EMA/601977/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: enoxaparin (except for biosimilars)

Procedure no.: PSUSA/00010560/201904

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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
LOVENOX 10 x 4.000 IE (10 x 40 mg) Injektionslösung im Fertigpen	AT/H/0756/001	1-21761	SANOFI-AVENTIS GMBH OSTERREICH	AT
Clexane 2.000 I. E. (20 mg)/0,2 ml Praxis Injektionslösung in einer Fertigspritze	DE/H/5213/001	32766.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Praxis Injektionslösung in einer Fertigspritze	DE/H/5213/002	32766.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
LOVENOX 10.000 IE (100 mg)/10 ml Injektionslösung in einer Durchstichflasche	AT/H/0755/001	1-21700	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 100.000 I. E. (1000 mg)/10 ml Injektionslösung	DE/H/5215/003	99842.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Qualiop Klinik 50.000 I. E. (500 mg)/5 ml Injektionslösung	DE/H/5215/002	99841.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 2.000 I. E. (20 mg)/0,2 ml Klinik Injektionslösung in einer Fertigspritze	DE/H/5212/001	16071.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane 4.000 I. E. (40 mg)/0,4 ml Klinik Injektionslösung in einer Fertigspritze	DE/H/5212/002	16071.01.00	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
CLEXANE	AT/H/0751/001	026966034 (PRE-FILLED)	SANOFI S.P.A	IT
CLEXANE T	AT/H/0751/003	029111046	SANOFI S.P.A	IT
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/001	RVG 121186	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE, 8000 RÜ (80 mg)/0,8 ml süstelahus süstlis	AT/H/0751/004	940617	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane 8000 UI (80 mg)/0,8 ml soluţie injectabilă	AT/H/0751/004	4928/2012/01	SANOFI ROMANIA SRL	RO
Klexane 6000 IE (60 mg)/0,6 ml injektionsvätska, lösning, förfylld spruta	AT/H/0751/003	56483	SANOFI AB	SE
CLEXANE 2.000 IU (20 mg)/0,2 mL ενέσιμο διάλυμα	AT/H/0751/001	19159	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE 6.000 IU (60 mg)/0,6 mL ενέσιμο διάλυμα	AT/H/0751/003	19744	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE 4.000 IU (40 mg)/0,4 mL ενέσιμο διάλυμα	AT/H/0751/002	19160	SANOFI-AVENTIS CYPRUS LTD	CY

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 4.000 IU (40 mg)/0,4 mL ενέσιμο διάλυμα	AT/H/0751/002	28078/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE 6.000 IU (60 mg)/0,6 mL ενέσιμο διάλυμα	AT/H/0751/003	9059/92/26-10-93	SANOFI-AVENTIS AEBE	GR
CLEXANE 10000 IU (100 mg)/1 ml injekčný roztok	AT/H/0751/005	16/0175/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 8.000 IU (80 mg)/0,8 mL ενέσιμο διάλυμα	AT/H/0751/004	9058/92/26-10-93	SANOFI-AVENTIS AEBE	GR
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 2000 NE (20 mg)/0,2 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/001	OGYI-T-4097/02	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 4000 NE (40 mg)/0,4 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/002	OGYI-T-4097/04	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

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 2.000 IU (20 mg)/0,2 mL ενέσιμο διάλυμα	AT/H/0751/001	28077/19-10-88	SANOFI-AVENTIS AEBE	GR
CLEXANE	AT/H/0751/001	9700003	SANOFI BULGARIA EOOD	BG
CLEXANE 8.000 IU (80 mg)/0,8 mL ενέσιμο διάλυμα	AT/H/0751/004	19161	SANOFI-AVENTIS CYPRUS LTD	CY
CLEXANE T 10.000 UI (100 mg)/1 ml soluzione iniettabile	AT/H/0751/005	029111135	SANOFI S.P.A	IT
CLEXANE	AT/H/0751/002	026966061 (SAFETY SYRINGE)	SANOFI S.P.A	IT
CLEXANE T	AT/H/0751/004	02911123 (SAFETY SYRINGE)	SANOFI S.P.A	IT
LOVENOX 2.000 IE (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	1-18663	SANOFI-AVENTIS GMBH OSTERREICH	AT
CLEXANE 2.000 UI (20 mg)/0,2 ml solución inyectable en jeringa precargada	AT/H/0751/001	58502	SANOFI-AVENTIS, S.A.	ES
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/001	BE144365	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
CLEXANE, 2000 RÜ (20 mg)/0,2 ml süstelahus süstlis	AT/H/0751/001	940317	SANOFI-AVENTIS ESTONIA OÜ	EE
Klexane 2000 IU (20 mg)/0,2 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/001	7507	SANOFI-AVENTIS NORGE AS	NO
Klexane, injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/001	13118	SANOFI-AVENTIS DENMARK A/S	DK
Clexane 2.000 I. E. (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0751/001	15854.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Klexane, injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/001	13118	SANOFI-AVENTIS DENMARK A/S	DK
Clexane 2000 IU (20 mg)/0,2 ml otopina za injekciju u napunjenoj štrcaljki	AT/H/0751/001	HR-H-099632462-10	SANOFI-AVENTIS CROATIA D.O.O.	HR
Clexane 2,000 IU (20 mg) /0.2 mL Solution for Injection in pre-filled syringes	AT/H/0751/001	PA 540/97/4	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
CLEXANE 2000 SV (20 mg)/0,2 ml šķīdums injekcijām pilnšļircē	AT/H/0751/001	98-0717	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 2000 TV (20 mg)/0,2 ml injekcinis tirpalas užpildytame švirkšte	AT/H/0751/001	LT/1/98/1560/022	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
Klexane 2000 IE (20 mg)/0,2 ml injektionsvätska, lösning, förfylld spruta	AT/H/0751/001	56481	SANOFI AB	SE
CLEXANE 2.000 i.e. (20 mg)/0,2 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0751/001	H/99/02469/002	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 2 000 IU (20 mg)/0,2 ml injekční roztok v předplněné injekční stříkačce	AT/H/0751/001	16/250/93-A/C	SANOFI-AVENTIS SRO	CZ
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie	AT/H/0751/002	BE144347	SANOFI BELGIUM	BE
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable en seringue préremplie	AT/H/0751/003	BE204565	SANOFI BELGIUM	BE
CLEXANE 8 000 UI (80 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0751/004	BE161953	SANOFI BELGIUM	BE
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable en seringue préremplie	AT/H/0751/005	BE161944	SANOFI BELGIUM	BE
Clexane 2,000 IU (20 mg) /0.2 mL Solution for Injection in pre-filled syringes	AT/H/0751/001	MA 082/00702	SANOFI MALTA LTD	MT
Klexane 2.000 a.e. (20 mg)/0,2 ml stungulyf, lausn í áfylltri sprautu	AT/H/0751/001	870238	SANOFI-AVENTIS NORGE AS	IS

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 2 000 UI (20 mg)/0,2 ml solution injectable en seringue préremplie	AT/H/0751/001	0861674	SANOFI BELGIUM	LU
CLEXANE 2.000 IE (20 mg)/0,2 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/001	RVG 121186	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 2000 IU (20 mg)/0,2 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0751/001	16/0108/96-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane® Syringes 2,000 IU (20 mg)/0.2 ml solution for injection in pre-filled syringes	AT/H/0751/001	PL 04425/0187	AVENTIS PHARMA LTD	UK
LOVENOX 4.000 IE (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	1-18662	SANOFI-AVENTIS GMBH OSTERREICH	AT
КЛЕКСАН 4 000 IU (40 mg)/0,4 ml инжекционен разтвор в предварително напълнени спринцовки	AT/H/0751/002	9700003	SANOFI BULGARIA EOOD	BG
Clexane 4000 IU (40 mg)/0,4 ml otopina za injekciju u napunjenoj štrcaljki	AT/H/0751/002	HR-H-990481301-15	SANOFI-AVENTIS CROATIA D.O.O.	HR
CLEXANE 4 000 IU (40 mg)/0,4 ml injekční roztok v předplněné injekční stříkačce	AT/H/0751/002	16/250/93-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, 4000 RÜ (40 mg)/0,4 ml süstelahus süstlis	AT/H/0751/002	940417	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane 4.000 I. E. (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	AT/H/0751/002	15854.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Klexane 4.000 a.e. (40 mg)/0,4 ml stungulyf, lausn í áfylltri sprautu	AT/H/0751/002	870238	SANOFI-AVENTIS NORGE AS	IS
CLEXANE 4000 SV (40 mg)/0,4 ml šķīdums injekcijām pilnšļircē	AT/H/0751/002	98-0718	SANOFI-AVENTIS LATVIA SIA	LV
CLEXANE 4000 TV (40 mg)/0,4 ml injekcinis tirpalas užpildytame švirkšte	AT/H/0751/002	LT/1/98/1560/046	UAB SANOFI-AVENTIS LIETUVA	LT
Clexane 4,000 IU (40 mg) /0.4 mL Solution for Injection in pre-filled syringes	AT/H/0751/002	PA 540/97/5	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
CLEXANE 4 000 UI (40 mg)/0,4 ml solution injectable en seringue préremplie	AT/H/0751/002	0861772	SANOFI BELGIUM	LU
Clexane 4,000 IU (40 mg) /0.4 mL Solution for Injection in pre-filled syringes	AT/H/0751/002	MA 082/00703	SANOFI MALTA LTD	MT
CLEXANE 4.000 IE (40 mg)/0,4 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/002	RVG 121187	SANOFI-AVENTIS NETHERLANDS B.V.	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
Klexane 4000 IU (40 mg)/0,4 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/002	17-11771	SANOFI-AVENTIS NORGE AS	NO
Clexane, 4000 j.m. (40 mg) /0,4 ml, roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0751/002	R/0484	SANOFI-AVENTIS FRANCE	PL
Clexane 4000 UI (40 mg)/0,4 ml soluţie injectabilă în seringă preumplută	AT/H/0751/002	4927/2012/31	SANOFI ROMANIA SRL	RO
Clexane 4000 UI (40 mg)/0,4 ml soluţie injectabilă în seringă preumplută	AT/H/0751/002	4927/2012/30	SANOFI ROMANIA SRL	RO
CLEXANE 4000 IU (40 mg)/0,4 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0751/002	16/0172/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 4.000 i.e. (40 mg)/0,4 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0751/002	H/99/02469/053	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 4.000 UI (40 mg)/0,4 ml solución inyectable en jeringa precargada	AT/H/0751/002	58503	SANOFI-AVENTIS, S.A.	ES
Klexane 4000 IE (40 mg)/0,4 ml injektionsvätska, lösning, förfylld spruta	AT/H/0751/002	56482	SANOFI AB	SE
Clexane® Syringes 4,000 IU (40 mg)/0.4 ml solution for injection in pre-filled syringes	AT/H/0751/002	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
LOVENOX 6.000 IE (60 mg)/0,6 ml Injektionslösung in einer Fertigspritze	AT/H/0751/003	1-21538	SANOFI-AVENTIS GMBH OSTERREICH	AT
КЛЕКСАН 6 000 IU (60 mg)/0,6 ml инжекционен разтвор в предварително напълнени спринцовки	AT/H/0751/003	20020197	SANOFI BULGARIA EOOD	BG
Clexane 6000 IU (60 mg)/0,6 ml otopina za injekciju u napunjenoj štrcaljki	AT/H/0751/003	HR-H-041381787-24	SANOFI-AVENTIS CROATIA D.O.O.	HR
Klexane,injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/003	13118	SANOFI-AVENTIS DENMARK A/S	DK
Clexane 6.000 I. E. (60 mg)/0,6 ml Injektionslösung in einer Fertigspritze	AT/H/0751/003	40759.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 6 000 IU (60 mg)/0,6 ml injekční roztok v předplněné injekční stříkačce	AT/H/0751/003	16/250/93-C/C	SANOFI-AVENTIS SRO	CZ
Klexane 6.000 a.e. (60 mg)/0,6 ml stungulyf, lausn í áfylltri sprautu	AT/H/0751/003	870238	SANOFI-AVENTIS NORGE AS	IS
CLEXANE, 6000 RÜ (60 mg)/0,6 ml süstelahus süstlis	AT/H/0751/003	940517	SANOFI-AVENTIS ESTONIA OÜ	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
CLEXANE 6000 SV (60 mg)/0,6 ml šķīdums injekcijām pilnšļircē	AT/H/0751/003	98-0719	SANOFI-AVENTIS LATVIA SIA	LV
Clexane 6,000 IU (60 mg) /0.6 mL Solution for Injection in pre-filled syringes	AT/H/0751/003	PA 540/97/6	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
CLEXANE 6000 TV (60 mg)/0,6 ml injekcinis tirpalas užpildytame švirkšte	AT/H/0751/003	LT/1/98/1560/060	UAB SANOFI-AVENTIS LIETUVA	LT
Clexane 6.000 IE (60 mg)/0,6 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/003	RVG 121188	SANOFI-AVENTIS NETHERLANDS B.V.	NL
CLEXANE 6 000 UI (60 mg)/0,6 ml solution injectable en seringue préremplie	AT/H/0751/003	0861819	SANOFI BELGIUM	LU
Klexane 6000 IU (60 mg)/0,6 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/003	17-11772	SANOFI-AVENTIS NORGE AS	NO
Klexane 6000 IU (60 mg)/0,6 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/003	17-11772	SANOFI-AVENTIS NORGE AS	NO
Clexane, 6000 j.m. (60 mg) /0,6 ml, roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0751/003	7748	SANOFI-AVENTIS FRANCE	PL
Clexane 6,000 IU (60 mg) /0.6 mL Solution for Injection in pre-filled syringes	AT/H/0751/003	MA 082/00704	SANOFI MALTA LTD	MT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 6000 IU (60 mg)/0,6 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0751/003	16/0173/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 6.000 UI (60 mg)/0,6 ml solución inyectable en jeringa precargada	AT/H/0751/003	62470	SANOFI-AVENTIS, S.A.	ES
CLEXANE 6.000 i.e. (60 mg)/0,6 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0751/003	H/99/02469/079	SANOFI-AVENTIS D.O.O.	SI
Clexane® Syringes 6,000 IU (60 mg)/0.6 ml solution for injection in pre-filled syringes	AT/H/0751/003	PL 04425/0187	AVENTIS PHARMA LTD	UK
LOVENOX 8.000 IE (80 mg)/0,8 ml Injektionslösung in einer Fertigspritze	AT/H/0751/004	1-21539	SANOFI-AVENTIS GMBH OSTERREICH	AT
CLEXANE 8 000 IU (80 mg)/0,8 ml injekční roztok v předplněné injekční stříkačce	AT/H/0751/004	16/250/93-D/C	SANOFI-AVENTIS SRO	CZ
Klexane, injektionsvæske, opløsning i fyldt injektionssprøjte	AT/H/0751/004	13118	SANOFI-AVENTIS DENMARK A/S	DK
Clexane 8.000 I. E. (80 mg)/0,8 ml Injektionslösung in einer Fertigspritze	AT/H/0751/004	40759.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
КЛЕКСАН 8 000 IU (80 mg)/0,8 ml инжекционен разтвор в предварително напълнени	AT/H/0751/004	20020196	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
спринцовки				
Clexane 8000 IU (80 mg)/0,8 ml otopina za injekciju u napunjenoj štrcaljki	AT/H/0751/004	HR-H-173049079-17	SANOFI-AVENTIS CROATIA D.O.O.	HR
Klexane 8.000 a.e. (80 mg)/0,8 ml stungulyf, lausn í áfylltri sprautu	AT/H/0751/004	870238	SANOFI-AVENTIS NORGE AS	IS
Clexane 8,000 IU (80 mg) /0.8 mL Solution for Injection in pre-filled syringes	AT/H/0751/004	PA 540/97/7	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Clexane® Syringes 8,000 IU (80 mg)/0.8 ml solution for injection in pre-filled syringes	AT/H/0751/004	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE 8000 SV (80 mg)/0,8 ml šķīdums injekcijām pilnšļircē	AT/H/0751/004	98-0720	SANOFI-AVENTIS LATVIA SIA	LV
Klexane 8000 IE (80 mg)/0,8 ml injektionsvätska, lösning, förfylld spruta	AT/H/0751/004	56484	SANOFI AB	SE
CLEXANE 8000 TV (80 mg)/0,8 ml injekcinis tirpalas užpildytame švirkšte	AT/H/0751/004	LT/1/98/1560/087	UAB SANOFI-AVENTIS LIETUVA	LT
CLEXANE 8.000 UI (80 mg)/0,8 ml solución inyectable en jeringa precargada	AT/H/0751/004	62471	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 8 000 UI (80 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0751/004	0656561	SANOFI BELGIUM	LU
CLEXANE 8.000 i.e. (80 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0751/004	H/99/02469/110	SANOFI-AVENTIS D.O.O.	SI
Clexane 8,000 IU (80 mg) /0.8 mL Solution for Injection in pre-filled syringes	AT/H/0751/004	MA 082/00705	SANOFI MALTA LTD	MT
CLEXANE 8.000 IE (80 mg)/0,8 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/004	RVG 121189	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Klexane 8000 IU (80 mg)/0,8 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/004	17-11773	SANOFI-AVENTIS NORGE AS	NO
Clexane, 8000 j.m. (80 mg) /0,8 ml, roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0751/004	7750	SANOFI-AVENTIS FRANCE	PL
CLEXANE 8000 IU (80 mg)/0,8 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0751/004	16/0174/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
Clexane 10,000 IU (100mg)/1ml Solution for Injection in pre-filled syringes	AT/H/0751/005	MA 082/00701	SANOFI MALTA LTD	MT
Clexane 10.000 IE (100 mg)/1 ml oplossing voor injectie in een voorgevulde spuit	AT/H/0751/005	RVG 121190	SANOFI-AVENTIS NETHERLANDS B.V.	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
Klexane 10 000 IU (100 mg)/1 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0751/005	17-11774	SANOFI-AVENTIS NORGE AS	NO
Clexane, 10 000 j.m. (100 mg) /1 ml, roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0751/005	7749	SANOFI-AVENTIS FRANCE	PL
CLEXANE 10.000 UI (100 mg)/1 ml solución inyectable en jeringa precargada	AT/H/0751/005	62472	SANOFI-AVENTIS, S.A.	ES
CLEXANE 10.000 i.e. (100 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0751/005	H/99/02469/156	SANOFI-AVENTIS D.O.O.	SI
Klexane 10000 IE (100 mg)/1 ml injektionsvätska, lösning, förfylld spruta	AT/H/0751/005	56485	SANOFI AB	SE
Clexane® Syringes 10,000 IU (100 mg)/1 ml solution for injection in pre-filled syringes	AT/H/0751/005	PL 04425/0187	AVENTIS PHARMA LTD	UK
CLEXANE 10 000 UI (100 mg)/1 ml solution injectable en seringue préremplie	AT/H/0751/005	0862011	SANOFI BELGIUM	LU
Clexane 10,000 IU (100mg)/1ml Solution for Injection in pre-filled syringes	AT/H/0751/005	PA 540/97/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
LOVENOX 10.000 IE (100 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0751/005	1-21540	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
Clexane 10 000 IU (100 mg)/1 ml otopina za injekciju u napunjenoj štrcaljki	AT/H/0751/005	HR-H-165030097-13	SANOFI-AVENTIS CROATIA D.O.O.	HR
KLEXANE	AT/H/0751/005	13118	SANOFI A/S	DK
CLEXANE, 10 000 RÜ (100 mg)/1 ml süstelahus süstlis	AT/H/0751/005	240398	SANOFI-AVENTIS ESTONIA OÜ	EE
Clexane 10.000 I. E. (100 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0751/005	40759.02.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
CLEXANE 10 000 IU (100 mg)/1 ml injekcní roztok v predplnené injekcní stríkacce	AT/H/0751/005	16/250/93-E/C	SANOFI-AVENTIS SRO	CZ
Klexane 10.000 a.e. (100 mg)/1 ml stungulyf, lausn í áfylltri sprautu	AT/H/0751/005	870238	SANOFI-AVENTIS NORGE AS	IS
Clexane 12.000 IE (120 mg)/0,8 ml oplossing voor injectie	AT/H/0754/001	RVG 121191	SANOFI-AVENTIS NETHERLANDS B.V.	NL
KLEXANE 10 000 IU (100 mg)/1 ml injektioneste, liuos, esitäytetty ruisku	AT/H/0751/001	10432	SANOFI OY	FI
LOVENOX 2 000 UI (20 mg)/0,2 ml, solution injectable en seringue préremplie	AT/H/0751/001	NL14582	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
KLEXANE 4 000 IU (40 mg)/0,4 ml injektioneste, liuos, esitäytetty ruisku	AT/H/0751/002	10432	SANOFI OY	FI
LOVENOX 4 000 UI (40 mg)/0,4 ml, solution injectable en seringue préremplie	AT/H/0751/002	NL14584	SANOFI-AVENTIS FRANCE	FR
Clexane, 2000 j.m. (20 mg) /0,2 ml, roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0751/001	R/0483	SANOFI-AVENTIS FRANCE	PL
LOVENOX 2 000 UI (20 mg)/0,2 ml solução injetável em seringa pré-cheia	AT/H/0751/001	AT/H/0751/001	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 4 000 UI (40 mg)/0,4 ml solução injetável em seringa pré-cheia	AT/H/0751/002	AT/H/0751/002	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Clexane 2000 UI (20 mg)/0,2 ml soluţie injectabilă în seringă preumplută	AT/H/0751/001	4926/2012/19	SANOFI ROMANIA SRL	RO
KLEXANE 8 000 IU (80 mg)/0,8 ml injektioneste, liuos, esitäytetty ruisku	AT/H/0751/004	10432	SANOFI OY	FI
KLEXANE 6 000 IU (60 mg)/0,6 ml injektioneste, liuos, esitäytetty ruisku	AT/H/0751/003	10432	SANOFI OY	FI
LOVENOX 6 000 UI (60 mg)/0,6 ml, solution injectable en seringue préremplie	AT/H/0751/003	NL18577	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 8 000 UI (80 mg)/0,8 ml, solution injectable en seringue préremplie	AT/H/0751/004	34009 301 538 7 1	SANOFI-AVENTIS FRANCE	FR
LOVENOX 8 000 UI (80 mg)/0,8 ml solução injetável em seringa pré-cheia	AT/H/0751/004	AT/H/0751/004	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 6 000 UI (60 mg)/0,6 ml solução injetável em seringa prá-cheia	AT/H/0751/003	AT/H/0751/003	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Clexane 6000 UI (60 mg)/0,6 ml soluţie injectabilă în seringă preumplută	AT/H/0751/003	4928/2012/34	SANOFI ROMANIA SRL	RO
KLEXANE 10 000 IU (100 mg)/1 ml injektioneste, liuos, esitäytetty ruisku	AT/H/0751/005	10432	SANOFI OY	FI
LOVENOX 10 000 UI (100 mg)/1 ml, solution injectable en seringue préremplie	AT/H/0751/005	NL18579	SANOFI-AVENTIS FRANCE	FR
LOVENOX 10 000 UI (100 mg)/1 ml solução injetável em seringa pré-cheia	AT/H/0751/005	AT/H/0751/005	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Enoxaparin Sanofi 4.000 I. E. (40 mg)/0,4 ml Injektionslösung in einer Fertigspritze	DE/H/5211/002	16349.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 2.000 I. E. (20 mg)/0,2 ml Injektionslösung in einer Fertigspritze	DE/H/5211/001	16349.00.00	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 6.000 I. E. (60 mg)/0,6 ml Injektionslösung in einer Fertigspritze	DE/H/5211/003	78635.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 8.000 I. E. (80 mg)/0,8 ml Injektionslösung in einer Fertigspritze	DE/H/5211/004	78636.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi 10.000 I. E. (100 mg)/1 ml Injektionslösung in einer Fertigspritze	DE/H/5211/005	78637.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
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
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
Clexane® Multidose Vial 30,000 IU (300 mg)/3 ml solution for injection	AT/H/0752/001	PL 04425/0186	AVENTIS PHARMA LTD	UK
LOVENOX 30.000 IE (300 mg)/3 ml Injektionslösung in einer Durchstichflasche	AT/H/0752/001	1-23838	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
Klexane 30000 IE (300 mg)/3 ml injektionsvätska, lösning	AT/H/0752/001	56486	SANOFI AB	SE
Klexane 100000 IE (1000 mg)/10 ml injektionsvätska, lösning	AT/H/0752/003	56487	SANOFI AB	SE
Klexane Cum Conservans	AT/H/0752/001	13568	SANOFI OY	FI
Klexane Cum Conservans	AT/H/0752/003	13568	SANOFI OY	FI
KLEXANE	AT/H/0752/001	19402	SANOFI A/S	DK
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 30000 IU (300 mg)/3 ml injekčný roztok	AT/H/0752/001	16/0176/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
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 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	OGYI-T-4097/11	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
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
LOVENOX 100.000 IE (1000 mg)/10ml Injektionslösung in einer Durchstichflasche	AT/H/0752/003	237480	SANOFI-AVENTIS GMBH OSTERREICH	AT
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 T	AT/H/0752/001	029111073	SANOFI S.P.A	IT
Clexane multidose 50.000 I.E. (500 mg)/5 ml Injektionslösung	AT/H/0752/002	99843.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 100.000 I.E. (1000 mg)/10 ml Injektionslösung	AT/H/0752/003	99844.00.00	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
Clexane multidose 50.000 I.E. (500 mg)/5 ml Injektionslösung	AT/H/0752/002	99843.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 100.000 I.E. (1000 mg)/10 ml Injektionslösung	AT/H/0752/003	99844.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
LOVENOX 50.000 IE (500 mg)/5 ml Injektionslösung in einer Durchstichflasche	AT/H/0752/002	237481	SANOFI-AVENTIS GMBH OSTERREICH	AT
Clexane 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	AT/H/0752/001	SANOFI-AVENTIS ZRT	HU
Clexane 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	OGYI-T-4097/11	SANOFI-AVENTIS ZRT	HU
Clexane 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	AT/H/0752/001	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 30000 IU (300 mg)/3 ml injekčný roztok	AT/H/0752/001	16/0176/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
LOVENOX 30 000 UI (300 mg)/3 ml solution injectable	AT/H/0752/001	34009 550 551 5 0	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 30 000 UI (300 mg)/3 ml solution injectable	AT/H/0752/001	34009 550 551 6 7	SANOFI-AVENTIS FRANCE	FR
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 (120 mg)/0,8 ml solução injetável	AT/H/0754/001	3136686	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
LOVENOX 12 000 UI (120 mg)/0,8 ml solução injetável	AT/H/0754/001	3136587	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
CLEXANE 12.000 UI (120 mg)/0,8 ml solución inyectable en jeringa precargada	AT/H/0754/001	63002	SANOFI-AVENTIS, S.A.	ES
CLEXANE FORTE 12 000 IU (120 mg)/0,8 ml injekční roztok v předplněné injekční stříkačce	AT/H/0754/001	16/338/01-A/C	SANOFI-AVENTIS SLOVAKIA SRO	CZ
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	0316489	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 15 000 UI (150 mg)/1 ml solution injectable	AT/H/0754/002	0656608	SANOFI BELGIUM	LU
CLEXANE FORTE	AT/H/0754/001	8914	AVENTIS PHARMA LTD	PL
CLEXANE 15.000 IU (150 mg)/1 mL ενέσιμο διάλυμα	AT/H/0754/002	59659/12-11-2003	SANOFI-AVENTIS AEBE	GR
CLEXANE 12.000 IU (120 mg)/0,8 mL ενέσιμο διάλυμα	AT/H/0754/001	59658/12-11-2003	SANOFI-AVENTIS AEBE	GR
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 2000 NE (20 mg)/0,2 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/001	OGYI-T-4097/02	SANOFI-AVENTIS ZRT	HU
Clexane 10 000 NE (100 mg)/1 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/005	OGYI-T-4097/10	SANOFI-AVENTIS ZRT	HU
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 4000 NE (40 mg)/0,4 ml oldatos injekció előretöltött fecskendőben	AT/H/0751/002	OGYI-T-4097/03	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
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 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 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
Klexane 15000 IE (150 mg) / 1 ml injektionsvätska, lösning, förfylld spruta	AT/H/0754/002	56489	SANOFI AB	SE
CLEXANE 12.000 IE (120 mg)/0,8 ml oplossing voor injectie	AT/H/0754/001	BE230176	SANOFI BELGIUM	BE
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 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	BE230185	SANOFI BELGIUM	BE
LOVENOX 12.000 IE (120 mg)/0,2 ml Injektionslösung in einer Fertigspritze	AT/H/0754/001	1-23836	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
Clexane Forte 12 000 NE (120 mg)/0,8 ml oldatos injekció előretöltött fecskendőben	AT/H/0754/001	AT/H/0754/001	SANOFI-AVENTIS ZRT	HU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0316492	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0862122	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0862136	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0656592	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0862105	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0555099	SANOFI BELGIUM	LU
CLEXANE 12 000 UI (120 mg)/0,8 ml solution injectable en seringue préremplie	AT/H/0754/001	0862119	SANOFI BELGIUM	LU
LOVENOX 15.000 IE (150 mg)/1 ml Injektionslösung in einer Fertigspritze	AT/H/0754/002	1-23837	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
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 FORTE 15 000 IU (150 mg)/1 ml injekční roztok v předplněné injekční stříkačce	AT/H/0754/002	16/338/01-B/C	SANOFI-AVENTIS SRO	CZ
Klexane 12 000 IU (120 mg)/0,8 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0754/001	99-3240	SANOFI-AVENTIS NORGE AS	NO
Clexane Forte, 12 000 j.m. (120 mg) /0,8 ml roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0754/001	8914	SANOFI-AVENTIS FRANCE	PL
Clexane Forte 15,000 IU (150mg)/1ml Solution for Injection in pre-filled syringes	AT/H/0754/002	540/97/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
LOVENOX 12 000 UI (120 mg)/0,8 ml solução injetável	AT/H/0754/001	AT/H/0754/001	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
CLEXANE FORTE 12000 IU (120 mg)/0,8 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0754/001	16/0033/02-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/008	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/011	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/019	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/009	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/003	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/005	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/014	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski	AT/H/0754/001	H/99/02470/001	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
brizgi				
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/004	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/013	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/015	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/016	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/017	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/002	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/021	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/010	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/007	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/006	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/012	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/018	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski	AT/H/0754/001	H/99/02470/024	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
brizgi				
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/020	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/023	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 12.000 i.e. (120 mg)/0,8 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/001	H/99/02470/022	SANOFI-AVENTIS D.O.O.	SI
Clexane 15.000 IE (150 mg)/1 ml oplossing voor injectie	AT/H/0754/002	RVG 121192	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Klexane 15 000 IU (150 mg)/1 ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	AT/H/0754/002	17-11775	SANOFI-AVENTIS NORGE AS	NO
Clexane Forte 15 000 j.m. (150 mg) /1 ml roztwór do wstrzykiwań w ampułko-strzykawkach	AT/H/0754/002	8915	SANOFI-AVENTIS FRANCE	PL
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0862167	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 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0862184	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0862171	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0555104	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0656608	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0862198	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0862203	SANOFI BELGIUM	LU
CLEXANE 15 000 UI (150 mg)/1 ml solution injectable en seringue préremplie	AT/H/0754/002	0316489	SANOFI BELGIUM	LU
Klexane 12000 IE (120 mg) / 0,8 ml injektionsvätska, lösning, förfylld spruta	AT/H/0754/001	56488	SANOFI AB	SE
Clexane® Forte Syringes 12,000 IU (120 mg)/0.8 ml solution for injection in pre-filled syringes	AT/H/0754/001	PL 04425/0185	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
LOVENOX 15 000 UI (150 mg)/1 ml solução injetável em seringa pré-cheia	AT/H/0754/002	AT/H/0754/002	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
CLEXANE FORTE 15000 IU (150 mg)/1 ml injekčný roztok v naplnenej injekčnej striekačke	AT/H/0754/002	16/0171/17-S	SANOFI-AVENTIS SLOVAKIA SRO	SK
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/044	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/040	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/047	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/048	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/033	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/030	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/037	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/028	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/034	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/031	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/046	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/039	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/032	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/027	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/041	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/035	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/036	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/043	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/029	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/025	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/026	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/042	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/038	SANOFI-AVENTIS D.O.O.	SI
CLEXANE 15.000 UI (150 mg)/1 ml solución inyectable en jeringa precargada	AT/H/0754/002	63000	SANOFI-AVENTIS, S.A.	ES
CLEXANE 15.000 i.e. (150 mg)/1 ml raztopina za injiciranje v napolnjeni injekcijski brizgi	AT/H/0754/002	H/99/02470/045	SANOFI-AVENTIS D.O.O.	SI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Clexane® Forte Syringes 15,000 IU (150 mg)/1 ml solution for injection in pre-filled syringes	AT/H/0754/002	PL 04425/0185	AVENTIS PHARMA LTD	UK
Clexane 30 000 NE (300 mg)/3 ml oldatos injekció	AT/H/0752/001	OGYI-T-4097/11	SANOFI-AVENTIS ZRT	HU
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 100.000 I. E. (1000 mg)/10 ml Praxis Injektionslösung	DE/H/5218/003	99850.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Clexane multidose 50.000 I. E. (500 mg)/5 ml Praxis Injektionslösung	DE/H/5218/002	99849.00.00	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 50.000 I. E. (500 mg)/5 ml Injektionslösung	DE/H/5216/002	99845.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Enoxaparin Sanofi multidose 100.000 I. E. (1000 mg)/10 ml Injektionslösung	DE/H/5216/003	99846.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE