22 July 2021
EMA/466570/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: Glatiramer

Procedure no.: PSUSA/00001529/202011

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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
Acetato de Glatirâmero Mepha 20 mg/ml Solução injetável em seringa pré-cheia	DE/H/5449/001	NOT PROVIDED BY HA	MEPHA LDA	PT
Acetato de Glatirâmero Mepha 40 mg/ml Solução injetável em seringa pré-cheia	DE/H/5449/002	5759121	MEPHA LDA	PT
Acetato de glatirâmero Mylan 20 mg/1 ml Solução injetável em seringa pré-cheia	NL/H/3213/001	5697271	MYLAN, LDA	PT
Brabio 20 mg/ml solution for injection, pre-filled syringe	NL/H/3211/001	PA0577/211/001	MCDERMOTT LABORATORIES LTD	IE
Brabio 20 mg/mL Solution for Injection, Pre-filled Syringe.	NL/H/3211/001	PL 04569/1725	GENERICS [UK] LIMITED	XI
Brabio 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3211/001	RVG 115980	SYNTHON BV	NL
Brabio 20 mg/ml, Solution for Injection, Pre-filled Syringe	NL/H/3211/001	MA1148/00101	MYLAN S.A.S	MT
Brabio 40 mg/ml solution for injection in pre-filled syringe	NL/H/3777/001	PA0577/212/001	MCDERMOTT LABORATORIES LTD	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Brabio 40 mg/ml solution for injection, pre-filled syringe	NL/H/3777/001	PL 04569/1756	GENERICS [UK] LIMITED	XI
CLIFT 20 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3213/001	92742.00.00	MYLAN GERMANY GMBH	DE
CLIFT 20 mg/mL, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	NL/H/3211/001	94837 / 13-12-2016	MYLAN S.A.S	GR
CLIFT 20 mg/mL, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα.	NL/H/3211/001	022666	MYLAN S.A.S	CY
CLIFT 40 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3777/001	97606.00.00	MYLAN GERMANY GMBH	DE
Clift 40 mg/ml solução injetável, seringa pré-cheia	NL/H/3777/001	5752829	MYLAN, LDA	PT
Copaxobene 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/001	138636	TEVA GMBH	AT
Copaxobene 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/002	138637	TEVA GMBH	AT
Copaxone ® 20 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/002	PL 10921/0023	TEVA PHARMACEUTICALS LTD	XI

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
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/002	TEVA GMBH	LT
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/004	TEVA GMBH	LT
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/003	TEVA GMBH	LT
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/005	TEVA GMBH	LT
Copaxone 20 mg/ml injekční roztok v předplněné injekční stříkačce	DE/H/5283/002	59/481/06-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Copaxone 20 mg/ml Injekčný roztok, predplnená injekčná striekačka	DE/H/5283/002	59/0463/06-S	TEVA GMBH	SK
Copaxone 20 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte.	DE/H/5283/002	03-2205	TEVA GMBH	NO
Copaxone 20 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5283/002	18786	TEVA GMBH	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Copaxone 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	1-25380	TEVA GMBH	AT
Copaxone 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	BE260881	TEVA GMBH	BE
Copaxone 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	52069.00.01	TEVA GMBH	DE
Copaxone 20 mg/ml injektionsvätska, lösning i förfylld spruta	DE/H/5283/002	18786	TEVA GMBH	FI
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/03	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/02	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 20 mg/ml oplossing voor injectie in een voorgevulde spuit.	DE/H/5283/002	BE260881	TEVA GMBH	BE
Copaxone 20 mg/ml otopina za injekciju u napunjenoj štrcaljki	DE/H/5283/002	HR-H-157986508	PLIVA HRVATSKA D.O.O.	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
Copaxone 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	DE/H/5283/002	H/07/00419/001	TEVA GMBH	SI
COPAXONE 20 mg/ml šķīdums injekcijām pilnšļircē	DE/H/5283/002	06-0274	TEVA GMBH	LV
Copaxone 20 mg/ml Solução Injetável	DE/H/5283/002	4937686	TEVA GMBH	PT
Copaxone 20 mg/ml Solución inyectable en jeringa precargada	DE/H/5283/002	65983	TEVA GMBH	ES
Copaxone 20 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/002	10755/2018/03	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 20 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/002	10755/2018/04	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 20 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/002	10755/2018/02	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 20 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/002	10755/2018/01	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 20 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/002	PA22579/001/001	TEVA GMBH	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
Copaxone 20 mg/ml solution injectable, seringue préremplie	DE/H/5283/002	2008069847	TEVA GMBH	LU
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418045	TEVA GMBH	IT
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418033	TEVA GMBH	IT
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418058	TEVA GMBH	IT
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418021	TEVA GMBH	IT
Copaxone 20 mg/ml voorgevulde spuit, oplossing voor injectie	DE/H/5283/002	RVG 30086	TEVA GMBH	NL
Copaxone 20 mg/ml, injektionsvätska, lösning, förfylld spruta	DE/H/5283/002	20043	TEVA GMBH	SE
Copaxone 20 mg/ml, solution injectable en seringe préremplie	DE/H/5283/002	BE260881	TEVA GMBH	BE
Copaxone 20 mg/ml, solution injectable en seringue préremplie	DE/H/5283/002	NL43730	TEVA SANTÉ	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
Copaxone 20 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/002	20187	TEVA GMBH	CY
Copaxone 20 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/002	58960/16.09.2008	TEVA GMBH	GR
Copaxone 40 mg, stungulyf, lausn í áfylltri sprautu	DE/H/5283/004	IS/1/14/096/01	TEVA GMBH	IS
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/008	TEVA GMBH	LT
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/007	TEVA GMBH	LT
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/006	TEVA GMBH	LT
Copaxone 40 mg/ml injekční roztok v předplněné injekční stříkačce	DE/H/5283/004	59/043/15-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
Copaxone 40 mg/ml injekčný roztok naplnený v injekčnej striekačke	DE/H/5283/004	59/0104/15-S	TEVA GMBH	SK
Copaxone 40 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	DE/H/5283/004	13-9600	TEVA GMBH	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
Copaxone 40 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5283/004	31602	TEVA GMBH	FI
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	135998	TEVA GMBH	AT
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	BE467902	TEVA GMBH	BE
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	90395.00.00	TEVA GMBH	DE
Copaxone 40 mg/ml injektionsvätska, lösning i förfylld spruta	DE/H/5283/004	31602	TEVA GMBH	FI
Copaxone 40 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/004	OGYI-T-9993/04	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 40 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/004	OGYI-T-9993/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 40 mg/ml oplossing voor injectie in een voorgevulde spuit.	DE/H/5283/004	BE467902	TEVA GMBH	BE
Copaxone 40 mg/ml otopina za injekciju u napunjenoj štrcaljki	DE/H/5283/004	HR-H-304476486	PLIVA HRVATSKA D.O.O.	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
Copaxone 40 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	DE/H/5283/004	H/07/00419/005	TEVA GMBH	SI
COPAXONE 40 mg/ml šķīdums injekcijām pilnšļircē	DE/H/5283/004	15-0026	TEVA GMBH	LV
Copaxone 40 mg/ml solução injetável, seringa pré-cheia.	DE/H/5283/004	5633136	TEVA GMBH	PT
Copaxone 40 mg/ml solución inyectable en jeringa precargada	DE/H/5283/004	79515	TEVA GMBH	ES
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	12772/2019/01	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	12772/2019/02	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	12772/2019/03	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/004	PA22579/001/002	TEVA GMBH	IE
Copaxone 40 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/004	PL 10921/0026	TEVA PHARMACEUTICALS LTD	XI

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
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418084	TEVA GMBH	IT
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418072	TEVA GMBH	IT
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418060	TEVA GMBH	IT
Copaxone 40 mg/ml voorgevulde spuit, oplossing voor injectie	DE/H/5283/004	RVG 113849	TEVA GMBH	NL
Copaxone 40 mg/ml ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/004	18611/17-02-2020	TEVA GMBH	GR
Copaxone 40 mg/ml, injektionsvätska, lösning, förfylld spruta	DE/H/5283/004	49557	TEVA GMBH	SE
Copaxone 40 mg/ml, solution injectable en seringue préremplie	DE/H/5283/004	BE467902	TEVA GMBH	BE
COPAXONE 40 mg/ml, solution injectable en seringue préremplie	DE/H/5283/004	NL46033	TEVA SANTÉ	FR
Copaxone 40 mg/ml, solution injectable en seringue préremplie	DE/H/5283/004	2015040057	TEVA GMBH	LU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Copaxone 40 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/004	22376	TEVA GMBH	CY
Copaxone Pen 40 mg injekční roztok v předplněném peru	DE/H/5283/005	59/048/20-C	TEVA PHARMACEUTICALS CR, S.R.O.	CZ
COPAXONE PEN 40 mg injekčný roztok v naplnenom pere	DE/H/5283/005	59/0164/20-S	TEVA GMBH	SK
COPAXONE PEN 40 mg Injektionslösung im Fertigpen	DE/H/5283/005	2201422.00.00	TEVA GMBH	DE
COPAXONE PEN 40 mg, solution injectable en stylo prérempli	DE/H/5283/005	NL52191	TEVA SANTÉ	FR
Copaxone Pen 40mg Solution injectable en stylo prerempli	DE/H/5283/005	2019050068	TEVA GMBH	LU
Copaxone, 20 mg/ml süstelahus süstlis	DE/H/5283/002	527006	TEVA GMBH	EE
Copaxone, 20 mg/ml, roztwór do wstrzykiwań, ampułkostrzykawki	DE/H/5283/002	12562	TEVA GMBH	PL
Copaxone, 40 mg/ml süstelahus süstlis	DE/H/5283/004	863414	TEVA GMBH	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
Copaxone, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	DE/H/5283/004	22364	TEVA GMBH	PL
Copaxone, injektionsvæske, opløsning, fyldt injektionssprøjte 20 mg/ml	DE/H/5283/002	35719	TEVA GMBH	DK
Copaxone, injektionsvæske, opløsning, fyldt injektionssprøjte 40 mg/ml	DE/H/5283/004	52806	TEVA GMBH	DK
Copaxone, stungulyf, lausn í áfylltri sprautu	DE/H/5283/002	IS/1/04/002/01	TEVA GMBH	IS
Copemyl 20 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte.	NL/H/3211/001	14-10261	MYLAN AB	NO
Copemyl 20 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3213/001	043860016	MYLAN S.P.A.	IT
Copemyl 20 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3213/001	043860028	MYLAN S.P.A.	IT
Copemyl 20 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3213/001	043860030	MYLAN S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Copemyl 20 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3213/001	043860042	MYLAN S.P.A.	IT
Copemyl 40 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	NL/H/3777/001	16-11184	MYLAN AB	NO
Copemyl, injektionsvæske, opløsning i fyldt injektionssprøjte 20 mg/ml	NL/H/3211/001	54650	MYLAN AB	DK
Copemyl, injektionsvæske, opløsning i fyldt injektionssprøjte 40 mg/ml	NL/H/3777/001	57826	MYLAN AB	DK
Copemyltri 40 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3777/001	045673011	MYLAN S.P.A.	IT
Copemyltri 40 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3777/002	045673023	MYLAN S.P.A.	IT
Copemyltri 40 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3777/003	045673035	MYLAN S.P.A.	IT
Galtipex 20 mg/mL Solution for Injection, Pre-filled Syringe	MA1243/00101	MA1243/00101	ASPEN HEALTHCARE MALTA LIMITED	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
Galtipex 40 mg/mL solution for injection, pre-filled syringe	MA1243/00201	MA1243/00201	ASPEN HEALTHCARE MALTA LIMITED	MT
Glataxon 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/001	BE535004	ACTAVIS GROUP PTC EHF.	BE
Glataxon 20 mg/ml oplossing voor injectie in een voorgevulde spuit	DE/H/5449/001	BE535004	ACTAVIS GROUP PTC EHF.	BE
Glataxon 20 mg/ml solution injectable en seringue préremplie	DE/H/5449/001	BE535004	ACTAVIS GROUP PTC EHF.	BE
Glataxon 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/002	BE535013	ACTAVIS GROUP PTC EHF.	BE
Glataxon 40 mg/ml oplossing voor injectie in een voorgevulde spuit	DE/H/5449/002	BE535013	ACTAVIS GROUP PTC EHF.	BE
Glataxon 40 mg/ml solution injectable en seringue préremplie	DE/H/5449/002	BE535013	ACTAVIS GROUP PTC EHF.	BE
Glatimyl 20 mg/ml injektioneste, liuos, esitäytetty ruisku	NL/H/3211/001	32543	MYLAN AB	FI
Glatimyl 20 mg/ml injektionsvätska, lösning i förfylld spruta	NL/H/3211/001	32543	MYLAN AB	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glatimyl 20 mg/ml, injektionsvätska, lösning, förfylld spruta	NL/H/3211/001	51654	MYLAN AB	SE
Glatimyl 40 mg/ml injektioneste, liuos, esitäytetty ruisku	NL/H/3777/001	34220	MYLAN AB	FI
Glatimyl 40 mg/ml, injektionsvätska, lösning i förfylld spruta	NL/H/3777/001	54808	MYLAN AB	SE
Glatirameeracetaat Alvogen 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3778/001	RVG 119321	ALVOGEN IPCO S.AR.L	NL
Glatirameeracetaat Mylan 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3213/001	RVG 115993	MYLAN B.V.	NL
Glatirameeracetaat Mylan 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3777/001	RVG 119319	MYLAN B.V.	NL
Glatiramer acetate ratiopharm 40 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5449/002	35737	RATIOPHARM GMBH	FI
Glatiramer acetate ratiopharm 40 mg/ml injektionsvätska, lösning i förfylld spruta	DE/H/5449/002	35737	RATIOPHARM GMBH	FI

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glatiramer acetate Teva, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	DE/H/5449/001	25040	TEVA GMBH	PL
Glatiramer acetate Teva, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	DE/H/5449/002	25041	TEVA GMBH	PL
Glatiramer G.L. 40 mg/ml-Injektionslösung in einer Fertigspritze	not available	139058	G.L. PHARMA GMBH	AT
GLATIRAMER MYLAN 20 mg/ml, solution injectable en seringue préremplie	NL/H/3213/001	NL 44923	MYLAN MEDICAL SAS	FR
GLATIRAMER MYLAN 40 mg/ml, solution injectable en seringue préremplie	NL/H/3777/001	NL47156	MYLAN MEDICAL SAS	FR
Glatiramer/Mylan 40 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	NL/H/3777/001	023037	MYLAN S.A.S	CY
Glatiramer/Mylan 40 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	NL/H/3777/01	40332/16/02-02-2018	MYLAN S.A.S	GR
Glatirameracetat AbZ 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/001	2201434.00.00	ABZ-PHARMA GMBH	DE
Glatirameracetat AbZ 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5449/002	2201435.00.00	ABZ-PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Glatirameracetat Pliva 20 mg/ml otopina za injekciju u napunjenoj štrcaljki	DE/H/5449/001	HR-H-336175671	PLIVA HRVATSKA D.O.O.	HR
Glatirameracetat Pliva 40 mg/ml otopina za injekciju u napunjenoj štrcaljki	DE/H/5449/002	HR-H-436335081	PLIVA HRVATSKA D.O.O.	HR
Glatirameracetát Teva 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	DE/H/5449/001	59/0356/18-S	TEVA GMBH	SK
Glatirameracetát Teva 40 mg/ml injekčný roztok naplnený v injekčnej striekačke	DE/H/5449/002	59/0357/18-S	TEVA GMBH	SK
Glatiramero Mylan 20 mg/ml solución inyectable en jeringa precargada	NL/H/3213/001	80933	MYLAN PHARMACEUTICALS S.L.	ES
Glatiramero Mylan 40 mg/ml solución inyectable en jeringa precargada	NL/H/3777/001	82703	MYLAN PHARMACEUTICALS S.L.	ES
Glatiramyl 20 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3213/001	BE497386	MYLAN BVBA/SPRL	BE
Glatiramyl 20 mg/ml oplossing voor injectie in een voorgevulde spuit	NL/H/3213/001	BE497386	MYLAN BVBA/SPRL	BE
Glatiramyl 20 mg/ml solution injectable en seringue préremplie	NL/H/3213/001	BE497386	MYLAN BVBA/SPRL	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
Glatiramyl 40 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3777/001	BE518524	MYLAN BVBA/SPRL	BE
Glatiramyl 40 mg/ml solution injectable en seringue préremplie	NL/H/3777/001	BE518524	MYLAN BVBA/SPRL	BE
Glatiramyl 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3777/001	BE518524	MYLAN BVBA/SPRL	BE
Glaxaton 20 mg/ 1 ml Solution injecatable en seringue préremplie	DE/H/5449/001	2019050080	ACTAVIS GROUP PTC EHF.	LU
Glaxaton 40 mg/ 1 ml Solution injecatable en seringue préremplie	DE/H/5449/002	2019050081	ACTAVIS GROUP PTC EHF.	LU
Marcyto 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3776/001	RVG 119315	SYNTHON BV	NL
Marcyto 40 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3776/001	2018060182	SYNTHON BV	LU
Perscleran 20 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3212/001	136854	G.L. PHARMA GMBH	AT
Perscleran 20 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3212/001	136854	G.L. PHARMA 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
Perscleran 40 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3779/001	137986	G.L. PHARMA GMBH	AT
Remurel 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3211/001	59/0285/16-S	ZENTIVA, K.S.	SK
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/001	ZENTIVA, K.S.	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/002	ZENTIVA, K.S.	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/003	ZENTIVA, K.S.	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/004	ZENTIVA, K.S.	LT
Remurel 20 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3211/001	59/222/16-C	ZENTIVA, K.S.	CZ
Remurel 20 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3211/001	OGYI-T-23013/01	ALVOGEN IPCO S.AR.L	HU
Remurel 20 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3211/001	OGYI-T-23013/02	ALVOGEN IPCO S.AR.L	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
Remurel 20 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3211/001	OGYI-T-23013/03	ALVOGEN IPCO S.AR.L	HU
Remurel 20 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3211/001	OGYI-T-23013/04	ALVOGEN IPCO S.AR.L	HU
Remurel 20 mg/ml otopina za injekciju u napunjenoj štrcaljki	NL/H/3211/001	HR-H-759520422	ALVOGEN IPCO S.AR.L	HR
Remurel 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3211/001	H/16/02156/001	ALVOGEN IPCO S.AR.L	SI
Remurel 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3211/001	H/16/02156/002	ALVOGEN IPCO S.AR.L	SI
Remurel 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3211/001	H/16/02156/003	ALVOGEN IPCO S.AR.L	SI
Remurel 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3211/001	H/16/02156/004	ALVOGEN IPCO S.AR.L	SI
Remurel 20 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3211/001	16-0080	ZENTIVA, K.S.	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
Remurel 20 mg/ml soluție injectabilă în seringă preumplută	NL/H/3211/001	9075/2016/01	ALVOGEN IPCO S.AR.L	RO
Remurel 20 mg/ml soluție injectabilă în seringă preumplută	NL/H/3211/001	9075/2016/02	ALVOGEN IPCO S.AR.L	RO
Remurel 20 mg/ml soluție injectabilă în seringă preumplută	NL/H/3211/001	9075/2016/03	ALVOGEN IPCO S.AR.L	RO
Remurel 20 mg/ml soluție injectabilă în seringă preumplută	NL/H/3211/001	9075/2016/04	ALVOGEN IPCO S.AR.L	RO
Remurel 20 mg/ml stungulyf, lausn í áfylltri sprautu	NL/H/3211/001	IS/1/16/034/01	ALVOGEN IPCO S.AR.L	IS
Remurel 40 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3778/001	59/0141/18-S	ZENTIVA, K.S.	SK
Remurel 40 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3778/001	LT/1/17/4164/001	ALVOGEN IPCO S.AR.L	LT
Remurel 40 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3778/001	LT/1/17/4164/002	ALVOGEN IPCO S.AR.L	LT
Remurel 40 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3778/001	LT/1/17/4164/003	ALVOGEN IPCO S.AR.L	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
Remurel 40 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3778/001	59/676/16-C	ALVOGEN IPCO S.AR.L	CZ
Remurel 40 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3778/001	OGYI-T-23013/05	ALVOGEN IPCO S.AR.L	HU
Remurel 40 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3778/001	OGYI-T-23013/06	ALVOGEN IPCO S.AR.L	HU
Remurel 40 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3778/001	OGYI-T-23013/07	ALVOGEN IPCO S.AR.L	HU
Remurel 40 mg/ml otopina za injekciju, u napunjenoj štrcaljki	NL/H/3778/001	HR-H-589516554	ALVOGEN IPCO S.AR.L	HR
Remurel 40 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3778/001	H/18/02435/001	ALVOGEN IPCO S.AR.L	SI
Remurel 40 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3778/001	H/18/02435/002	ALVOGEN IPCO S.AR.L	SI
Remurel 40 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	NL/H/3778/001	H/18/02435/003	ALVOGEN IPCO S.AR.L	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
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ZENTIVA, K.S.	LV
Remurel 40 mg/ml soluţie injectabilă în seringă preumplută	NL/H/3778/001	10601/2018/01	ALVOGEN IPCO S.AR.L	RO
Remurel 40 mg/ml soluţie injectabilă în seringă preumplută	NL/H/3778/001	10601/2018/02	ALVOGEN IPCO S.AR.L	RO
Remurel 40 mg/ml soluţie injectabilă în seringă preumplută	NL/H/3778/001	10601/2018/03	ALVOGEN IPCO S.AR.L	RO
Remurel 40 mg/ml, stungulyf, lausn í áfylltri sprautu	NL/H/3778/001	IS/1/17/085/01	ALVOGEN IPCO S.AR.L	IS
Remurel, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3211/001	23282	ZENTIVA, K.S.	PL
Remurel, 20 mg/ml, süstelahus süstlis	NL/H/3211/001	909016	ZENTIVA, K.S.	EE
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko strzykawce	NL/H/3778/001	24402	ZENTIVA, K.S.	PL
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ZENTIVA, K.S.	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
Sclerthon 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3212/001	RVG 115987	SYNTHON BV	NL
Sclerthon 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3212/001	RVG 115987	SYNTHON BV	NL
Sclerthon 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3779/001	RVG 119322	SYNTHON BV	NL
Копаксон 20 mg/ml инжекционен разтвор, предварително напълнена спринцовка	not available	20040256	TEVA B.V	BG
Копаксон 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	not available	20160001	TEVA B.V	BG
Ремурел 20 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3211/001	20160170	ALVOGEN IPCO S.AR.L	BG
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3778/001	20170354	ALVOGEN IPCO S.AR.L	BG