26 January 2023
EMA/101192/2023
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: levomethadone

Procedure no.: PSUSA/00001855/202205

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
Ellepalmiron 10 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/002	RVG 124250	DNE PHARMA AS	NL
Ellepalmiron 10 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/002	2203490.00.00	DNE PHARMA AS	DE
Ellepalmiron 10 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/002	999020	DNE PHARMA AS	EE
Ellepalmiron 15 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/003	RVG 124251	DNE PHARMA AS	NL
Ellepalmiron 15 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/003	2203491.00.00	DNE PHARMA AS	DE
Ellepalmiron 15 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/003	999120	DNE PHARMA AS	EE
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711050	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711062	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711074	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO 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
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711086	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711098	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ellepalmiron 2,5 mg/ml soluzione orale	DE/H/3805/003	043711100	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	IT
Ellepalmiron 20 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/004	RVG 124252	DNE PHARMA AS	NL
Ellepalmiron 20 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/004	2203492.00.00	DNE PHARMA AS	DE
Ellepalmiron 20 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/004	999220	DNE PHARMA AS	EE
Ellepalmiron 25 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/005	RVG 124253	DNE PHARMA AS	NL
Ellepalmiron 25 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/005	2203493.00.00	DNE PHARMA AS	DE
Ellepalmiron 25 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/005	999320	DNE PHARMA AS	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Ellepalmiron 30 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/006	RVG 124254	DNE PHARMA AS	NL
Ellepalmiron 30 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/006	2203494.00.00	DNE PHARMA AS	DE
Ellepalmiron 30 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/006	999420	DNE PHARMA AS	EE
Ellepalmiron 35 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/007	RVG 124555	DNE PHARMA AS	NL
Ellepalmiron 35 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/007	2203495.00.00	DNE PHARMA AS	DE
Ellepalmiron 35 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/007	999520	DNE PHARMA AS	EE
Ellepalmiron 40 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/008	RVG 124256	DNE PHARMA AS	NL
Ellepalmiron 40 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/008	2203496.00.00	DNE PHARMA AS	DE
Ellepalmiron 40 mg suukaudne lahus üheannuselises	NO/H/0301/008	999620	DNE PHARMA AS	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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Ellepalmiron 45 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/009	RVG 124257	DNE PHARMA AS	NL
Ellepalmiron 45 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/009	2203497.00.00	DNE PHARMA AS	DE
Ellepalmiron 45 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/009	999720	DNE PHARMA AS	EE
Ellepalmiron 5 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/001	RVG 124245	DNE PHARMA AS	NL
Ellepalmiron 5 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/001	2203489.00.00	DNE PHARMA AS	DE
Ellepalmiron 5 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/001	998920	DNE PHARMA AS	EE
Ellepalmiron 50 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/010	RVG 124258	DNE PHARMA AS	NL
Ellepalmiron 50 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/010	2203498.00.00	DNE PHARMA AS	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
Ellepalmiron 50 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/010	999820	DNE PHARMA AS	EE
Ellepalmiron 55 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/011	RVG 124259	DNE PHARMA AS	NL
Ellepalmiron 55 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/011	2203499.00.00	DNE PHARMA AS	DE
Ellepalmiron 55 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/011	999920	DNE PHARMA AS	EE
Ellepalmiron 60 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/012	RVG 124261	DNE PHARMA AS	NL
Ellepalmiron 60 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/012	2203500.00.00	DNE PHARMA AS	DE
Ellepalmiron 60 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/012	1000020	DNE PHARMA AS	EE
Ellepalmiron 65 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/013	RVG 124262	DNE PHARMA AS	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
Ellepalmiron 65 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/013	2203501.00.00	DNE PHARMA AS	DE
Ellepalmiron 65 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/013	1000120	DNE PHARMA AS	EE
Ellepalmiron 70 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/014	RVG 124263	DNE PHARMA AS	NL
Ellepalmiron 70 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/014	2203502.00.00	DNE PHARMA AS	DE
Ellepalmiron 70 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/014	1000220	DNE PHARMA AS	EE
Ellepalmiron 75 mg drank in verpakking voor éénmalig gebruik	NO/H/0301/015	RVG 124266	DNE PHARMA AS	NL
Ellepalmiron 75 mg Lösung zum Einnehmen im Einzeldosisbehältnis	NO/H/0301/015	2203503.00.00	DNE PHARMA AS	DE
Ellepalmiron 75 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/015	1000320	DNE PHARMA AS	EE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levometadon G.L. 10 mg tabletter	DK/H/2991/003	18-12539	G.L. PHARMA GMBH	NO
Levometadon G.L. 2,5 mg tabletter	DK/H/2991/001	18-12537	G.L. PHARMA GMBH	NO
Levometadon G.L. 20 mg tabletter	DK/H/2991/004	18-12540	G.L. PHARMA GMBH	NO
Levometadon G.L. 30 mg tabletter	DK/H/2991/005	18-12541	G.L. PHARMA GMBH	NO
Levometadon G.L. 5 mg tabletter	DK/H/2991/002	18-12538	G.L. PHARMA GMBH	NO
Levomethadon "G.L. Pharma", tabletter	DK/H/2991/001	61841	G.L. PHARMA GMBH	DK
Levomethadon "G.L. Pharma", tabletter	DK/H/2991/002	61842	G.L. PHARMA GMBH	DK
Levomethadon "G.L. Pharma", tabletter	DK/H/2991/003	61843	G.L. PHARMA GMBH	DK
Levomethadon "G.L. Pharma", tabletter	DK/H/2991/004	61844	G.L. PHARMA GMBH	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
Levomethadon "G.L. Pharma", tabletter	DK/H/2991/005	61845	G.L. PHARMA GMBH	DK
Levomethadon Aristo 2,5 mg Tabletten	DE/H/4849/001	98636.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Levomethadon Aristo 20 mg Tabletten	DE/H/4849/003	98638.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Levomethadon Aristo 30 mg Tabletten	DE/H/4849/004	98639.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Levomethadon Aristo 5 mg Tabletten	DE/H/4849/002	98637.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Levomethadon Aristo 7,5 mg Tabletten	DE/H/4849/005	2203449.00.00	ARISTO PHARMA GMBH (ART 57)	DE
Levomethadon Molteni 2,5 mg/ml Lösung zum Einnehmen	DE/H/3805/003	2202684.00.00	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	DE
Levomethadone G.L. 10 mg tabletter	DK/H/2991/003	58451	G.L. PHARMA GMBH	SE
Levomethadone G.L. 2,5 mg tabletter	DK/H/2991/001	58449	G.L. PHARMA GMBH	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
Levomethadone G.L. 20 mg tabletter	DK/H/2991/004	58452	G.L. PHARMA GMBH	SE
Levomethadone G.L. 30 mg tabletter	DK/H/2991/005	58453	G.L. PHARMA GMBH	SE
Levomethadone G.L. 5 mg tabletter	DK/H/2991/002	58450	G.L. PHARMA GMBH	SE
Levomethadone G.L. Pharma 10 mg tabletti	DK/H/2991/003	468700	G.L. PHARMA GMBH	FI
Levomethadone G.L. Pharma 2,5 mg tabletti	DK/H/2991/001	468698	G.L. PHARMA GMBH	FI
Levomethadone G.L. Pharma 20 mg tabletti	DK/H/2991/004	468701	G.L. PHARMA GMBH	FI
Levomethadone G.L. Pharma 30 mg tabletti	DK/H/2991/005	468702	G.L. PHARMA GMBH	FI
Levomethadone G.L. Pharma 5 mg tabletti	DK/H/2991/002	468699	G.L. PHARMA GMBH	FI
Levomethadone hydrochloride dne 10 mg mikstur, oppløsning i endosebeholder	NO/H/0301/002	18-12610	DNE PHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levomethadone hydrochloride dne 15 mg mikstur, oppløsning i endosebeholder	NO/H/0301/003	18-12611	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 20 mg mikstur, oppløsning i endosebeholder	NO/H/0301/004	18-12612	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 25 mg mikstur, oppløsning i endosebeholder	NO/H/0301/005	18-12613	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 30 mg mikstur, oppløsning i endosebeholder	NO/H/0301/006	18-12614	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 35 mg mikstur, oppløsning i endosebeholder	NO/H/0301/007	18-12615	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 40 mg mikstur, oppløsning i endosebeholder	NO/H/0301/008	18-12616	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 45 mg mikstur, oppløsning i endosebeholder	NO/H/0301/009	18-12617	DNE PHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levomethadone hydrochloride dne 5 mg mikstur, oppløsning i endosebeholder	NO/H/0301/001	18-12609	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 50 mg mikstur, oppløsning i endosebeholder	NO/H/0301/010	18-12618	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 55 mg mikstur, oppløsning i endosebeholder	NO/H/0301/011	18-12619	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 60 mg mikstur, oppløsning i endosebeholder	NO/H/0301/012	18-12620	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 65 mg mikstur, oppløsning i endosebeholder	NO/H/0301/013	18-12621	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 70 mg mikstur, oppløsning i endosebeholder	NO/H/0301/014	18-12622	DNE PHARMA AS	NO
Levomethadone hydrochloride dne 75 mg mikstur, oppløsning i endosebeholder	NO/H/0301/015	18-12623	DNE PHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levomethadone Hydrochloride Molteni, 2,5 mg/ml, roztwór doustny	DE/H/3805/003	25518	L. MOLTENI AND C. DEI F.LLI ALITTI SOCIETÀ DI ESERCIZIO S.P.A.	PL
Levo-Methasan 10 mg Tabletten	DE/H/5966/003	2203047.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 10 mg Tabletten	DE/H/5966/003	2020040098	G.L. PHARMA GMBH	LU
Levo-Methasan 2,5 mg Tabletten	DE/H/5966/001	2203045.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 2,5 mg Tabletten	DE/H/5966/001	2020040096	G.L. PHARMA GMBH	LU
Levo-Methasan 20 mg Tabletten	DE/H/5966/004	2203048.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 20 mg Tabletten	DE/H/5966/004	2020040099	G.L. PHARMA GMBH	LU
Levo-Methasan 30 mg Tabletten	DE/H/5966/005	2203049.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 30 mg Tabletten	DE/H/5966/005	20200400100	G.L. PHARMA GMBH	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
Levo-Methasan 5 mg Tabletten	DE/H/5966/002	2203046.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 5 mg Tabletten	DE/H/5966/002	2020040097	G.L. PHARMA GMBH	LU
Levo-Methasan® 10 mg-Tabletten	not available	140531	G.L. PHARMA GMBH	AT
Levo-Methasan® 2,5 mg-Tabletten	not available	140528	G.L. PHARMA GMBH	AT
Levo-Methasan® 30 mg-Tabletten	not available	140529	G.L. PHARMA GMBH	AT
Levo-Methasan® 20 mg-Tabletten		11428169	G.L. PHARMA GMBH	AT
Levo-Methasan® 5 mg-Tabletten		11428264	G.L. PHARMA GMBH	AT
Levopidon 10 mg mikstur, oppløsning	NO/H/0256/002	17-12068	DNE PHARMA AS	NO
Levopidon 10 mg oraaliliuos	NO/H/0256/002	35915	DNE PHARMA AS	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
Levopidon 10 mg oral lösning	NO/H/0256/002	57334	DNE PHARMA AS	SE
Levopidon 10 mg oral lösning i endosbehållare	NO/H/0256/002	35915	DNE PHARMA AS	FI
Levopidon 15 mg mikstur, oppløsning	NO/H/0256/003	17-12047	DNE PHARMA AS	NO
Levopidon 15 mg oraaliliuos	NO/H/0256/003	35844	DNE PHARMA AS	FI
Levopidon 15 mg oral lösning	NO/H/0256/003	57310	DNE PHARMA AS	SE
Levopidon 15 mg oral lösning i endosbehållare	NO/H/0256/003	35844	DNE PHARMA AS	FI
Levopidon 20 mg mikstur, oppløsning	NO/H/0256/004	17-12069	DNE PHARMA AS	NO
Levopidon 20 mg oraaliliuos	NO/H/0256/004	35916	DNE PHARMA AS	FI
Levopidon 20 mg oral lösning	NO/H/0256/004	57335	DNE PHARMA AS	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
Levopidon 20 mg oral lösning i endosbehållare	NO/H/0256/004	35916	DNE PHARMA AS	FI
Levopidon 25 mg mikstur, oppløsning	NO/H/0256/005	17-12070	DNE PHARMA AS	NO
Levopidon 25 mg oraaliliuos	NO/H/0256/005	35917	DNE PHARMA AS	FI
Levopidon 25 mg oral lösning	NO/H/0256/005	57336	DNE PHARMA AS	SE
Levopidon 25 mg oral lösning i endosbehållare	NO/H/0256/005	35917	DNE PHARMA AS	FI
Levopidon 30 mg mikstur, oppløsning	NO/H/0256/006	16-11328	DNE PHARMA AS	NO
Levopidon 30 mg oraaliliuos	NO/H/0256/006	34499	DNE PHARMA AS	FI
Levopidon 30 mg oral lösning	NO/H/0256/006	55302	DNE PHARMA AS	SE
Levopidon 30 mg oral lösning i endosbehållare	NO/H/0256/006	34499	DNE PHARMA AS	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
Levopidon 35 mg mikstur, oppløsning	NO/H/0256/007	16-11329	DNE PHARMA AS	NO
Levopidon 35 mg oraaliliuos	NO/H/0256/007	34500	DNE PHARMA AS	FI
Levopidon 35 mg oral lösning	NO/H/0256/007	55303	DNE PHARMA AS	SE
Levopidon 35 mg oral lösning i endosbehållare	NO/H/0256/007	34500	DNE PHARMA AS	FI
Levopidon 40 mg mikstur, oppløsning	NO/H/0256/008	16-11330	DNE PHARMA AS	NO
Levopidon 40 mg oraaliliuos	NO/H/0256/008	34501	DNE PHARMA AS	FI
Levopidon 40 mg oral lösning	NO/H/0256/008	55304	DNE PHARMA AS	SE
Levopidon 40 mg oral lösning i endosbehållare	NO/H/0256/008	34501	DNE PHARMA AS	FI
Levopidon 45 mg mikstur, oppløsning	NO/H/0256/009	17-11965	DNE PHARMA AS	NO

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Levopidon 45 mg oraaliliuos	NO/H/0256/009	35695	DNE PHARMA AS	FI
Levopidon 45 mg oral lösning	NO/H/0256/009	57025	DNE PHARMA AS	SE
Levopidon 45 mg oral lösning i endosbehållare	NO/H/0256/009	35695	DNE PHARMA AS	FI
Levopidon 5 mg mikstur, oppløsning	NO/H/0256/001	17-12046	DNE PHARMA AS	NO
Levopidon 5 mg oraaliliuos	NO/H/0256/001	35845	DNE PHARMA AS	FI
Levopidon 5 mg oral lösning	NO/H/0256/001	57309	DNE PHARMA AS	SE
Levopidon 5 mg oral lösning i endosbehållare	NO/H/0256/001	35845	DNE PHARMA AS	FI
Levopidon 50 mg mikstur, oppløsning	NO/H/0256/010	17-11966	DNE PHARMA AS	NO
Levopidon 50 mg oraaliliuos	NO/H/0256/010	35696	DNE PHARMA AS	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
Levopidon 50 mg oral lösning	NO/H/0256/010	57026	DNE PHARMA AS	SE
Levopidon 50 mg oral lösning i endosbehållare	NO/H/0256/010	35696	DNE PHARMA AS	FI
Levopidon 55 mg mikstur, oppløsning	NO/H/0256/011	17-11967	DNE PHARMA AS	NO
Levopidon 55 mg oraaliliuos	NO/H/0256/011	35697	DNE PHARMA AS	FI
Levopidon 55 mg oral lösning	NO/H/0256/011	57027	DNE PHARMA AS	SE
Levopidon 55 mg oral lösning i endosbehållare	NO/H/0256/011	35697	DNE PHARMA AS	FI
Levopidon 60 mg mikstur, oppløsning	NO/H/0256/012	17-11968	DNE PHARMA AS	NO
Levopidon 60 mg oraaliliuos	NO/H/0256/012	35698	DNE PHARMA AS	FI
Levopidon 60 mg oral lösning	NO/H/0256/012	57028	DNE PHARMA AS	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
Levopidon 60 mg oral lösning i endosbehållare	NO/H/0256/012	35698	DNE PHARMA AS	FI
Levopidon 65 mg mikstur, oppløsning	NO/H/0256/013	17-11969	DNE PHARMA AS	NO
Levopidon 65 mg oraaliliuos	NO/H/0256/013	35699	DNE PHARMA AS	FI
Levopidon 65 mg oral lösning	NO/H/0256/013	57029	DNE PHARMA AS	SE
Levopidon 65 mg oral lösning i endosbehållare	NO/H/0256/013	35699	DNE PHARMA AS	FI
Levopidon 70 mg mikstur, oppløsning	NO/H/0256/014	17-11970	DNE PHARMA AS	NO
Levopidon 70 mg oraaliliuos	NO/H/0256/014	35700	DNE PHARMA AS	FI
Levopidon 70 mg oral lösning	NO/H/0256/014	57030	DNE PHARMA AS	SE
Levopidon 70 mg oral lösning i endosbehållare	NO/H/0256/014	35700	DNE PHARMA AS	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
Levopidon 75 mg mikstur, oppløsning	NO/H/0256/015	17-11971	DNE PHARMA AS	NO
Levopidon 75 mg oraaliliuos	NO/H/0256/015	35701	DNE PHARMA AS	FI
Levopidon 75 mg oral lösning	NO/H/0256/015	57031	DNE PHARMA AS	SE
Levopidon 75 mg oral lösning i endosbehållare	NO/H/0256/015	35701	DNE PHARMA AS	FI
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/001	60729	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/002	60730	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/003	60731	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/004	60732	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/005	60733	DNE PHARMA AS	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
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/006	60734	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/007	60735	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/008	60736	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/009	60737	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/010	60738	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/011	60739	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/012	60740	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/013	60741	DNE PHARMA AS	DK
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/014	60742	DNE PHARMA AS	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
Levopidon, oral opløsning i enkeltdosisbeholder	NO/H/0256/015	60743	DNE PHARMA AS	DK
L-Poladdict 20 mg Tabletten	DE/H/4600/003	96251.00.00	HEXAL AG	DE
L-Poladdict 30 mg Tabletten	DE/H/4600/004	96252.00.00	HEXAL AG	DE
L-Poladdict 5 mg Tabletten	DE/H/4600/002	96250.00.00	HEXAL AG	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	1-31939	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND 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
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 20 mg Tabletten	DE/H/3528/003	86818.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	1-31938	SANOFI-AVENTIS DEUTSCHLAND GMBH	AT
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND 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
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE