25 June 2026
EMADOC-1700519818-3394728
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): terbinafine

EURD list No. PSUSA/00002896/202509

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Lamisil 1% - Creme	not available	1-19581	KARO HEALTHCARE AB	AT
Terbinafin Moberg Pharma 98 mg/ml Lösung zur Anwendung auf der Haut	SE/H/2249/001	141820	MOBERG PHARMA AB	AT
Terbinafin Moberg Pharma 98 mg/ml Lösung zur Anwendung auf der Haut	SE/H/2249/001	141820	MOBERG PHARMA AB	AT
Terbinafin Moberg Pharma 98 mg/ml Lösung zur Anwendung auf der Haut	SE/H/2249/001	141820	MOBERG PHARMA AB	AT
Terbinafin Moberg Pharma 98 mg/ml Lösung zur Anwendung auf der Haut	SE/H/2249/001	141820	MOBERG PHARMA AB	AT
LamisilOnce 1 % Lösung zur einmaligen Anwendung auf der Haut	SE/H/0992/004	1-26665	KARO HEALTHCARE AB	AT
Lamisil 250 mg Tabletten	not available	BE155705	NOVARTIS PHARMA N.V.	BE
Lamisil 250 mg comprimés	not available	BE155705	NOVARTIS PHARMA N.V.	BE
Lamisil 250 mg tabletten	not available	BE155705	NOVARTIS PHARMA N.V.	BE
Lamisil 1% Creme	not available	BE457351	KARO HEALTHCARE AB	BE
Lamisil 1% Creme	not available	BE159092	KARO HEALTHCARE AB	BE
Lamisil 1% crème	not available	BE159092	KARO HEALTHCARE AB	BE
Lamisil 1% crème	not available	BE457351	KARO HEALTHCARE AB	BE
Lamisil 1% crème	not available	BE159092	KARO HEALTHCARE AB	BE
Lamisil 1% crème	not available	BE457351	KARO HEALTHCARE AB	BE
Terbinafine Moberg Pharma 98 mg/ml solution pour application cutanée	SE/H/2249/001	BE662330	MOBERG PHARMA AB	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
Terbinafine Moberg Pharma 98 mg/ml solution pour application cutanée	SE/H/2249/001	BE662329	MOBERG PHARMA AB	BE
Terbinafine Moberg Pharma 98 mg/ml solution pour application cutanée	SE/H/2249/001	BE662330	MOBERG PHARMA AB	BE
Terbinafine Moberg Pharma 98 mg/ml solution pour application cutanée	SE/H/2249/001	BE662329	MOBERG PHARMA AB	BE
LAMISIL ONCE 1 % , Lösung zur Anwendung auf der Haut	SE/H/0992/004	BE291322	KARO HEALTHCARE AB	BE
Lamisil Once 1 %, solution pour application cutanée	SE/H/0992/004	BE291322	KARO HEALTHCARE AB	BE
Lamisil Once 1%, oplossing voor cutaan gebruik.	SE/H/0992/004	BE291322	KARO HEALTHCARE AB	BE
Lamisil Dermgel 1%, gel	not available	BE283464	KARO HEALTHCARE AB	BE
Lamisil Dermgel 1%, gel	not available	BE283464	KARO HEALTHCARE AB	BE
LAMISIL DERMGEL 1%, Gel	not available	BE283464	KARO HEALTHCARE AB	BE
Lamisil Dermgel 1%, gel	not available	BE474035	KARO HEALTHCARE AB	BE
Lamisil Dermgel 1%, gel	not available	BE474035	KARO HEALTHCARE AB	BE
LAMISIL DERMGEL 1%, Gel	not available	BE474035	KARO HEALTHCARE AB	BE
Terbinafine Viatris 1% crème	NL/H/4613/001	BE365346	VIATRIS GX	BE
Terbinafine Viatris 1% crème	NL/H/4613/001	BE365346	VIATRIS GX	BE
Terbinafine Viatris 1% Creme	NL/H/4613/001	BE365346	VIATRIS GX	BE
ЛАМИЗИЛ 1% Спрей за	not available	II-6055/23.10.2009	KARO HEALTHCARE AB	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
кожа, разтвор				
Exoterbyn	DE/H/6404/001	20200234	SANDOZ PHARMACEUTICALS D.D.	BG
ЛАМИЗИЛ 1% Крем	not available	20020380	KARO HEALTHCARE AB	BG
Terbinafin Sandoz 78 mg/ml léčivý lak na nehty	DE/H/6404/001	26/271/19-C	SANDOZ S.R.O.	CZ
Terbinafin Moberg Pharma 98 mg/ml kožní roztok	SE/H/2249/001	26/111/22-C 0260514	MOBERG PHARMA AB	CZ
Terbinafin Moberg Pharma 98 mg/ml kožní roztok	SE/H/2249/001	26/111/22-C 0260517	MOBERG PHARMA AB	CZ
Terbinafin Moberg Pharma 98 mg/ml kožní roztok	SE/H/2249/001	26/111/22-C 0260514	MOBERG PHARMA AB	CZ
Terbinafin Moberg Pharma 98 mg/ml kožní roztok	SE/H/2249/001	26/111/22-C 0280150	MOBERG PHARMA AB	CZ
Lamisil 10 mg/g krém	not available	26/417/91-S/C	KARO HEALTHCARE AB	CZ
LAMISIL 250 mg tablety	not available	26/418/91-B/C	NOVARTIS S.R.O.,	CZ
LAMISIL 125 mg tablety	not available	26/418/91-A/C	NOVARTIS S.R.O.,	CZ
LAMISIL 250 mg tablety	not available	26/418/91-B/C	NOVARTIS S.R.O.,	CZ
Terbiderm Creme 10 mg/g Creme	DE/H/6469/001	83083.00.00	DERMAPHARM AG	DE
Terbiderm Creme	DE/H/6469/001/DC	83083.00.00	DERMAPHARM AG	DE
Terbinafin - 1 A Pharma Nagellack gegen Nagelpilz 78,22 mg/ml wirkstoffhaltiger Nagellack	DE/H/6404/001	2204412.00.00	1 A PHARMA GMBH	DE
Terbiderm Spray, 10 mg/g für Erwachsene	not available	66032.00.00	DERMAPHARM AG	DE
Terbiderm Spray, 10 mg/g für Erwachsene	not available	66032.00.00	DERMAPHARM AG	DE
Lamisil Spray 1 % Spray	SE/H/0992/002	41418.00.00	KARO HEALTHCARE AB	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
zur Anwendung auf der Haut, Lösung				
Lamisil Once 1 % Lösung zur Anwendung auf der Haut	SE/H/0992/004	65563.00.00	KARO HEALTHCARE AB	DE
Lamisil Creme 10 mg/g	not available	26373.00.00	KARO HEALTHCARE AB	DE
Terbiderm Gel	not available	66031.00.00	DERMAPHARM AG	DE
Lamisil, creme	not available	14096	KARO HEALTHCARE AB	DK
Lamisil, gel	SE/H/0992/003	31237	KARO HEALTHCARE AB	DK
Terbinafin "Moberg Pharma", kutanopløsning	SE/H/2249/001	67637	MOBERG PHARMA AB	DK
Terbinafin "Moberg Pharma", kutanopløsning	SE/H/2249/001	67637	MOBERG PHARMA AB	DK
Terbinafin "Moberg Pharma", kutanopløsning	SE/H/2249/001	67637	MOBERG PHARMA AB	DK
Terbinafin "Moberg Pharma", kutanopløsning	SE/H/2249/001	67637	MOBERG PHARMA AB	DK
Terclara, kutanopløsning	SE/H/2250/001	67638	MOBERG PHARMA AB	DK
Terclara, kutanopløsning	SE/H/2250/001	67638	MOBERG PHARMA AB	DK
Terclara, kutanopløsning	SE/H/2250/001	67638	MOBERG PHARMA AB	DK
Terclara, kutanopløsning	SE/H/2250/001	67638	MOBERG PHARMA AB	DK
Lamisil Once, kutanopløsning	SE/H/0992/004	39088	KARO HEALTHCARE AB	DK
Exotafin 78,22 mg/ml ravimküünelakk	DE/H/6404/001	1039021	SANDOZ PHARMACEUTICALS D.D.	EE
Lamisil DermGel 10 mg/g geel	SE/H/0992/003	266699	KARO HEALTHCARE AB	EE
Lamisil Uno, 1% nahalahus	SE/H/0992/004	520606	KARO HEALTHCARE AB	EE
Lamisil, 10 mg/g kreem	not available	215998	KARO HEALTHCARE AB	EE
Lamisil 250 mg comprimidos	not available	59.435	NOVARTIS FARMACÉUTICA S.A.	ES
Lamisil 250 mg comprimidos	not available	59.435	NOVARTIS FARMACÉUTICA S.A.	ES
Lamisil 10 mg/g crema	not available	59.434	NOVARTIS FARMACÉUTICA	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
			S.A.	
Lamisil 10 mg/g solución para pulverización cutánea	not available	62.783	NOVARTIS FARMACÉUTICA S.A.	ES
Terbinafina Moberg Pharma 98 mg/ml solución cutánea	SE/H/2249/001	89103	MOBERG PHARMA AB	ES
Terbinafina Moberg Pharma 98 mg/ml solución cutánea	SE/H/2249/001	89103	MOBERG PHARMA AB	ES
Terbinafina Moberg Pharma 98 mg/ml solución cutánea	SE/H/2249/001	89103	MOBERG PHARMA AB	ES
Terbinafina Moberg Pharma 98 mg/ml solución cutánea	SE/H/2249/001	89103	MOBERG PHARMA AB	ES
Lamisil DermGel 1 % geeli	SE/H/0992/003	13752	KARO HEALTHCARE AB	FI
Lamisil DermGel 1 % gel	SE/H/0992/003	13752	KARO HEALTHCARE AB	FI
Terbinafin Sandoz 78,22 mg/ml lääkekynsilakka	DE/H/6404/001	37279	SANDOZ A/S	FI
Lamisil 1 % sumute iholle, liuos	SE/H/0992/002	12945	KARO HEALTHCARE AB	FI
Lamisil 1 % kutan spray, lösning	SE/H/0992/002	12945	KARO HEALTHCARE AB	FI
Terbinafin Moberg Pharma 98 mg/ml liuos iholle	SE/H/2249/001	40845	MOBERG PHARMA AB	FI
Terbinafin Moberg Pharma 98 mg/ml liuos iholle	SE/H/2249/001	40845	MOBERG PHARMA AB	FI
Terbinafin Moberg Pharma 98 mg/ml liuos iholle	SE/H/2249/001	40845	MOBERG PHARMA AB	FI
Terbinafin Moberg Pharma 98 mg/ml liuos	SE/H/2249/001	40845	MOBERG PHARMA 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
iholle				
Terclara 98 mg/ml liuos iholle	SE/H/2250/001	40843	MOBERG PHARMA AB	FI
Terclara 98 mg/ml liuos iholle	SE/H/2250/001	40843	MOBERG PHARMA AB	FI
Terclara 98 mg/ml liuos iholle	SE/H/2250/001	40843	MOBERG PHARMA AB	FI
Terclara 98 mg/ml liuos iholle	SE/H/2250/001	40843	MOBERG PHARMA AB	FI
Lamisil Solo 1 % liuos iholle	SE/H/0992/004	21966	KARO HEALTHCARE AB	FI
Lamisil Solo 1 %, kutan lösning	SE/H/0992/004	21966	KARO HEALTHCARE AB	FI
Lamisil 1 % emulsiovoide	not available	11046	KARO HEALTHCARE AB	FI
Lamisil 1 % kräm	not available	11046	KARO HEALTHCARE AB	FI
LAMISILDERMGEL 1 %, gel	SE/H/0992/003	34009 350 016 3 4	KARO HEALTHCARE AB	FR
LAMISILDERMGEL 1 %, gel	SE/H/0992/003	34009 350 018 6 3	KARO HEALTHCARE AB	FR
LAMISILDERMGEL 1 %, gel	SE/H/0992/003	34009 350 019 2 4	KARO HEALTHCARE AB	FR
LAMISILATE 1 %, crème	not available	34009 357 029 3 7	KARO HEALTHCARE AB	FR
LAMISILATE 1 %, crème	not available	34009 334 962 5 8	KARO HEALTHCARE AB	FR
LAMISILATE 1 %, crème	not available	34009 301 589 2 0	KARO HEALTHCARE AB	FR
LAMISILATE 1 %, crème	not available	34009 301 589 4 4	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 334 959 4 7	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 334 960 2 9	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 334 961 9 7	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 301 588 9 0	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 301 589 0 6	KARO HEALTHCARE AB	FR
LAMISIL 1 %, crème	not available	34009 301 589 1 3	KARO HEALTHCARE AB	FR
LAMISIL 1 %, solution pour pulvérisation cutanée	SE/H/0992/002	34009 355 855 3 0	KARO HEALTHCARE AB	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
LAMISIL 1 %, solution pour pulvérisation cutanée	SE/H/0992/002	34009 345 681 2 1	KARO HEALTHCARE AB	FR
TERBINAFINE MOBERG PHARMA 98 mg/mL, solution pour application cutanée	SE/H/2249/001	34009 302 767 8 5	MOBERG PHARMA AB	FR
TERBINAFINE MOBERG PHARMA 98 mg/mL, solution pour application cutanée	SE/H/2249/001	34009 302 767 9 2	MOBERG PHARMA AB	FR
TERBINAFINE MOBERG PHARMA 98 mg/mL, solution pour application cutanée	SE/H/2249/001	34009 302 767 9 2	MOBERG PHARMA AB	FR
TERBINAFINE MOBERG PHARMA 98 mg/mL, solution pour application cutanée	SE/H/2249/001	34009 302 767 7 8	MOBERG PHARMA AB	FR
LAMISIL 250 mg, comprimé sécable	not available	34009 334 955 9 6	NOVARTIS PHARMA S.A.S.	FR
LAMISIL 250 mg, comprimé sécable	not available	34009 334 956 5 7	NOVARTIS PHARMA S.A.S.	FR
LAMISIL 250 mg, comprimé sécable	not available	34009 334 957 1 8	NOVARTIS PHARMA S.A.S.	FR
LAMISILATE MONODOSE 1%, solution pour application cutanée	SE/H/0992/004	34009 377 068 4 1	KARO HEALTHCARE AB	FR
TERBINAFINE SANDOZ 250 mg, comprimé sécable	not available	34009 498 907 7 4	SANDOZ	FR
TERBINAFINE SANDOZ 250 mg, comprimé sécable	not available	34009 498 908 3 5	SANDOZ	FR
TERBINAFINE SANDOZ 250 mg, comprimé	not available	34009 498 906 0 6	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
sécable				
TERBINAFINE VIATRIS 1 %, crème	NL/H/4613/001	34009 387 098 3 4	VIATRIS SANTE	FR
TERBINAFINE VIATRIS 1 %, crème	NL/H/4613/001	34009 387 100 8 3	VIATRIS SANTE	FR
TERBINAFINE VIATRIS 1 %, crème	NL/H/4613/001	34009 387 097 7 3	VIATRIS SANTE	FR
LAMISIL DermGel 1% γέλη	SE/H/0992/003	38722/01-04-2024	KARO HEALTHCARE AB	GR
LAMISIL 1% Δερματικό εκνέφωμα, διάλυμα	SE/H/0992/002	38721/01-04-2024	KARO HEALTHCARE AB	GR
Lamisil ONCE 1% δερματικό διάλυμα	SE/H/0992/004	38723/01-04-2024	KARO HEALTHCARE AB	GR
LAMISIL 1% κρέμα	not available	35486/01-04-2024	KARO HEALTHCARE AB	GR
Lamisil 250 mg tablete	not available	HR-H-798097379-01	NOVARTIS HRVATSKA D.O.O.	HR
Lamisil 10 mg/g krema	not available	HR-H-374155033	KARO HEALTHCARE AB	HR
Exoterbyn 78,22 mg/ml ljekoviti lak za nokte	DE/H/6404/001	HR-H-110905071	SANDOZ D.O.O.	HR
Atere 10 mg/g krema	not available	HR-H-664829608	MIBE PHARMACEUTICALS D.O.O.	HR
Lamisil DermGel 10 mg/g gel	not available	HR-H-344287404	KARO HEALTHCARE AB	HR
Lamisil Derma 1% krém	not available	OGYI-T-1867/01	KARO HEALTHCARE AB	HU
Lamisil Derma 1% krém	not available	OGYI-T-1867/02	KARO HEALTHCARE AB	HU
Lamisil Derma 1% krém	not available	OGYI-T-1867/07	KARO HEALTHCARE AB	HU
Lamisil Derma 1% krém	not available	OGYI-T-1867/10	KARO HEALTHCARE AB	HU
Lamisil Derma 1% krém	not available	OGYI-T-1867/11	KARO HEALTHCARE AB	HU
Lamisil Derma 1% krém	not available	OGYI-T-1867/12	KARO HEALTHCARE AB	HU
Exoterbyn gyógyszeres körömlakk	DE/H/6404/001	OGYI-T-23769/02	SANDOZ HUNGÁRIA KFT	HU
Exoterbyn gyógyszeres körömlakk	DE/H/6404/001	OGYI-T-23769/01	SANDOZ HUNGÁRIA KFT	HU
Terbinafin Moberg Pharma 98 mg/ml	SE/H/2249/001	OGYI-T-24270/01	MOBERG PHARMA AB	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
külsöleges oldat				
Terbinafin Moberg Pharma 98 mg/ml külsöleges oldat	SE/H/2249/001	OGYI-T-24270/03	MOBERG PHARMA AB	HU
Terbinafin Moberg Pharma 98 mg/ml külsőleges oldat	SE/H/2249/001	OGYI-T-24270/05	MOBERG PHARMA AB	HU
Terbinafin Moberg Pharma 98 mg/ml külsőleges oldat	SE/H/2249/001	OGYI-T-24270/04	MOBERG PHARMA AB	HU
Lamisil 1% w/w Cream	not available	PA22650/009/001	KARO PHARMA AB	IE
Lamisil AT Cream	not available	PA22650/009/002	KARO PHARMA AB	IE
Lamisil 250 mg Tablets	not available	PA0896/015/001	NOVARTIS IRELAND LIMITED	IE
Terbza 98 mg/ml cutaneous solution	SE/H/2249/001	PA23368/001/001	MOBERG PHARMA AB	IE
Terbza 98 mg/ml cutaneous solution	SE/H/2249/001	PA23368/001/001	MOBERG PHARMA AB	IE
Terbza 98 mg/ml cutaneous solution	SE/H/2249/001	PA23368/001/001	MOBERG PHARMA AB	IE
Terbza 98 mg/ml cutaneous solution	SE/H/2249/001	PA23368/001/001	MOBERG PHARMA AB	IE
Lamisil Once 1% cutaneous solution	SE/H/0992/004	PA22650/009/003	KARO PHARMA AB	IE
Lamisil Once 10 mg/g húðlausn.	SE/H/0992/004	IS/1/06/017/01	KARO HEALTHCARE AB	IS
Lamisil 10 mg/g krem.	not available	910178	KARO HEALTHCARE AB	IS
LAMISIL 1% crema	not available	028176129	NOVARTIS FARMA S.P.A.	IT
LAMISIL 1% crema	not available	028176042	NOVARTIS FARMA S.P.A.	IT
Onicoter	DE/H/6404/001	048131027	SANDOZ S.P.A.	IT
Onicoter	DE/H/6404/001	048131015	SANDOZ S.P.A.	IT
Terbinafina Moberg Pharma 98 mg/mL soluzione cutanea	SE/H/2249/001	050747017	MOBERG PHARMA AB	IT
Terbinafina Moberg	SE/H/2249/001	050747031	MOBERG PHARMA AB	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
Pharma 98 mg/mL soluzione cutanea				
Terbinafina Moberg Pharma 98 mg/mL soluzione cutanea	SE/H/2249/001	050747056	MOBERG PHARMA AB	IT
Terbinafina Moberg Pharma 98 mg/mL soluzione cutanea	SE/H/2249/001	050747043	MOBERG PHARMA AB	IT
LAMISIL 250 mg Compresse	not available	028176028	NOVARTIS FARMA S.P.A.	IT
LAMISIL 250 mg Compresse	not available	028176105	NOVARTIS FARMA S.P.A.	IT
LAMISIL 10 mg/g kremas	not available	LT/1/94/0101/002	KARO HEALTHCARE AB	LT
Posamyk 78,22 mg/ml vaistinis nagų lakas	DE/H/6404/001	LT/1/21/4822/001	SANDOZ PHARMACEUTICALS D.D.	LT
Posamyk 78,22 mg/ml vaistinis nagų lakas	DE/H/6404/001	LT/1/21/4822/002	SANDOZ PHARMACEUTICALS D.D.	LT
LAMISIL UNO 1 % odos tirpalas	SE/H/0992/004	LT/1/06/0586/001	KARO HEALTHCARE AB	LT
Lamisil DermGel 10 mg/g gelis	not available	LT/1/01/3040/001	KARO HEALTHCARE AB	LT
Lamisil 1% crème	not available	2009050331	KARO HEALTHCARE AB	LU
Lamisil 1% crème	not available	2009050331	KARO HEALTHCARE AB	LU
Lamisil Once 1 %, solution pour application cutanée.	SE/H/0992/004	2007060046	KARO HEALTHCARE AB	LU
Lamisil Dermgel 1%, gel	not available	2021040058	KARO HEALTHCARE AB	LU
Lamisil Dermgel 1%, gel	not available	2021040058	KARO HEALTHCARE AB	LU
Lamisil Dermgel 1%, gel	not available	2021040058	KARO HEALTHCARE AB	LU
Lamisil DermGel 10 mg/g gels	not available	98-0252	KARO HEALTHCARE AB	LV
Exotafin 78,22 mg/ml ārstnieciskā nagu laka	DE/H/6404/001	21-0126	SANDOZ PHARMACEUTICALS D.D.	LV
Lamisil Uno 1% uz ādas lietojams šķīdums	SE/H/0992/004	06-0178	KARO HEALTHCARE AB	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
Lamisil 10 mg/g krēms	not available	93-0550	KARO HEALTHCARE AB	LV
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Terbinafine Aurobindo	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
125 mg tablets			(MALTA) LIMITED	
Terbinafine Aurobindo 125 mg tablets	NL/H/2399/001	MA807/04701	AUROBINDO PHARMA (MALTA) LIMITED	MT
Lamisil voetschimmelcrème 10 mg/g, hydrofiele crème	not available	RVG 113799	KARO HEALTHCARE AB	NL
Lamisil DermGel 1%, gel.	SE/H/0992/003	RVG 24945	KARO HEALTHCARE AB	NL
Terbinafine Moberg Pharma 98 mg/ml oplossing voor cutaan gebruik	SE/H/2249/001	RVG 129854	MOBERG PHARMA AB	NL
Terbinafine Moberg Pharma 98 mg/ml oplossing voor cutaan gebruik	SE/H/2249/001	RVG 129854	MOBERG PHARMA AB	NL
Terbinafine Moberg Pharma 98 mg/ml oplossing voor cutaan gebruik	SE/H/2249/001	RVG 129854	MOBERG PHARMA AB	NL
Terbinafine Moberg Pharma 98 mg/ml oplossing voor cutaan gebruik	SE/H/2249/001	RVG 129854	MOBERG PHARMA AB	NL
Lamisil Spray 1%, huidspray, oplossing	not available	RVG 21006	KARO HEALTHCARE AB	NL
Lamisil Lotion 1%, oplossing voor cutaan gebruik	not available	RVG 21005	KARO HEALTHCARE AB	NL
Lamisil Once 1%, oplossing voor cutaan gebruik 10mg/g	SE/H/0992/004	RVG 33714	KARO HEALTHCARE AB	NL
Terbinafine HCl Viatris 10 mg/g, creme	NL/H/4613/001	RVG 101608	VIATRIS LIMITED	NL
Lamisil huidschimmelcrème 10	not available	RVG 14843	KARO HEALTHCARE AB	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
mg/g, hydrofiele crème				
Lamisil 10 mg/ml hudspray, oppløsning	not available	96-3559	KARO HEALTHCARE AB	NO
Terbinafin Moberg Pharma 98 mg/ml liniment, oppløsning	SE/H/2249/001	22-14691	MOBERG PHARMA AB	NO
Terbinafin Moberg Pharma 98 mg/ml liniment, oppløsning	SE/H/2249/001	22-14691	MOBERG PHARMA AB	NO
Terbinafin Moberg Pharma 98 mg/ml liniment, oppløsning	SE/H/2249/001	22-14691	MOBERG PHARMA AB	NO
Terbinafin Moberg Pharma 98 mg/ml liniment, oppløsning	SE/H/2249/001	22-14691	MOBERG PHARMA AB	NO
Terclara 98 mg/ml liniment, oppløsning	SE/H/2250/001	22-14684	MOBERG PHARMA AB	NO
Terclara 98 mg/ml liniment, oppløsning	SE/H/2250/001	22-14684	MOBERG PHARMA AB	NO
Terclara 98 mg/ml liniment, oppløsning	SE/H/2250/001/DC	22-14684	MOBERG PHARMA AB	NO
Terclara 98 mg/ml liniment, oppløsning	SE/H/2250/001/DC	22-14684	MOBERG PHARMA AB	NO
Lamisil 1% liniment, oppløsning	SE/H/0992/004	06-3985	KARO HEALTHCARE AB	NO
Lamisil DermGel 1% gel	not available	97-3304	KARO HEALTHCARE AB	NO
Lamisil 10 mg/g krem	not available	7963	KARO HEALTHCARE AB	NO
Terbinafinhydroklorid Norfri 10 mg/g krem	not available	17-11768	EVOLAN PHARMA AB	NO
LAMISILATT 10 mg/g, krem	not available	R/1191	KARO HEALTHCARE AB	PL
Tersilat, 10 mg/g, krem	DE/H/6469/001	18989	SUN-FARM SP. Z.O.O.	PL
Tersilat, 10 mg/g, krem	DE/H/6469/001/DC	18989	SUN-FARM SP. Z.O.O.	PL
Tersilat, 10 mg/g, aerozol na skórê, roztwór	not available	22036	SUN-FARM 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
Lamisil 10 mg/g solução para pulverização cutânea	SE/H/0992/002	2690386	KARO HEALTHCARE AB	PT
Lamisil 10 mg/g solução para pulverização cutânea	SE/H/0992/002	3532686	KARO HEALTHCARE AB	PT
LAMISIL, 10 mg/g Creme	not available	2135184	KARO HEALTHCARE AB	PT
Lamisil 1 10 mg/g solução cutânea	SE/H/0992/004	5929781	KARO HEALTHCARE AB	PT
Lamisil 250 mg Comprimidos	not available	2134989	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Lamisil 250 mg Comprimidos	not available	2135085	NOVARTIS FARMA - PRODUTOS FARMACÊUTICOS S.A.	PT
Exoterfyn 78,22 mg/ml Lac de unghii medicamentos	DE/H/6404/001	16306/2025/01	SANDOZ PHARMACEUTICALS S.R.L.	RO
Exoterfyn 78,22 mg/ml Lac de unghii medicamentos	DE/H/6404/001	16306/2025/02	SANDOZ PHARMACEUTICALS S.R.L.	RO
Lamisil 10 mg/g cremă	not available	11028/2018/01	KARO HEALTHCARE AB	RO
Lamisil 10 mg/g cremă	not available	11028/2018/02	KARO HEALTHCARE AB	RO
Lamisil 1% kutan lösning	SE/H/0992/001	13759	KARO HEALTHCARE AB	SE
Terbinafin Apofri 10 mg/g kräm	not available	46305	EVOLAN PHARMA AB	SE
Terbinafin ABECE 10 mg/g kräm	not available	56084	EVOLAN PHARMA AB	SE
Lamisil 1% kräm	not available	11523	KARO HEALTHCARE AB	SE
Lamisil Dermgel 1%, gel	SE/H/0992/003	14867	KARO HEALTHCARE AB	SE
Lamisil 1% kutan spray, lösning	SE/H/0992/002	13760	KARO HEALTHCARE AB	SE
Terbinafin Moberg Pharma 98 mg/ml kutan lösning	SE/H/2249/001	63250	MOBERG PHARMA AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Terbinafin Moberg Pharma 98 mg/ml kutan lösning	SE/H/2249/001	63250	MOBERG PHARMA AB	SE
Terbinafin Moberg Pharma 98 mg/ml kutan lösning	SE/H/2249/001	63250	MOBERG PHARMA AB	SE
Terbinafin Moberg Pharma 98 mg/ml kutan lösning	SE/H/2249/001	63250	MOBERG PHARMA AB	SE
Terclara 98 mg/ml kutan lösning	SE/H/2250/001	63251	MOBERG PHARMA AB	SE
Terclara 98 mg/ml kutan lösning	SE/H/2250/001	63251	MOBERG PHARMA AB	SE
Terclara 98 mg/ml kutan lösning	SE/H/2250/001	63251	MOBERG PHARMA AB	SE
Terclara 98 mg/ml kutan lösning	SE/H/2250/001	63251	MOBERG PHARMA AB	SE
Lamisil Singeldos 1 %, kutan lösning	SE/H/0992/004	23495	KARO HEALTHCARE AB	SE
Lamisil DermGel 10 mg/g gel	not available	H/93/00872/007	KARO HEALTHCARE AB	SI
Lamisil DermGel 10 mg/g gel	not available	H/93/00872/008	KARO HEALTHCARE AB	SI
Lamisil DermGel 10 mg/g gel	not available	H/93/00872/009	KARO HEALTHCARE AB	SI
Lamisil DermGel 10 mg/g gel	not available	H/93/00872/010	KARO HEALTHCARE AB	SI
Lamisil DermGel 10 mg/g gel	not available	H/93/00872/011	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g dermalno pršilo, raztopina	not available	H/93/00872/005	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g dermalno pršilo, raztopina	not available	H/93/00872/006	KARO HEALTHCARE AB	SI
Exoter 78,22 mg/ml	DE/H/6404/001	H/21/02800/001	LEK PHARMACEUTICALS	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
zdravilni lak za nohte			D.D. LJUBLJANA	
Exoter 78,22 mg/ml zdravilni lak za nohte	DE/H/6404/001	H/21/02800/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Lamisil 250 mg tablete	not available	H/93/02072/004	NOVARTIS EUROPHARM LIMITED	SI
Lamisil 125 mg tablete	not available	H/93/02072/001	NOVARTIS EUROPHARM LIMITED	SI
Lamisil 250 mg tablete	not available	H/93/02072/003	NOVARTIS EUROPHARM LIMITED	SI
Lamisil 125 mg tablete	not available	H/93/02072/002	NOVARTIS EUROPHARM LIMITED	SI
Lamisil 10 mg/g krema	not available	H/93/00872/001	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g krema	not available	H/93/00872/002	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g krema	not available	H/93/00872/003	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g krema	not available	H/93/00872/004	KARO HEALTHCARE AB	SI
Lamisil 10 mg/g krém	not available	26/0417/91-S	KARO HEALTHCARE AB	SK
Exoterin	DE/H/6404/001	26/0184/21-S	SANDOZ PHARMACEUTICALS D.D.	SK
Terbinafine 250mg Tablets	not available	PL 42765/0012	RENATA (UK) LIMITED	XI
Terbinafine 250 mg Tablets	not available	PL 21880/0099	MEDREICH PLC	XI
Terbinafine Hydrochloride 1% Cream	not available	PL 28444/0040	ACTIVASE PHARMACEUTICALS LIMITED	XI
Terbinafine 250 mg tablets	not available	PL 36687/0369	TORRENT PHARMA (UK) LTD.	XI
Terbinafine 1 % Cream	NL/H/4613/001	PL 04569/0889	GENERICS [UK] LIMITED	XI