23 February 2023
EMA/164678/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): Ropinirole

Procedure No. PSUSA/00002661/202207

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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
ADARTREL 0,25 mg comprimidos recubiertos con película	FR/H/258/001	67.919	GLAXOSMITHKLINE, S.A.	ES
Adartrel 0,25 mg comprimidos revestidos por película	FR/H/258/001	5825583	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 0,25 mg filmdragerade tabletter	FR/H/258/001	21478	GLAXOSMITHKLINE AB	SE
ADARTREL 0,25 mg filmom obalené tablety	FR/H/258/001	27/0218/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
ADARTREL 0,25 mg Filmtabletten	FR/H/0258/001	60916.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
ADARTREL 0,25 mg, comprimé pelliculé	FR/H/258/001	NL29199	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel 0,5 mg comprimidos revestidos por película	FR/H/258/002	5825682	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 0,5 mg filmdragerade tabletter.	FR/H/258/002	21479	GLAXOSMITHKLINE AB	SE
ADARTREL 0,5 mg filmom obalené tablety	FR/H/258/002	27/0219/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
ADARTREL 0,5 mg Filmtabletten	FR/H/0258/002	60917.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
ADARTREL 0,50 mg comprimidos recubiertos con película	FR/H/258/002	67.921	GLAXOSMITHKLINE S.A.	ES
ADARTREL 0,50 mg, comprimé pelliculé	FR/H/0258/002	NL29200	LABORATOIRE GLAXOSMITHKLINE	FR
ADARTREL 2 mg comprimidos recubiertos con película	FR/H/258/004	67.922	GLAXOSMITHKLINE 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
Adartrel 2 mg comprimidos revestidos por película	FR/H/258/004	5825880	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 2 mg filmdragerade tabletter	FR/H/258/004	21481	GLAXOSMITHKLINE AB	SE
ADARTREL 2 mg Filmtabletten	FR/H/0258/004	60919.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
ADARTREL 2 mg, comprimé pelliculé	FR/H/0258/004	NL29202	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel, 0,25 mg, tabletki powlekane	FR/H/0258/001	12423	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Adartrel, 0,5 mg, tabletki powlekane	FR/H/0258/002	12424	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Adartrel, 2 mg, tabletki powlekane	FR/H/0258/004	12426	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Raponer, prolonged-release tablets	DK/H/2043/004	47992	ACTAVIS GROUP PTC EHF.	DK
Requip 5 mg film-coated tablets	FR/H/0111/005	MA 192/00105	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletes	FR/H/0255/006	LT/1/08/1077/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	FR/H/0255/006	LT/1/08/1077/003	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	FR/H/0255/006	LT/1/08/1077/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletes	FR/H/0255/008	LT/1/08/1077/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletes	FR/H/0255/008	LT/1/08/1077/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletes	FR/H/0255/009	LT/1/08/1077/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	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
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletes	FR/H/0255/009	LT/1/08/1077/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
Requip 0,25 mg - Filmtabletten	FR/H/0111/001	1-21708	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 0,25 mg compresse rivestite con film	FR/H/0111/001	032261063	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 0,25 mg compresse rivestite con film	FR/H/0111/001	032261099	LABORATOIRE GLAXOSMITHKLINE	IT
REQUIP 0,25 mg comprimidos recubiertos con película	FR/H/111/001	61.464	GLAXOSMITHKLINE S.A.	ES
Requip 0,25 mg filmdragerade tabletter.	FR/H/111/001	13222	GLAXOSMITHKLINE AB	SE
REQUIP 0,25 mg filmsko obložene tablete	not available	H/02/01344/001	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP 0,25 mg filmsko obložene tablete	not available	H/02/01344/002	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Requip 0,25 mg filmtabletta	not available	OGYI-T-6200/08	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP 0,25 mg Filmtabletten	FR/H/111/001	38707.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP 0,25 mg tabletki powlekane	FR/H/111/001	11115	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP 0,25 mg, comprimé pelliculé	FR/H/0111/001	NL21562	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 0,25, filmomhulde tabletten	FR/H/0111/001	RVG 20761	GLAXOSMITHKLINE B.V.	NL
Requip 0,5 mg compresse rivestite con film	FR/H/0111/002	032261101	LABORATOIRE GLAXOSMITHKLINE	IT
REQUIP 0,5 mg comprimidos recubiertos	FR/H/111/002	61.465	GLAXOSMITHKLINE 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
con película				
Requip 0,5 mg filmtabletta	not available	OGYI-T-6200/10	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip 0,5 mg filmtabletta	not available	OGYI-T-6200/12	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP 0,5 mg Filmtabletten	FR/H/111/002	38707.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP 0,5 mg tabletki powlekane	FR/H/0111/002	11116	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP 0,50 mg, comprimé pelliculé	FR/H/0111/002	NL21563	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 0.25 mg film-coated tablets	FR/H/0111/001	PA 1077/037/001	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Requip 0.25 mg film-coated tablets	FR/H/0111/001	MA 192/00101	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 1 mg - Filmtabletten	FR/H/0111/003	1-21710	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 1 mg compresse rivestite con film	FR/H/0111/003	032261125	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 1 mg comprimés pelliculés	FR/H/0111/003	BE182314	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip 1 mg comprimés pelliculés	FR/H/0111/003	2009 08 0539	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 1 mg comprimidos recubiertos con película	FR/H/111/003	61.466	GLAXOSMITHKLINE S.A.	ES
Requip 1 mg film-coated tablets	FR/H/0111/003	PA 1077/037/003	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Requip 1 mg film-coated tablets	FR/H/0111/003	MA 192/00103	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 1 mg filmdragerade tabletter.	FR/H/111/003	13224	GLAXOSMITHKLINE AB	SE
Requip 1 mg filmomhulde tabletten.	FR/H/0111/003	BE182314	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 1 mg filmsko obložene tablete	not available	H/02/01344/004	GLAXOSMITHKLINE TRADING SERVICES	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
			LIMITED	
Requip 1 mg filmtabletta	not available	OGYI-T-6200/15	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip 1 mg Filmtabletten	FR/H/0111/003	BE182314	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 1 mg Filmtabletten	FR/H/0111/003	38707.02.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Requip 1 mg Filmtabletten	FR/H/0111/003	2009 08 0539	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 1 mg tabletki powlekane	FR/H/111/003	11122	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP 1 mg, comprimé pelliculé	FR/H/111/003	NL21564	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 1, filmomhulde tabletten	FR/H/0111/003	RVG 20763	GLAXOSMITHKLINE B.V.	NL
Requip 2 mg - Filmtabletten	FR/H/0111/004	1-21711	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 2 mg compresse a rilascio prolungato	FR/H/0111/006	032261190	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg compresse a rilascio prolungato	FR/H/0111/006	032261202	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg compresse a rilascio prolungato	FR/H/0111/006	032261214	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg compresse rivestite con film	FR/H/0111/004	032261149	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg comprimés pelliculés	FR/H/0111/004	BE182305	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip 2 mg comprimés pelliculés	FR/H/111/004	2009 08 0540	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 2 mg comprimidos recubiertos con película	FR/H/111/004	61.468	GLAXOSMITHKLINE S.A.	ES
Requip 2 mg film-coated tablets	FR/H/0111/004	PA 1077/037/004	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Requip 2 mg film-coated tablets	FR/H/0111/004	MA 192/00104	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 2 mg	FR/H/111/004	13225	GLAXOSMITHKLINE AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
filmdragerade tabletter.				
Requip 2 mg filmomhulde tabletten	FR/H/0111/004	BE182305	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip 2 mg filmtabletta	not available	OGYI-T-6200/17	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip 2 mg Filmtabletten	FR/H/0111/004	BE182305	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 2 mg Filmtabletten	FR/H/0111/004	38707.03.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Requip 2 mg Filmtabletten	FR/H/0111/004	2009 08 0540	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 2 mg tabletki powlekane	FR/H/111/004	11113	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP 2 mg, comprimé pelliculé	FR/H/0111/004	NL21565	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 2, filmomhulde tabletten	FR/H/0111/004	RVG 20764	GLAXOSMITHKLINE B.V.	NL
Requip 4 mg compresse a rilascio prolungato	FR/H/0111/008	032261240	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 4 mg compresse a rilascio prolungato	FR/H/0111/008	032261253	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 5 mg - Filmtabletten	FR/H/111/005	1-21712	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 5 mg compresse rivestite con film	FR/H/0111/005	032261164	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 5 mg comprimés pelliculés	FR/H/111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip 5 mg comprimés pelliculés	FR/H/111/005	2009 08 0541	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 5 mg comprimidos recubiertos con película	FR/H/111/005	61.469	GLAXOSMITHKLINE S.A.	ES
Requip 5 mg film-coated tablets	FR/H/0111/005	PA 1077/037/005	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
Requip 5 mg filmdragerade tabletter	FR/H/111/005	13226	GLAXOSMITHKLINE AB	SE
Requip 5 mg	FR/H/111/005	BE184502	GLAXOSMITHKLINE	BE

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filmomhulde tabletten			PHARMACEUTICALS SA	
Requip 5 mg filmtabletta	not available	OGYI-T-6200/19	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip 5 mg Filmtabletten	FR/H/111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 5 mg Filmtabletten	FR/H/0111/005	38707.04.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
Requip 5 mg Filmtabletten	FR/H/0111/005	2009 08 0541	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP 5 mg tabletki powlekane	FR/H/0111/005	11114	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP 5 mg, comprimé pelliculé	FR/H/0111/005	NL21566	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 5, filmomhulde tabletten	FR/H/0111/005	RVG 20765	GLAXOSMITHKLINE B.V.	NL
Requip 8 mg compresse a rilascio prolungato	FR/H/0111/009	032261265	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 8 mg compresse a rilascio prolungato	FR/H/0111/009	032261277	LABORATOIRE GLAXOSMITHKLINE	IT
Requip Depot 2 mg depottablett	FR/H/0111/006	23934	GLAXOSMITHKLINE OY	FI
Requip Depot 2 mg depottabletter	not available	06-3959	GLAXOSMITHKLINE AS	NO
Requip Depot 2 mg depottabletter	FR/H/111/006	25843	GLAXOSMITHKLINE AB	SE
Requip Depot 2 mg depottabletti	FR/H/0111/006	23934	GLAXOSMITHKLINE OY	FI
Requip Depot 2 mg forðatöflur	not available	IS/1/07/164/01	GLAXOSMITHKLINE PHARMA A/S	IS
Requip Depot 4 mg depottabletter	FR/H/0111/008	23936	GLAXOSMITHKLINE OY	FI
Requip Depot 4 mg depottabletter	not available	06-3961	GLAXOSMITHKLINE AS	NO
Requip Depot 4 mg depottabletter	FR/H/0111/008	25845	GLAXOSMITHKLINE AB	SE
Requip Depot 4 mg depottabletti	FR/H/0111/008	23936	GLAXOSMITHKLINE OY	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
Requip Depot 4 mg forðatöflur	not available	IS/1/07/164/03	GLAXOSMITHKLINE PHARMA A/S	IS
Requip Depot 8 mg depottabletter	FR/H/0111/009	23937	GLAXOSMITHKLINE OY	FI
Requip Depot 8 mg depottabletter	not available	06-3962	GLAXOSMITHKLINE AS	NO
Requip Depot 8 mg depottabletter	FR/H/0111/009	25846	GLAXOSMITHKLINE AB	SE
Requip Depot 8 mg depottabletti	FR/H/0111/009	23937	GLAXOSMITHKLINE OY	FI
Requip Depot 8 mg forðatöflur	not available	IS/1/07/164/04	GLAXOSMITHKLINE PHARMA A/S	IS
Requip LP 2 mg comprimidos de libertação prolongada	FR/H/111/006	5080767	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Requip LP 2 mg comprimidos de libertação prolongada	FR/H/0111/006	5073911	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
REQUIP LP 2 mg, comprimé à libération prolongée	FR/H/0111/006	NL32489	LABORATOIRE GLAXOSMITHKLINE	FR
Requip LP 4 mg comprimidos de libertação prolongada	FR/H/0111/008	5080809	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Requip LP 4 mg comprimidos de libertação prolongada	FR/H/111/008	5073929	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
REQUIP LP 4 mg, comprimé à libération prolongée	FR/H/111/008	NL32491	LABORATOIRE GLAXOSMITHKLINE	FR
Requip LP 8 mg comprimidos de libertação prolongada	FR/H/0111/009	5080817	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT

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Requip LP 8 mg comprimidos de libertação prolongada	FR/H/111/009	5073937	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
REQUIP LP 8 mg, comprimé à libération prolongée	FR/H/0111/009	NL32492	LABORATOIRE GLAXOSMITHKLINE	FR
Requip Modutab 2 mg tablete s produljenim oslobađanjem	not available	HR-H-575488109	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Requip Modutab 2 mg tablety s prodlouženým uvolňováním	not available	27/461/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 4 mg tablete s produljenim oslobađanjem	not available	HR-H-723056317	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Requip Modutab 4 mg tablety s prodlouženým uvolňováním	not available	27/463/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 8 mg tablete s produljenim oslobađanjem	not available	HR-H-864556067	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HR
Requip Modutab 8 mg tablety s prodlouženým uvolňováním	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip XL 2 mg d?s??a pa?atetaμ???? ap?d?sμe?s??.	FR/H/0111/006	2330606	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Requip XL 4 mg δισκία παρατεταμένης αποδέσμευσης.	FR/H/0111/008	2330608	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Requip XL 8 mg δισκία παρατεταμένης αποδέσμευσης.	FR/H/0111/009	2330609	GLAXOSMITHKLINE SINGLE MEMBER A.E.B.E.	GR
Requip-Modutab 2 mg - Retardtabletten	FR/H/0111/006	1-27360	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 2 mg comprimate cu eliberare	not available	13169/2020/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO

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prelungita				
Requip-Modutab 2 mg comprimate cu eliberare prelungita	not available	13169/2020/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/07	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
REQUIP-MODUTAB 2 mg comprimés à libération prolongée.	FR/H/0111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP-MODUTAB 2 mg comprimés à libération prolongée.	FR/H/111/006	2008 06 0006	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg prolonged-release tablets	FR/H/0111/006	PA 1077/037/006	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REQUIP-MODUTAB 2 mg prolonged-release tablets	FR/H/255/006	MA192/00106	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU

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Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/20	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP-MODUTAB 2 mg Retardtabletten	FR/H/111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP-MODUTAB 2 mg Retardtabletten	FR/H/0111/006	69867.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP-MODUTAB 2 mg Retardtabletten	FR/H/0111/006	2008 06 0006	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 2 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-Modutab 2 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/010	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-MODUTAB 2 mg tabletten met verlengde afgifte	FR/H/111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip-Modutab 2 mg tabletten met verlengde afgifte	FR/H/0111/006	RVG 100571	GLAXOSMITHKLINE B.V.	NL
REQUIP-MODUTAB 2 mg tablety s predlženým uvolnovaním	not available	27/0311/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
REQUIP-MODUTAB 2 mg tablety s predlženým uvolnovaním	not available	27/0311/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Requip-Modutab 2 mg toimeainet prolongeeritult vabastavad tabletid	not available	532306	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Requip-Modutab 4 mg - Retardtabletten	FR/H/0111/008	1-27362	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 4 mg comprimate cu eliberare	not available	13170/2020/02	GLAXOSMITHKLINE (IRELAND) LIMITED	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
prelungită				
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
REQUIP-MODUTAB 4 mg comprimés à libération prolongée	FR/H/0111/008	2008 06 0008	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-MODUTAB 4 mg comprimés à libération prolongée.	FR/H/0111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip-Modutab 4 mg ilgstošas darbibas tabletes	not available	06-0263	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 4 mg prolonged-release tablets	FR/H/0111/008	PA 1077/037/008	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REQUIP-MODUTAB 4 mg prolonged-release tablets	FR/H/255/008	MA192/00108	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip-Modutab 4 mg retard filmtabletta	not available	OGYI-T-6200/05	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip-Modutab 4 mg retard filmtabletta	not available	OGYI-T-6200/21	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/111/008	69869.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/0111/008	2008 06 0008	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	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
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-MODUTAB 4 mg tabletten met verlengde afgifte	FR/H/111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip-Modutab 4 mg tabletten met verlengde afgifte	FR/H/0111/008	RVG 100574	GLAXOSMITHKLINE B.V.	NL
REQUIP-MODUTAB 4 mg tablety s predlženým uvolnovaním	not available	27/0313/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
REQUIP-MODUTAB 4 mg tablety s predlženým uvolnovaním	not available	27/0313/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Requip-Modutab 4 mg toimeainet prolongeeritult vabastavad tabletid	not available	532106	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Requip-Modutab 8 mg - Retardtabletten	FR/H/0111/009	1-27364	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	13171/2020/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	13171/2020/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	13171/2020/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	13171/2020/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
REQUIP-MODUTAB 8 mg comprimés à libération prolongée	FR/H/0111/009	BE316565	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP-MODUTAB 8 mg comprimés à libération	FR/H/0111/009	2008 06 0004	GLAXOSMITHKLINE PHARMACEUTICALS SA	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
prolongée				
Requip-Modutab 8 mg ilgstošas darbibas tabletes	not available	06-0264	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 8 mg prolonged-release tablets	FR/H/0111/009	PA 1077/037/009	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REQUIP-MODUTAB 8 mg prolonged-release tablets	FR/H/255/009	MA192/00109	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip-Modutab 8 mg retard filmtabletta	not available	OGYI-T-6200/22	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip-Modutab 8 mg retard filmtabletta	not available	OGYI-T-6200/06	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP-MODUTAB 8 mg Retardtabletten	FR/H/111/009	BE 316565	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP-MODUTAB 8 mg Retardtabletten	FR/H/0111/009	69870.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP-MODUTAB 8 mg Retardtabletten	FR/H/0111/009	2008 06 0004	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 8 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-Modutab 8 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/012	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-MODUTAB 8 mg tabletten met verlengde afgifte	FR/H/111/009	BE316565	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
Requip-Modutab 8 mg tabletten met verlengde afgifte	FR/H/0111/009	RVG 100576	GLAXOSMITHKLINE B.V.	NL
REQUIP-MODUTAB 8 mg tablety s predlženým uvolnovaním	not available	27/0314/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
REQUIP-MODUTAB 8 mg tablety s predlženým uvolnovaním	not available	27/0314/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Requip-Modutab 8 mg toimeainet prolongeeritult vabastavad tabletid	not available	532006	GLAXOSMITHKLINE (IRELAND) LIMITED	EE
Requip-Modutab, 2 mg, tabletki o przedluzonym uwalnianiu	FR/H/0255/006	14577	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Requip-Modutab, 4 mg, tabletki o przedłużonym uwalnianiu	FR/H/0255/008	14579	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
Requip-Modutab, 8 mg, tabletki o przedłużonym uwalnianiu	FR/H/255/009	14580	GLAXOSMITHKLINE (IRELAND) LIMITED	PL
REQUIP-PROLIB 2 mg comprimidos de liberación prolongada.	FR/H/111/006	69.700	GLAXOSMITHKLINE S.A.	ES
REQUIP-PROLIB 4 mg comprimidos de liberación prolongada	FR/H/111/008	69.701	GLAXOSMITHKLINE S.A.	ES
REQUIP-PROLIB 8 mg comprimidos de liberación prolongada	FR/H/0111/009	69.702	GLAXOSMITHKLINE S.A.	ES
Ropinirol - 1 A Pharma 3 mg Filmtabletten	not available	71161.00.00	1 A PHARMA GMBH	DE
Ropinirol - 1 A Pharma 4 mg Filmtabletten	not available	71162.00.00	1 A PHARMA GMBH	DE
Ropinirol "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099304	DECOMED FARMACÊUTICA, LDA.	PT
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099437	DECOMED FARMACÊUTICA, LDA.	PT
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099270	DECOMED FARMACÊUTICA, LDA.	PT
Ropinirol Genedec 4 mg	not available	5099320	DECOMED FARMACÊUTICA,	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
comprimidos revestidos por película			LDA.	
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099452	DECOMED FARMACÊUTICA, LDA.	PT
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099460	DECOMED FARMACÊUTICA, LDA.	PT
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099312	DECOMED FARMACÊUTICA, LDA.	PT
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	NL34106	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	NL34107	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	NL34108	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	NL34109	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE PAUCOURT LP 2 mg, comprimé à libération prolongée	FR/H/0255/006	NL32485	LABORATOIRES PAUCOURT	FR
ROPINIROLE PAUCOURT LP 4 mg, comprimé à libération prolongée	FR/H/0255/008	NL32487	LABORATOIRES PAUCOURT	FR
ROPINIROLE PAUCOURT LP 8 mg, comprimé à libération prolongée	FR/H/0255/009	NL32488	LABORATOIRES PAUCOURT	FR
Ropinirol-neuraxpharm 3 mg Filmtabletten	not available	68843.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Ropinirol-neuraxpharm 4 mg Filmtabletten	not available	68844.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE