29 January 2026
EMADOC-1700519818-2971691
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): levomethadone

EURD list No. PSUSA/00001855/202505

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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
Agovadin	DE/H/6794/001	049742075	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742190	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742202	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742214	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742226	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742238	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742240	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742253	G.L. PHARMA GMBH	IT
Agovadin 10 mg compresse	DE/H/6794/003	049742265	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742012	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742024	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742036	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742048	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742051	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742063	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742087	G.L. PHARMA GMBH	IT
Agovadin 2,5 mg compresse	DE/H/6794/001	049742099	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742277	G.L. PHARMA GMBH	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
Agovadin 20 mg compresse	DE/H/6794/004	049742289	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742291	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742303	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742315	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742327	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742339	G.L. PHARMA GMBH	IT
Agovadin 20 mg compresse	DE/H/6794/004	049742341	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742354	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742366	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742378	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742380	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742392	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742404	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742416	G.L. PHARMA GMBH	IT
Agovadin 30 mg compresse	DE/H/6794/005	049742428	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742101	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742113	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742125	G.L. PHARMA GMBH	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
Agovadin 5 mg compresse	DE/H/6794/002	049742137	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742149	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742152	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742164	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742176	G.L. PHARMA GMBH	IT
Agovadin 5 mg compresse	DE/H/6794/002	049742188	G.L. PHARMA GMBH	IT
Ellepalmiron 10 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/002	999020	DNE PHARMA AS	EE
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
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	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
			ESERCIZIO S.P.A.	
Ellepalmiron 20 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/004	999220	DNE PHARMA AS	EE
Ellepalmiron 25 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/005	999320	DNE PHARMA AS	EE
Ellepalmiron 30 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/006	999420	DNE PHARMA AS	EE
Ellepalmiron 35 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/007	999520	DNE PHARMA AS	EE
Ellepalmiron 40 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/008	999620	DNE PHARMA AS	EE
Ellepalmiron 45 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/009	999720	DNE PHARMA AS	EE
Ellepalmiron 5 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/001	998920	DNE PHARMA AS	EE
Ellepalmiron 50 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/010	999820	DNE PHARMA AS	EE
Ellepalmiron 55 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/011	999920	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 60 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/012	1000020	DNE PHARMA AS	EE
Ellepalmiron 65 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/013	1000120	DNE PHARMA AS	EE
Ellepalmiron 70 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/014	1000220	DNE PHARMA AS	EE
Ellepalmiron 75 mg suukaudne lahus üheannuselises konteineris	NO/H/0301/015	1000320	DNE PHARMA AS	EE
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 G.L. Pharma 10 mg Tabletten	DE/H/6794/003	7000726.00.00	G.L. PHARMA GMBH	DE
Levomethadon G.L. Pharma 2,5 mg Tabletten	DE/H/6794/001	7000724.00.00	G.L. PHARMA GMBH	DE
Levomethadon G.L. Pharma 20 mg Tabletten	DE/H/6794/004	7000727.00.00	G.L. PHARMA GMBH	DE
Levomethadon G.L. Pharma 30 mg Tabletten	DE/H/6794/005	7000728.00.00	G.L. PHARMA GMBH	DE
Levomethadon G.L. Pharma 5 mg Tabletten	DE/H/6794/002	7000725.00.00	G.L. PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
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 hydrochloride dne 10 mg mikstur, oppløsning i endosebeholder	NO/H/0301/002	18-12610	DNE PHARMA AS	NO
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
Levomethadone hydrochloride dne 5 mg	NO/H/0301/001	18-12609	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
mikstur, oppløsning i endosebeholder				
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
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 2,5 mg Tabletten	DE/H/5966/001	2203045.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 20 mg Tabletten	DE/H/5966/004	2203048.00.00	G.L. PHARMA GMBH	DE
Levo-Methasan 30 mg	DE/H/5966/005	2203049.00.00	G.L. PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tabletten				
Levo-Methasan 5 mg Tabletten	DE/H/5966/002	2203046.00.00	G.L. PHARMA GMBH	DE
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
Levopidon 10 mg mikstur, oppløsning i endosebeholder	NO/H/0256/002	17-12068	DNE PHARMA AS	NO
Levopidon 10 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/002	35915	DNE PHARMA AS	FI
Levopidon 10 mg oral lösning i endosbehållare	NO/H/0256/002	35915	DNE PHARMA AS	FI
Levopidon 10 mg oral lösning i endosbehållare	NO/H/0256/002	57334	DNE PHARMA AS	SE
Levopidon 10 mg tabletter	DK/H/2991/003	36513	DNE PHARMA AS	FI
Levopidon 10 mg tabletter	DK/H/2991/003	58451	DNE PHARMA AS	SE
Levopidon 10 mg tabletti	DK/H/2991/003	36513	DNE PHARMA AS	FI
Levopidon 15 mg mikstur, oppløsning i endosebeholder	NO/H/0256/003	17-12047	DNE PHARMA AS	NO
Levopidon 15 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/003	35844	DNE PHARMA AS	FI
Levopidon 15 mg oral lösning i endosbehållare	NO/H/0256/003	35844	DNE PHARMA AS	FI
Levopidon 15 mg oral lösning i endosbehållare	NO/H/0256/003	57310	DNE PHARMA AS	SE
Levopidon 2,5 mg tabletter	DK/H/2991/001	36511	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 2,5 mg tabletter	DK/H/2991/001	58449	DNE PHARMA AS	SE
Levopidon 2,5 mg tabletti	DK/H/2991/001	36511	DNE PHARMA AS	FI
Levopidon 20 mg mikstur, oppløsning i endosebeholder	NO/H/0256/004	17-12069	DNE PHARMA AS	NO
Levopidon 20 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/004	35916	DNE PHARMA AS	FI
Levopidon 20 mg oral lösning i endosbehållare	NO/H/0256/004	35916	DNE PHARMA AS	FI
Levopidon 20 mg oral lösning i endosbehållare	NO/H/0256/004	57335	DNE PHARMA AS	SE
Levopidon 20 mg tabletter	DK/H/2991/004	36514	DNE PHARMA AS	FI
Levopidon 20 mg tabletter	DK/H/2991/004	58452	DNE PHARMA AS	SE
Levopidon 20 mg tabletti	DK/H/2991/004	36514	DNE PHARMA AS	FI
Levopidon 25 mg mikstur, oppløsning i endosebeholder	NO/H/0256/005	17-12070	DNE PHARMA AS	NO
Levopidon 25 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/005	35917	DNE PHARMA AS	FI
Levopidon 25 mg oral lösning i endosbehållare	NO/H/0256/005	35917	DNE PHARMA AS	FI
Levopidon 25 mg oral lösning i endosbehållare	NO/H/0256/005	57336	DNE PHARMA AS	SE
Levopidon 30 mg mikstur, oppløsning i endosebeholder	NO/H/0256/006	16-11328	DNE PHARMA AS	NO
Levopidon 30 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/006	34499	DNE PHARMA AS	FI
Levopidon 30 mg oral	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
lösning i endosbehållare				
Levopidon 30 mg oral lösning i endosbehållare	NO/H/0256/006	55302	DNE PHARMA AS	SE
Levopidon 30 mg tabletter	DK/H/2991/005	36515	DNE PHARMA AS	FI
Levopidon 30 mg tabletter	DK/H/2991/005	58453	DNE PHARMA AS	SE
Levopidon 30 mg tabletti	DK/H/2991/005	36515	DNE PHARMA AS	FI
Levopidon 35 mg mikstur, oppløsning i endosebeholder	NO/H/0256/007	16-11329	DNE PHARMA AS	NO
Levopidon 35 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/007	34500	DNE PHARMA AS	FI
Levopidon 35 mg oral lösning i endosbehållare	NO/H/0256/007	34500	DNE PHARMA AS	FI
Levopidon 35 mg oral lösning i endosbehållare	NO/H/0256/007	55303	DNE PHARMA AS	SE
Levopidon 40 mg mikstur, oppløsning i endosebeholder	NO/H/0256/008	16-11330	DNE PHARMA AS	NO
Levopidon 40 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/008	34501	DNE PHARMA AS	FI
Levopidon 40 mg oral lösning i endosbehållare	NO/H/0256/008	34501	DNE PHARMA AS	FI
Levopidon 40 mg oral lösning i endosbehållare	NO/H/0256/008	55304	DNE PHARMA AS	SE
Levopidon 45 mg mikstur, oppløsning i endosebeholder	NO/H/0256/009	17-11965	DNE PHARMA AS	NO
Levopidon 45 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/009	35695	DNE PHARMA AS	FI
Levopidon 45 mg oral lösning i endosbehållare	NO/H/0256/009	35695	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 45 mg oral lösning i endosbehållare	NO/H/0256/009	57025	DNE PHARMA AS	SE
Levopidon 5 mg mikstur, oppløsning i endosebeholder	NO/H/0256/001	17-12046	DNE PHARMA AS	NO
Levopidon 5 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/001	35845	DNE PHARMA AS	FI
Levopidon 5 mg oral lösning i endosbehållare	NO/H/0256/001	35845	DNE PHARMA AS	FI
Levopidon 5 mg oral lösning i endosbehållare	NO/H/0256/001	57309	DNE PHARMA AS	SE
Levopidon 5 mg tabletter	DK/H/2991/002	36512	DNE PHARMA AS	FI
Levopidon 5 mg tabletter	DK/H/2991/002	58450	DNE PHARMA AS	SE
Levopidon 5 mg tabletti	DK/H/2991/002	36512	DNE PHARMA AS	FI
Levopidon 50 mg mikstur, oppløsning i endosebeholder	NO/H/0256/010	17-11966	DNE PHARMA AS	NO
Levopidon 50 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/010	35696	DNE PHARMA AS	FI
Levopidon 50 mg oral lösning i endosbehållare	NO/H/0256/010	35696	DNE PHARMA AS	FI
Levopidon 50 mg oral lösning i endosbehållare	NO/H/0256/010	57026	DNE PHARMA AS	SE
Levopidon 55 mg mikstur, oppløsning i endosebeholder	NO/H/0256/011	17-11967	DNE PHARMA AS	NO
Levopidon 55 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/011	35697	DNE PHARMA AS	FI
Levopidon 55 mg oral lösning i endosbehållare	NO/H/0256/011	35697	DNE PHARMA AS	FI
Levopidon 55 mg oral lösning i endosbehållare	NO/H/0256/011	57027	DNE PHARMA AS	SE
Levopidon 60 mg	NO/H/0256/012	17-11968	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
mikstur, oppløsning i endosebeholder				
Levopidon 60 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/012	35698	DNE PHARMA AS	FI
Levopidon 60 mg oral lösning i endosbehållare	NO/H/0256/012	35698	DNE PHARMA AS	FI
Levopidon 60 mg oral lösning i endosbehållare	NO/H/0256/012	57028	DNE PHARMA AS	SE
Levopidon 65 mg mikstur, oppløsning i endosebeholder	NO/H/0256/013	17-11969	DNE PHARMA AS	NO
Levopidon 65 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/013	35699	DNE PHARMA AS	FI
Levopidon 65 mg oral lösning i endosbehållare	NO/H/0256/013	35699	DNE PHARMA AS	FI
Levopidon 65 mg oral lösning i endosbehållare	NO/H/0256/013	57029	DNE PHARMA AS	SE
Levopidon 70 mg mikstur, oppløsning i endosebeholder	NO/H/0256/014	17-11970	DNE PHARMA AS	NO
Levopidon 70 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/014	35700	DNE PHARMA AS	FI
Levopidon 70 mg oral lösning i endosbehållare	NO/H/0256/014	35700	DNE PHARMA AS	FI
Levopidon 70 mg oral lösning i endosbehållare	NO/H/0256/014	57030	DNE PHARMA AS	SE
Levopidon 75 mg mikstur, oppløsning i endosebeholder	NO/H/0256/015	17-11971	DNE PHARMA AS	NO
Levopidon 75 mg oraaliliuos, kerta-annospakkaus	NO/H/0256/015	35701	DNE PHARMA AS	FI
Levopidon 75 mg oral lösning i endosbehållare	NO/H/0256/015	35701	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 oral lösning i endosbehållare	NO/H/0256/015	57031	DNE PHARMA AS	SE
Levopidon, tabletter	DK/H/2991/001	61841	DNE PHARMA AS	DK
Levopidon, tabletter	DK/H/2991/002	61842	DNE PHARMA AS	DK
Levopidon, tabletter	DK/H/2991/003	61843	DNE PHARMA AS	DK
Levopidon, tabletter	DK/H/2991/004	61844	DNE PHARMA AS	DK
Levopidon, tabletter	DK/H/2991/005	61845	DNE PHARMA AS	DK
Levopidon. 10 mg tabletter	DK/H/2991/003	18-12539	DNE PHARMA AS	NO
Levopidon. 2,5 mg tabletter	DK/H/2991/001	18-12537	DNE PHARMA AS	NO
Levopidon. 20 mg tabletter	DK/H/2991/004	18-12540	DNE PHARMA AS	NO
Levopidon. 30 mg tabletter	DK/H/2991/005	18-12541	DNE PHARMA AS	NO
Levopidon. 5 mg tabletter	DK/H/2991/002	18-12538	DNE PHARMA AS	NO
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 40 mg Tabletten	not available	7007265.00.00	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	86818.00.00	HEXAL AG	DE
L-Polamidon 5 mg Tabletten	DE/H/3528/002	86817.00.00	HEXAL AG	DE
L-Polamidon 5 mg/ml Lösung zum Einnehmen	not available	45583.00.00	HEXAL AG	DE
L-Polamidon 5 mg/ml Tropfen zum Einnehmen, Lösung	not available	6196782.00.00	HEXAL AG	DE