10 June 2021
EMA/333431/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: lidocaine

Procedure no.: PSUSA/00001861/202009

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
Xylocaine 10 mg/razpršek pršilo, raztopina	not available	H/94/01682/001	ASPEN PHARMA TRADING LIMITED	SI
Actilogic 700 mg apósito adhesivo medicamentoso	DE/H/5699/001	83398	GRÜNENTHAL PHARMA S.A.	ES
Actilogic 700 mg wirkstoffhaltiges Pflaster	DE/H/5699/001	98976.00.00	GRÜNENTHAL GMBH	DE
Asensil 40mg/g Crema	IE/H/0727/001	043742016	LOGOFARMA S.R.L.	IT
Asensil 40mg/g Crema	IE/H/0727/001	043742028	LOGOFARMA S.R.L.	IT
Asensil 40mg/g Crema	IE/H/0727/001	043742030	LOGOFARMA S.R.L.	IT
Asensil 40mg/g Crema	IE/H/0727/001	043742042	LOGOFARMA S.R.L.	IT
Asensil 40mg/g Crema	IE/H/0727/001	043742055	LOGOFARMA S.R.L.	IT
Calmotop® 700 mg wirkstoffhaltiges Pflaster	DE/H/5703/001	97540.00.00	GRÜNENTHAL GMBH	DE
Dolocopin 40mg/g Cream	IE/H/0727/001	PA22869/001/001	FERNDALE LABORATORIES LIMITED	IE
Dynexan 2 %, crème buccale	not available	3400935279897	CHEMISCHE FABRIK KREUSSLER & CO. GMBH	FR
Dynexangival 1 %, crème buccale	not available	34009 498 755 2 8	CHEMISCHE FABRIK KREUSSLER & CO. GMBH	FR
Endor 700 mg wirkstoffhaltiges Pflaster	DE/H/5695/001	98972.00.00	GRÜNENTHAL GMBH	DE
Esliva 700 mg lääkelaastari	DE/H/5696/001	34621	GRÜNENTHAL GMBH	FI
Esliva 700 mg wirkstoffhaltiges Pflaster	DE/H/5696/001	98973.00.00	GRÜNENTHAL GMBH	DE
Instillagel 20,9 mg gel	not available	10132	FARCO-PHARMA GMBH	SE
Intermed XYLOJELL® Γέλη 2% w/w	not available	10876/15/12.10.2018	IOULIA AND IRENE TSETI PHARMACEUTICAL LABORATORIES S.A.	GR
Intermed XYLOJELL® Εκνέφωμα 10% w/v	not available	28568/15/12.10.2018	IOULIA AND IRENE TSETI PHARMACEUTICAL LABORATORIES S.A.	GR
LAMBDALINA 40 mg/g	SE/H/0699/001	69.789	ISDIN, 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
crema				
LAMBDALINA 40 mg/g crema	SE/H/0699/001	69.789	ISDIN, S.A.	ES
LAMBDALINA 40 mg/g creme	SE/H/0699/001	5095369	ISDIN LDA	PT
LAMBDALINA 40 mg/g creme	SE/H/0699/001	5095377	ISDIN LDA	PT
LAMBDALINA 40 mg/g creme	SE/H/0699/001	5095369	ISDIN LDA	PT
LAMBDALINA 40 mg/g creme	SE/H/0699/001	5095377	ISDIN LDA	PT
Licain PUREN 10 mg/ml Injektionslösung	PT/H/1585/001	96439.00.00	PUREN PHARMA GMBH & CO. KG	DE
Lidbree	NL/H/4625/001	61872	GEDEON RICHTER PLC.	DK
LIDBREE 42 mg/ml gel cu cedare intrauterină	NL/H/4625/001	13518/2020/01	GEDEON RICHTER ROMÂNIA S.A.	RO
Lidbree 42 mg/ml gel intra-utérin	NL/H/4625/001	BE567146	GEDEON RICHTER PLC.	BE
Lidbree 42 mg/ml gel intra-utérin	NL/H/4625/001	2020080203	GEDEON RICHTER PLC.	LU
Lidbree 42 mg/ml gel intrauterino	NL/H/4625/001	85483	GEDEON RICHTER PLC.	ES
Lidbree 42 mg/mL gel intrauterino	NL/H/4625/001	047309012	GEDEON RICHTER ROMÂNIA S.A.	IT
Lidbree 42 mg/ml gel voor intra-uterien gebruik	NL/H/4625/001	BE567146	GEDEON RICHTER PLC.	BE
Lidbree 42 mg/ml gel voor intra-uterien gebruik	NL/H/4625/001	RVG 124015	GEDEON RICHTER PLC.	NL
Lidbree 42 mg/ml Gel zur intrauterinen Anwendung	NL/H/4625/001	BE567146	GEDEON RICHTER PLC.	BE
Lidbree 42 mg/ml Gel	NL/H/4625/001	2203302.00.00	GEDEON RICHTER PLC.	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 intrauterinen Anwendung				
Lidbree 42 mg/ml gimdos ertmės gelis	NL/H/4625/001	LT/1/20/4598/001	GEDEON RICHTER PLC.	LT
Lidbree 42 mg/ml intrauteriingeel	NL/H/4625/001	1009320	GEDEON RICHTER PLC.	EE
Lidbree 42 mg/ml intrauterin gel	NL/H/4625/001	58469	GEDEON RICHTER PLC.	SE
Lidbree 42 mg/ml intrauterin gél	NL/H/4625/001	OGYI-T-23713/01	GEDEON RICHTER PLC.	HU
Lidbree 42 mg/ml intrauterīnais gels	NL/H/4625/001	20-0089	GEDEON RICHTER PLC.	LV
Lidbree 42 mg/mL intrauterine gel	NL/H/4625/001	PL 04854/0166	GEDEON RICHTER PLC.	UK
Lidbree 42 mg/mL intrauterine gel	NL/H/4625/001	PA1330/025/001	GEDEON RICHTER PLC.	IE
Lidbree 42 mg/mL intrauterine gel	NL/H/4625/001	MA1031/00601	GEDEON RICHTER PLC.	MT
Lidbree 42 mg/ml intrauterini gel	NL/H/4625/001	H/20/02742/001	GEDEON RICHTER PLC.	SI
Lidbree 42 mg/ml intrauterini gel	NL/H/4625/001	HR-H-772844274	GEDEON RICHTER PLC.	HR
Lidbree 42 mg/ml intrauterinní gel	NL/H/4625/001	01/467/18-C	GEDEON RICHTER PLC.	CZ
Lidbree 42 mg/ml intrauterinný gél	NL/H/4625/001	01/0163/20-S	GEDEON RICHTER PLC.	SK
Lidbree 42 mg/ml leghlaup	NL/H/4625/001	IS/1/20/079/01	GEDEON RICHTER PLC.	IS
LIDBREE 42 mg/mL, gel intra-utérin	NL/H/4625/001	34009 302 121 2 7	GEDEON RICHTER PLC.	FR
Lidbree® 42 mg/ml Gel zur intrauterinen Anwendung	NL/H/4625/001	140367	GEDEON RICHTER PLC.	AT
LIDIAQ 40 mg/g cremă	PL/H/0643/001	8940/2016/01	FERNDALE LABORATORIES	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
			LIMITED	
LIDIAQ 40 mg/g dermálny krém	PL/H/0643/001	01/0463/15-S	FERNDALE LABORATORIES LIMITED	SK
LIDIQ 40mg/g krema	PL/H/0643/001	H/16/02099/001	FERNDALE LABORATORIES LIMITED	SI
Lidocain Grünenthal 700 mg wirkstoffhaltiges Pflaster	DE/H/5734/001	96944.00.00	GRÜNENTHAL GMBH	DE
Lidocain Walter Ritter 100 mg/ml Spray zur Anwendung auf der Haut, Lösung	AT/H/0689/001/DC	137843	WALTER RITTER GMBH + CO. KG	AT
Lidocain Walter Ritter 100 mg/ml Spray zur Anwendung auf der Haut, Lösung	AT/H/0689/001/DC	137843	WALTER RITTER GMBH + CO. KG	LI
Lidocaína 1% Braun 10 mg/ml solução injetável	not available	5184171	B. BRAUN MEDICAL LDA.	PT
Lidocaína 1% Braun 10 mg/ml solução injetável	not available	3699188	B. BRAUN MEDICAL LDA.	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g crema	IE/H/0727/001/DC	82.257	ISDIN, S.A.	ES
Lidocaína Ferndale Pharmaceuticals 40 mg/g crema	IE/H/0727/001/DC	82.257	ISDIN, S.A.	ES
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5717632	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715511	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g	IE/H/0727/001	5715545	ISDIN 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
creme				
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715602	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715537	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715560	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715503	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715479	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715529	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715552	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5717632	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715511	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715545	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715602	ISDIN 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
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715537	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715560	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715503	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715479	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715529	ISDIN LDA	PT
Lidocaína Ferndale Pharmaceuticals 40 mg/g creme	IE/H/0727/001	5715552	ISDIN LDA	PT
Lidocaine 5% m/m Ointment	not available	PL 00289/0761	TEVA UK LIMITED	XI
LIDOCAINE CHLORHYDRATE RENAUDIN 20 mg/mL, solution pour injection/perfusion	NL/H/4574/001	34009 302 082 5 0	LABORATOIRE RENAUDIN	FR
LIDOCAINE CHLORHYDRATE RENAUDIN 20 mg/mL, solution pour injection/perfusion	NL/H/4574/001	34009 302 082 6 7	LABORATOIRE RENAUDIN	FR
LIDOCAINE CHLORHYDRATE RENAUDIN 20 mg/mL, solution pour	NL/H/4574/001	34009 302 082 7 4	LABORATOIRE RENAUDIN	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
injection/perfusion				
Lidocaine Grünenthal 700 mg lääkelaastari	DE/H/5734/001	33896	GRÜNENTHAL GMBH	FI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v	not available	PL 03265/0005R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v	not available	PL 03265/0005R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v	not available	PL 03265/0005R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v	not available	PL 03265/0005R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v	not available	PL 03265/0005R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Hydrochloride Injection BP Minijet 1% w/v solution for injection.	not available	PA22684/008/001	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 1% w/v solution for injection.	not available	PA22684/008/001	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 1% w/v solution for injection.	not available	PA22684/008/001	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 1% w/v solution for injection.	not available	PA22684/008/001	DLRC PHARMA SERVICES LIMITED	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**
Lidocaine Hydrochloride Injection BP Minijet 2% w/v Solution for Injection	not available	PA22684/008/002	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 2% w/v Solution for Injection	not available	PA22684/008/002	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 2% w/v Solution for Injection	not available	PA22684/008/002	DLRC PHARMA SERVICES LIMITED	IE
Lidocaine Hydrochloride Injection BP Minijet 2% w/v Solution for Injection	not available	PA22684/008/002	DLRC PHARMA SERVICES LIMITED	IE
Lidocaïne Hydrochloride Renaudin 20 mg/ml, oplossing voor injectie/infusie	NL/H/4574/001	RVG 123705	LABORATOIRE RENAUDIN	NL
Lidocaine Injection BP Minijet 2% w/v	not available	PL 03265/0006R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Injection BP Minijet 2% w/v	not available	PL 03265/0006R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Injection BP Minijet 2% w/v	not available	PL 03265/0006R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Injection BP Minijet 2% w/v	not available	PL 03265/0006R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
Lidocaine Injection BP Minijet 2% w/v	not available	PL 03265/0006R	INTERNATIONAL MEDICATION SYSTEMS (UK) LIMITED	XI
LIDOFAST 10 mg/g gel	not available	034478014	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F.	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
			S.P.A.	
LIDOFAST 10 mg/g gel	not available	034478014	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
LIDOFAST 25 mg/g gel uretrale	not available	034478026	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
LIDOFAST 25 mg/g gel uretrale	not available	034478026	AZIENDE CHIMICHE RIUNITE ANGELINI FRANCESCO - A.C.R.A.F. S.P.A.	IT
LidoGalen 40 mg/g Creme	not available	91519.00.00	GALENPHARMA GMBH	DE
Lidokain Isdin 40 mg/g kräm	SE/H/699/001	21239	ISDIN, S.A.	SE
Lidokain Isdin 40 mg/g kräm	SE/H/699/001	21239	ISDIN, S.A.	SE
Lidoposterin 50 mg/g tepalas	not available	LT/1/02/2982/001	DR. KADE PHARMAZEUTISCHE FABRIK GMBH	LT
LIDOPOSTERIN 50 mg/g, maść	not available	11628	DR. KADE PHARMAZEUTISCHE FABRIK GMBH	PL
Lidotec® 700 mg wirkstoffhaltiges Pflaster	DE/H/5701/001	96984.00.00	GRÜNENTHAL GMBH	DE
Linisol 1 %, Injektionslösung	not available	BE166695	B.BRAUN MELSUNGEN AG	BE
Linisol 1 %, oplossing voor injectie	not available	BE166695	B.BRAUN MELSUNGEN AG	BE
Linisol 1 %, solution injectable	not available	BE166695	B.BRAUN MELSUNGEN AG	BE
Linisol 2 % oplossing	not available	BE166704	B.BRAUN MELSUNGEN AG	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
voor injectie				
Linisol 2 %, Injektionslösung	not available	BE166704	B.BRAUN MELSUNGEN AG	BE
Linisol 2 %, solution injectable	not available	BE166704	B.BRAUN MELSUNGEN AG	BE
Lonioban 700 mg medicated plaster	DE/H/5702/001	PL 21727/0073	GRÜNENTHAL LTD.	XI
Lonioban 700 mg medicinskt plåster	DE/H/5702/001	54619	GRÜNENTHAL GMBH	SE
Lonioban 700 mg medisinert plaster	DE/H/5702/001	16-11135	GRÜNENTHAL GMBH	NO
Lonioban, medicinsk plaster	DE/H/5702/001	57581	GRÜNENTHAL GMBH	DK
Lonioban® 700 mg wirkstoffhaltiges Pflaster	DE/H/5702/001	97539.00.00	GRÜNENTHAL GMBH	DE
Luan 1% gel	not available	005638022	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Luan 2,5% gel	not available	005638 010	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Maxilene 40 mg/g kräm	IE/H/0727/001	50469	FERNDALE LABORATORIES LIMITED	SE
MESOCAINE 25 mg/5ml solution injectable	not available	34009 331 077 0 3	LABORATOIRES PHARMY II	FR
MESOCAINE 25 mg/5ml solution injectable	not available	34009 331 077 0 3	LABORATOIRES PHARMY II	FR
MESOCAINE 50 mg/5ml solution injectable	not available	34009 331 080 1 4	LABORATOIRES PHARMY II	FR
MESOCAINE 50 mg/5ml solution injectable	not available	34009 331 080 1 4	LABORATOIRES PHARMY II	FR
Nanolipo, 40 mg/g, krem	UK/H/5616/001	23052	ADAMED PHARMA S.A.	PL
Nydriar 700 mg cerotto medicato	DE/H/5697/001	045702014	GRÜNENTHAL ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Nydriar 700 mg cerotto medicato	DE/H/5697/001	045702026	GRÜNENTHAL ITALIA S.R.L.	IT
Nydriar 700 mg cerotto medicato	DE/H/5697/001	045702038	GRÜNENTHAL ITALIA S.R.L.	IT
Nydriar 700 mg cerotto medicato	DE/H/5697/001	045702040	GRÜNENTHAL ITALIA S.R.L.	IT
Nydriar 700 mg cerotto medicato	DE/H/5697/001	045702053	GRÜNENTHAL ITALIA S.R.L.	IT
ORTODERMINA crema al 5%	not available	005556028	SOFAR S.P.A.	IT
ORTODERMINA crema al 5%	not available	005556016	SOFAR S.P.A.	IT
ORTODERMINA crema al 5%	not available	005556030	SOFAR S.P.A.	IT
ORTODERMINA crema al 5%	not available	005556042	SOFAR S.P.A.	IT
Posterisan® akut 50 mg/g Rektalsalbe	not available	6077267.00.00	DR. KADE PHARMAZEUTISCHE FABRIK GMBH	DE
Posterisan® akut 50 mg/g Rektalsalbe mit Analdehner	not available	6077267.00.00	DR. KADE PHARMAZEUTISCHE FABRIK GMBH	DE
Premjact Desensitizing Spray for Men	MT/H/510/001	PL 02294/5000R	POUND INTERNATIONAL LTD	UK
Premjact Desensitizing Spray for Men, 9.6% w/w, cutaneous spray	MT/H/510/001	PL 02294/5000R	POUND INTERNATIONAL LTD	UK
Ralvo 700 mg medicated plaster	UK/H/6362/001	PL 21727/0075	GRÜNENTHAL LTD.	XI
Severtos 700 mg emplâtre médicamenteux	DE/H/5702/001	BE505786	SA GRÜNENTHAL N.V.	BE
Severtos 700 mg emplâtre médicamenteux	DE/H/5702/001	2017070246	SA GRÜNENTHAL N.V.	LU
Severtos 700 mg pleister	DE/H/5702/001	BE505786	SA GRÜNENTHAL 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
Severtos 700 mg wirkstoffhaltiges Pflaster	DE/H/5698/001	98975.00.00	GRÜNENTHAL GMBH	DE
STUD 100 Desensitizing Spray for Men	MT/H/510/001	MA526/00201	POUND INTERNATIONAL LTD	MT
STUD 100 Desensitizing Spray for Men	MT/H/510/001	PL 02294/5000R	POUND INTERNATIONAL LTD	UK
Truvinut 700 mg cerotto medicato	DE/H/5696/001	045701012	GRÜNENTHAL ITALIA S.R.L.	IT
Truvinut 700 mg cerotto medicato	DE/H/5696/001	045701024	GRÜNENTHAL ITALIA S.R.L.	IT
Truvinut 700 mg cerotto medicato	DE/H/5696/001	045701036	GRÜNENTHAL ITALIA S.R.L.	IT
Truvinut 700 mg cerotto medicato	DE/H/5696/001	045701048	GRÜNENTHAL ITALIA S.R.L.	IT
Truvinut 700 mg cerotto medicato	DE/H/5696/001	045701051	GRÜNENTHAL ITALIA S.R.L.	IT
Verira 700 mg apósito adhesivo medicamentoso	DE/H/5698/001	83397	GRÜNENTHAL PHARMA S.A.	ES
Versatis 5 % liečivá náplasť	DE/H/5733/001	01/0054/10-S	GRÜNENTHAL GMBH	SK
Versatis 5% medicinski flaster	not available	UP/I-530-09/13-02/166	GRÜNENTHAL GMBH	HR
Versatis 700 mg apósito adhesivo medicamentoso	DE/H/5733/001	71848	GRÜNENTHAL PHARMA S.A.	ES
Versatis 700 mg ārstnieciskais plāksteris	DE/H/5733/001	10-0176	GRÜNENTHAL GMBH	LV
Versatis 700 mg cerotto medicato	DE/H/5733/001	040335010	GRÜNENTHAL ITALIA S.R.L.	IT
Versatis 700 mg cerotto medicato	DE/H/5733/001	040335046	GRÜNENTHAL ITALIA S.R.L.	IT
Versatis 700 mg cerotto medicato	DE/H/5733/001	040335022	GRÜNENTHAL ITALIA S.R.L.	IT
Versatis 700 mg cerotto medicato	DE/H/5733/001	040335034	GRÜNENTHAL ITALIA S.R.L.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Versatis 700 mg cerotto medicato	DE/H/5733/001	040335059	GRÜNENTHAL ITALIA S.R.L.	IT
Versatis 700 mg emplastru medicamentos	DE/H/5733/001	11189/2018/02	GRÜNENTHAL GMBH	RO
Versatis 700 mg emplastru medicamentos	DE/H/5733/001	11189/2018/03	GRÜNENTHAL GMBH	RO
Versatis 700 mg emplastru medicamentos	DE/H/5733/001	11189/2018/04	GRÜNENTHAL GMBH	RO
Versatis 700 mg emplastru medicamentos	DE/H/5733/001	11189/2018/05	GRÜNENTHAL GMBH	RO
Versatis 700 mg emplastru medicamentos	DE/H/5733/001	11189/2018/01	GRÜNENTHAL GMBH	RO
Versatis 700 mg emplâtre médicamenteux	DE/H/5733/001	BE312462	SA GRÜNENTHAL N.V.	BE
Versatis 700 mg emplâtre médicamenteux	DE/H/5733/001	2008040037	SA GRÜNENTHAL N.V.	LU
Versatis 700 mg gyógyszeres tapasz	DE/H/5733/001	OGYI-T-21784/02	GRÜNENTHAL GMBH	HU
Versatis 700 mg gyógyszeres tapasz	DE/H/5733/001	OGYI-T-21784/05	GRÜNENTHAL GMBH	HU
Versatis 700 mg gyógyszeres tapasz	DE/H/5733/001	OGYI-T-21784/03	GRÜNENTHAL GMBH	HU
Versatis 700 mg gyógyszeres tapasz	DE/H/5733/001	OGYI-T-21784/01	GRÜNENTHAL GMBH	HU
Versatis 700 mg gyógyszeres tapasz	DE/H/5733/001	OGYI-T-21784/04	GRÜNENTHAL GMBH	HU
VERSATIS 700 mg léčivá náplast	DE/H/5733/001	01/284/10-C	GRÜNENTHAL GMBH	CZ
Versatis 700 mg lyfjaplástur	DE/H/5733/001	IS/1/10/033/01	GRÜNENTHAL GMBH	IS
Versatis 700 mg medicated plaster	DE/H/5733/001	PA 2242/7/1	GRÜNENTHAL PHARMA LTD.	IE
Versatis 700 mg medicated plaster	DE/H/5733/001	MA840/00201	GRÜNENTHAL GMBH	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
Versatis 700 mg medicated plaster	DE/H/5733/001	PL 21727/0016	GRÜNENTHAL LTD.	XI
Versatis 700 mg medicinskt plåster	DE/H/5733/001	25314	GRÜNENTHAL GMBH	SE
Versatis 700 mg medisinert plaster	DE/H/5733/001	10-7564	GRÜNENTHAL GMBH	NO
Versatis 700 mg pleister	DE/H/5733/001	BE312462	SA GRÜNENTHAL N.V.	BE
Versatis 700 mg pleister	DE/H/5733/001	RVG 113654	GRÜNENTHAL B.V.	NL
Versatis 700 mg vaistinis pleistras	DE/H/5733/001	LT/1/10/2279/001	GRÜNENTHAL GMBH	LT
Versatis 700 mg vaistinis pleistras	DE/H/5733/001	LT/1/10/2279/003	GRÜNENTHAL GMBH	LT
Versatis 700 mg vaistinis pleistras	DE/H/5733/001	LT/1/10/2279/002	GRÜNENTHAL GMBH	LT
Versatis 700 mg vaistinis pleistras	DE/H/5733/001	LT/1/10/2279/005	GRÜNENTHAL GMBH	LT
Versatis 700 mg vaistinis pleistras	DE/H/5733/001	LT/1/10/2279/004	GRÜNENTHAL GMBH	LT
Versatis 700 mg wirkstoffhaltiges Pflaster	DE/H/5733/001	1-29025	GRÜNENTHAL GES. M.B.H.	AT
Versatis 700 mg wirkstoffhaltiges Pflaster	DE/H/5733/001	69088.00.00	GRÜNENTHAL GMBH	DE
Versatis 700 mg zdravilni obliž	DE/H/5733/001	H/07/01632/002	GRÜNENTHAL GMBH	SI
Versatis 700 mg zdravilni obliž	DE/H/5733/001	H/07/01632/003	GRÜNENTHAL GMBH	SI
Versatis 700 mg zdravilni obliž	DE/H/5733/001	H/07/01632/004	GRÜNENTHAL GMBH	SI
Versatis 700 mg zdravilni obliž	DE/H/5733/001	H/07/01632/005	GRÜNENTHAL GMBH	SI
Versatis 700 mg zdravilni obliž	DE/H/5733/001	H/07/01632/001	GRÜNENTHAL GMBH	SI
Versatis 700 mg έμπλαστρο φαρμακούχο	DE/H/5733/001	22834	GRÜNENTHAL GMBH	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
Versatis 700 mg ἐμπλαστρο φαρμακούχο	DE/H/5733/001	79655/24-09-2019	GRÜNENTHAL GMBH	GR
VERSATIS 700 mg, emplâtre médicamenteux	DE/H/5733/001	34009 382 853 8 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
VERSATIS 700 mg, emplâtre médicamenteux	DE/H/5733/001	34009 382 855 0 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
VERSATIS 700 mg, emplâtre médicamenteux	DE/H/5733/001	34009 382 854 4 4	LABORATOIRES GRÜNENTHAL S.A.S.	FR
VERSATIS 700 mg, emplâtre médicamenteux	DE/H/5733/001	34009 382 852 1 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
VERSATIS 700 mg, emplâtre médicamenteux	DE/H/5733/001	34009 382 856 7 3	LABORATOIRES GRÜNENTHAL S.A.S.	FR
Versatis, 700 mg ravimplaaster	DE/H/5733/001	675610	GRÜNENTHAL GMBH	EE
Versatis, 700 mg, plaster leczniczy	DE/H/5733/001	17841	GRÜNENTHAL GMBH	PL
Versatis, medicinsk plaster	DE/H/5733/001	46834	GRÜNENTHAL GMBH	DK
Vessatis 700 mg emplastro medicamentoso	DE/H/5733/001	5669478	GRÜNENTHAL S.A.	PT
Vessatis 700 mg emplastro medicamentoso	DE/H/5733/001	5281472	GRÜNENTHAL S.A.	PT
Vessatis 700 mg emplastro medicamentoso	DE/H/5733/001	5281514	GRÜNENTHAL S.A.	PT
Vessatis 700 mg emplastro medicamentoso	DE/H/5733/001	5281506	GRÜNENTHAL S.A.	PT
Vobolon 700 mg apósito adhesivo medicamentoso	DE/H/5697/001	83399	GRÜNENTHAL PHARMA S.A.	ES
Vobolon 700 mg medicated plaster	DE/H/5697/001	PA 2242/001/001	GRÜNENTHAL PHARMA LTD.	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Vobolon 700 mg wirkstoffhaltiges Pflaster	DE/H/5697/001	98974.00.00	GRÜNENTHAL GMBH	DE
Xylocain - viscös oral 2 %	not available	8.840	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 1% - Durchstechflasche	not available	16.720	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 10 mg/ml injektionsvätska, lösning	not available	3841	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 10 mg/ml stungulyf, lausn	not available	IS/1/02/134/01	ASPEN PHARMA TRADING LIMITED	IS
Xylocain 10% - Pumpspray	not available	12.979	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 100 mg/ml kutan spray, lösning	not available	11258	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 100 mg/ml spray	not available	4112	ASPEN PHARMA TRADING LIMITED	NO
Xylocain 2 % - Gel	not available	8.456	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 2% - Ampullen	not available	16.722	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 2% - Durchstechflasche	not available	16.721	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 2% gel med konserveringsmedel	not available	4114	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 2% gel utan konserveringsmedel	not available	10445	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 2% hlaup	not available	843388	ASPEN PHARMA TRADING LIMITED	IS
Xylocain 2% hlaup	not available	640474	ASPEN PHARMA TRADING LIMITED	IS
Xylocain 20 mg/ml injektionsvätska, lösning	not available	3553	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 20 mg/ml stungulyf, lausn	not available	IS/1/02/134/02	ASPEN PHARMA TRADING LIMITED	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
Xylocain 5 mg/ml injektionsvätska, lösning	not available	3840	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 5% Salbe	not available	8.498	ASPEN PHARMA TRADING LIMITED	AT
Xylocain 5% salva	not available	3601	ASPEN PHARMA TRADING LIMITED	SE
Xylocain 5% smyrsli	not available	IS/1/03/126/01	ASPEN PHARMA TRADING LIMITED	IS
Xylocain injeksjonsvæske, oppløsning 10 mg/ml	not available	3605	ASPEN PHARMA TRADING LIMITED	NO
Xylocain injeksjonsvæske, oppløsning 20 mg/ml	not available	2162	ASPEN PHARMA TRADING LIMITED	NO
Xylocain liniment 3 %	not available	6271	ASPEN PHARMA TRADING LIMITED	NO
Xylocain salve 5 %	not available	03-1921	ASPEN PHARMA TRADING LIMITED	NO
Xylocain utan konserveringsmedel 10 mg/ml injektionsvätska, lösning	not available	10598	ASPEN PHARMA TRADING LIMITED	SE
Xylocain utan konserveringsmedel 20 mg/ml injektionsvätska, lösning	not available	10705	ASPEN PHARMA TRADING LIMITED	SE
Xylocain Viskös 20 mg/ml oral lösning	not available	4546	ASPEN PHARMA TRADING LIMITED	SE
Xylocain, 100 mg/ml, húðúði, lausn	not available	640471	ASPEN PHARMA TRADING LIMITED	IS
Xylocain, gel	not available	9467	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, gel	not available	9467	ASPEN PHARMA TRADING 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
Xylocain, injektionsvæske, opløsning	not available	01109	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, injektionsvæske, opløsning	not available	01109	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, injektionsvæske, opløsning	not available	09469	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, injektionsvæske, opløsning	not available	09469	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, kutanopløsning	not available	14449	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, kutanspray, opløsning	not available	3352	ASPEN PHARMA TRADING LIMITED	DK
Xylocain, salve	not available	9470	ASPEN PHARMA TRADING LIMITED	DK
Xylocain® 1 % Injektionslösung	not available	6077072.01.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® 100 mg/ml kutan spray, lösning	not available	10384	ASPEN PHARMA TRADING LIMITED	FI
Xylocain® 100 mg/ml sumute iholle, liuos	not available	10384	ASPEN PHARMA TRADING LIMITED	FI
Xylocain® 2 % gel	not available	9058	ASPEN PHARMA TRADING LIMITED	FI
Xylocain® 2 % gel	not available	2779	ASPEN PHARMA TRADING LIMITED	NO
Xylocain® 2 % Injektionslösung	not available	6077072.02.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® 2 % Injektionslösung	not available	6225422.00.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® 2% geeli	not available	9058	ASPEN PHARMA TRADING 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
Xylocain® Gel 2 %	not available	6077215.00.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® Pumpspray 10 mg/Sprühstoß Spray zur Anwendung auf der Haut, Lösung	not available	6084764.00.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® Pumpspray dental 10 mg/Sprühstoß Spray zur Anwendung in der Mundhöhle	not available	6077184.00.00	ASPEN PHARMA TRADING LIMITED	DE
Xylocain® Viscös 2 % Lösung zum Einnehmen	not available	6077149.00.00	ASPEN PHARMA TRADING LIMITED	DE
XYLOCAINA 20 mg/ml soluzione iniettabile	not available	004535213	ASPEN PHARMA TRADING LIMITED	IT
XYLOCAINA 20 mg/ml soluzione iniettabile	not available	004535011	ASPEN PHARMA TRADING LIMITED	IT
XYLOCAINE 2% GEL (en tube), gel urétral	not available	2006048534	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 5%, pommade	not available	2006048535	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 1 %, solution injectable	not available	BE052631	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 1 %, solution injectable	not available	1995123279	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 1%, Injektionslösung	not available	BE052631	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 1%, Injektionslösung	not available	1995123279	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 1%, oplossing voor injectie	not available	BE052631	ASPEN PHARMA TRADING LIMITED	BE
Xylocaine 10 mg Spray	not available	PL 39699/0086	ASPEN PHARMA TRADING LIMITED	XI
Xylocaine 10 mg/delivered dose	not available	1691/028/001	ASPEN PHARMA TRADING LIMITED	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
Mucosal Spray				
Xylocaine 10 mg/delivered dose Mucosal Spray	not available	MA955/01201	ASPEN PHARMA TRADING LIMITED	MT
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 560 066 7 0	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 342 239 7 6	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 086 8 5	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 551 658 2 8	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 561 455 7 7	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 087 4 6	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 085 1 7	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 311 531 8 4	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 10 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 347 920 4 5	ASPEN PHARMA TRADING LIMITED	FR
Xylocaine 10% (pump spray) 10 mg/dose εκνέφωμα	not available	0057007	ASPEN PHARMA TRADING LIMITED	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
XYLOCAINE 10% , Spray, Lösung	not available	BE052796	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 10% , Spray, Lösung	not available	2006048536	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 10%, solution pour pulvérisation	not available	BE 052796	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 10%, solution pour pulvérisation	not available	2006048536	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 10%, spray oplossing	not available	BE 052796	ASPEN PHARMA TRADING LIMITED	BE
Xylocaine 100 mg/ml Spray, spray	not available	RVG 07831	ASPEN PHARMA TRADING LIMITED	NL
XYLOCAINE 2 %, gel urétral en seringue préremplie	not available	34009 363 446 1 7	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 2 %, gel urétral en seringue préremplie	not available	34009 565 324 4 5	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 2 %, solution injectable	not available	BE052735	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2 %, solution injectable	not available	1995123281	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 2% GEL (en seringue), gel urétral	not available	BE 149843	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2% GEL (en seringue), gel urétral	not available	0173/06048534	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 2% GEL (en tube), gel urétral	not available	BE052595	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2% GEL (in einer Spritze), Uretralgel	not available	BE149843	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2% GEL (in einer Spritze), Uretralgel	not available	2006048534	ASPEN PHARMA TRADING LIMITED	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
XYLOCAINE 2% GEL (in einer Tube), Uretralgel	not available	BE052595	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2% GEL (in einer Tube), Uretralgel	not available	0173/06048534	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 2% GEL (in spuit), gel voor urethraal gebruik	not available	BE149843	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2% GEL (in tube), gel voor urethraal gebruik	not available	BE052595	ASPEN PHARMA TRADING LIMITED	BE
Xylocaine 2% w/w γέλη	not available	0057004	ASPEN PHARMA TRADING LIMITED	GR
Xylocaine 2%, 20 mg/ml, roztwór do wstrzykiwań	not available	R/1309	ASPEN PHARMA TRADING LIMITED	PL
XYLOCAINE 2%, Injektionslösung	not available	BE052735	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 2%, Injektionslösung	not available	1995123281	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 2%, oplossing voor injectie	not available	BE052735	ASPEN PHARMA TRADING LIMITED	BE
Xylocaine 20 mg/g gel	not available	H/94/01682/002	ASPEN PHARMA TRADING LIMITED	SI
Xylocaine 20 mg/g Gel, catheterslijm	not available	RVG 07830	ASPEN PHARMA TRADING LIMITED	NL
Xylocaine 20 mg/ml raztopina za injiciranje	not available	H/94/01682/003	ASPEN PHARMA TRADING LIMITED	SI
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 083 9 5	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 082 2 7	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 20 mg/ml SANS CONSERVATEUR,	not available	34009 347 919 6 3	ASPEN PHARMA TRADING LIMITED	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
solution injectable				
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 561 454 0 9	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 342 240 5 8	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 560 067 3 1	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 20 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 084 5 6	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 560 065 0 2	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 089 7 5	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 347 921 0 6	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 342 150 6 3	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 561 456 3 8	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable	not available	34009 348 088 0 7	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE 5 mg/ml SANS CONSERVATEUR, solution injectable.	not available	34009 348 090 5 7	ASPEN PHARMA TRADING LIMITED	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
XYLOCAINE 5%, pommade	not available	BE 052561	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 5%, Salbe	not available	BE052561	ASPEN PHARMA TRADING LIMITED	BE
XYLOCAINE 5%, Salbe	not available	2006048535	ASPEN PHARMA TRADING LIMITED	LU
XYLOCAINE 5%, zalf	not available	BE052561	ASPEN PHARMA TRADING LIMITED	BE
Xylocaine 50 mg/g Zalf, hydrofiele zalf	not available	RVG 01553	ASPEN PHARMA TRADING LIMITED	NL
XYLOCAINE PUMP SPRAY 10%, 10 mg/annuses, nahasprei, lahus	not available	287099	ASPEN PHARMA TRADING LIMITED	EE
XYLOCAINE VISQUEUSE 2 %, gel oral	not available	34009 311 537 6 4	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE VISQUEUSE 2 %, gel oral	not available	34009 551 645 8 6	ASPEN PHARMA TRADING LIMITED	FR
Xylocaine® 10 mg/ml, oplossing voor injectie	not available	RVG 07828	ASPEN PHARMA TRADING LIMITED	NL
Xylocaine® 20 mg/ml, oplossing voor injectie	not available	RVG 07829	ASPEN PHARMA TRADING LIMITED	NL
XYLOCAINE® 5 POUR CENT NEBULISEUR, solution pour pulvérisation buccale	not available	34009 322 993 8 6	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE® 5 POUR CENT NEBULISEUR, solution pour pulvérisation buccale	not available	34009 555 788 8 8	ASPEN PHARMA TRADING LIMITED	FR
XYLOCAINE® 5 POUR CENT NEBULISEUR, solution pour pulvérisation buccale	not available	34009 555 760 6 8	ASPEN PHARMA TRADING LIMITED	FR
XYLOCARD 100, 20	not available	BE052832	ASPEN PHARMA TRADING	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
mg/ml, Injektionslösung			LIMITED	
XYLOCARD 100, 20 mg/ml, Injektionslösung	not available	2006028435	ASPEN PHARMA TRADING LIMITED	LU
XYLOCARD 100, 20 mg/ml, oplossing voor injectie	not available	BE052832	ASPEN PHARMA TRADING LIMITED	BE
XYLOCARD 100, 20 mg/ml, solution injectable	not available	BE 052832	ASPEN PHARMA TRADING LIMITED	BE
XYLOCARD 100, 20 mg/ml, solution injectable	not available	2006028435	ASPEN PHARMA TRADING LIMITED	LU
Xylocard 20 mg/ml injektionsvätska, lösning, glasampull	not available	22566	ASPEN PHARMA TRADING LIMITED	SE
Xylocard 20 mg/ml stungulyf, lausn, glerlykja	not available	711443	ASPEN PHARMA TRADING LIMITED	IS
XYLOCARD® 20 mg/ml INTRAVEINEUX, solution injectable	not available	374 970-9	ASPEN PHARMA TRADING LIMITED	FR
XYLOCARD® 20 mg/ml INTRAVEINEUX, solution injectable	not available	377 810-2	ASPEN PHARMA TRADING LIMITED	FR
XYLOCARD® 50mg/ml, solution injectable pour perfusion	not available	34009 550 941 2 8	ASPEN PHARMA TRADING LIMITED	FR
Xylonor Spray N 15%, Dentallösung	not available	53314.00.00	SEPTODONT GMBH	DE
Версатис 5% лечебен пластир	DE/H/5733/001	20100719	GRÜNENTHAL GMBH	BG
ЛИДБРИ 42 mg/ml вътрематочен гел	NL/H/4625/001	20200188	GEDEON RICHTER PLC.	BG