12 July 2018
EMA/537197/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): glatiramer

Procedure No.: PSUSA/00001529/201711

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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
Perscleran 40 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3779/001	137986	G.L. PHARMA GMBH	AT
Sclerthon 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3779/001	RVG 119322	SYNTHON BV	NL
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
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 Alvogen 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3778/001	RVG 119321	ALVOGEN IPCO S.AR.L	NL
Glatirameeracetaat Alvogen 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3778/001	RVG 119321	ALVOGEN IPCO S.AR.L	NL
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 otopina za injekciju u napunjenoj štrcaljki	NL/H/3778/001	HR-H-589516554	ALVOGEN IPCO S.AR.L	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
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 40 mg/ml, stungulyf, lausn í áfylltri sprautu	NL/H/3778/001	IS/1/17/085/01	ALVOGEN IPCO S.AR.L	IS
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 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 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
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	BG
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	BG
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	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
спринцовка				
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
Remurel 40 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3778/001	17-0229	ALVOGEN IPCO S.AR.L	LV
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 Alvogen 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3778/001	RVG 119321	ALVOGEN IPCO S.AR.L	NL
Glatirameeracetaat Alvogen 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3778/001	RVG 119321	ALVOGEN IPCO S.AR.L	NL
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 otopina za injekciju u	NL/H/3778/001	HR-H-589516554	ALVOGEN IPCO S.AR.L	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
napunjenoj štrcaljki				
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 40 mg/ml, stungulyf, lausn í áfylltri sprautu	NL/H/3778/001	IS/1/17/085/01	ALVOGEN IPCO S.AR.L	IS
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 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 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
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Remurel, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3778/001	24402	ALVOGEN IPCO S.AR.L	PL
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	BG
Ремурел 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	BG
Ремурел 40 mg/ml инжекционен	NL/H/3778/001	II-39829	ALVOGEN IPCO S.AR.L	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
разтвор в предварително напълнена спринцовка				
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Remurel, 40 mg/ml, süstelahus süstlis	NL/H/3778/001	955417	ALVOGEN IPCO S.AR.L	EE
Copaxone 20 mg/ml Injekčný roztok, predplnená injekčná striekačka	DE/H/5283/002	59/0463/06-S	TEVA PHARMACEUTICALS CR, S.R.O.	SK
Copaxone 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	1-25380	TEVA GMBH	AT
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/01	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone® 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	52069.00.01	TEVA GMBH	DE
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418045	TEVA PHARMACEUTICALS LTD	IT
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, roztwór do wstrzykiwań, ampułkostrzykawki	DE/H/5283/002	12562	TEVA PHARMACEUTICALS LTD	PL
Copaxone 20 mg/ml Solução Injetável	DE/H/5283/002	4937686	TEVA PHARMACEUTICALS LTD	PT
Copaxone 20 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte.	DE/H/5283/002	03-2205	TEVA PHARMACEUTICALS LTD	NO
Copaxone 20 mg/ml voorgevulde spuit, oplossing voor injectie	DE/H/5283/002	RVG 30086	TEVA PHARMACEUTICALS LTD	NL
Copaxone ® 20 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/002	PL 10921/0023	TEVA PHARMACEUTICALS LTD	UK
Copaxone 20 mg/ml Solución inyectable	DE/H/5283/002	65.983	TEVA PHARMACEUTICALS	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
en jeringa precargada			LTD	
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418033	TEVA PHARMACEUTICALS LTD	IT
Copaxone 20 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5283/002	18786	TEVA PHARMACEUTICALS LTD	FI
Copaxone 20 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/002	58960/16.09.2008	TEVA PHARMACEUTICALS LTD	GR
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418058	TEVA PHARMACEUTICALS LTD	IT
Copaxone 20 mg/ml, injektionsvätska, lösning, förfylld spruta	DE/H/5283/002	20043	TEVA PHARMACEUTICALS LTD	SE
Copaxone 20mg/ml, solution injectable en seringue préremplie	UK/H/0453/002	2008069847	TEVA PHARMACEUTICALS LTD	LU
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/002	TEVA PHARMACEUTICALS LTD	LT
Copaxone 20 mg, stungulyf, lausn í áfylltri sprautu	DE/H/5283/002	IS/1/04/002/01	TEVA PHARMACEUTICALS LTD	IS
Copaxone 20 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	DE/H/5283/002	H/07/00419/001	TEVA PHARMACEUTICALS LTD	SI
Copaxone 20 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/002	035418021	TEVA PHARMACEUTICALS LTD	IT
COPAXONE 20 mg/ml šķīdums injekcijām pilnšļircē	DE/H/5283/002	06-0274	TEVA PHARMACEUTICALS LTD	LV
Copaxone 20 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/002	MA504/00102	TEVA PHARMACEUTICALS LTD	MT
Copaxone 20 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/002	7380/2015/01-04	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 20 mg/ml otopina za injekciju u	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
napunjenoj štrcaljki				
Copaxone 20 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/002	20187	TEVA PHARMACEUTICALS LTD	CY
Copaxone, injektionsvæske, opløsning, fyldt injektionssprøjte 20 mg/ml	DE/H/5283/002	35719	TEVA PHARMACEUTICALS LTD	DK
Copaxone 20mg/ml, solution injectable en seringue préremplie	DE/H/5283/002	BE260881	TEVA PHARMACEUTICALS LTD	BE
Copaxone, 20 mg/ml süstelahus süstlis	DE/H/5283/002	527006	TEVA PHARMACEUTICALS LTD	EE
Copaxone, 40 mg/ml süstelahus süstlis	DE/H/5283/004	863414	TEVA PHARMACEUTICALS LTD	EE
Copaxone 40 mg/ml injekčný roztok naplnený v injekčnej striekačke	DE/H/5283/004	59/0104/15-S	TEVA PHARMACEUTICALS CR, S.R.O.	SK
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	90395.00.00	TEVA GMBH	DE
Copaxone 40 mg/ml otopina za injekciju u napunjenoj štrcaljki	DE/H/5283/004	HR-H-304476486	PLIVA HRVATSKA D.O.O.	HR
Copaxone 40 mg/ml solution injectable, seringue préremplie	DE/H/5283/004	2015040057	TEVA PHARMACEUTICALS LTD	LU
Copaxone 40 mg/ml voorgevulde spuit, oplossing voor injectie	DE/H/5283/004	RVG 113849	TEVA PHARMACEUTICALS LTD	NL
Copaxone 40 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/004	PL 10921/0026	TEVA PHARMACEUTICALS LTD	UK
Copaxone, 40 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	DE/H/5283/004	22364	TEVA PHARMACEUTICALS LTD	PL
Copaxone 40 mg/ml, solution injectable en seringue préremplie	DE/H/5283/004	BE467902	TEVA PHARMACEUTICALS LTD	BE
Copaxone 40 mg/ml oldatos injekció	DE/H/5283/004	OGYI-T-9993/04	TEVA GYÓGYSZERGYÁR 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
előretöltött fecskendőben				
Copaxone 40 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5283/004	31602	TEVA PHARMACEUTICALS LTD	FI
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 Solution for Injection, Pre-filled Syringe	DE/H/5283/004	PA 1014/001/004	TEVA PHARMACEUTICALS LTD	IE
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418084	TEVA PHARMACEUTICALS LTD	IT
Copaxone, injektionsvæske, opløsning, fyldt injektionssprøjte 40 mg/ml	DE/H/5283/004	52806	TEVA PHARMACEUTICALS LTD	DK
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418072	TEVA PHARMACEUTICALS LTD	IT
Copaxone 40 mg/ml soluzione iniettabile, siringa preriempita	DE/H/5283/004	035418060	TEVA PHARMACEUTICALS LTD	IT
Copaxone 40 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/004	OGYI-T-9993/05	TEVA GYÓGYSZERGYÁR ZRT	HU
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	135998	TEVA GMBH	AT
Copaxone 40 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/004	MA 504/00104	TEVA PHARMACEUTICALS LTD	MT
COPAXONE 40 mg/ml šķīdums injekcijām pilnšļircē	DE/H/5283/004	15-0026	TEVA PHARMACEUTICALS LTD	LV
Copaxone 40 mg, stungulyf, lausn í áfylltri sprautu	DE/H/5283/004	IS/1/14/096/01	TEVA PHARMACEUTICALS LIMITED	IS
Copaxone 40 mg/ml solución inyectable en jeringa precargada	DE/H/5283/004	79515	TEVA PHARMACEUTICALS LTD	ES
Copaxone 40 mg/ml, injektionsvätska,	DE/H/5283/004	49557	TEVA PHARMACEUTICALS	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
lösning, förfylld spruta			LTD	
Copaxone 40 mg/ml solução injetável, seringa pré-cheia	DE/H/5283/004	5633136	TEVA PHARMACEUTICALS LTD	PT
Copaxone 40 mg/ml raztopina za injiciranje v napolnjeni injekcijski brizgi	DE/H/5283/004	H/07/00419/005	TEVA PHARMACEUTICALS LTD	SI
Copaxone 40 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	DE/H/5283/004	13-9600	TEVA PHARMACEUTICALS LTD	NO
Copaxone 20 mg/ml Solution for Injection, Pre-filled Syringe	DE/H/5283/002	PA 1014/001/002	TEVA PHARMACEUTICALS LTD	IE
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/03	TEVA PHARMACEUTICAL WORKS LTD	HU
Copaxone 20 mg/ml oldatos injekció előretöltött fecskendőben	DE/H/5283/002	OGYI-T-9993/02	TEVA PHARMACEUTICAL WORKS LTD	HU
Copaxone 20 mg/ml injektioneste, liuos, esitäytetty ruisku	DE/H/5283/002	18786	TEVA PHARMACEUTICALS LTD	FI
Copaxone 20 mg/ml oplossing voor injectie in een voorgevulde spuit.	DE/H/5283/002	BE260881	TEVA PHARMACEUTICALS LTD	BE
Copaxone 40 mg/ml injektionsvätska, lösning, förfylld spruta	DE/H/5283/004	31602	TEVA PHARMACEUTICALS LTD	FI
Copaxone 40 mg/ml oplossing voor injectie in een voorgevulde spuit.	DE/H/5283/004	BE467902	TEVA PHARMACEUTICALS LTD	BE
Copaxone 20 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/002	BE260881	TEVA PHARMACEUTICALS LTD	BE
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	8504/2016/01	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml Injektionslösung in einer Fertigspritze	DE/H/5283/004	BE467902	TEVA PHARMACEUTICALS LTD	BE
COPAXONE 20 mg/ml injekcinis tirpalas	DE/H/5283/002	LT/1/05/0240/004	TEVA PHARMACEUTICALS	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
užpildytame švirkšte			LTD	
COPAXONE 40 mg/ml, solution injectable en seringue préremplie	DE/H/5283/004	NL46033	TEVA SANTÉ	FR
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/003	TEVA PHARMACEUTICALS LTD	LT
Copaxone 40 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/004	81367/16-11-2015	TEVA PHARMA GMBH	GR
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	8504/2016/03	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml soluție injectabilă în seringă preumplută	DE/H/5283/004	8504/2016/02	TEVA PHARMACEUTICALS S.R.L	RO
Copaxone 40 mg/ml, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	DE/H/5283/004	22376	TEVA PHARMACEUTICALS LTD	CY
COPAXONE 20 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/002	LT/1/05/0240/005	TEVA PHARMACEUTICALS LTD	LT
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/008	TEVA PHARMACEUTICALS LTD	LT
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/007	TEVA PHARMACEUTICALS LTD	LT
COPAXONE 40 mg/ml injekcinis tirpalas užpildytame švirkšte	DE/H/5283/004	LT/1/05/0240/006	TEVA PHARMACEUTICALS LTD	LT
Copaxone 20 mg/ml, solution injectable en seringue préremplie	DE/H/5283/002	NL43730	TEVA SANTÉ	FR
Glatirameeracetaat Mylan 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3213/001	RVG 115993	MYLAN B.V.	NL
CLIFT 20 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3213/001	92742.00.00	MYLAN DURA 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
Glatiramyl 20 mg/ml Injektionslösung in einer Fertigspritze	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
Glatiramyl 20 mg/ml oplossing voor injectie in een voorgevulde spuit	NL/H/3213/001	BE497386	MYLAN BVBA/SPRL	BE
Glatiramero Mylan 20 mg/ml solución inyectable en jeringa precargada	NL/H/3213/001	80933	MYLAN PHARMACEUTICALS S.L.	ES
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
Copemyl 20 mg/ml soluzione iniettabile in siringa preriempita	NL/H/3213/001	043860042	MYLAN S.P.A.	IT
Acetato de glatirâmero Mylan 20 mg/1 ml Solução injetável em seringa pré-cheia	NL/H/3213/001	5697271	MYLAN, LDA	PT
GLATIRAMER MYLAN 20 mg/ml, solution injectable en seringue préremplie	NL/H/3213/001	NL 44923	MYLAN S.A.S	FR
Perscleran 20 mg/ml-Injektionslösung in einer Fertigspritze	NL/H/3212/001	136854	G.L. PHARMA GMBH	AT
Sclerthon 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3212/001	RVG 115987	SYNTHON BV	NL
Marcyto 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3776/001	RVG 119315	SYNTHON BV	NL
Копаксон 20 mg/ml инжекционен разтвор, предварително напълнена	not available	20040256	TEVA PHARMACEUTICALS 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
спринцовка				
Копаксон 40 mg/ml инжекционен разтвор в предварително напълнена спринцовка	not available	20160001	TEVA PHARMACEUTICALS BULGARIA EOOD	BG
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/001	ALVOGEN IPCO S.AR.L	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/002	ALVOGEN IPCO S.AR.L	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/003	ALVOGEN IPCO S.AR.L	LT
Remurel 20 mg/ml injekcinis tirpalas užpildytame švirkšte	NL/H/3211/001	LT/1/16/3917/004	ALVOGEN IPCO S.AR.L	LT
Remurel, 20 mg/ml, süstelahus süstlis	NL/H/3211/001	909016	ALVOGEN IPCO S.AR.L	EE
Remurel, 20 mg/ml, süstelahus süstlis	NL/H/3211/001	909016	ALVOGEN IPCO S.AR.L	EE
Remurel, 20 mg/ml, süstelahus süstlis	NL/H/3211/001	909016	ALVOGEN IPCO S.AR.L	EE
Remurel, 20 mg/ml, süstelahus süstlis	NL/H/3211/001	909016	ALVOGEN IPCO S.AR.L	EE
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 otopina za injekciju, u napunjenoj štrcaljki	NL/H/3211/001	HR-H-759520422	ALVOGEN IPCO S.AR.L	HR
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 otopina za injekciju, u napunjenoj štrcaljki	NL/H/3211/001	HR-H-759520422	ALVOGEN IPCO S.AR.L	HR
Brabio 20 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3211/001	RVG 115980	SYNTHON BV	NL
Remurel 20 mg/ml oldatos injekció előretöltött fecskendőben	NL/H/3211/001	OGYI-T-23013/01	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-23013/02	ALVOGEN IPCO S.AR.L	HU
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, stungulyf, lausn í áfylltri sprautu	NL/H/3211/001	IS/1/16/034/01	ALVOGEN IPCO S.AR.L	IS
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 20 mg/ml, stungulyf, lausn í áfylltri sprautu	NL/H/3211/001	IS/1/16/034/01	ALVOGEN IPCO S.AR.L	IS
Remurel 20 mg/ml, stungulyf, lausn í áfylltri sprautu	NL/H/3211/001	IS/1/16/034/01	ALVOGEN IPCO S.AR.L	IS
Copemyl, injektionsvæske, opløsning i fyldt injektionssprøjte 20 mg/ml	NL/H/3211/001	54650	MYLAN AB	DK
Glatimyl 20 mg/ml, injektionsvätska, lösning, förfylld spruta	NL/H/3211/001	51654	MYLAN AB	SE
Remurel 20 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3211/001	16-0080	ALVOGEN IPCO S.AR.L	LV
Remurel 20 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3211/001	16-0080	ALVOGEN IPCO S.AR.L	LV
Remurel 20 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3211/001	16-0080	ALVOGEN IPCO S.AR.L	LV
Remurel 20 mg/ml šķīdums injekcijām pilnšļircē	NL/H/3211/001	16-0080	ALVOGEN IPCO S.AR.L	LV
Remurel, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3211/001	23282	ALVOGEN IPCO S.AR.L	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
Remurel, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3211/001	23282	ALVOGEN IPCO S.AR.L	PL
Remurel, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3211/001	23282	ALVOGEN IPCO S.AR.L	PL
Remurel, 20 mg/ml, roztwór do wstrzykiwań w ampułko-strzykawce	NL/H/3211/001	23282	ALVOGEN IPCO S.AR.L	PL
Ремурел 20 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3211/001	II-33545	ALVOGEN IPCO S.AR.L	BG
Ремурел 20 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3211/001	II-33545	ALVOGEN IPCO S.AR.L	BG
Ремурел 20 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3211/001	II-33545	ALVOGEN IPCO S.AR.L	BG
Ремурел 20 mg/ml инжекционен разтвор в предварително напълнена спринцовка	NL/H/3211/001	II-33545	ALVOGEN IPCO S.AR.L	BG
Remurel 20 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3211/001	59/222/16-C	ALVOGEN IPCO S.AR.L	CZ
Remurel 20 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3211/001	59/222/16-C	ALVOGEN IPCO S.AR.L	CZ
Remurel 20 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3211/001	59/222/16-C	ALVOGEN IPCO S.AR.L	CZ
Remurel 20 mg/ml injekční roztok v předplněné injekční stříkačce	NL/H/3211/001	59/222/16-C	ALVOGEN IPCO S.AR.L	CZ
Remurel 20 mg/ml soluție injectabilă în seringă preumplută	NL/H/3211/001	9075/2016/01	ALVOGEN IPCO S.AR.L	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
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
Brabio 20 mg/ml, Solution for Injection, Pre-filled Syringe	NL/H/3211/001	MA1148/00101	SYNTHON BV	MT
Remurel 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3211/001	59/0285/16-S	ALVOGEN IPCO S.AR.L	SK
Remurel 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3211/001	59/0285/16-S	ALVOGEN IPCO S.AR.L	SK
Remurel 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3211/001	59/0285/16-S	ALVOGEN IPCO S.AR.L	SK
Remurel 20 mg/ml injekčný roztok naplnený v injekčnej striekačke	NL/H/3211/001	59/0285/16-S	ALVOGEN IPCO S.AR.L	SK
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
Copemyl 20 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte.	NL/H/3211/001	14-10261	MYLAN AB	NO
Brabio 20 mg/ml solution for injection, pre-filled syringe	NL/H/3211/001	PA0405/099/001	GENERICS [UK] LIMITED	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
CLIFT 20 mg/mL, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	NL/H/3211/001	94837 / 13-12-2016	MYLAN S.A.S	GR
Brabio 20 mg/mL solution for injection, pre-filled syringe	NL/H/3211/001	PL 04569/1725	GENERICS [UK] LIMITED	UK
Brabio 20 mg/mL, ενέσιμο διάλυμα σε προγεμισμένη σύριγγα	NL/H/3211/001	022466	MYLAN S.A.S	CY
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
CLIFT 40 mg/ml Injektionslösung in einer Fertigspritze	NL/H/3777/001	97606.00.00	MYLAN DURA GMBH	DE
Glatirameeracetaat Mylan 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3777/001	RVG 119319	MYLAN B.V.	NL
Glatiramyl 40 mg/ml, oplossing voor injectie in een voorgevulde spuit	NL/H/3777/001	BE518524	MYLAN BVBA/SPRL	BE
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
Brabio 40 mg/ml solution for injection, pre-filled syringe	NL/H/3777/001	PA0405/101/001	GENERICS [UK] LIMITED	IE
Brabio 40 mg/ml solution for injection, pre-filled syringe	NL/H/3777/001	PL 04569/1756	GENERICS [UK] LIMITED	UK
Glatiramero Mylan 40 mg/ml solución inyectable en jeringa precargada	NL/H/3777/001	82703	MYLAN PHARMACEUTICALS S.L.	ES
GLATIRAMER MYLAN 40 mg/ml, solution	NL/H/3777/001	NL47156	MYLAN S.A.S	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
injectable en seringue préremplie				
Glatimyl 40 mg/ml, injektionsvätska, lösning i förfylld spruta	NL/H/3777/001	54808	MYLAN AB	SE
Copemyl 40 mg/ml injeksjonsvæske, oppløsning i ferdigfylt sprøyte	NL/H/3777/001	16-11184	MYLAN AB	NO
Glatimyl 40 mg/ml injektioneste, liuos, esitäytetty ruisku	NL/H/3777/001	34220	MYLAN AB	FI
Copemyl, injektionsvæske, opløsning i fyldt injektionssprøjte 40 mg/ml	NL/H/3777/001	57826	MYLAN AB	DK