30 November 2017
EMA/83934/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): clarithromycin

Procedure No.: PSUSA/00000788/201704

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Product Name (in authorisation country)	MRP / DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Biclar 125 mg/5 ml, granulaat voor orale suspensie	not available	BE165575	MYLAN EPD SPRL	BE
Biclar 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	BE165575	MYLAN EPD SPRL	BE
Biclar 125 mg/5 ml, granulés pour suspension buvable	not available	BE165575	MYLAN EPD SPRL	BE
Biclar 125 mg/5 ml, granulés pour suspension buvable	not available	1994110818	MYLAN EPD SPRL	LU
Biclar 250 mg, comprimés enrobés	not available	BE154086	MYLAN EPD SPRL	BE
Biclar 250 mg, comprimés enrobés	not available	1997124275	MYLAN EPD SPRL	LU
Biclar 250 mg, omhulde tabletten	not available	BE154086	MYLAN EPD SPRL	BE
Biclar 250 mg, überzogene Tabletten	not available	BE154086	MYLAN EPD SPRL	BE
Biclar Forte 500 mg, comprimés enrobés	not available	BE173485	MYLAN EPD SPRL	BE
Biclar Forte 500 mg, comprimés enrobés	not available	1996060324	MYLAN EPD SPRL	LU
Biclar Forte 500 mg, omhulde tabletten	not available	BE173485	MYLAN EPD SPRL	BE
Biclar Forte 500 mg, überzogene Tabletten	not available	BE173485	MYLAN EPD SPRL	BE
Biclar I.V. 500 mg, poeder voor oplossing voor infusie	not available	BE173476	MYLAN EPD SPRL	BE
Biclar I.V. 500 mg, poudre pour solution pour perfusion	not available	BE173476	MYLAN EPD SPRL	BE
Biclar I.V. 500 mg,	not available	1996060326	MYLAN EPD SPRL	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
poudre pour solution pour perfusion				
Biclar I.V. 500 mg, Pulver zur Herstellung einer Infusionslösung	not available	BE173476	MYLAN EPD SPRL	BE
Biclar Kids 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	BE215092	MYLAN EPD SPRL	BE
Biclar Kids 250 mg/5ml, granulaat voor orale suspensie	not available	BE215092	MYLAN EPD SPRL	BE
Biclar Kids 250 mg/5ml, granulés pour suspension buvable	not available	BE215092	MYLAN EPD SPRL	BE
Biclar Kids 250 mg/5ml, granulés pour suspension buvable	not available	2001010001	MYLAN EPD SPRL	LU
BICLAR UNO 500 mg – Filmtabletten mit veränderter Wirkstofffreisetzung	IE/H/0105/001	BE187144	MYLAN EPD SPRL	BE
BICLAR UNO 500 mg – Filmtabletten mit veränderter Wirkstofffreisetzung	IE/H/0105/001	2005106594	MYLAN EPD SPRL	LU
Biclar Uno 500 mg tabletten met gereguleerde afgifte	IE/H/0105/001	BE187144	MYLAN EPD SPRL	BE
Biclar Uno 500mg, comprimés à libération modifiée	IE/H/0105/001	BE187144	MYLAN EPD SPRL	BE
Biclar Uno 500mg, comprimés à libération modifiée	IE/H/0105/001	2005106594	MYLAN EPD SPRL	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
BREVIL OD® 200 mg ?a????a pa?atetaµ???? ap?d?sµe?s??, s?????	FI/H/0624/001	62178/14/09/2010	MEDITRINA PHARMACEUTICAL LIMITED	GR
Clarithomycin 125mg/5ml Suspension	NL/H/2144/001	PL 04416/0609	SANDOZ LTD	UK
Clarithomycin 250mg/5ml Suspension	NL/H/2144/002	PL 04416/0610	SANDOZ LTD	UK
Clarithromycin 1A Pharma 500 mg - Filmtabletten	not available	1-26245	1A PHARMA GMBH	AT
Clarithromycin 250 mg Film-Coated Tablets	not available	PL 11311/0499	TILLOMED LABORATORIES LTD	UK
Clarithromycin 250mg Film-coated Tablets	UK/H/0782/001	PL 04416/0641	SANDOZ LTD	UK
Clarithromycin 500 mg Film-Coated Tablets	not available	PL 11311/0500	TILLOMED LABORATORIES LTD	UK
Clarithromycin Hexal 500 mg – Filmtabletten	not available	1-26248	HEXAL PHARMA GMBH	AT
Clarithromycin Sandoz 125 mg/5 ml – Granulat für orale Suspension	NL/H/2144/001	1-25595	SANDOZ GMBH	AT
Clarithromycin Sandoz 250 mg/5 ml – Granulat für orale Suspension	NL/H/2144/002	1-25596	SANDOZ GMBH	AT
Clarithromycin Sandoz 500 mg - Filmtabletten	not available	1-26244	SANDOZ GMBH	AT
CLARITHROMYCINE BIPHAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345 670 0 1	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345 668 6 8	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE	not available	34009 334 172 4 6	MYLAN MEDICAL 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
BIPHAR 250 mg, comprimé pelliculé				
CLARITHROMYCINE BIPHAR 250 mg, comprimé pelliculé	not available	34009 347 828 0 0	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 250 mg, comprimé pelliculé	not available	34009 347 829 7 8	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 250 mg, comprimé pelliculé	not available	34009 337 224 5 6	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 50 mg/ml, granulés pour suspension buvable	not available	34009 266 833 4 4	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 50 mg/ml, granulés pour suspension buvable	not available	34009 266 834 0 5	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 564 0 6	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 347 830 5 0	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 340 974 1 6	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 347 831 1 1	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 565 7 4	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE	not available	34009 337 566 3 5	MYLAN MEDICAL 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
BIPHAR 500 mg, comprimé pelliculé				
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 568 6 4	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34008 340 975 8 4	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 569 2 5	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 570 0 7	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 571 7 5	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 337 572 3 6	MYLAN MEDICAL SAS	FR
CLARITHROMYCINE BIPHAR 500 mg, comprimé pelliculé	not available	34009 348 736 2 1	MYLAN MEDICAL SAS	FR
Clarithromycine Mylan EPD 125 mg/5 ml, granulaat voor orale suspensie	not available	BE198387	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	BE198387	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD 125 mg/5 ml, granulés pour	not available	BE198387	MYLAN EPD 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
suspension buvable				
Clarithromycine Mylan EPD 125 mg/5 ml, granulés pour suspension buvable	not available	1998090053	MYLAN EPD SPRL	LU
Clarithromycine Mylan EPD 250 mg, comprimés enrobés	not available	BE198396	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD 250 mg, comprimés enrobés	not available	1998090052	MYLAN EPD SPRL	LU
Clarithromycine Mylan EPD 250 mg, filmomhulde tabletten	not available	BE198396	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD 250 mg, überzogene Tabletten	not available	BE198396	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Forte 500 mg, comprimés enrobés	not available	BE198405	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Forte 500 mg, comprimés enrobés	not available	1998090051	MYLAN EPD SPRL	LU
Clarithromycine Mylan EPD Forte 500 mg, filmomhulde tabletten	not available	BE198405	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Forte 500 mg, überzogene Tabletten	not available	BE198405	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD I.V. 500 mg, poeder voor oplossing voor intraveneuze infusie	not available	BE198414	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD I.V. 500 mg, poudre	not available	BE198414	MYLAN EPD 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
pour solution pour perfusion				
Clarithromycine Mylan EPD I.V. 500 mg, poudre pour solution pour perfusion	not available	1998090050	MYLAN EPD SPRL	LU
Clarithromycine Mylan EPD I.V. 500 mg, Pulver zur Herstellung einer Infusionslösung	not available	BE198414	MYLAN EPD SPRL	BE
CLARITHROMYCINE MYLAN EPD UNO 500 mg – Filmtabletten mit veränderter Wirkstofffreisetzung	not available	BE187153	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Uno 500 mg comprimés à libération modifiée	not available	BE187153	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Uno 500 mg tabletten met gereguleerde afgifte	not available	BE187153	MYLAN EPD SPRL	BE
Clarithromycine Mylan EPD Uno 500 mg, comprimés à libération modifiée	not available	2002076594	MYLAN EPD SPRL	LU
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 542 8 3	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 543 4 4	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
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 544 0 5	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 545 7 3	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 546 3 4	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 548 6 3	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 549 2 4	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 550 0 6	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 551 7 4	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à libération modifiée	not available	34009 565 960 8 9	MYLAN S.A.S	FR
CLARITHROMYCINE MYLAN PHARMA 500 mg, comprimé pelliculé à	not available	34009 565 961 4 0	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
libération modifiée				
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258504	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258470	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258462	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258454	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258447	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258439	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258421	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Farmoz OD 500 mg comprimidos de libertação modificada	not available	5258413	FARMOZ - SOCIEDADE TÉCNICO MEDICINAL, S.A.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5248612	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5248620	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg	not available	5250204	TETRAFARMA- 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
comprimidos de libertação modificada				
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250212	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250220	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250238	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250246	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Claritromicina Tetrafarma OD 500 mg comprimidos de libertação modificada	not available	5250253	TETRAFARMA- PRODUTOS FARMACÊUTICOS, LDA.	PT
Clarium 200 mg depotkapseli, kova	FI/H/0624/001	19747	LABORATOIRES SMB S.A.	FI
Heliclar 500 mg, comprimés enrobés	not available	BE173451	MYLAN EPD SPRL	BE
Heliclar 500 mg, comprimés enrobés	not available	1996060325	MYLAN EPD SPRL	LU
Heliclar 500 mg, filmomhulde tabletten	not available	BE173451	MYLAN EPD SPRL	BE
Heliclar 500 mg, überzogene Tabletten	not available	BE173451	MYLAN EPD SPRL	BE
KLACID - Lactobionat 0,5 g - Trockensubstanz zur Infusionsbereitung	not available	1-20013	BGP PRODUCTS 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
KLACID - Lactobionat 0,5 g - Trockensubstanz zur Infusionsbereitung	not available	1-20013	BGP PRODUCTS GES. M. B. H.	AT
KLACID 125 mg / 5 ml - Granulat für orale Suspension	not available	1-19911	BGP PRODUCTS GES. M. B. H.	AT
KLACID 125 mg / 5 ml - Granulat für orale Suspension	not available	1-19911	BGP PRODUCTS GES. M. B. H.	AT
Klacid 125 mg/5 ml granulas iekšķīgi lietojamas suspensijas pagatavošanai	not available	94-0308	BGP PRODUCTS SIA	LV
KLACID 125 mg/5 ml granulát na perorálnu suspenziu	not available	15/0283/97-S	BGP PRODUCTS S.R.O.	SK
KLACID 125 mg/5 ml Granulato per sospensione orale	not available	027370067	BGP PRODUCTS S.R.L.	IT
Klacid 125 mg/5 ml granulátum 100 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/02	MYLAN EPD KFT.	HU
Klacid 125 mg/5 ml granule pentru suspensie orală	not available	8386/2015/01	BGP PRODUCTS S.R.L.	RO
Klacid 125 mg/5 ml granule pentru suspensie orală	not available	8386/2015/02	BGP PRODUCTS S.R.L.	RO
KLACID 125 mg/5 ml Granule pro perorální suspenzi	not available	15/634/94-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
Klacid 125 mg/5 ml granulės geriamajai suspensijai	not available	LT/1/98/0264/005	BGP PRODUCTS SIA	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
Klacid 125 mg/5 ml granulės geriamajai suspensijai	not available	LT/1/98/0264/004	BGP PRODUCTS SIA	LT
Klacid 125 mg/5 ml, suukaudse suspensiooni graanulid	not available	268799	BGP PRODUCTS SIA	EE
Klacid 25 mg/ml granulado para suspensión oral	not available	59462	BGP PRODUCTS OPERATIONS SL	ES
Klacid 25 mg/ml granulat til mikstur	not available	94-1401	BGP PRODUCTS AB	NO
Klacid 25 mg/ml granulat till oral suspension	not available	11987	BGP PRODUCTS AB	SE
KLACID 250 mg - Filmtabletten	not available	1-19163	BGP PRODUCTS GES. M. B. H.	AT
KLACID 250 mg - Filmtabletten	not available	1-19163	BGP PRODUCTS GES. M. B. H.	AT
KLACID 250 mg / 5 ml - Granulat für orale Suspension	not available	1-21047	BGP PRODUCTS GES. M. B. H.	AT
KLACID 250 mg / 5 ml - Granulat für orale Suspension	not available	1-21047	BGP PRODUCTS GES. M. B. H.	AT
Klacid 250 mg apvalkotas tabletes	not available	94-0307	BGP PRODUCTS SIA	LV
KLACID 250 mg Coated Tablets	not available	MA 1064/00901	BGP PRODUCTS LTD.	MT
KLACID 250 mg Compresse rivestite	not available	027370055	BGP PRODUCTS S.R.L.	IT
Klacid 250 mg comprimate filmate	not available	8383/2015/01	BGP PRODUCTS S.R.L.	RO
Klacid 250 mg comprimate filmate	not available	8383/2015/02	BGP PRODUCTS S.R.L.	RO
Klacid 250 mg comprimido revestido	not available	5546684	BGP PRODUCTS UNIPESSOAL, 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
Klacid 250 mg Film-coated Tablets	not available	PA 2010/4/1	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid 250 mg filmtabletta	not available	OGYI-T-2200/05	MYLAN EPD KFT.	HU
Klacid 250 mg filmtabletta	not available	OGYI-T-2200/04	MYLAN EPD KFT.	HU
KLACID 250 mg Granulato per sospensione orale	not available	027370093	BGP PRODUCTS S.R.L.	IT
Klacid 250 mg plėvele dengtos tabletės	not available	LT/1/98/0264/006	BGP PRODUCTS SIA	LT
Klacid 250 mg tabletter	not available	11372	BGP PRODUCTS AB	SE
Klacid 250 mg tabletti, kalvopäällysteinen	not available	11451	BGP PRODUCTS AB	FI
Klacid 250 mg, õhukese polümeerikattega tabletid	not available	268699	BGP PRODUCTS SIA	EE
KLACID 250 mg/5 ml granulát na perorálnu suspenziu	not available	15/0179/99-S	BGP PRODUCTS S.R.O.	SK
KLACID 250 mg/5 ml Granulato per sospensione orale	not available	027370117	BGP PRODUCTS S.R.L.	IT
KLACID 250 mg/5 ml Granule pro perorální suspenzi	not available	15/355/98-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
KLACID 250 Potahované tablety	not available	15/1157/94-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
Klacid 250mg/5ml Granules for Oral Suspension	not available	PA 2010/4/6	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid 50 mg/ml granulado para suspensión oral	not available	60516	BGP PRODUCTS OPERATIONS SL	ES
Klacid 50 mg/ml granulat	not available	97-01645	BGP PRODUCTS AB	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
til mikstur				
Klacid 50 mg/ml por oldatos infúzióhoz való koncentrátumhoz	not available	OGYI-T-2200/10	MYLAN EPD KFT.	HU
Klacid 50 mg/ml rakeet oraalisuspensiota varten	not available	13089	BGP PRODUCTS AB	FI
KLACID 500 mg - Filmtabletten	not available	1-21527	BGP PRODUCTS GES. M. B. H.	AT
KLACID 500 mg - Filmtabletten	not available	1-21527	BGP PRODUCTS GES. M. B. H.	AT
Klacid 500 mg apvalkotās tabletes	not available	10-0609	BGP PRODUCTS SIA	LV
KLACID 500 mg Coated Tablets	not available	MA 1138/00802	MYLAN PRODUCTS LIMITED	MT
KLACID 500 mg Compresse rivestite	not available	027370129	BGP PRODUCTS S.R.L.	IT
Klacid 500 mg comprimido revestido	not available	5546783	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid 500 mg comprimidos recubiertos con película	not available	60515	BGP PRODUCTS OPERATIONS SL	ES
KLACID 500 mg filmom obalené tablety	not available	15/0215/00-S	BGP PRODUCTS S.R.O.	SK
Klacid 500 mg filmtabletta	not available	OGYI-T-2200/06	MYLAN EPD KFT.	HU
Klacid 500 mg filmuhúðaðar töflur	not available	930158 (IS)	BGP PRODUCTS AB	IS
Klacid 500 mg granulado para suspensión oral	not available	60518	BGP PRODUCTS OPERATIONS SL	ES
KLACID 500 mg Granulato per sospensione orale	not available	027370105	BGP PRODUCTS S.R.L.	IT
Klacid 500 mg infuusiokuiva-aine, liuosta varten	not available	14371	BGP PRODUCTS AB	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
Klacid 500 mg liofilizado para solución para perfusión	not available	60202	BGP PRODUCTS OPERATIONS SL	ES
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/009	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/010	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/011	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/012	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/013	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/014	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/015	BGP PRODUCTS SIA	LT
Klacid 500 mg plėvele dengtos tabletės	not available	LT/1/98/0264/008	BGP PRODUCTS SIA	LT
Klacid 500 mg tabletter	not available	12164	BGP PRODUCTS AB	SE
Klacid 500 mg tabletti, kalvopäällysteinen	not available	11923	BGP PRODUCTS AB	FI
KLACID 500 mg/10 ml Polvere e solvente per soluzione per infusione	not available	027370042	BGP PRODUCTS S.R.L.	IT
KLACID 500 Potahované tablety	not available	15/374/97-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
Klacid Baby 125 mg/5 ml granulátum 60 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/01	MYLAN EPD KFT.	HU
Klacid Forte 500mg Film-coated Tablets	not available	PA 2010/4/2	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid i.v. 500 mg milteliai infuziniam	not available	LT/1/98/0264/007	BGP PRODUCTS SIA	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
tirpalui				
KLACID I.V. 500 mg Prášek pro infuzní roztok	not available	15/236/99-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
KLACID I.V. 500 mg prášok na infúzny roztok	not available	15/0010/99-S	BGP PRODUCTS S.R.O.	SK
Klacid i.v. 500 mg pulveris infūziju šķīduma pagatavošanai	not available	94-0306	BGP PRODUCTS SIA	LV
Klacid i.v., 500 mg infusioonilahuse pulber	not available	312800	BGP PRODUCTS SIA	EE
Klacid I.V.500 mg pulbere pentru soluţie perfuzabilă	not available	8385/2015/01	MYLAN MEDICAL SAS	RO
Klacid IV 500mg Powder for Concentrate for Solution for Infusion	not available	PA 2010/4/3	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid Kid 250 mg/5 ml granulátum 70 ml belsőleges szuszpenzióhoz	not available	OGYI-T-2200/03	MYLAN EPD KFT.	HU
Klacid LA 500mg Modified release tablets	IE/H/0105/001	PA 2010/4/4	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518784	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519386	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055687	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055182	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg	IE/H/0105/001	4055281	BGP PRODUCTS	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos de libertação modificada			UNIPESSOAL, LDA.	
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519287	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519188	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627586	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055083	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056289	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056081	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4056180	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055380	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519485	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4054987	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519584	BGP PRODUCTS UNIPESSOAL, 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
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055984	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055885	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518883	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3518982	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627487	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519089	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	2627685	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519683	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519782	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	3519881	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055489	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de	IE/H/0105/001	2627784	BGP PRODUCTS UNIPESSOAL, 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
libertação modificada				
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055588	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg comprimidos de libertação modificada	IE/H/0105/001	4055786	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid OD 500 mg depottabletter	not available	97-2350	BGP PRODUCTS AB	NO
Klacid OD 500 mg säädellysti vapauttava tabletti	IE/H/0105/001	12843	BGP PRODUCTS AB	FI
Klacid Paediatric Suspension 125mg/5ml, Granules for Oral Suspension	not available	PA 2010/4/5	MYLAN IRE HEALTHCARE LIMITED	IE
Klacid Pediátrico 25 mg/ml granulado para suspensão oral	not available	2146983	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid Pediátrico 50 mg/ml granulado para suspensão oral	not available	2525384	BGP PRODUCTS UNIPESSOAL, LDA.	PT
Klacid RM 500 mg Compresse a rilascio modificato	not available	027370143	BGP PRODUCTS S.R.L.	IT
Klacid Saft® Forte 250 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	40373.00.00	MYLAN HEALTHCARE GMBH	DE
Klacid Saft®, 125 mg/5 ml, Granulat zur Herstellung einer Suspension zum Einnehmen	not available	19219.00.01	MYLAN HEALTHCARE 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
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/01	MYLAN PRODUCTS LIMITED	RO
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/02	MYLAN PRODUCTS LIMITED	RO
Klacid SR 500 mg comprimate cu eliberare prelungită	not available	8384/2015/03	MYLAN PRODUCTS LIMITED	RO
Klacid SR 500 mg ilgstošās darbības tabletes	not available	98-0691	BGP PRODUCTS SIA	LV
Klacid SR 500 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/98/0264/002	BGP PRODUCTS SIA	LT
Klacid SR 500 mg modifikuoto atpalaidavimo tabletės	not available	LT/1/98/0264/003	BGP PRODUCTS SIA	LT
KLACID SR 500 mg tablety s riadeným uvoľňovaním	not available	15/0141/99-S	BGP PRODUCTS S.R.O.	SK
KLACID SR 500 mg Tablety s řízeným uvolňováním	not available	15/062/98-C	BGP PRODUCTS CZECH REPUBLIC S.R.O.	CZ
Klacid SR 500 mg, toimeainet modifitseeritult vabastavad tabletid	not available	268999	BGP PRODUCTS SIA	EE
Klacid SR, 500 mg tablet met gereguleerde afgifte	IE/H/0105/001	RVG 21555	BGP PRODUCTS B.V.	NL
Klacid unidía 500 mg comprimidos de liberación modificada	not available	63380	BGP PRODUCTS OPERATIONS SL	ES
Klacid Uno	IE/H/0105/001	18861	BGP PRODUCTS AB	DK
Klacid Uno	IE/H/0105/001	18861	BGP PRODUCTS AB	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
Klacid Uno – Filmtabletten mit veränderter Wirkstoff-Freisetzung	IE/H/0105/001	1-22085	BGP PRODUCTS GES. M. B. H.	AT
Klacid UNO 500 mg retard filmtabletta	not available	OGYI-T-2200/08	MYLAN EPD KFT.	HU
Klacid Uno, 500 mg, tabletki o zmodyfikowanym uwalnianiu	not available	7930	BGP PRODUCTS POLAND SP. Z.O.O.	PL
KLACID XL 500 mg Modified-Release Coated Tablets	not available	MA 1064/00903	BGP PRODUCTS LTD.	MT
Klacid, 125 mg/5 ml, granulat do sporządzania zawiesiny doustnej	not available	R/3317	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Klacid, 250 mg, tabletki powlekane	not available	R/3318	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Klacid, 250 mg/5 ml, granulat do sporządzania zawiesiny doustnej	not available	7806	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Klacid, 500 mg õhukese polümeerikattega tabletid	not available	751011	BGP PRODUCTS SIA	EE
Klacid, 500 mg, proszek do sporządzania roztworu do infuzji	not available	R/3757	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Klacid, 500 mg, tabletki powlekane	not available	R/7194	BGP PRODUCTS POLAND SP. Z.O.O.	PL
Klacid, filmovertrukne tabletter	not available	15527	BGP PRODUCTS AB	DK
Klacid, filmovertrukne tabletter	not available	13941	BGP PRODUCTS AB	DK
Klacid, granulat til oral suspension	not available	18018	BGP PRODUCTS AB	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
Klacid, granulat til oral suspension	not available	15382	BGP PRODUCTS AB	DK
Klacid, pulver til infusionsvæske, opløsning	not available	30742	BGP PRODUCTS AB	DK
Klacid® 500 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung	not available	46110.00.00	MYLAN HEALTHCARE GMBH	DE
Klacid® 500 mg Pulver und Lösungsmittel für ein Konzentrat zur Herstellung einer Infusionslösung	not available	93335.00.00	MYLAN HEALTHCARE GMBH	DE
Klacid® Filmtabletten, 250 mg	not available	19219.00.00	MYLAN HEALTHCARE GMBH	DE
Klacid® Pro, 250 mg Filmtabletten	not available	28401.00.00	MYLAN HEALTHCARE GMBH	DE
Klacid® Uno, 500 mg Retardtablette	not available	61478.00.00	MYLAN HEALTHCARE GMBH	DE
Klaricid 250 mg Tablets	not available	PL 46302/0016	MYLAN PRODUCTS LIMITED	UK
Klaricid 500 mg Tablets	not available	PL 46302/0017	MYLAN PRODUCTS LIMITED	UK
Klaricid Adult Sachet 250mg	not available	PL 46302/0015	MYLAN PRODUCTS LIMITED	UK
Klaricid IV 500 mg	not available	PL 46302/0018	MYLAN PRODUCTS LIMITED	UK
Klaricid Paediatric Suspension 125mg/5ml	not available	PL 46302/0014	MYLAN PRODUCTS LIMITED	UK
Klaricid Paediatric Suspension 250mg/5ml	not available	PL 46302/0019	MYLAN PRODUCTS LIMITED	UK
Klaricid XL 500 mg Tablets	not available	PL 46302/0020	MYLAN PRODUCTS LIMITED	UK
KLARICID®	not available	36466/10/18-03-2011	BGP PRODUCTS LTD (GREECE)	GR
KLARICID®	not available	36469/10/18-03-2011	BGP PRODUCTS LTD (GREECE)	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
KLARICID®	not available	36464/10/18-03-2011	BGP PRODUCTS LTD (GREECE)	GR
KLARICID®	not available	36468/10/18-03-2011	BGP PRODUCTS LTD (GREECE)	GR
KLARICID® OD	not available	36472/10/18-03-2011	BGP PRODUCTS LTD (GREECE)	GR
Kofron 25 mg/ml granulado para suspensión oral	not available	59464	GUIDOTTI FARMA, S.L.U.	ES
Kofron 50 mg/ml granulado para suspensión oral	not available	60592	GUIDOTTI FARMA, S.L.U.	ES
Kofron 500 mg granulado para suspensión oral	not available	60593	GUIDOTTI FARMA, S.L.U.	ES
Kofron unidía 500 mg comprimidos de liberación modificada	not available	63.383	GUIDOTTI FARMA, S.L.U.	ES
LEKOKLAR 250 mg/5 ml, granule pentru suspensie orală	NL/H/2099/002	4386/2012/01	S.C. SANDOZ S.R.L.	RO
LEKOKLAR 250 mg/5 ml, granule pentru suspensie orală	NL/H/2099/002	4386/2012/02	S.C. SANDOZ S.R.L.	RO
LEKOKLAR 250 mg/5 ml, granule pentru suspensie orală	NL/H/2099/002	4386/2012/03	S.C. SANDOZ S.R.L.	RO
LEKOKLAR 250 mg/5 ml, granule pentru suspensie orală	NL/H/2099/002	4386/2012/04	S.C. SANDOZ S.R.L.	RO
Lekoklar 500 mg film-coated tablets	NL/H/2087/002	20110649	SANDOZ PHARMACEUTICALS D.D.	BG
MACLADIN 125 mg/5 ml Granulato per sospensione orale	not available	027530068	LABORATORI GUIDOTTI 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
MACLADIN 250 mg Compresse rivestite	not available	027530056	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 250 mg Granulato per sospensione orale	not available	027530094	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 250 mg/5 ml Granulato per sospensione orale	not available	027530120	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg Compresse rivestite	not available	027530118	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg Granulato per sospensione orale	not available	027530106	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN 500 mg/10 ml Polvere e solvente per soluzione per infusione	not available	027530043	LABORATORI GUIDOTTI S.P.A.	IT
MACLADIN RM 500 mg Compresse a rilascio modificato	not available	027530144	LABORATORI GUIDOTTI S.P.A.	IT
Maclar 500 mg, comprimés enrobés	not available	BE173467	MYLAN EPD SPRL	BE
Maclar 500 mg, comprimés enrobés	not available	1996060327	MYLAN EPD SPRL	LU
Maclar 500 mg, filmomhulde tabletten	not available	BE173467	MYLAN EPD SPRL	BE
Maclar 500 mg, überzogene Tabletten	not available	BE173467	MYLAN EPD SPRL	BE
MAKCIN 500 mg filmom obložene tablete	not available	HR-H-696573111	BELUPO D.D.	HR
MAKCIN SR 500 mg tablete s produljenim oslobađanjem	not available	UP/I-530-09/12-02/458	BELUPO D.D.	HR
Monoclarium 200 mg Hartkapsel, retardiert	FI/H/0624/001	BE312417	LABORATOIRES SMB S.A.	BE
Monoclarium 200 mg	FI/H/0624/001	2008110062	LABORATOIRES SMB 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
Hartkapsel, retardiert				
Monoclarium 200 mg, gélule à libération prolongée	FI/H/0624/001	BE312417	LABORATOIRES SMB S.A.	BE
Monoclarium 200 mg, gélule à libération prolongée	FI/H/0624/001	2008110062	LABORATOIRES SMB S.A.	LU
Monoclarium 200 mg, harde capsules met vertraagde afgifte	FI/H/0624/001	BE312417	LABORATOIRES SMB S.A.	BE
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 533 9 2	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 565 957 7 8	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 565 958 3 9	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 539 7 2	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 537 4 3	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 534 5 3	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 535 1 4	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 536 8 2	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg,	not available	34009 365 541 1 5	MYLAN MEDICAL 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
comprimé pelliculé à libération modifiée				
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 540 5 4	MYLAN MEDICAL SAS	FR
MONOZECLAR 500 mg, comprimé pelliculé à libération modifiée	not available	34009 365 538 0 4	MYLAN MEDICAL SAS	FR
SORICLAR 125 mg/5 ml Granulato per sospensione orale	not available	037456011	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 125 mg/5 ml Granulato per sospensione orale	not available	037456011	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg Compresse rivestite	not available	037456050	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg Compresse rivestite	not available	037456050	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg Granulato per sospensione orale	not available	037456086	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg Granulato per sospensione orale	not available	037456086	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg/5 ml Granulato per sospensione orale	not available	037456023	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 250 mg/5 ml Granulato per sospensione orale	not available	037456023	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 500 mg Compresse rivestite	not available	037456062	ABIOGEN PHARMA S.P.A.	IT
SORICLAR 500 mg Compresse rivestite	not available	037456062	ABIOGEN PHARMA S.P.A.	IT
SORICLAR RM 500 mg	not available	037456098	ABIOGEN PHARMA 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
Compresse a rilascio modificato				
SORICLAR RM 500 mg Compresse a rilascio modificato	not available	037456098	ABIOGEN PHARMA S.P.A.	IT
VECLAM 125 mg/5 ml Granulato per sospensione orale	not available	027529041	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg Compresse rivestite	not available	027529054	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg Granulato per sospensione orale	not available	027529080	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 250 mg/5 ml Granulato per sospensione orale	not available	027529104	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 500 mg Compresse rivestite	not available	027529116	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 500 mg Granulato per sospensione orale	not available	027529092	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM 500 mg/10 ml Polvere e solvente per soluzione per infusione	not available	027529039	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
VECLAM RM 500 mg Compresse a rilascio modificato	not available	027529130	MALESCI ISTITUTO FARMACOBIOLOGICO - S.P.A.	IT
WINCLAR 250 mg/5 mL granulato per sospensione orale	not available	037446022	I.B.N. SAVIO S.R.L.	IT
WINCLAR 500 mg Compresse rivestite	not available	037446061	I.B.N. SAVIO S.R.L.	IT
ZECLAR 0,5 g, poudre	not available	34009 3558435 9	MYLAN MEDICAL 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
pour solution pour perfusion				
ZECLAR 0,5 g, poudre pour solution pour perfusion	not available	34009 3464390 3	MYLAN MEDICAL SAS	FR
ZECLAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345653 9 7	MYLAN MEDICAL SAS	FR
ZECLAR 25 mg/ml, granulés pour suspension buvable	not available	34009 345655 1 9	MYLAN MEDICAL SAS	FR
Zeclar 250 mg tabletti, kalvopäällysteinen	not available	12341	ORION OYJ	FI
Zeclar 250 mg tabletti, kalvopäällysteinen	not available	12341	ORION OYJ	FI
ZECLAR 250 mg, comprimé pelliculé	not available	34009 3448445 2	MYLAN MEDICAL SAS	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 3448451 3	MYLAN MEDICAL SAS	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 3340311 9	MYLAN MEDICAL SAS	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 3448468 1	MYLAN MEDICAL SAS	FR
ZECLAR 250 mg, comprimé pelliculé	not available	34009 3448474 2	MYLAN MEDICAL SAS	FR
Zeclar 50 mg/ml rakeet oraalisuspensiota varten	not available	13090	ORION OYJ	FI
Zeclar 50 mg/ml rakeet oraalisuspensiota varten	not available	13090	ORION OYJ	FI
ZECLAR 50 mg/ml, granulés pour suspension buvable	not available	34009 347 855 8 0	MYLAN MEDICAL SAS	FR
ZECLAR 50 mg/ml, granulés pour suspension buvable	not available	34009 347 853 5 1	MYLAN MEDICAL 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
Zeclar 500 mg tabletti, kalvopäällysteinen	not available	12342	ORION OYJ	FI
Zeclar 500 mg tabletti, kalvopäällysteinen	not available	12342	ORION OYJ	FI
ZECLAR 500 mg, comprimé pelliculé	not available	34009 346 497 0 7	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 280 6 6	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 281 2 7	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 013 8 0	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 303 6 6	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 207 3 5	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 346 498 7 5	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 205 0 6	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 338 279 8 4	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 204 4 5	MYLAN MEDICAL SAS	FR
ZECLAR 500 mg, comprimé pelliculé	not available	34009 337 206 7 4	MYLAN MEDICAL SAS	FR
Zeclar OD 500 mg säädellysti vapauttavat tabletit	not available	12844	ORION CORPORATION	FI
Zeclar OD 500 mg säädellysti vapauttavat tabletit	not available	12844	ORION CORPORATION	FI
Клацид 125 mg/5 ml гранули за перорална суспензия	not available	20010617	MYLAN PRODUCTS LIMITED	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
Клацид 250 mg филмирани таблетки	not available	20010734	MYLAN PRODUCTS LIMITED	BG
Клацид 250 mg/5 ml гранули за перорална суспензия	not available	20050512	MYLAN PRODUCTS LIMITED	BG
Клацид 500 mg прах за инфузионен разтвор	not available	9700113	MYLAN PRODUCTS LIMITED	BG
Клацид SR 500 mg таблетки с изменено освобождаване	not available	20010517	MYLAN PRODUCTS LIMITED	BG