31 October 2018
EMA/792944/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): enoxaparin (except for biosimilars)

Procedure No.: PSUSA/00010560/201804

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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
CLEXANE	AT/H/0751/001	9700003	SANOFI BULGARIA EOOD	BG
CLEXANE	AT/H/0751/001	9700003	SANOFI BULGARIA EOOD	BG
CLEXANE	AT/H/0751/004	19161	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE	AT/H/0751/001	19159	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE	AT/H/0751/003	19744	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE	AT/H/0751/002	19160	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE	AT/H/0751/001	16/250/93-A/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/002	16/250/93-B/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/001	16/250/93-A/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/004	16/250/93-D/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/002	16/250/93-B/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/002	16/250/93-B/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/001	16/250/93-A/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/003	16/250/93-C/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/003	16/250/93-C/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/004	16/250/93-D/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/004	16/250/93-D/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/005	16/250/93-E/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/005	16/250/93-E/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/005	16/250/93-E/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/003	16/250/93-C/C	SANOFI-AVENTIS SRO	CZ
CLEXANE	AT/H/0751/002	28078/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/003	9059/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/004	9058/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/001	28077/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/002	28078/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/002	28078/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/003	9059/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/004	9058/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/001	28077/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/004	9058/92/26-10-93	SANOFI-AVENTIS AEBE	GR

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CLEXANE	AT/H/0751/001	28077/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/003	9059/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/001	28077/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/002	28078/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0754/001	59658/12-11-2003	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0754/002	59659/12-11-03	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0754/001	59658/12-11-2003	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0754/002	59659/12-11-03	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/002	026966046 (PRE-FILLED)	SANOFI SPA	IT
CLEXANE	AT/H/0751/001	026966034 (PRE-FILLED)	SANOFI SPA	IT
CLEXANE	AT/H/0752/001	MA082/00706	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/001	MA082/00702	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/002	MA082/00703	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/003	MA082/00704	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/005	MA082/00701	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/004	MA082/00705	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/004	MA082/00705	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/004	MA082/00705	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/005	MA082/00701	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/005	MA082/00701	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/001	MA082/00702	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/001	MA082/00702	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/002	MA082/00703	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/002	MA082/00703	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/003	MA082/00704	SANOFI MALTA LTD	MT
CLEXANE	AT/H/0751/003	MA082/00704	SANOFI MALTA LTD	MT
Clexane ® Multidose Vial 30,000 IU (300 mg)/3 ml solution for injection	AT/H/0752/001	PL 04425/0186	AVENTIS PHARMA LTD	UK
Clexane ® Multidose Vial 30,000 IU (300 mg)/3 ml solution for injection	AT/H/0752/001	PL 04425/0186	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 10,000 IU (100 mg)/1 ml	AT/H/0751/005	PL 04425/0187	AVENTIS PHARMA LTD	UK

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 for injection				
Clexane ® Syringes 10,000 IU (100 mg)/1 ml solution for injection	AT/H/0751/005	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 4,000 IU (40 mg)/0.4 ml solution for injection	AT/H/0751/002	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 4,000 IU (40 mg)/0.4 ml solution for injection	AT/H/0751/002	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 6,000 IU (60 mg)/0.6 ml solution for injection	AT/H/0751/003	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 6,000 IU (60 mg)/0.6 ml solution for injection	AT/H/0751/003	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 8,000 IU (80 mg)/0.8 ml solution for injection	AT/H/0751/004	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane ® Syringes 8,000 IU (80 mg)/0.8 ml solution for injection	AT/H/0751/004	PL 04425/0187	AVENTIS PHARMA LTD	UK
Clexane 10 000 IU (100 mg)/1 ml otopina za injekciju	AT/H/0751/005	HR-H-165030097-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 10 000 IU (100 mg)/1 ml otopina za injekciju	AT/H/0751/005	HR-H-165030097-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 10 000 NE (100 mg)/1 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/005	OGYI-T-4097/09	SANOFI-AVENTIS ZRT	HU
Clexane 10 000 NE (100 mg)/1 ml oldatos injekció előretöltött	AT/H/0751/005	OGYI-T-4097/10	SANOFI-AVENTIS ZRT	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
fecskendőben				
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	0555085	SANOFI BELGIUM	LU
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	0656589	SANOFI BELGIUM	LU
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	0190485	SANOFI BELGIUM	LU
Clexane 10,000 IU (100mg)/1ml Solution for Injection	AT/H/0751/005	540/97/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 10,000 IU (100mg)/1ml Solution for Injection	AT/H/0751/005	540/97/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 10,000 IU (100mg)/1ml Solution for Injection	AT/H/0751/005	540/97/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 10.000 i.e. (100 mg)/1 ml raztopina za injiciranje	AT/H/0751/005	H/99/00399/005	SANOFI-AVENTIS D.O.O.	SI
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 10.000 I.E. (100	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS	DE

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mg)/1,0 ml Injektionslösung			DEUTSCHLAND GMBH	
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 10.000 I.E. (100 mg)/1,0 ml Injektionslösung	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 10.000 IE (100 mg)/1 ml oplossing voor injectie	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
CLEXANE 10.000 IE (100 mg)/1 ml oplossing voor injectie	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
CLEXANE 10.000 IE (100 mg)/1 ml oplossing voor injectie	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
Clexane 10.000 IE (100 mg)/1 ml oplossing voor injectie	AT/H/0751/005	RVG 121190	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 10.000 UI (100 mg)/1 ml solución inyectable	AT/H/0751/005	62472	SANOFI-AVENTIS, S.A.	ES
CLEXANE 10.000 UI (100 mg)/1 ml solución inyectable	AT/H/0751/005	62472	SANOFI-AVENTIS, 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
CLEXANE 10.000 UI (100 mg)/1 ml solución inyectable	AT/H/0751/005	62472	SANOFI-AVENTIS, S.A.	ES
CLEXANE 10000 IU (100 mg)/1 ml injekčný roztok	AT/H/0751/005	16/0175/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 10000 IU (100 mg)/1 ml injekčný roztok	AT/H/0751/005	16/0175/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 10000 IU (100 mg)/1 ml injekčný roztok	AT/H/0751/005	16/0175/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	0316492	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	0656592	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable	AT/H/0754/001	0555099	SANOFI BELGIUM	LU
Clexane 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje	AT/H/0754/001	H/99/00399/006	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 IE (120 mg)/0,8 ml oplossing voor injectie	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
CLEXANE 12.000 IE (120 mg)/0,8 ml oplossing	AT/H/0754/001	BE230176	SANOFI BELGIUM	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				
CLEXANE 12.000 IE (120 mg)/0,8 ml oplossing voor injectie	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
Clexane 12.000 IE (120 mg)/0,8 ml oplossing voor injectie	AT/H/0754/001	RVG 121191	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 12.000 UI (120 mg)/0,8 ml solución inyectable	AT/H/0754/001	63002	SANOFI-AVENTIS, S.A.	ES
CLEXANE 12.000 UI (120 mg)/0,8 ml solución inyectable	AT/H/0754/001	63002	SANOFI-AVENTIS, S.A.	ES
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	0555104	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	0316489	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	0656608	SANOFI BELGIUM	LU
Clexane 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje	AT/H/0754/002	H/99/00399/007	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 IE (150	AT/H/0754/002	BE230185	SANOFI BELGIUM	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)/1 ml oplossing voor injectie				
CLEXANE 15.000 IE (150 mg)/1 ml oplossing voor injectie	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
CLEXANE 15.000 IE (150 mg)/1 ml oplossing voor injectie	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
Clexane 15.000 IE (150 mg)/1 ml oplossing voor injectie	AT/H/0754/002	RVG 121192	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 15.000 UI (150 mg)/1 ml solución inyectable	AT/H/0754/002	63000	SANOFI-AVENTIS, S.A.	ES
CLEXANE 15.000 UI (150 mg)/1 ml solución inyectable	AT/H/0754/002	63000	SANOFI-AVENTIS, S.A.	ES
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	0555037	SANOFI BELGIUM	LU
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	0173128	SANOFI BELGIUM	LU

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CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	0829114	SANOFI BELGIUM	LU
Clexane 2,000 IU (20 mg) /0.2 mL Solution for Injection	AT/H/0751/001	540/97/4	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 2,000 IU (20 mg) /0.2 mL Solution for Injection	AT/H/0751/001	540/97/4	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 2,000 IU (20 mg) /0.2 mL Solution for Injection	AT/H/0751/001	540/97/4	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE

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Injektionslösung				
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I.E. (20 mg)/0,2 ml Injektionslösung	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 i.e. (20 mg)/0,2 ml raztopina za injiciranje	AT/H/0751/001	H/99/00399/001	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
CLEXANE 2.000 IE (20	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE

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mg)/0,2 ml oplossing voor injectie				
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie	AT/H/0751/001	BE144365	SANOFI BELGIUM	BE
Clexane 2.000 IE (20 mg)/0,2 ml oplossing voor injectie	AT/H/0751/001	RVG 121186	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 2.000 UI (20 mg)/0,2 ml solución inyectable	AT/H/0751/001	58502	SANOFI-AVENTIS, S.A.	ES
CLEXANE 2.000 UI (20 mg)/0,2 ml solución inyectable	AT/H/0751/001	58502	SANOFI-AVENTIS, S.A.	ES
CLEXANE 2.000 UI (20 mg)/0,2 ml solución inyectable	AT/H/0751/001	58502	SANOFI-AVENTIS, S.A.	ES
CLEXANE 2.000 UI (20 mg)/0,2 ml soluzione iniettabile	AT/H/0751/001	026966059	SANOFI S.P.A	IT
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 20 MG KLINIK	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 2000 IU (20 mg)/0,2 ml injekčný roztok	AT/H/0751/001	16/0108/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
CLEXANE 2000 IU (20 mg)/0,2 ml injekčný roztok	AT/H/0751/001	16/0108/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 2000 IU (20 mg)/0,2 ml injekčný roztok	AT/H/0751/001	16/0108/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane 2000 IU (20 mg)/0,2 ml otopina za injekciju	AT/H/0751/001	HR-H-099632462-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 2000 IU (20 mg)/0,2 ml otopina za injekciju	AT/H/0751/001	HR-H-099632462-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 2000 NE (20 mg)/0,2 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/001	OGYI-T-4097/01	SANOFI-AVENTIS ZRT	HU
Clexane 2000 NE (20 mg)/0,2 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/001	OGYI-T-4097/02	SANOFI-AVENTIS ZRT	HU
CLEXANE 2000 SV (20 mg)/0,2 ml šķīdums injekcijām	AT/H/0751/001	98-0717	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 2000 SV (20 mg)/0,2 ml šķīdums injekcijām	AT/H/0751/001	98-0717	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 2000 SV (20 mg)/0,2 ml šķīdums injekcijām	AT/H/0751/001	98-0717	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 2000 TV (20 mg)/0,2 ml injekcinis tirpalas	AT/H/0751/001	LT/1/98/1560/006	UAB SANOFI-AVENTIS LIETUVA	LT
CLEXANE 2000 TV (20 mg)/0,2 ml injekcinis tirpalas	AT/H/0751/001	LT/1/98/1560/005	UAB SANOFI-AVENTIS LIETUVA	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Clexane 2000 UI (20 mg)/0,2 ml soluţie injectabilă	AT/H/0751/001	4926/2012/01	SANOFI ROMANIA SRL	RO
Clexane 2000 UI (20 mg)/0,2 ml soluţie injectabilă	AT/H/0751/001	4926/2012/02	SANOFI ROMANIA SRL	RO
Clexane 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	OGYI-T-4097/11	SANOFI-AVENTIS ZRT	HU
CLEXANE 30000 IU (300 mg)/3 ml injekčný roztok	AT/H/0752/001	16/0176/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	0173145	SANOFI BELGIUM	LU
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	0555041	SANOFI BELGIUM	LU
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	0829128	SANOFI BELGIUM	LU
Clexane 4,000 IU (40 mg) /0.4 mL Solution for Injection	AT/H/0751/002	540/97/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	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
Clexane 4,000 IU (40 mg) /0.4 mL Solution for Injection	AT/H/0751/002	540/97/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 4,000 IU (40 mg) /0.4 mL Solution for Injection	AT/H/0751/002	540/97/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis	DE/H/5213/002	32766.01.00	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
Injektionslösung				
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I.E. (40 mg)/0,4 ml Injektionslösung	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 i.e. (40 mg)/0,4 ml raztopina za injiciranje	AT/H/0751/002	H/99/00399/002	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
Clexane 4.000 IE (40	AT/H/0751/002	RVG 121187	SANOFI-AVENTIS	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)/0,4 ml oplossing voor injectie			NETHERLANDS B.V.	
CLEXANE 4.000 UI (40 mg)/0,4 ml solución inyectable	AT/H/0751/002	58503	SANOFI-AVENTIS, S.A.	ES
CLEXANE 4.000 UI (40 mg)/0,4 ml solución inyectable	AT/H/0751/002	58503	SANOFI-AVENTIS, S.A.	ES
CLEXANE 4.000 UI (40 mg)/0,4 ml solución inyectable	AT/H/0751/002	58503	SANOFI-AVENTIS, S.A.	ES
CLEXANE 4.000 UI (40 mg)/0,4 ml solución inyectable	AT/H/0751/002	58503	SANOFI-AVENTIS, S.A.	ES
CLEXANE 4.000 UI (40 mg)/0,4 ml soluzione iniettabile	AT/H/0751/002	026966061	SANOFI S.P.A	IT
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 40 MG KLINIK	DE/H/5212/002	16071.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 4000 IU (40 mg)/0,4 ml injekčný roztok	AT/H/0751/002	16/0172/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 4000 IU (40 mg)/0,4 ml injekčný roztok	AT/H/0751/002	16/0172/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
CLEXANE 4000 IU (40 mg)/0,4 ml injekčný roztok	AT/H/0751/002	16/0172/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane 4000 IU (40 mg)/0,4 ml otopina za injekciju	AT/H/0751/002	HR-H-990481301-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 4000 IU (40 mg)/0,4 ml otopina za injekciju	AT/H/0751/002	HR-H-990481301-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 4000 NE (40 mg)/0,4 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/002	OGYI-T-4097/04	SANOFI-AVENTIS ZRT	HU
Clexane 4000 NE (40 mg)/0,4 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/002	OGYI-T-4097/03	SANOFI-AVENTIS ZRT	HU
CLEXANE 4000 SV (40 mg)/0,4 ml šķīdums injekcijām	AT/H/0751/002	98-0718	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 4000 SV (40 mg)/0,4 ml šķīdums injekcijām	AT/H/0751/002	98-0718	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 4000 SV (40 mg)/0,4 ml šķīdums injekcijām	AT/H/0751/002	98-0718	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 4000 TV (40 mg)/0,4 ml injekcinis tirpalas	AT/H/0751/002	LT/1/98/1560/008	UAB SANOFI-AVENTIS LIETUVA	LT
CLEXANE 4000 TV (40 mg)/0,4 ml injekcinis tirpalas	AT/H/0751/002	LT/1/98/1560/007	UAB SANOFI-AVENTIS LIETUVA	LT
Clexane 4000 UI (40 mg)/0,4 ml soluţie injectabilă	AT/H/0751/002	4927/2012/02	SANOFI ROMANIA SRL	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
Clexane 4000 UI (40 mg)/0,4 ml soluţie injectabilă	AT/H/0751/002	4927/2012/01	SANOFI ROMANIA SRL	RO
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	0267613	SANOFI BELGIUM	LU
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	0555054	SANOFI BELGIUM	LU
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	0656558	SANOFI BELGIUM	LU
Clexane 6,000 IU (60 mg) /0.6 mL Solution for Injection	AT/H/0751/003	540/97/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 6,000 IU (60 mg) /0.6 mL Solution for Injection	AT/H/0751/003	540/97/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 6,000 IU (60 mg) /0.6 mL Solution for Injection	AT/H/0751/003	540/97/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 I.E. (60 mg)/0,6 ml	AT/H/0751/003	40759.00.00	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
Injektionslösung				
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 I.E. (60 mg)/0,6 ml Injektionslösung	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 6.000 i.e. (60 mg)/0,6 ml raztopina za injiciranje	AT/H/0751/003	H/99/00399/003	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 6.000 IE (60 mg)/0,6 ml oplossing voor injectie	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 6.000 IE (60 mg)/0,6 ml oplossing voor injectie	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 6.000 IE (60 mg)/0,6 ml oplossing voor injectie	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
Clexane 6.000 IE (60 mg)/0,6 ml oplossing voor injectie	AT/H/0751/003	RVG 121188	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 6.000 IU (60 mg)/0,6 mL ενέσιμο διάλυμα	AT/H/0751/003	19744	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE 6.000 UI (60	AT/H/0751/003	62470	SANOFI-AVENTIS, 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
mg)/0,6 ml solución inyectable				
CLEXANE 6.000 UI (60 mg)/0,6 ml solución inyectable	AT/H/0751/003	62470	SANOFI-AVENTIS, S.A.	ES
CLEXANE 6.000 UI (60 mg)/0,6 ml solución inyectable	AT/H/0751/003	62470	SANOFI-AVENTIS, S.A.	ES
CLEXANE 6000 IU (60 mg)/0,6 ml injekčný roztok	AT/H/0751/003	16/0173/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 6000 IU (60 mg)/0,6 ml injekčný roztok	AT/H/0751/003	16/0173/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 6000 IU (60 mg)/0,6 ml injekčný roztok	AT/H/0751/003	16/0173/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane 6000 IU (60 mg)/0,6 ml otopina za injekciju	AT/H/0751/003	HR-H-041381787-01	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 6000 IU (60 mg)/0,6 ml otopina za injekciju	AT/H/0751/003	HR-H-041381787-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 6000 NE (60 mg)/0,6 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/003	OGYI-T-4097/06	SANOFI-AVENTIS ZRT	HU
Clexane 6000 NE (60 mg)/0,6 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/003	OGYI-T-4097/05	SANOFI-AVENTIS ZRT	HU
CLEXANE 6000 SV (60 mg)/0,6 ml šķīdums injekcijām	AT/H/0751/003	98-0719	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 6000 SV (60	AT/H/0751/003	98-0719	SANOFI-AVENTIS LATVIA	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
mg)/0,6 ml šķīdums injekcijām			SIA	
CLEXANE 6000 SV (60 mg)/0,6 ml šķīdums injekcijām	AT/H/0751/003	98-0719	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 6000 TV (60 mg)/0,6 ml injekcinis tirpalas	AT/H/0751/003	LT/1/98/1560/009	UAB SANOFI-AVENTIS LIETUVA	LT
CLEXANE 6000 TV (60 mg)/0,6 ml injekcinis tirpalas	AT/H/0751/003	LT/1/98/1560/010	UAB SANOFI-AVENTIS LIETUVA	LT
Clexane 6000 UI (60 mg)/0,6 ml soluţie injectabilă	AT/H/0751/003	4928/2012/02	SANOFI ROMANIA SRL	RO
Clexane 6000 UI (60 mg)/0,6 ml soluţie injectabilă	AT/H/0751/003	4928/2012/01	SANOFI ROMANIA SRL	RO
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	0190454	SANOFI BELGIUM	LU
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	0555068	SANOFI BELGIUM	LU
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	0656561	SANOFI BELGIUM	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
Clexane 8,000 IU (80 mg) /0.8 mL Solution for Injection	AT/H/0751/004	540/97/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 8,000 IU (80 mg) /0.8 mL Solution for Injection	AT/H/0751/004	540/97/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 8,000 IU (80 mg) /0.8 mL Solution for Injection	AT/H/0751/004	540/97/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 I.E. (80 mg)/0,8 ml Injektionslösung	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 8.000 i.e. (80 mg)/0,8 ml raztopina za injiciranje	AT/H/0751/004	H/99/00399/004	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 8.000 IE (80 mg)/0,8 ml oplossing	AT/H/0751/004	BE161953	SANOFI BELGIUM	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				
CLEXANE 8.000 IE (80 mg)/0,8 ml oplossing voor injectie	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
CLEXANE 8.000 IE (80 mg)/0,8 ml oplossing voor injectie	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
Clexane 8.000 IE (80 mg)/0,8 ml oplossing voor injectie	AT/H/0751/004	RVG 121189	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 8.000 IU (80 mg)/0,8 mL ενέσιμο διάλυμα	AT/H/0751/004	19161	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE 8.000 UI (80 mg)/0,8 ml solución inyectable	AT/H/0751/004	62471	SANOFI-AVENTIS, S.A.	ES
CLEXANE 8.000 UI (80 mg)/0,8 ml solución inyectable	AT/H/0751/004	62471	SANOFI-AVENTIS, S.A.	ES
CLEXANE 8.000 UI (80 mg)/0,8 ml solución inyectable	AT/H/0751/004	62471	SANOFI-AVENTIS, S.A.	ES
CLEXANE 8000 IU (80 mg)/0,8 ml injekčný roztok	AT/H/0751/004	16/0174/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 8000 IU (80 mg)/0,8 ml injekčný roztok	AT/H/0751/004	16/0174/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 8000 IU (80 mg)/0,8 ml injekčný roztok	AT/H/0751/004	16/0174/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane 8000 IU (80 mg)/0,8 ml otopina za injekciju	AT/H/0751/004	HR-H-173049079-02	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 8000 IU (80	AT/H/0751/004	HR-H-173049079-01	SANOFI-AVENTIS CROATIA	HR

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)/0,8 ml otopina za injekciju			D.O.O.	
Clexane 8000 NE (80 mg)/0,8 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/004	OGYI-T-4097/08	SANOFI-AVENTIS ZRT	HU
Clexane 8000 NE (80 mg)/0,8 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/004	OGYI-T-4097/07	SANOFI-AVENTIS ZRT	HU
CLEXANE 8000 SV (80 mg)/0,8 ml šķīdums injekcijām	AT/H/0751/004	98-0720	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 8000 SV (80 mg)/0,8 ml šķīdums injekcijām	AT/H/0751/004	98-0720	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 8000 SV (80 mg)/0,8 ml šķīdums injekcijām	AT/H/0751/004	98-0720	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 8000 TV (80 mg)/0,8 ml injekcinis tirpalas	AT/H/0751/004	LT/1/98/1560/012	UAB SANOFI-AVENTIS LIETUVA	LT
CLEXANE 8000 TV (80 mg)/0,8 ml injekcinis tirpalas	AT/H/0751/004	LT/1/98/1560/011	UAB SANOFI-AVENTIS LIETUVA	LT
Clexane 8000 UI (80 mg)/0,8 ml soluţie injectabilă	AT/H/0751/004	4929/2012/02	SANOFI ROMANIA SRL	RO
Clexane 8000 UI (80 mg)/0,8 ml soluţie injectabilă	AT/H/0751/004	4928/2012/01	SANOFI ROMANIA SRL	RO
CLEXANE FORTE	AT/H/0754/002	16/338/01-B/C	SANOFI-AVENTIS SRO	CZ
CLEXANE FORTE	AT/H/0754/001	16/338/01-A/C	SANOFI-AVENTIS SLOVAKIA SRO	CZ
CLEXANE FORTE	AT/H/0754/002	16/338/01-B/C	SANOFI-AVENTIS SRO	CZ

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
CLEXANE FORTE	AT/H/0754/001	16/338/01-A/C	SANOFI-AVENTIS SLOVAKIA SRO	CZ
CLEXANE FORTE	AT/H/0754/002	8915	AVENTIS PHARMA LTD	PL
CLEXANE FORTE	AT/H/0754/001	8914	AVENTIS PHARMA LTD	PL
CLEXANE FORTE 12 000 IU (120 mg)/0,8 ml injekčný roztok	AT/H/0754/001	16/0033/02-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE FORTE 12 000 IU (120 mg)/0,8 ml injekčný roztok	AT/H/0754/001	16/0033/02-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane Forte 12 000 NE (120 mg)/0,8 ml oldatos injekció előretöltött fecskendőben	AT/H/0754/001	OGYI-T-4097/12	SANOFI-AVENTIS ZRT	HU
Clexane Forte 12 000 NE (120 mg)/0,8 ml oldatos injekció előretöltött fecskendőben	AT/H/0754/001	OGYI-T-4097/13	SANOFI-AVENTIS ZRT	HU
Clexane Forte 12,000 IU (120 mg) / 0.8 mL Solution for Injection	AT/H/0754/001	540/97/8	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 12,000 IU (120 mg) / 0.8 mL Solution for Injection	AT/H/0754/001	540/97/8	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 12,000 IU (120 mg) / 0.8 mL Solution for Injection	AT/H/0754/001	540/97/8	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 12,000 IU (120 mg) / 0.8 mL Solution for Injection	AT/H/0754/001	540/97/8	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
CLEXANE FORTE 15 000 IU (150 mg)/1 ml injekčný roztok	AT/H/0754/002	19/0171/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE FORTE 15 000 IU (150 mg)/1 ml	AT/H/0754/002	16/0171/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK

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
injekčný roztok				
Clexane Forte 15 000 NE (150 mg)/1 ml oldatos injekció előretöltött fecskendőben	AT/H/0754/002	OGYI-T-4097/14	SANOFI-AVENTIS ZRT	HU
Clexane Forte 15 000 NE (150 mg)/1 ml oldatos injekció előretöltött fecskendőben	AT/H/0754/002	OGYI-T-4097/15	SANOFI-AVENTIS ZRT	HU
Clexane Forte 15,000 IU (150mg)/1ml Solution for Injection	AT/H/0754/002	540/97/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 15,000 IU (150mg)/1ml Solution for Injection	AT/H/0754/002	540/97/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 15,000 IU (150mg)/1ml Solution for Injection	AT/H/0754/002	540/97/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane Forte 15,000 IU (150mg)/1ml Solution for Injection	AT/H/0754/002	540/97/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
CLEXANE FORTE SYRINGES	AT/H/0754/002	PL 04425/0185	AVENTIS PHARMA LTD	UK
CLEXANE FORTE SYRINGES	AT/H/0754/002	PL 04425/0185	AVENTIS PHARMA LTD	UK
CLEXANE FORTE SYRINGES	AT/H/0754/002	PL 04425/0185	AVENTIS PHARMA LTD	UK
CLEXANE FORTE SYRINGES	AT/H/0754/001	PL 04425/0185	AVENTIS PHARMA LTD	UK
CLEXANE FORTE SYRINGES	AT/H/0754/001	PL 04425/0185	AVENTIS PHARMA LTD	UK
CLEXANE FORTE SYRINGES	AT/H/0754/001	PL 04425/0185	AVENTIS PHARMA LTD	UK
Clexane multidose 30.000 I. E. (300 mg)/3	DE/H/5218/001	50480.00.00	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
ml Praxis Injektionslösung				
Clexane multidose 30.000 I. E. (300 mg)/3 ml Praxis Injektionslösung	DE/H/5218/001	50480.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 30.000 I. E. (300 mg)/3 ml Praxis Injektionslösung	DE/H/5218/001	50480.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 30.000 I. E. (300 mg)/3 ml Praxis Injektionslösung	DE/H/5218/001	50480.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 30.000 I.E. (300 mg)/3 ml Injektionslösung	AT/H/0752/001	40758.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 30.000 I.E. (300 mg)/3 ml Injektionslösung	AT/H/0752/001	40758.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 30.000 I.E. (300 mg)/3 ml Injektionslösung	AT/H/0752/001	40758.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE MULTIDOSE VIAL	AT/H/0752/001	PL 04425/0186	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/002	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/003	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/005	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/001	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/001	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/001	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE SYRINGES	AT/H/0751/004	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE T	AT/H/0752/001	029111073	SANOFI SPA	IT
CLEXANE T	AT/H/0751/003	029111085 (PRE-FILLED)	SANOFI SPA	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
CLEXANE T	AT/H/0751/004	029111059	SANOFI SPA	IT
CLEXANE T	AT/H/0751/004	029111097 (PRE-FILLED)	SANOFI SPA	IT
CLEXANE T	AT/H/0751/003	029111046	SANOFI SPA	IT
CLEXANE T	AT/H/0751/005	029111061	SANOFI SPA	IT
CLEXANE T	AT/H/0751/005	029111109 (PRE-FILLED)	SANOFI SPA	IT
CLEXANE T 10.000 UI (100 mg)/1 ml soluzione iniettabile	AT/H/0751/005	029111135	SANOFI S.P.A	IT
CLEXANE T 6.000 UI (60 mg)/0,6 ml soluzione iniettabile	AT/H/0751/003	029111111	SANOFI S.P.A	IT
CLEXANE T 8.000 UI (80 mg)/0,8 ml soluzione iniettabile	AT/H/0751/004	029111123	SANOFI S.P.A	IT
Clexane, 10 000 j.m. (100 mg) /1 ml, roztwór do wstrzykiwań	AT/H/0751/005	7749	SANOFI-AVENTIS FRANCE	PL
Clexane, 10 000 j.m. (100 mg) /1 ml, roztwór do wstrzykiwań	AT/H/0751/005	7749	SANOFI-AVENTIS FRANCE	PL
Clexane, 10 000 j.m. (100 mg) /ml (30 000 j.m. (300 mg) /3 ml), roztwór do wstrzykiwań	AT/H/0752/001	4561	SANOFI-AVENTIS FRANCE	PL
Clexane, 10 000 j.m. (100 mg) /ml (30 000 j.m. (300 mg) /3 ml), roztwór do wstrzykiwań	AT/H/0752/001	4561	SANOFI-AVENTIS FRANCE	PL
CLEXANE, 10 000 RÜ (100 mg)/1 ml süstelahus	AT/H/0751/005	240398	SANOFI-AVENTIS ESTONIA OÜ	EE
CLEXANE, 10 000 RÜ (100 mg)/1 ml süstelahus	AT/H/0751/005	240398	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane, 2000 j.m. (20	AT/H/0751/001	R/0483	SANOFI-AVENTIS FRANCE	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
mg) /0,2 ml, roztwór do wstrzykiwań				
CLEXANE, 2000 RÜ (20 mg)/0,2 ml süstelahus	AT/H/0751/001	940317	SANOFI-AVENTIS ESTONIA OÜ	EE
CLEXANE, 2000 RÜ (20 mg)/0,2 ml süstelahus	AT/H/0751/001	940317	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane, 4000 j.m. (40 mg) /0,4 ml, roztwór do wstrzykiwań	AT/H/0751/002	R/0484	SANOFI-AVENTIS FRANCE	PL
CLEXANE, 4000 RÜ (40 mg)/0,4 ml süstelahus	AT/H/0751/002	940417	SANOFI-AVENTIS ESTONIA OÜ	EE
CLEXANE, 4000 RÜ (40 mg)/0,4 ml süstelahus	AT/H/0751/002	940417	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane, 6000 j.m. (60 mg) /0,6 ml, roztwór do wstrzykiwań	AT/H/0751/003	7748	SANOFI-AVENTIS FRANCE	PL
Clexane, 6000 j.m. (60 mg) /0,6 ml, roztwór do wstrzykiwań	AT/H/0751/003	7748	SANOFI-AVENTIS FRANCE	PL
CLEXANE, 6000 RÜ (60 mg)/0,6 ml süstelahus	AT/H/0751/003	940517	SANOFI-AVENTIS ESTONIA OÜ	EE
CLEXANE, 6000 RÜ (60 mg)/0,6 ml süstelahus	AT/H/0751/003	940517	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane, 8000 j.m. (80 mg) /0,8 ml, roztwór do wstrzykiwań	AT/H/0751/004	7750	SANOFI-AVENTIS FRANCE	PL
Clexane, 8000 j.m. (80 mg) /0,8 ml, roztwór do wstrzykiwań	AT/H/0751/004	7750	SANOFI-AVENTIS FRANCE	PL
CLEXANE, 8000 RÜ (80 mg)/0,8 ml süstelahus	AT/H/0751/004	940617	SANOFI-AVENTIS ESTONIA OÜ	EE
CLEXANE, 8000 RÜ (80 mg)/0,8 ml süstelahus	AT/H/0751/004	940617	SANOFI-AVENTIS ESTONIA OÜ	EE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1	DE/H/5211/005	78637.00.00	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**
ml Injektionslösung				
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung	DE/H/5211/001	16349.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung	DE/H/5211/001	16349.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung	DE/H/5211/001	16349.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung	DE/H/5211/001	16349.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung	DE/H/5211/001	16349.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 4.000	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS	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
I. E. (40 mg)/0,4 ml Injektionslösung			DEUTSCHLAND GMBH	
Enoxaparin Sanofi 4.000 I. E. (40 mg)/0,4 ml Injektionslösung	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 4.000 I. E. (40 mg)/0,4 ml Injektionslösung	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 4.000 I. E. (40 mg)/0,4 ml Injektionslösung	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 4.000 I. E. (40 mg)/0,4 ml Injektionslösung	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 6.000 I. E. (60 mg)/0,6 ml Injektionslösung	DE/H/5211/003	78635.00.00	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
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E.	DE/H/5216/001	50479.00.00	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
(300 mg)/3 ml Injektionslösung				
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5216/001	50479.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
KLEXANE	AT/H/0751/001	13118	SANOFI-AVENTIS DENMARK A/S	DK
KLEXANE	AT/H/0751/003	13118	SANOFI-AVENTIS DENMARK A/S	DK
KLEXANE	AT/H/0751/002	10432	SANOFI OY	FI
KLEXANE	AT/H/0751/004	10432	SANOFI OY	FI
KLEXANE	AT/H/0751/003	10432	SANOFI OY	FI
KLEXANE	AT/H/0751/001	10432	SANOFI OY	FI
KLEXANE	AT/H/0751/005	10432	SANOFI OY	FI
Klexane 10.000 a.e. (100 mg)/1 ml stungulyf, lausn	AT/H/0751/005	870238	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 10000 IU (100 MG)/1 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/005	17-11774	SANOFI-AVENTIS NORGE AS	NO
Klexane 100000 IE	AT/H/0752/003	56487	SANOFI 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
(1000 mg)/10 ml injektionsvätska, lösning				
KLEXANE 12000 IU (120 MG)/0,8 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0754/001	99-3240	SANOFI-AVENTIS NORGE AS	NO
KLEXANE 15000 IU (150 MG)/1 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0754/002	17-11775	SANOFI-AVENTIS NORGE AS	NO
Klexane 2.000 a.e. (20 mg)/0,2 ml stungulyf, lausn	AT/H/0751/001	870238	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 2000 IU (20 MG)/0,2 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/001	7507	SANOFI-AVENTIS NORGE AS	NO
KLEXANE 2000 IU (20 MG)/0,2 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/001	7507	SANOFI-AVENTIS NORGE AS	NO
Klexane 30.000 a.e. (300 mg)/3 ml stungulyf, lausn	AT/H/0752/001	970454	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 30000 IU (300 MG)/3 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0752/001	05-3487	SANOFI-AVENTIS NORGE AS	NO
Klexane 4.000 a.e. (40 mg)/0,4 ml stungulyf, lausn	AT/H/0751/002	870238	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 4000 IU (40 MG)/0,4 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/002	17-11771	SANOFI-AVENTIS NORGE 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
KLEXANE 4000 IU (40 MG)/0,4 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/002	17-11771	SANOFI-AVENTIS NORGE AS	NO
Klexane 6.000 a.e. (60 mg)/0,6 ml stungulyf, lausn	AT/H/0751/003	870238	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 6000 IU (60 MG)/0,6 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/003	17-11772	SANOFI-AVENTIS NORGE AS	NO
Klexane 8.000 a.e. (80 mg)/0,8 ml stungulyf, lausn	AT/H/0751/004	870238	SANOFI-AVENTIS NORGE AS	IS
KLEXANE 8000 IU (80 MG)/0,8 ML INJEKSJONSVAESKE, OPPLOSNING	AT/H/0751/004	17-11773	SANOFI-AVENTIS NORGE AS	NO
KLEXANE cum conservans 100 000 IU (1 000 mg)/10 ml injektioneste, liuos	AT/H/0752/003	13568	SANOFI OY	FI
KLEXANE cum conservans 30 000 IU (300 mg)/3 ml injektioneste, liuos	AT/H/0752/001	13568	SANOFI OY	FI
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/001	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/004	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/001	13118	SANOFI-AVENTIS DENMARK A/S	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
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/002	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/002	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/005	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/002	13118	SANOFI-AVENTIS DENMARK A/S	DK
Klexane,injektionsvæske, opløsning i hætteglas	AT/H/0752/001	19402	SANOFI-AVENTIS DENMARK A/S	DK
LOVENOX 10 000 UI (100 mg)/1 ml solução injetável	AT/H/0751/005	2842185	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 10 000 UI (100 mg)/1 ml solução injetável	AT/H/0751/005	2529782	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 10 000 UI (100 mg)/1 ml solução injetável	AT/H/0751/005	2842284	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	34009 300 142 7 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	34009 364 688 9 4	SANOFI-AVENTIS FRANCE	FR
LOVENOX 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	34009 336 062 1 3	SANOFI-AVENTIS FRANCE	FR
LOVENOX 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	34009 364 689 5 5	SANOFI-AVENTIS FRANCE	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
LOVENOX 10 000 UI (100 mg)/1 ml solution injectable	AT/H/0751/005	34009 336 061 5 2	SANOFI-AVENTIS FRANCE	FR
LOVENOX 10 x 4.000 IE (10 x 40 mg) Injektionslösung in einem Fertigpen	AT/H/0756/001	1-21761	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 10.000 IE (100 mg)/1 ml Injektionslösung in einer Ampulle	AT/H/0753/001	1-18753	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 10.000 IE (100 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0751/005	1-21540	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 10.000 IE (100 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0751/005	1-21540	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 10.000 IE (100 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0751/005	1-21540	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 10.000 IE (100 mg)/10 ml Injektionslösung in einer Durchstichflasche	AT/H/0755/001	1-21700	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 100.000 IE (1000 mg)/10ml Injektionslösung in einer Durchstichflasche	AT/H/0752/003	237480	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 12 000 UI (120 mg)/0,8 ml solução injetável	AT/H/0754/001	3136488	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 12 000 UI	AT/H/0754/001	3136686	SANOFI - PRODUTOS	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
(120 mg)/0,8 ml solução injetável			FARMACEUTICOS LDA	
LOVENOX 12 000 UI (120 mg)/0,8 ml solução injetável	AT/H/0754/001	3136587	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 12.000 IE (120 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0754/001	1-23836	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 12.000 IE (120 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0754/001	1-23836	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 12.000 IE (120 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0754/001	1-23836	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 15 000 UI (150 mg)/1 ml solução injetável	AT/H/0754/002	3136280	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 15 000 UI (150 mg)/1 ml solução injetável	AT/H/0754/002	3136389	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 15 000 UI (150 mg)/1 ml solução injetável	AT/H/0754/002	3136181	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 15.000 IE (150 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0754/002	1-23837	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 15.000 IE (150 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0754/002	1-23837	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 15.000 IE	AT/H/0754/002	1-23837	SANOFI-AVENTIS 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
(150 mg)/1 ml Injektionslösung in einer Fertigspritze			OSTERREICH	
LOVENOX 2 000 UI (20 mg)/0,2 ml solução injetável	AT/H/0751/001	2308682	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 2 000 UI (20 mg)/0,2 ml solução injetável	AT/H/0751/001	2263283	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 2 000 UI (20 mg)/0,2 ml solução injetável	AT/H/0751/001	2046688	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 556 142 4 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 364 684 3 6	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 334 351 6 5	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 329 487 0 3	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 364 683 7 5	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2 000 UI (20 mg)/0,2 ml solution injectable	AT/H/0751/001	34009 300 142 2 6	SANOFI-AVENTIS FRANCE	FR
LOVENOX 2.000 IE (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	1-18663	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 2.000 IE (20	AT/H/0751/001	1-18663	SANOFI-AVENTIS 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
mg)/0,2 ml Injektionslösung in einer Fertigspritze			OSTERREICH	
LOVENOX 2.000 IE (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	1-18663	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 2.000 IE (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	1-18663	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 2.000 IE (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	1-18663	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 30 000 UI (300 mg)/3 ml solução injetável	AT/H/0752/001	5349956	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 30 000 UI (300 mg)/3 ml solution injectable	AT/H/0752/001	34009 561 070 8 7	SANOFI-AVENTIS FRANCE	FR
LOVENOX 30.000 IE (300 mg)/3 ml Injektionslösung in einer Durchstichflasche	AT/H/0752/001	1-23838	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 4 000 UI (40 mg)/0,4 ml solução injetável	AT/H/0751/002	2046787	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 4 000 UI (40 mg)/0,4 ml solução injetável	AT/H/0751/002	2308781	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 4 000 UI (40 mg)/0,4 ml solução injetável	AT/H/0751/002	2263382	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 4 000 UI (40	AT/H/0751/002	34009 364 695 5 6	SANOFI-AVENTIS FRANCE	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
mg)/0,4 ml solution injectable				
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 348 564 7 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 329 486 4 2	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 556 141 8 0	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 364 686 6 5	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 364 687 2 6	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 334 352 2 6	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4 000 UI (40 mg)/0,4 ml solution injectable	AT/H/0751/002	34009 300 142 3 3	SANOFI-AVENTIS FRANCE	FR
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	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
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 6 000 UI (60 mg)/0,6 ml solução injetável	AT/H/0751/003	2841781	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 6 000 UI (60 mg)/0,6 ml solução injetável	AT/H/0751/003	2841880	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 6 000 UI (60 mg)/0,6 ml solução injetável	AT/H/0751/003	2529584	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	34009 364 692 6 6	SANOFI-AVENTIS FRANCE	FR
LOVENOX 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	34009 336 058 4 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	34009 336 057 8 0	SANOFI-AVENTIS FRANCE	FR
LOVENOX 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	34009 364 690 3 7	SANOFI-AVENTIS FRANCE	FR
LOVENOX 6 000 UI (60 mg)/0,6 ml solution injectable	AT/H/0751/003	34009 300 142 4 0	SANOFI-AVENTIS FRANCE	FR
LOVENOX 6.000 IE (60 mg)/0,6 ml Injektionslösung in einer	AT/H/0751/003	1-21538	SANOFI-AVENTIS GMBH OSTERREICH	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
Fertigspritze				
LOVENOX 6.000 IE (60 mg)/0,6 ml Injektionslösung in einer Fertigspritze	AT/H/0751/003	1-21538	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 6.000 IE (60 mg)/0,6 ml Injektionslösung in einer Fertigspritze	AT/H/0751/003	1-21538	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 8 000 UI (80 mg)/0,8 ml solução injetável	AT/H/0751/004	2841989	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 8 000 UI (80 mg)/0,8 ml solução injetável	AT/H/0751/004	2842086	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 8 000 UI (80 mg)/0,8 ml solução injetável	AT/H/0751/004	2529683	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	34009 336 060 9 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	34009 300 142 6 4	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	34009 364 693 2 7	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	34009 364 694 9 5	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8 000 UI (80 mg)/0,8 ml solution injectable	AT/H/0751/004	34009 336 059 0 2	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8.000 IE (80 mg)/0,8 ml	AT/H/0751/004	1-21539	SANOFI-AVENTIS GMBH OSTERREICH	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
Injektionslösung in einer Fertigspritze				
LOVENOX 8.000 IE (80 mg)/0,8 ml Injektionslösung in einer Fertigspritze	AT/H/0751/004	1-21539	SANOFI-AVENTIS GMBH OSTERREICH	AT
LOVENOX 8.000 IE (80 mg)/0,8 ml Injektionslösung in einer Fertigspritze	AT/H/0751/004	1-21539	SANOFI-AVENTIS GMBH OSTERREICH	AT
Qualiop Klinik 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5215/001	16071.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Qualiop Klinik 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5215/001	16071.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Qualiop Klinik 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5215/001	16071.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Qualiop Klinik 30.000 I. E. (300 mg)/3 ml Injektionslösung	DE/H/5215/001	16071.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
КЛЕКСАН 4 000 IU (40 mg)/0,4 ml инжекционен разтвор	AT/H/0751/002	9700003	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 4 000 IU (40 mg)/0,4 ml инжекционен разтвор	AT/H/0751/002	9700003	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 4 000 IU (40 mg)/0,4 ml инжекционен разтвор	AT/H/0751/002	9700003	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 6 000 IU (60 mg)/0,6 ml инжекционен разтвор	AT/H/0751/003	20020197	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 6 000 IU (60	AT/H/0751/003	20020197	SANOFI BULGARIA EOOD	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
mg)/0,6 ml инжекционен разтвор				
КЛЕКСАН 6 000 IU (60 mg)/0,6 ml инжекционен разтвор	AT/H/0751/003	20020197	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 8 000 IU (80 mg)/0,8 ml инжекционен разтвор	AT/H/0751/004	20020196	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 8 000 IU (80 mg)/0,8 ml инжекционен разтвор	AT/H/0751/004	20020196	SANOFI BULGARIA EOOD	BG
КЛЕКСАН 8 000 IU (80 mg)/0,8 ml инжекционен разтвор	AT/H/0751/004	20020196	SANOFI BULGARIA EOOD	BG