Vancomicina Labesfal 1000 mg Pó para concentrado para solução para perfusão.	not available	3211497	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Vancomicina Labesfal 1000 mg Pó para concentrado para solução para perfusão.	not available	5164033	LABESFAL LABORATORIOS ALMIRO, 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
Vancomicina Labesfal 500 mg Pó para concentrado para solução para perfusão.	not available	3211190	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Vancomicina Labesfal 500 mg Pó para concentrado para solução para perfusão.	not available	3211299	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Vancomicina Labesfal 500 mg Pó para concentrado para solução para perfusão.	not available	5164025	LABESFAL LABORATORIOS ALMIRO, S.A.	PT
Vancomicina MIP 1000 mg polvo para solución para perfusión EFG	FI/H/0889/002	78671	MIP PHARMA GMBH	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
Vancomicina MIP 500 mg polvo para solución para perfusión EFG	FI/H/0889/001	78672	MIP PHARMA GMBH	ES
Vancomicina Mylan 1 g polvere per soluzione per infusione	CZ/H/0351/002	041220082/M	MYLAN S.P.A.	IT
Vancomicina Mylan 1 g polvere per soluzione per infusione	CZ/H/0351/002	041220070/M	MYLAN S.P.A.	IT
Vancomicina Mylan 1 g polvere per soluzione per infusione	CZ/H/0351/002	041220068/M	MYLAN S.P.A.	IT
Vancomicina Mylan 1 g polvere per soluzione per infusione	CZ/H/0351/002	41220056/M	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomicina Mylan 1000 mg pó para concentrado para solução para perfusão.	SE/H/1158/002	5648928	MYLAN, LDA	PT
Vancomicina Mylan 1000 mg pó para concentrado para solução para perfusão.	SE/H/1158/002	5648944	MYLAN, LDA	PT
Vancomicina Mylan 1000 mg pulbere pentru concentrat pentru soluţie perfuzabilă	SE/H/1158/002	9921/2017/01	MYLAN S.A.S	RO
Vancomicina Mylan 500 mg pó para concentrado para solução para perfusão.	SE/H/1158/001	5648910	MYLAN, 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
Vancomicina Mylan 500 mg polvere per soluzione per infusione	CZ/H/0351/001	041220017/M	MYLAN S.P.A.	IT
Vancomicina Mylan 500 mg polvere per soluzione per infusione	CZ/H/0351/001	041220029/M	MYLAN S.P.A.	IT
Vancomicina Mylan 500 mg polvere per soluzione per infusione	CZ/H/0351/001	041220031/M	MYLAN S.P.A.	IT
Vancomicina Mylan 500 mg polvere per soluzione per infusione	CZ/H/0351/001	041220043	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomicina Mylan 500 mg pulbere pentru concentrat pentru soluţie perfuzabilă	SE/H/1158/001	9920/2017/01	MYLAN S.A.S	RO
Vancomicina Pfizer 1.000 mg polvo para concentrado para solución para perfusión EFG.	AT/H/0876/002	73785	PFIZER, S.L.	ES
Vancomicina Pfizer 500 mg polvo para concentrado para solución para perfusión EFG	AT/H/0876/001	73784	PFIZER, S.L.	ES
Vancomicina Quimedical, 1000 mg, Pó para concentrado para solução para perfusão	PT/H/1401/002	5623624	LABORATÓRIOS AZEVEDOS - INDÚSTRIA 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
Vancomicina Quimedical, 1000 mg, Pó para concentrado para solução para perfusão	PT/H/1401/002	5623632	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Quimedical, 1000 mg, Pó para concentrado para solução para perfusão	PT/H/1401/002	5623640	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Quimedical, 1000 mg, Pó para concentrado para solução para perfusão	PT/H/1401/002	5623657	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Quimedical, 500 mg, Pó para concentrado para solução para perfusão	PT/H/1401/001	5623368	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Quimedical, 500 mg, Pó para concentrado para solução para perfusão	PT/H/1401/001	5623376	LABORATÓRIOS AZEVEDOS - INDÚSTRIA 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
Vancomicina Quimedical, 500 mg, Pó para concentrado para solução para perfusão	PT/H/1401/001	5623400	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
Vancomicina Quimedical, 500 mg, Pó para concentrado para solução para perfusão	PT/H/1401/001	5623418	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	PT
VANCOMICINA SALA 1 g, Polvo para solución inyectable EFG	not available	67.283	LABORATORIO REIG JOFRE, S.A.	ES
VANCOMICINA SALA 500 mg, Polvo para solución inyectable	not available	67.282	LABORATORIO REIG JOFRE, 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
Vancomimylan 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	SE/H/1158/002	BE434147	MYLAN BVBA/SPRL	BE
Vancomimylan 1000 mg, poeder voor concentraat voor oplossing voor infusie	SE/H/1158/002	BE434147	MYLAN BVBA/SPRL	BE
Vancomimylan 1000 mg, poudre pour solution à diluer pour solution pour perfusion	SE/H/1158/002	BE434147	MYLAN BVBA/SPRL	BE
Vancomimylan 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	SE/H/1158/001	BE434131	MYLAN BVBA/SPRL	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
Vancomimylan 500 mg, poeder voor concentraat voor oplossing voor infusie	SE/H/1158/001	BE434131	MYLAN BVBA/SPRL	BE
Vancomimylan 500 mg, poudre pour solution à diluer pour solution pour perfusion	SE/H/1158/001	BE434131	MYLAN BVBA/SPRL	BE
Vancomycin CP 1,0 g Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.03.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin CP 500 mg Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.02.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Vancomycin "Lederle" 1000 mg; Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	not available	36654.01.00	RIEMSER PHARMA GMBH	DE
Vancomycin "Lederle" 500 mg; Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	not available	36654.00.00	RIEMSER PHARMA GMBH	DE
Vancomycin "Reig Jofre", pulver til koncentrat til infusionsvæske, opløsning	not available	53803	LABORATORIO REIG JOFRE, S.A.	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
Vancomycin "Reig Jofre", pulver til koncentrat til infusionsvæske, opløsning	not available	53804	LABORATORIO REIG JOFRE, S.A.	DK
Vancomycin "Sandoz"	NL/H/4445/001	42478	SANDOZ A/S	DK
Vancomycin "Sandoz"	NL/H/4445/002	42479	SANDOZ A/S	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/002	57500	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
Vancomycin "Strides", hårde kapsler	DK/H/2629/001	57499	STRIDES PHARMA (CYPRUS) LIMITED	DK
VANCOMYCIN / ΦΟΙΝΙΞΦΑΡΜ (Βανκομυκίνη CP, χρωματογραφικά κεκαθαρμένη βανκομυκίνη)	not available	18136/21-12-2011	FINIXFARM LTD	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
Vancomycin ”Fresenius Kabi”, pulver til koncentrat til infusionsvæske, opløsning	PT/H/2179/001	45259	FRESENIUS KABI AB	DK
Vancomycin ”Fresenius Kabi”, pulver til koncentrat til infusionsvæske, opløsning	PT/H/2179/002	45260	FRESENIUS KABI AB	DK
Vancomycin ”Mylan”, pulver til koncentrat til infusionsvæske, opløsning	SE/H/1158/001	50506	MYLAN IRELAND LIMITED	DK
Vancomycin ”Mylan”, pulver til koncentrat til infusionsvæske, opløsning	SE/H/1158/002	50507	MYLAN IRELAND LIMITED	DK
Vancomycin ”Orion”, pulver til koncentrat til infusionsvæske, opløsning	FI/H/0845/001	53549	ORION CORPORATION	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
Vancomycin ”Orion”, pulver til koncentrat til infusionsvæske, opløsning	FI/H/0845/002	53550	ORION CORPORATION	DK
Vancomycin 1000 mg powder for concentrate for solution for infusion	PT/H/1401/002	PA 1852/004/002	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	IE
Vancomycin 1000 mg Powder for concentrate for solution for infusion	PT/H/0771/002	PA1217/009/002	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IE
Vancomycin 1000 mg Powder for concentrate for solution for infusion	PT/H/0771/002	PA1217/009/002	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IE
Vancomycin 1000 mg Powder for concentrate for solution for infusion	PT/H/2179/002	PA 2059/066/002	FRESENIUS KABI DEUTSCHLAND GMBH	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
Vancomycin 1000 mg Powder for concentrate for solution for infusion and oral solution	IE/H/0706/002	PA1122/008/002	NORIDEM ENTERPRISES LTD	IE
Vancomycin 1000 mg Powder for concentrate for solution for infusion and oral solution	IE/H/0706/002	PL 24598/0016	NORIDEM ENTERPRISES LTD	UK
Vancomycin 1000 mg Powder for concentrate for solution for infusion.	PT/H/2179/002	PL 08828/0238	FRESENIUS KABI LIMITED	UK
Vancomycin 1000 mg powder for concentrate for solution for infusion.	AT/H/0876/002	PL 00057/1169	PFIZER LIMITED	UK
Vancomycin 1000 mg, Powder for concentrate for solution for infusion	PT/H/1401/002	PL 24065/0006 - 0001	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin 1000 mg, powder for concentrate for solution for infusion	SE/H/1158/002	PL 04569/1603	GENERICS [UK] LIMITED	UK
Vancomycin 1000 mg, Powder for solution for infusion	FI/H/0889/002	PL 26928/0016	MIP PHARMA GMBH	UK
Vancomycin 1000, 1000 mg Pulver zur Herstellung einer Infusionslösung	DE/H/0368/002	39981.01.00	MIP PHARMA GMBH	DE
Vancomycin 125 mg hard capsules	UK/H/5577/001	PL 13606/0196	STRIDES PHARMA UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin 125 mg hard capsules	UK/H/5577/001	PL 13606/0196	STRIDES PHARMA UK LIMITED	UK
Vancomycin 125mg Capsules, Hard.	UK/H/2508/001	PL 20117/0149	MORNINGSIDE HEALTHCARE LTD	UK
Vancomycin 1g Powder for Solution for Infusion	not available	29831/0322	WOCKHARDT UK LTD	UK
Vancomycin 1g Powder for Solution for Infusion	not available	29831/0322	WOCKHARDT UK LTD	UK
Vancomycin 1g Powder for Solution for Infusion	not available	PL 44095/0023	REIG JOFRE UK LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin 250 mg hard capsules	UK/H/5577/002	PL 13606/0197	STRIDES PHARMA UK LIMITED	UK
Vancomycin 250 mg hard capsules	UK/H/5577/002	PL 13606/0197	STRIDES PHARMA UK LIMITED	UK
Vancomycin 500 mg powder for concentrate for solution for infusion	PT/H/1401/001	PA 1852/004/001	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	IE
Vancomycin 500 mg Powder for concentrate for solution for infusion	PT/H/0771/001	PA1217/009/001	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	IE
Vancomycin 500 mg Powder for concentrate for solution for infusion	PT/H/0771/001	PA1217/009/001	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	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
Vancomycin 500 mg Powder for concentrate for solution for infusion and oral solution	IE/H/0706/001	PA 1122/008/001	NORIDEM ENTERPRISES LTD	IE
Vancomycin 500 mg Powder for concentrate for solution for infusion and oral solution	IE/H/0706/001	PL 24598/0015	NORIDEM ENTERPRISES LTD	UK
Vancomycin 500 mg Powder for concentrate for solution for infusion.	PT/H/2179/001	PA 2059/066/001	FRESENIUS KABI DEUTSCHLAND GMBH	IE
Vancomycin 500 mg Powder for concentrate for solution for infusion.	PT/H/2179/001	PL 08828/0237	FRESENIUS KABI LIMITED	UK
Vancomycin 500 mg powder for concentrate for solution for infusion.	AT/H/0876/001	PL 00057/1168	PFIZER LIMITED	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin 500 mg, Powder for concentrate for solution for infusion	PT/H/1401/001	PL 24065/0005 - 0001	LABORATÓRIOS AZEVEDOS - INDÚSTRIA FARMACÊUTICA, S.A.	UK
Vancomycin 500 mg, powder for concentrate for solution for infusion	SE/H/1158/001	PL 04569/1602	GENERICS [UK] LIMITED	UK
Vancomycin 500 mg, Powder for solution for infusion	FI/H/0889/001	PL 26928/0015	MIP PHARMA GMBH	UK
Vancomycin 500, 500 mg Pulver zur Herstellung einer Infusionslösung	DE/H/0368/001	42565.00.00	MIP PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin 500mg Powder for Solution for Infusion	not available	PL 44095/0022	REIG JOFRE UK LIMITED	UK
Vancomycin 500mg Powder for Solution for Infusion	UK/H/7133/001/MR	29831/0319	WOCKHARDT UK LTD	UK
Vancomycin 500mg Powder for Solution for Infusion	UK/H/7133/001/MR	29831/0319	WOCKHARDT UK LTD	UK
Vancomycin AB 1000 mg poeder voor concentraat voor oplossing voor infusie	NL/H/4181/002	BE469120	AUROBINDO PHARMA B.V.	BE
Vancomycin AB 1000 mg poudre pour solution à diluer pour perfusion	NL/H/4181/002	BE469120	AUROBINDO PHARMA B.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin AB 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	NL/H/4181/002	BE469120	AUROBINDO PHARMA B.V.	BE
Vancomycin AB 500 mg poeder voor concentraat voor oplossing voor infusie	NL/H/4181/001	BE469111	AUROBINDO PHARMA B.V.	BE
Vancomycin AB 500 mg poudre pour solution à diluer pour perfusion	NL/H/4181/001	BE469111	AUROBINDO PHARMA B.V.	BE
Vancomycin AB 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	NL/H/4181/001	BE469111	AUROBINDO PHARMA B.V.	BE
Vancomycin Actavis 1.000 mg stofn fyrir innrennslisþykkni, lausn	SE/H/1733/002	IS/1/15/019/02	ACTAVIS GROUP PTC EHF.	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
Vancomycin Actavis 1000 mg pulver till koncentrat till infusionsvätska, lösning	SE/H/1733/002	49322	ACTAVIS GROUP PTC EHF.	SE
Vancomycin Actavis 500 mg pulver till koncentrat till infusionsvätska, lösning	SE/H/1733/001	49321	ACTAVIS GROUP PTC EHF.	SE
Vancomycin Actavis 500 mg stofn fyrir innrennslisþykkni, lausn	SE/H/1733/001	IS/1/15/019/01	ACTAVIS GROUP PTC EHF.	IS
Vancomycin Capsules 125 mg	not available	PL 17815/0041	XELLIA PHARMACEUTICALS APS	UK
Vancomycin Capsules 250mg	not available	PL 17815/0042	XELLIA PHARMACEUTICALS APS	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin CP 1,0 g Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.03.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Vancomycin CP 1,0 g Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.03.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Vancomycin CP 1,0 g Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.03.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin CP 500 mg Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.02.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Vancomycin CP 500 mg Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.02.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE
Vancomycin CP 500 mg Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen Wirkstoff: Vancomycinhydrochlorid	not available	16803.02.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Dr. Eberth 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	AT/H/0569/002	92809.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	DE
Vancomycin Dr. Eberth 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder zur Herstellung einer Lösung zum Einnehmen	AT/H/0569/002	137151	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT
Vancomycin Dr. Eberth 125 mg Hartkapseln	DK/H/2629/001	138014	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT
Vancomycin Dr. Eberth 125 mg Hartkapseln	DK/H/2629/001	138014	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Dr. Eberth 125 mg Hartkapseln	DE/H/5908/001	90867.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	DE
Vancomycin Dr. Eberth 250 mg Hartkapseln	DK/H/2629/002	138015	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT
Vancomycin Dr. Eberth 250 mg Hartkapseln	DK/H/2629/002	138015	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT
Vancomycin Dr. Eberth 250 mg Hartkapseln	DE/H/5908/002	90868.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL 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
Vancomycin Dr. Eberth 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	AT/H/0569/001	92808.00.00	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	DE
Vancomycin Dr. Eberth 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder zur Herstellung einer Lösung zum Einnehmen	AT/H/0569/001	137150	DR. FRIEDRICH EBERTH ARZNEIMITTEL GMBH	AT
Vancomycin ENTEROCAPS 125 mg Hartkapseln	not available	95248.00.00	RIEMSER PHARMA GMBH	DE
VANCOMYCIN ENTEROCAPS® 250 mg Kapseln	not available	1-19057	RIEMSER PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin ENTEROCAPS® 250 mg, Hartkapseln	not available	7422.01.00	RIEMSER PHARMA GMBH	DE
Vancomycin FarmaPlus 1000 mg prášok na infúzny koncentrát	SE/H/1158/002	15/0177/14-S	MYLAN S.A.S	SK
Vancomycin Fresenius Kabi 1 g pulver til konsentrat til infusjonsvæske, oppløsning.	PT/H/2179/002	09-6842	FRESENIUS KABI NORGE AS	NO
Vancomycin Fresenius Kabi 1.000 mg stofn fyrir innrennslisþykkni, lausn.	PT/H/2179/002	IS/1/11/015/02	FRESENIUS KABI AB	IS
Vancomycin Fresenius Kabi 500 mg pulver til konsentrat til infusjonsvæske, oppløsning.	PT/H/2179/001	09-6841	FRESENIUS KABI NORGE 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
Vancomycin Fresenius Kabi 500 mg stofn fyrir innrennslisþykkni, lausn.	PT/H/2179/001	IS/1/11/015/01	FRESENIUS KABI AB	IS
Vancomycin Hikma 1000 mg - Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/0771/002	1-31670	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Vancomycin Hikma 1000 mg - Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/0771/002	1-31670	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Vancomycin Hikma 1000 mg por oldatos infúzióhoz	PT/H/0339/002	OGYI-T-21716/03	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	HU
Vancomycin Hikma 1000 mg por oldatos infúzióhoz	PT/H/0339/002	OGYI-T-21716/04	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	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
Vancomycin Hikma 500 mg - Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/0771/001	1-31669	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Vancomycin Hikma 500 mg - Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/0771/001	1-31669	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Vancomycin Hikma 500 mg por oldatos infúzióhoz	PT/H/0339/001	OGYI-T-21716/01	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	HU
Vancomycin Hikma 500 mg por oldatos infúzióhoz	PT/H/0339/001	OGYI-T-21716/02	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	HU
Vancomycin Hydrochloride 1 g Powder for Concentrate for Infusion	not available	PL 04515/0053	HOSPIRA UK LTD	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Hydrochloride 500 mg Powder for Concentrate for Infusion	not available	PL 04515/0053	HOSPIRA UK LTD	UK
Vancomycin Kabi 1 000 mg prášok na infúzny koncentrát	PT/H/2179/002	15/0325/11-S	FRESENIUS KABI S.R.O.	SK
Vancomycin Kabi 1 g Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/2179/002	78732.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Vancomycin Kabi 1 g Pulver zur Herstellung eines Konzentrats für eine Infusionslösung	PT/H/2179/002	2012100184	FRESENIUS KABI DEUTSCHLAND GMBH	LU
Vancomycin Kabi 1000 mg milteliai infuzinio tirpalo koncentratui	PT/H/2179/002	LT/1/11/2505/004	FRESENIUS KABI POLSKA SP. Z O.O.	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
Vancomycin Kabi 1000 mg milteliai infuzinio tirpalo koncentratui	PT/H/2179/002	LT/1/11/2505/002	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Vancomycin Kabi 1000 mg por oldatos infúzióhoz való koncentrátumhoz	PT/H/2179/002	OGYI-T-21953/04	FRESENIUS KABI HUNGARY KFT.	HU
Vancomycin Kabi 1000 mg por oldatos infúzióhoz való koncentrátumhoz	PT/H/2179/002	OGYI-T-21953/02	FRESENIUS KABI HUNGARY KFT.	HU
Vancomycin Kabi 1000 mg prášek pro koncentrát pro infuzní roztok	PT/H/2179/002	15/374/11-C	FRESENIUS KABI S.R.O.	CZ
Vancomycin Kabi 1000 mg pulveris infūziju šķīduma koncentrāta pagatavošanai.	PT/H/2179/002	11-0221	FRESENIUS KABI POLSKA SP. Z O.O.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Kabi 1000 mg, infusioonilahuse kontsentraadi pulber	PT/H/2179/002	733211	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Vancomycin Kabi 500 mg milteliai infuzinio tirpalo koncentratui	PT/H/2179/001	LT/1/11/2505/003	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Vancomycin Kabi 500 mg milteliai infuzinio tirpalo koncentratui	PT/H/2179/001	LT/1/11/2505/001	FRESENIUS KABI POLSKA SP. Z O.O.	LT
Vancomycin Kabi 500 mg por oldatos infúzióhoz való koncentrátumhoz	PT/H/2179/001	OGYI-T-21953/03	FRESENIUS KABI HUNGARY KFT.	HU
Vancomycin Kabi 500 mg por oldatos infúzióhoz való koncentrátumhoz	PT/H/2179/001	OGYI-T-21953/01,03	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
Vancomycin Kabi 500 mg prášek pro koncentrát pro infuzní roztok	PT/H/2179/001	15/373/111-C	FRESENIUS KABI S.R.O.	CZ
Vancomycin Kabi 500 mg prášok na infúzny koncentrát	PT/H/2179/001	15/0324/11-S	FRESENIUS KABI S.R.O.	SK
Vancomycin Kabi 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	PT/H/2179/001	78731.00.00	FRESENIUS KABI DEUTSCHLAND GMBH	DE
Vancomycin Kabi 500 mg Pulver zur Herstellung eines Konzentrats für eine Infusionslösung	PT/H/2179/001	2012100183	FRESENIUS KABI DEUTSCHLAND GMBH	LU
Vancomycin Kabi 500 mg pulveris infūziju šķīduma koncentrāta pagatavošanai.	PT/H/2179/001	11-0222	FRESENIUS KABI POLSKA SP. Z O.O.	LV

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Kabi 500 mg, infusioonilahuse kontsentraadi pulber	PT/H/2179/001	733111	FRESENIUS KABI POLSKA SP. Z O.O.	EE
Vancomycin Kabi, 1000 mg, proszek do sporządzania koncentratu roztworu do infuzji	PT/H/2179/002	18691	FRESENIUS KABI POLSKA SP. Z O.O.	PL
Vancomycin Kabi, 500 mg, proszek do sporządzania koncentratu roztworu do infuzji	PT/H/2179/001	18690	FRESENIUS KABI POLSKA SP. Z O.O.	PL
Vancomycin Lyomark 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	not available	99760.00.00	LYOMARK PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Lyomark 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung oder einer Lösung zum Einnnehmen	not available	6614009.00.00	LYOMARK PHARMA GMBH	DE
Vancomycin MIP 1 000 mg prášok na infúzny roztok	FI/H/0889/002	15/0265/15-S	MIP PHARMA GMBH	SK
Vancomycin MIP 1 000 mg pulver till infusionsvätska, lösning	DE/H/0368/002	21112	MIP PHARMA GMBH	SE
Vancomycin MIP 1000 mg por oldatos infúzióhoz	FI/H/0889/002	OGYI-T-22761/03	MIP PHARMA GMBH	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
Vancomycin MIP 1000 mg por oldatos infúzióhoz	FI/H/0889/002	OGYI-T-22761/04	MIP PHARMA GMBH	HU
Vancomycin MIP 1000 mg pulver til infusjonsvæske, oppløsning	DE/H/0368/002	04-2720	MIP PHARMA GMBH	NO
Vancomycin MIP 500 mg por oldatos infúzióhoz	FI/H/0889/001	OGYI-T-22761/01	MIP PHARMA GMBH	HU
Vancomycin MIP 500 mg por oldatos infúzióhoz	FI/H/0889/001	OGYI-T-22761/02	MIP PHARMA GMBH	HU
Vancomycin MIP 500 mg prášok na infúzny roztok	FI/H/0889/001	15/0264/15-S	MIP PHARMA GMBH	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
Vancomycin MIP 500 mg pulver til infusjonsvæske, oppløsning	DE/H/0368/001	04-2718	MIP PHARMA GMBH	NO
Vancomycin MIP 500 mg pulver till infusionsvätska, lösning	DE/H/0368/001	21111	MIP PHARMA GMBH	SE
Vancomycin MIP Pharma 1000 mg prášek pro infuzní roztok	FI/H/0889/002	15/028/15-C	MIP PHARMA GMBH	CZ
Vancomycin MIP Pharma 1000 mg, Infuusiokuiva-aine, liuosta varten	FI/H/0889/002	28524	MIP PHARMA GMBH	FI
Vancomycin MIP Pharma 1000 mg, pulver till infusionsvätska, lösning	FI/H/0889/002	28524	MIP PHARMA GMBH	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin MIP Pharma 500 mg prášek pro infuzní roztok	FI/H/0889/001	15/027/15-C	MIP PHARMA GMBH	CZ
Vancomycin MIP Pharma 500 mg, Infuusiokuiva-aine, liuosta varten	FI/H/0889/001	28523	MIP PHARMA GMBH	FI
Vancomycin MIP Pharma 500 mg, pulver till infusionsvätska, lösning	FI/H/0889/001	28523	MIP PHARMA GMBH	FI
Vancomycin Mylan 1 000 mg kuiva-aine välikonsentraatiksi infuusionestettä varten, liuos	SE/H/1158/002	30628	MYLAN IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Mylan 1 g prášok na infúzny roztok	CZ/H/0351/002	15/0365/11-S	MYLAN S.A.S	SK
Vancomycin Mylan 1 g, powder for solution for infusion	CZ/H/0351/002	PA0577/163/002	MCDERMOTT LABORATORIES LTD	IE
Vancomycin Mylan 1000 mg milteliai infuzinio tirpalo koncentratui	SE/H/1158/002	LT/1/13/3204/002	MYLAN S.A.S	LT
Vancomycin Mylan 1000 mg prášek pro přípravu infuzního roztoku	CZ/H/0351/002	15/355/11-C	MYLAN S.A.S	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Mylan 1000 mg pulver til konsentrat til infusjonsvæske, oppløsning	SE/H/1158/002	12-8969	MYLAN IRELAND LIMITED	NO
Vancomycin Mylan 1000 mg pulver till koncentrat till infusionsvätska, lösning	SE/H/1158/002	47797	MYLAN IRELAND LIMITED	SE
Vancomycin Mylan 1000 mg pulveris infūziju šķīduma koncentrāta pagatavošanai	SE/H/1158/002	13-0041	MYLAN S.A.S	LV
Vancomycin Mylan 1000 mg, pulver till koncentrat till infusionsvätska, lösning	SE/H/1158/002	30628	MYLAN IRELAND LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VANCOMYCIN MYLAN 1000 mg, κόνις για διάλυμα προς έγχυση	CZ/H/0351/002	21307	MYLAN S.A.S	CY
Vancomycin Mylan 500 mg kuiva-aine välikonsentraatiksi infuusionestettä varten, liuos	SE/H/1158/001	30627	MYLAN IRELAND LIMITED	FI
Vancomycin Mylan 500 mg milteliai infuzinio tirpalo koncentratui	SE/H/1158/001	LT/1/13/3204/001	MYLAN S.A.S	LT
Vancomycin Mylan 500 mg prášek pro infuzní roztok	CZ/H/0351/001	15/354/11-C	MYLAN S.A.S	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Mylan 500 mg prášok na infúzny roztok	CZ/H/0351/001	15/0364/11-S	MYLAN S.A.S	SK
Vancomycin Mylan 500 mg pulver til konsentrat til infusjonsvæske, oppløsning	SE/H/1158/001	12-8968	MYLAN IRELAND LIMITED	NO
Vancomycin Mylan 500 mg pulver till koncentrat till infusionsvätska, lösning	SE/H/1158/001	10413	MYLAN IRELAND LIMITED	SE
Vancomycin Mylan 500 mg pulveris infūziju šķīduma koncentrāta pagatavošanai	SE/H/1158/001	13-0042	MYLAN S.A.S	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
Vancomycin Mylan 500 mg, powder for solution for infusion	CZ/H/0351/001	PA0577/163/001	MCDERMOTT LABORATORIES LTD	IE
Vancomycin Mylan 500 mg, pulver till koncentrat till infusionsvätska, lösning	SE/H/1158/001	30627	MYLAN IRELAND LIMITED	FI
VANCOMYCIN MYLAN 500 mg, κόνις για διάλυμα προς έγχυση	CZ/H/0351/001	21307	MYLAN S.A.S	CY
Vancomycin Noridem 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung und Lösung zum Einnehmen	IE/H/0706/002	1-30790	NORIDEM ENTERPRISES LTD	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
Vancomycin Noridem 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung und Lösung zum Einnehmen	IE/H/0706/002	73936.00.00	NORIDEM ENTERPRISES LTD	DE
Vancomycin Noridem 500 mg Pulver fur ein Konzentrat zur Herstellung einer Infusionslosung und Losung zum Einnehmen	IE/H/0706/001	73935.00.00	NORIDEM ENTERPRISES LTD	DE
Vancomycin Noridem 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung und Lösung zum Einnehmen	IE/H/0706/001	1-30789	NORIDEM ENTERPRISES LTD	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
Vancomycin Olikla 1000 mg prášek pro koncentrát pro infuzní roztok	not available	15/793/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 1000 mg prášek pro koncentrát pro infuzní roztok	not available	15/793/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 1000 mg prášek pro koncentrát pro infuzní roztok	not available	15/793/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 1000 mg prášek pro koncentrát pro infuzní roztok	not available	15/793/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 500 mg prášek pro koncentrát pro infuzní roztok	not available	15/792/16-C	CZ PHARMA 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
Vancomycin Olikla 500 mg prášek pro koncentrát pro infuzní roztok	not available	15/792/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 500 mg prášek pro koncentrát pro infuzní roztok	not available	15/792/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Olikla 500 mg prášek pro koncentrát pro infuzní roztok	not available	15/792/16-C	CZ PHARMA S.R.O.	CZ
Vancomycin Orion 1 000 mg pulver till koncentrat till infusionsvätska, lösning	FI/H/0845/002	31991	ORION CORPORATION	FI
Vancomycin Orion 1000 mg kuiva-aine välikonsentraatiksi infuusionestettä varten, liuos	FI/H/0845/002	31991	ORION CORPORATION	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Orion 1000 mg pulver till koncentrat till infusionsvätska, lösning	FI/H/0845/002	50603	ORION CORPORATION	SE
Vancomycin Orion 500 mg kuiva-aine välikonsentraatiksi infuusionestettä varten, liuos	FI/H/0845/001	31990	ORION CORPORATION	FI
Vancomycin Orion 500 mg pulver till koncentrat till infusionsvätska, lösning	FI/H/0845/001	31990	ORION CORPORATION	FI
Vancomycin Orion 500 mg pulver till koncentrat till infusionsvätska, lösning	FI/H/0845/001	50602	ORION CORPORATION	SE
Vancomycin Pfizer 1000 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	AT/H/0876/002	1-30925	PFIZER CORPORATION AUSTRIA GESELLSCHAFT 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
Vancomycin Pfizer 1000 mg pulver til konsentrat til infusjonsvæske, oppløsning.	AT/H/0876/002	09-6844	PFIZER AS	NO
Vancomycin Pfizer 1000 mg pulveris infūziju šķīduma koncentrāta pagatavošanai.	AT/H/0876/002	11-0170	PFIZER EUROPE MA EEIG	LV
Vancomycin Pfizer 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	AT/H/0876/001	1-30924	PFIZER CORPORATION AUSTRIA GESELLSCHAFT M.B.H.	AT
Vancomycin Pfizer 500 mg pulver til konsentrat til infusjonsvæske, oppløsning.	AT/H/0876/001	09-6843	PFIZER AS	NO
Vancomycin Pfizer 500 mg pulveris infūziju šķīduma koncentrāta pagatavošanai.	AT/H/0876/001	11-0171	PFIZER EUROPE MA EEIG	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
Vancomycin Sandoz 1000 mg infuusiokuiva-aine, liuosta varten	NL/H/4445/002	24472	SANDOZ A/S	FI
Vancomycin Sandoz 1000 mg powder for solution for infusion	NL/H/4445/002	20100578	SANDOZ PHARMACEUTICALS D.D.	BG
Vancomycin Sandoz 1000 mg, infusioonilahuse pulber	NL/H/4445/002	658609	SANDOZ PHARMACEUTICALS D.D.	EE
VANCOMYCIN SANDOZ 1000 MG, PROSZEK DO SPORZĄDZANIA ROZTWORU DO INFUZJI	NL/H/4445/002	17021	SANDOZ GMBH	PL
Vancomycin Sandoz 500 mg infuusiokuiva-aine, liuosta varten	NL/H/4445/001	24471	SANDOZ A/S	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VANCOMYCIN SANDOZ 500 MG, PROSZEK DO SPORZĄDZANIA ROZTWORU DO INFUZJI	NL/H/4445/001	17020	SANDOZ GMBH	PL
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	34093	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 125 mg hårda kapslar	DK/H/2629/001	54527	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 125 mg harde kapsler	DK/H/2629/001	16-11120	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	34094	STRIDES PHARMA (CYPRUS) LIMITED	FI
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 250 mg hårda kapslar	DK/H/2629/002	54528	STRIDES PHARMA (CYPRUS) LIMITED	SE
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	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**
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	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
Vancomycin Strides 250 mg harde kapsler	DK/H/2629/002	16-11121	STRIDES PHARMA (CYPRUS) LIMITED	NO
Vancomycin Xellia 125 mg hårda kapslar	not available	12243	XELLIA PHARMACEUTICALS APS	SE
Vancomycin Xellia 125 mg hörð hylki.	not available	940198	XELLIA PHARMACEUTICALS APS	IS
Vancomycin Xellia 125 mg kapsel, hard	not available	94-1817	XELLIA PHARMACEUTICALS APS	NO
Vancomycin Xellia 125 mg, kapseli, kova	not available	11660	XELLIA PHARMACEUTICALS APS	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin Xellia 250 mg hårda kapslar	not available	12244	XELLIA PHARMACEUTICALS APS	SE
Vancomycin Xellia 250 mg hörð hylki.	not available	940199	XELLIA PHARMACEUTICALS APS	IS
Vancomycin Xellia 250 mg, kapseli, kova	not available	11661	XELLIA PHARMACEUTICALS APS	FI
Vancomycin, 1000 mg Powder for concentrate for solution for infusion	PT/H/0771/002	PL 15413/0008	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	UK
Vancomycin, 1000 mg Powder for concentrate for solution for infusion	PT/H/0771/002	PL 15413/0008	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	UK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycin, 500 mg Powder for concentrate for solution for infusion	PT/H/0771/001	PL 15413/0001	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	UK
Vancomycin, 500 mg Powder for concentrate for solution for infusion	PT/H/0771/001	PL 15413/0001	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	UK
VANCOMYCIN/GENERICS 1 g, κόνις για διάλυμα προς έγχυση	CZ/H/0351/002	73631/ 12-10-2016	GENERICS PHARMA HELLAS LTD	GR
VANCOMYCIN/GENERICS 500 mg, κόνις για διάλυμα προς έγχυση	CZ/H/0351/001	110870/ 14 / 12-10-2016	GENERICS PHARMA HELLAS LTD	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
Vancomycin/Kabi 1000mg κόνις για πυκνό σκεύασμα για παρασκευή διαλύματος προς έγχυση	PT/H/2179/002	68431/15-09-2016	FRESENIUS KABI HELLAS A.E.	GR
Vancomycin/Kabi 500 mg κόνις για πυκνό σκεύασμα για παρασκευή διαλύματος προς έγχυση	PT/H/2179/001	67587/15-09-2016	FRESENIUS KABI HELLAS A.E.	GR
Vancomycin/Noridem 1000 mg Κόνις για πυκνό σκεύασμα για Παρασκευή Διαλύματος προς Έγχυση.	IE/H/0706/002	2827902	NORIDEM ENTERPRISES LTD	GR
Vancomycin/Noridem 500 mg Κόνις για πυκνό σκεύασμα για Παρασκευή Διαλύματος προς Έγχυση	IE/H/0706/001	2827901	NORIDEM ENTERPRISES LTD	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
Vancomycin/Norma® pd.sol.inf. 500mg/vial	not available	44651/12/26-02-2013	NORMA HELLAS S.A.	GR
Vancomycin/Vocate: 1g/Vial κόνις για διάλυμα προς έγχυση	not available	94853/17/01-01-2018	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
Vancomycin/Vocate: 500mg/Vial κόνις για διάλυμα προς έγχυση	not available	93679/16/09-01-2018	VOCATE ΦΑΡΜΑΚΕΥΤΙΚΗ ΑΕ	GR
Vancomycin-CNP 1000 mg pulbere pentru solutie perfuzabila	not available	7508/2015/01	CNP PHARMA GMBH	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
Vancomycin-CNP 1000 mg pulbere pentru solutie perfuzabila	not available	7508/2015/02	CNP PHARMA GMBH	RO
Vancomycin-CNP 500 mg pulbere pentru soluţie perfuzabilă	not available	7507/2015/01	CNP PHARMA GMBH	RO
Vancomycin-CNP 500 mg pulbere pentru soluţie perfuzabilă	not available	7507/2015/02	CNP PHARMA GMBH	RO
Vancomycine Aurobindo 1000 mg, poeder voor concentraat voor oplossing voor infusie	NL/H/4181/002	RVG 113555	AUROBINDO PHARMA 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
Vancomycine Aurobindo 500 mg, poeder voor concentraat voor oplossing voor infusie	NL/H/4181/001	RVG 113830	AUROBINDO PHARMA B.V.	NL
Vancomycine Fresenius Kabi 1000 mg poeder voor concentraat voor oplossing voor infusie	PT/H/2179/002	BE398526	FRESENIUS KABI NV/SA	BE
Vancomycine Fresenius Kabi 1000 mg poeder voor concentraat voor oplossing voor infusie.	PT/H/2179/002	RVG 107977	FRESENIUS KABI NEDERLAND B.V.	NL
Vancomycine Fresenius Kabi 1000 mg poudre pour solution à diluer pour perfusion	PT/H/2179/002	BE398526	FRESENIUS KABI NV/SA	BE
Vancomycine Fresenius Kabi 1000 mg Pulver zur Herstellung eines Konzentrats für eine Infusionslösung	PT/H/2179/002	BE398526	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
Vancomycine Fresenius Kabi 500 mg poeder voor concentraat voor oplossing voor infusie	PT/H/2179/001	BE398517	FRESENIUS KABI NV/SA	BE
Vancomycine Fresenius Kabi 500 mg poeder voor concentraat voor oplossing voor infusie.	PT/H/2179/001	RVG 107976	FRESENIUS KABI NEDERLAND B.V.	NL
Vancomycine Fresenius Kabi 500 mg poudre pour solution à diluer pour perfusion	PT/H/2179/001	BE398517	FRESENIUS KABI NV/SA	BE
Vancomycine Fresenius Kabi 500 mg Pulver zur Herstellung eines Konzentrats für eine Infusionslösung	PT/H/2179/001	BE398517	FRESENIUS KABI NV/SA	BE
VANCOMYCINE HIKMA 1000 mg, Poudre pour solution à diluer pour perfusion	PT/H/0771/002	34009 585 364 10	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	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
VANCOMYCINE HIKMA 1000 mg, Poudre pour solution à diluer pour perfusion	PT/H/0771/002	34009 585 365 88	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
VANCOMYCINE HIKMA 500 mg, Poudre pour solution à diluer pour perfusion	PT/H/0771/001	34009 585 366 4 9	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
VANCOMYCINE HIKMA 500 mg, Poudre pour solution à diluer pour perfusion	PT/H/0771/001	34009 585 367 00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	FR
Vancomycine Hikma, 1000 mg, Poeder voor concentraat voor oplossing voor infusie	PT/H/0771/002	RVG 110476	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Vancomycine Hikma, 1000 mg, Poeder voor concentraat voor oplossing voor infusie	PT/H/0771/002	RVG 110476	HIKMA FARMACÊUTICA (PORTUGAL), 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
Vancomycine Hikma, 500 mg, Poeder voor concentraat voor oplossing voor infusie	PT/H/0771/001	RVG 110475	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Vancomycine Hikma, 500 mg, Poeder voor concentraat voor oplossing voor infusie	PT/H/0771/001	RVG 110475	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
VANCOMYCINE KABI 1 g, poudre pour solution injectable.	not available	34009 563 783 1 9	FRESENIUS KABI FRANCE S.A.S.	FR
VANCOMYCINE KABI 500 mg, poudre pour solution injectable.	not available	34009 563 782 5 8	FRESENIUS KABI FRANCE S.A.S.	FR
VANCOMYCINE MEDIPHA SANTE 1 g, poudre pour solution à diluer pour perfusion	not available	34009 550 451 9 9	MEDIPHA SANTE	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
VANCOMYCINE MEDIPHA SANTE 1 g, poudre pour solution à diluer pour perfusion	not available	34009 550 452 0 5	MEDIPHA SANTE	FR
VANCOMYCINE MEDIPHA SANTE 500 mg, poudre pour solution à diluer pour perfusion	not available	34009 550 451 6 8	MEDIPHA SANTE	FR
VANCOMYCINE MEDIPHA SANTE 500 mg, poudre pour solution à diluer pour perfusion	not available	34009 550 451 7 5	MEDIPHA SANTE	FR
VANCOMYCINE MEDIPHA SANTE 500 mg, poudre pour solution à diluer pour perfusion	not available	34009 550 451 8 2	MEDIPHA SANTE	FR
Vancomycine MIP 1000 mg, poeder voor oplossing voor infusie	FI/H/0889/002	RVG 114472	MIP PHARMA GMBH	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
Vancomycine MIP 1000 mg, poudre pour solution pour perfusion	not available	NL 39083, CIS : 6 500 547 9	MIP PHARMA GMBH	FR
Vancomycine MIP 500 mg, poeder voor oplossing voor infusie	FI/H/0889/001	RVG 114471	MIP PHARMA GMBH	NL
Vancomycine MIP 500 mg, poudre pour solution pour perfusion	not available	NL 39082, CIS : 6 862 482 1	MIP PHARMA GMBH	FR
Vancomycine Mylan 1 g poeder voor oplossing voor infusie	CZ/H/0351/002	BE395437	MYLAN BVBA/SPRL	BE
Vancomycine Mylan 1 g poudre pour solution pour perfusion	CZ/H/0351/002	BE395437	MYLAN BVBA/SPRL	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
Vancomycine Mylan 1 g Pulver zur Herstellung einer Infusionslösung	CZ/H/0351/002	BE395437	MYLAN BVBA/SPRL	BE
Vancomycine Mylan 1 g, poudre pour solution pour perfusion	not available	NL 22242	MYLAN S.A.S	FR
VANCOMYCINE MYLAN 125 mg, poudre pour solution à diluer pour perfusion	not available	NL 20620	MYLAN S.A.S	FR
VANCOMYCINE MYLAN 250 mg, poudre pour solution à diluer pour perfusion	not available	NL 20621	MYLAN S.A.S	FR
Vancomycine Mylan 500 mg poeder voor oplossing voor infusie	CZ/H/0351/001	BE395421	MYLAN BVBA/SPRL	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
Vancomycine Mylan 500 mg poudre pour solution pour perfusion	CZ/H/0351/001	BE395421	MYLAN BVBA/SPRL	BE
Vancomycine Mylan 500 mg Pulver zur Herstellung einer Infusionslösung	CZ/H/0351/001	BE395421	MYLAN BVBA/SPRL	BE
VANCOMYCINE MYLAN 500 mg, poudre pour solution à diluer pour perfusion ou pour solution buvable	not available	NL 17209	MYLAN S.A.S	FR
VANCOMYCINE MYLAN PHARMA 1000 mg, poudre pour solution à diluer pour perfusion	SE/H/1158/002	NL42604	MYLAN S.A.S	FR
VANCOMYCINE MYLAN PHARMA 500 mg, poudre pour solution à diluer pour perfusion	SE/H/1158/001	NL42603	MYLAN 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
Vancomycine Pfizer 1000 mg poeder voor concentraat voor oplossing voor infusie.	AT/H/0876/002	RVG 105372	PFIZER B.V.	NL
Vancomycine Pfizer 500 mg poeder voor concentraat voor oplossing voor infusie.	AT/H/0876/001	RVG 105369	PFIZER B.V.	NL
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 559 330 6 9	SANDOZ	FR
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 560 804 8 9	SANDOZ	FR
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 563 234 8 7	SANDOZ	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 563 235 4 8	SANDOZ	FR
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 566 134 4 1	SANDOZ	FR
VANCOMYCINE SANDOZ 1 g, poudre pour solution à diluer pour perfusion	not available	34009 566 140 4 2	SANDOZ	FR
Vancomycine Sandoz 1000 mg poeder voor oplossing voor infusie	NL/H/4445/002	BE361663	SANDOZ N.V.	BE
Vancomycine Sandoz 1000 mg, poeder voor oplossing voor infusie	NL/H/4445/002	RVG 101509	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
VANCOMYCINE SANDOZ 125 mg, poudre pour solution à diluer pour perfusion	not available	34009 559 334 1 0	SANDOZ	FR
VANCOMYCINE SANDOZ 125 mg, poudre pour solution à diluer pour perfusion	not available	34009 566 131 5 1	SANDOZ	FR
VANCOMYCINE SANDOZ 250 mg, poudre pour solution à diluer pour perfusion	not available	34009 566 132 1 2	SANDOZ	FR
VANCOMYCINE SANDOZ 250 mg, poudre pour solution à diluer pour perfusion	not available	34009 559 332 9 8	SANDOZ	FR
Vancomycine Sandoz 500 mg poeder voor oplossing voor infusie	NL/H/4445/001	BE361654	SANDOZ N.V.	BE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancomycine Sandoz 500 mg, poeder voor oplossing voor infusie	NL/H/4445/001	RVG 101505	SANDOZ B.V.	NL
VANCOMYCINE SANDOZ 500 mg, poudre pour solution pour perfusion	not available	34009 556 527 3 1	SANDOZ	FR
VANCOMYCINE SANDOZ 500 mg, poudre pour solution pour perfusion	not available	34009 566 133 8 0	SANDOZ	FR
Vancomycin-HUMAN 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-7807/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Vancomycin-HUMAN 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-7807/01	TEVA GYÓGYSZERGYÁR ZRT	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
Vancomycin-HUMAN 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-7807/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Vancomycin-HUMAN 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-7807/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Vancomycin-MIP 1000 mg Pulver zur Herstellung einer Infusionslösung	DE/H/0368/002	1-26310	MIP PHARMA AUSTRIA GMBH	AT
Vancomycin–MIP 1000, 1 g, proszek do sporzadzania roztworu do infuzji i roztworu doustnego	not available	10214	MIP PHARMA 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
Vancomycin-MIP 500 mg Pulver zur Herstellung einer Infusionslösung	DE/H/0368/001	1-26309	MIP PHARMA AUSTRIA GMBH	AT
Vancomycin–MIP 500, 500 mg, proszek do sporzadzania roztworu do infuzji i roztworu doustnego	not available	10213	MIP PHARMA POLSKA SP. Z O.O.	PL
Vancomycin-Teva 1000 mg por oldatos infúzióhoz való koncentrátumhoz	DE/H/5737/002	OGYI-T-23255/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Vancomycin-Teva 500 mg por oldatos infúzióhoz való koncentrátumhoz	DE/H/5737/001	OGYI-T-23255/01	TEVA GYÓGYSZERGYÁR ZRT	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
Vanco-ratiopharm® 1000 mg Pulver zur Herstellung einer Infusionslösung	DE/H/5737/002	82752.00.00	RATIOPHARM GMBH	DE
Vanco-ratiopharm® 500 mg Pulver zur Herstellung einer Infusionslösung	DE/H/5737/001	82751.00.00	RATIOPHARM GMBH	DE
Vanco-saar® 1 g Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	not available	49101.01.00	MIP PHARMA GMBH	DE
Vanco-saar® 500 mg Pulver zur Herstellung einer Infusionslösung oder einer Lösung zum Einnehmen	not available	39981.00.00	MIP PHARMA GMBH	DE
Vancosan 1000 mg milteliai infuziniam tirpalui	FI/H/0882/002	LT/1/11/2499/004	MIP PHARMA GMBH	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
Vancosan 1000 mg milteliai infuziniam tirpalui	FI/H/0882/002	LT/1/11/2499/006	MIP PHARMA GMBH	LT
Vancosan 1000 mg milteliai infuziniam tirpalui	FI/H/0882/002	LT/1/11/2499/003	MIP PHARMA GMBH	LT
Vancosan 1000 mg Pulver zur Herstellung einer Infusionslösung	FI/H/0882/002	81575.00.00	CNP PHARMA GMBH	DE
Vancosan 1000 mg pulveris infūziju šķīduma pagatavošanai	FI/H/0882/002	11-0190	MIP PHARMA GMBH	LV
Vancosan 1000 mg, infusioonilahuse pulber	FI/H/0882/002	740011	MIP PHARMA GMBH	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
Vancosan 1000 mg, Infuusiokuiva-aine, liuosta varten	FI/H/0882/002	28526	MIP PHARMA GMBH	FI
Vancosan 1000 mg, pulver till infusionsvätska, lösning	FI/H/0882/002	28526	MIP PHARMA GMBH	FI
Vancosan 500 mg milteliai infuziniam tirpalui	FI/H/0882/001	LT/1/11/2499/002	MIP PHARMA GMBH	LT
Vancosan 500 mg milteliai infuziniam tirpalui	FI/H/0882/001	LT/1/11/2499/005	MIP PHARMA GMBH	LT
Vancosan 500 mg milteliai infuziniam tirpalui	FI/H/0882/001	LT/1/11/2499/001	MIP PHARMA GMBH	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
Vancosan 500 mg Pulver zur Herstellung einer Infusionslösung	FI/H/0882/001	81574.00.00	CNP PHARMA GMBH	DE
Vancosan 500 mg pulveris infūziju šķīduma pagatavošanai	FI/H/0882/001	11-0191	MIP PHARMA GMBH	LV
Vancosan 500 mg, infusioonilahuse pulber	FI/H/0882/001	740111	MIP PHARMA GMBH	EE
Vancosan 500 mg, Infuusiokuiva-aine, liuosta varten	FI/H/0882/001	28525	MIP PHARMA GMBH	FI
Vancosan 500 mg, pulver till infusionsvätska, lösning	FI/H/0882/001	28525	MIP PHARMA GMBH	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vancosan oral 500 mg	not available	49101.00.00	MIP PHARMA GMBH	DE
VANCOTEN 500 mg, Κόνις για διάλυμα προς έγχυση	PT/H/1371/001	76812/19-10-2016	ANFARM HELLAS SA	GR
VANCOTEX 500 mg polvere per soluzione per infusione endovenosa e per soluzione orale	not available	034632036	PHARMATEX ITALIA SRL	IT
VANCOTEX 1 g polvere per soluzione per infusione endovenosa e per soluzione orale 1 flaconcino	not available	034632024	PHARMATEX ITALIA SRL	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
VANCOTEX 500 mg polvere per soluzione per infusione endovenosa e per soluzione orale 1 flaconcino	not available	034632012	PHARMATEX ITALIA SRL	IT
Vangrotin, 500 mg, pó para solução para perfusão	PT/H/1371/001	PT/H/1371/001	ANFARM HELLAS SA	PT
Vankomicin Apta 1000 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/002	H/16/02192/005	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Apta 1000 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/002	H/16/02192/006	APTA MEDICA INTERNACIONAL D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vankomicin Apta 1000 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/002	H/16/02192/007	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Apta 1000 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/002	H/16/02192/008	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Apta 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/001	H/16/02192/001	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Apta 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/001	H/16/02192/002	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Apta 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/001	H/16/02192/003	APTA MEDICA INTERNACIONAL D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vankomicin Apta 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/1401/001	H/16/02192/004	APTA MEDICA INTERNACIONAL D.O.O.	SI
Vankomicin Kabi 1 g prašek za koncentrat za raztopino za infundiranje	PT/H/2179/002	H/11/01613/003	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Vankomicin Kabi 1 g prašek za koncentrat za raztopino za infundiranje	PT/H/2179/002	H/11/01613/004	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Vankomicin Kabi 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/2179/001	H/11/01613/001	FRESENIUS KABI DEUTSCHLAND GMBH	SI
Vankomicin Kabi 500 mg prašek za koncentrat za raztopino za infundiranje	PT/H/2179/001	H/11/01613/002	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
Vankomicin Lek 1000 mg prašek za raztopino za infundiranje	NL/H/4445/002	H/10/01611/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 1000 mg prašek za raztopino za infundiranje	NL/H/4445/002	H/10/01611/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 1000 mg prašek za raztopino za infundiranje	NL/H/4445/002	H/10/01611/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 1000 mg prašek za raztopino za infundiranje	NL/H/4445/002	H/10/01611/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 500 mg prašek za raztopino za infundiranje	NL/H/4445/001	H/10/01611/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Vankomicin Lek 500 mg prašek za raztopino za infundiranje	NL/H/4445/001	H/10/01611/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 500 mg prašek za raztopino za infundiranje	NL/H/4445/001	H/10/01611/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin Lek 500 mg prašek za raztopino za infundiranje	NL/H/4445/001	H/10/01611/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Vankomicin MIP 1000 mg prašak za otopinu za infuziju	FI/H/0889/002	HR-H-126592758	MIP PHARMA CROATIA D.O.O.	HR
Vankomicin MIP 500 mg prašak za otopinu za infuziju	FI/H/0889/001	HR-H-949322881	MIP PHARMA CROATIA 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
Vankomicin Mylan 1000 mg prašek za raztopino za infundiranje	CZ/H/0351/002	H/11/01614/001	MYLAN S.A.S	SI
Vankomicin Mylan 1000 mg prašek za raztopino za infundiranje	CZ/H/0351/002	H/11/01614/002	MYLAN S.A.S	SI
Vankomicin Mylan 1000 mg prašek za raztopino za infundiranje	CZ/H/0351/002	H/11/01614/003	MYLAN S.A.S	SI
Vankomicin Mylan 1000 mg prašek za raztopino za infundiranje	CZ/H/0351/002	H/11/01614/004	MYLAN S.A.S	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
Vankomicin Mylan Pharma 1000 mg prašek za koncentrat za raztopino za infundiranje	SE/H/1158/002	H/13/01612/002	MYLAN S.A.S	SI
Vankomicin Mylan Pharma 500 mg prašek za koncentrat za raztopino za infundiranje	SE/H/1158/001	H/13/01612/001	MYLAN S.A.S	SI
Vankomicin Pfizer 1000 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/002	H/11/01615/005	PFIZER LUXEMBOURG SARL	SI
Vankomicin Pfizer 1000 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/002	H/11/01615/004	PFIZER LUXEMBOURG SARL	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
Vankomicin Pfizer 1000 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/002	H/11/01615/006	PFIZER LUXEMBOURG SARL	SI
Vankomicin Pfizer 500 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/001	H/11/01615/003	PFIZER LUXEMBOURG SARL	SI
Vankomicin Pfizer 500 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/001	H/11/01615/001	PFIZER LUXEMBOURG SARL	SI
Vankomicin Pfizer 500 mg prašek za koncentrat za raztopino za infundiranje	AT/H/0876/001	H/11/01615/002	PFIZER LUXEMBOURG SARL	SI
VONCON® 500 mg Κόνις για διάλυμα προς έγχυση	not available	58858/13/21-11-2014	PHARMASERVE-LILLY SACI	GR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
VONDEM	not available	38055/10	DEMO ABEE	GR
VONDEM	not available	2694202	DEMO ABEE	GR
VOXIN®	not available	AA721/00501	VIANEX S.A.	MT
VOXIN® 500mg/VIAL Κόνις για διάλυμα προς έγχυση	not available	66706/17-12-2019	VIANEX S.A.	GR
VOXIN® Κόνις για διάλυμα προς έγχυση 1g/VIAL	not available	66375/17-12-2019	VIANEX S.A.	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
ZENGAC “500 mg Polvere per soluzione per infusione e per uso orale”	not available	034634030	FISIOPHARMA SRL	IT
ZENGAC 1 g Polvere per soluzione per infusione e per uso orale	not available	034634028	FISIOPHARMA SRL	IT
Ванкомицин Бендалис 1000 mg прах за инфузионен разтвор	not available	20130237	BENDALIS GMBH	BG
Ванкомицин Бендалис 500 mg прах за инфузионен разтвор	not available	20130238	BENDALIS GMBH	BG
Ванкомицин Каби 1000 mg прах за концентрат за инфузионен разтвор	PT/H/2179/002	20130138	FRESENIUS KABI BULGARIA EOOD	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
Ванкомицин Каби 500 mg прах за концентрат за инфузионен разтвор	PT/H/2179/001	20130137	FRESENIUS KABI BULGARIA EOOD	BG
Ванкомицин-MIP 1000 mg прах за инфузионен разтвор	not available	20060707	MIP PHARMA GMBH	BG
Ванкомицин-MIP 500 mg прах за инфузионен разтвор	not available	20060706	MIP PHARMA GMBH	BG
Ванкоцин СР 1 g прах за инфузионен разтвор	not available	20000363	TCHAIKAPHARMA HIGH QUALITY MEDICINES, INC.	BG