26 March 2020
EMA/249485/2020
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: ropinirole

Procedure no.: PSUSA00002661201907

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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 filmdrasjerte tabletter	FR/H/258/001	04-2893	GLAXOSMITHKLINE AS	NO
ADARTREL 0,25 mg filmom obalené tablety	FR/H/258/001	27/0218/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ADARTREL 0,25 mg Filmtabletten	FR/H/0258/001	60916.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
ADARTREL 0,25 mg filmuhúðaðar töflur	FR/H/0258/001	IS/1/04/057/01	GLAXOSMITHKLINE PHARMA A/S	IS
ADARTREL 0,25 mg, comprimé pelliculé	FR/H/258/001	NL29199	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel 0,25 mg, filmomhulde tabletten 0,25 mg	FR/H/258/001	RVG 31670	GLAXOSMITHKLINE 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
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 filmdrasjerte tabletter	FR/H/0258/002	04-2895	GLAXOSMITHKLINE AS	NO
ADARTREL 0,5 mg filmom obalené tablety	FR/H/258/002	27/0219/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ADARTREL 0,5 mg Filmtabletten	FR/H/0258/002	60917.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
ADARTREL 0,5 mg filmuhúðaðar töflur	FR/H/0258/002	IS/1/04/057/02	GLAXOSMITHKLINE PHARMA A/S	IS
Adartrel 0,5 mg, filmomhulde tabletten 0,5 mg	FR/H/258/002	RVG 31671	GLAXOSMITHKLINE B.V.	NL
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

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 film-coated tablets	FR/H/258/001	PL 19494/0033	GLAXOSMITHKLINE UK LIMITED	UK
ADARTREL 0.5 mg film-coated tablets.	FR/H/0258/002	PL 19494/0034	GLAXOSMITHKLINE UK LIMITED	UK
ADARTREL 2 mg comprimidos recubiertos con película	FR/H/258/004	67.922	GLAXOSMITHKLINE S.A.	ES
Adartrel 2 mg comprimidos revestidos por película	FR/H/258/004	5825880	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 2 mg film-coated tablets.	FR/H/0258/004	PL 19494/0036	GLAXOSMITHKLINE UK LIMITED	UK
ADARTREL 2 mg filmdragerade tabletter	FR/H/258/004	21481	GLAXOSMITHKLINE AB	SE
ADARTREL 2 mg filmdrasjerte tabletter	FR/H/258/004	04-2897	GLAXOSMITHKLINE AS	NO
ADARTREL 2 mg filmom obalené tablety	FR/H/258/004	27/0221/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	SK
ADARTREL 2 mg Filmtabletten	FR/H/0258/004	60919.00.00	GLAXOSMITHKLINE GMBH & CO. KG	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
ADARTREL 2 mg filmuhúðaðar töflur.	FR/H/0258/004	IS/1/04/057/04	GLAXOSMITHKLINE PHARMA A/S	IS
ADARTREL 2 mg, comprimé pelliculé	FR/H/0258/004	NL29202	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel 2 mg, filmomhulde tabletten 2 mg	FR/H/258/004	RVG 31673	GLAXOSMITHKLINE B.V.	NL
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
Ipinnia XL 2 mg prolonged-release tablets	not available	PL 01883/0326	MACARTHYS LABORATORIES LTD	UK
Ipinnia XL 3 mg prolonged-release tablets	not available	PL 01883/0327	MACARTHYS LABORATORIES LTD	UK
Ipinnia XL 4 mg prolonged-release tablets	not available	PL 01883/0328	MACARTHYS LABORATORIES 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
Ipinnia XL 6 mg prolonged-release tablets	not available	PL 01883/0329	MACARTHYS LABORATORIES LTD	UK
Ipinnia XL 8 mg prolonged-release tablets	not available	PL 01883/0330	MACARTHYS LABORATORIES LTD	UK
Raponer PR 6 mg prolonged-release tablets	MT/H/0352/004	MA1341/00304	SIGILLATA LIMITED	MT
Raponer, depottabletter	DK/H/2043/004	47992	ACTAVIS GROUP PTC EHF.	DK
Requip	not available	PL 10592/0088	SMITHKLINE BEECHAM LTD	UK
Requip	not available	PL 10592/0089	SMITHKLINE BEECHAM LTD	UK
Requip	not available	PL 10592/0087	SMITHKLINE BEECHAM LTD	UK
Requip	not available	PL 10592/0085	SMITHKLINE BEECHAM LTD	UK
Requip 5 mg film-coated tablets	FR/H/0111/005	MA 192/00105	GLAXOSMITHKLINE (IRELAND) 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
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	FR/H/0255/006	LT/1/08/1077/002	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	FR/H/0255/006	LT/1/08/1077/003	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	FR/H/0255/006	LT/1/08/1077/004	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletės	FR/H/0255/008	LT/1/08/1077/007	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletės	FR/H/0255/008	LT/1/08/1077/008	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletes	FR/H/0255/009	LT/1/08/1077/009	GLAXOSMITHKLINE LIETUVA UAB	LT
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletės	FR/H/0255/009	LT/1/08/1077/010	GLAXOSMITHKLINE LIETUVA UAB	LT
Requip 0,25 filmdragerade tabletter	FR/H/111/001	13222	GLAXOSMITHKLINE AB	SE
Requip 0,25 mg - Filmtabletten	FR/H/0111/001	1-21708	GLAXOSMITHKLINE 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
Requip 0,25 mg compresse rivestite con film	FR/H/111/001	032261063	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 0,25 mg compresse rivestite con film	FR/H/111/001	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 filmsko obložene tablete	not available	H/02/01344/001	GLAXOSMITHKLINE D.O.O.	SI
REQUIP 0,25 mg filmsko obložene tablete	not available	H/02/01344/002	GLAXOSMITHKLINE D.O.O.	SI
Requip 0,25 mg filmtabletta	not available	OGYI-T-6200/08	GLAXOSMITHKLINE KFT.	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

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 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 con película	FR/H/111/002	61.465	GLAXOSMITHKLINE S.A.	ES
REQUIP 0,5 mg filmsko obložene tablete	not available	H/02/01344/003	GLAXOSMITHKLINE D.O.O.	SI
Requip 0,5 mg filmtabletta	not available	OGYI-T-6200/10	GLAXOSMITHKLINE KFT.	HU
Requip 0,5 mg filmtabletta	not available	OGYI-T-6200/12	GLAXOSMITHKLINE KFT.	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

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 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 filmdragerade tabletter	FR/H/111/003	13224	GLAXOSMITHKLINE AB	SE
Requip 1 mg - Filmtabletten	FR/H/0111/003	1-21710	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 1 mg compresse rivestite con film	FR/H/111/003	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

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 1 mg film-coated tablets	FR/H/0111/003	MA 192/00103	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
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 D.O.O.	SI
Requip 1 mg filmtabletta	not available	OGYI-T-6200/15	GLAXOSMITHKLINE KFT.	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	LABORATOIRES PAUCOURT	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
Requip 1, filmomhulde tabletten	FR/H/0111/003	RVG 20763	GLAXOSMITHKLINE B.V.	NL
Requip 2 filmdragerade tabletter	FR/H/111/004	13225	GLAXOSMITHKLINE AB	SE
Requip 2 mg - Filmtabletten	FR/H/0111/004	1-21711	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip 2 mg compresse a rilascio prolungato	FR/H/111/006	032261190	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg compresse a rilascio prolungato	FR/H/0111/006	032261214	LABORATOIRE GLAXOSMITHKLINE	IT
Requip 2 mg compresse a rilascio prolungato.	FR/H/0111/006	032261202	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

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 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 filmomhulde tabletten	FR/H/0111/004	BE182305	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 2 mg filmsko obložene tablete	not available	H/02/01344/005	GLAXOSMITHKLINE D.O.O.	SI
Requip 2 mg filmtabletta	not available	OGYI-T-6200/17	GLAXOSMITHKLINE KFT.	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

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 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/111/005	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

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 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 filmomhulde tabletten	FR/H/111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS SA	BE
REQUIP 5 mg filmsko obložene tablete	not available	H/02/01344/006	GLAXOSMITHKLINE D.O.O.	SI
Requip 5 mg filmtabletta	not available	OGYI-T-6200/18	GLAXOSMITHKLINE KFT.	HU
Requip 5 mg filmtabletta	not available	OGYI-T-6200/19	GLAXOSMITHKLINE KFT.	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

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 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

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 2 mg forðatöflur	not available	IS/1/07/164/01	GLAXOSMITHKLINE PHARMA A/S	IS
Requip Depot 4 mg depottablett	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
Requip Depot 4 mg forðatöflur	not available	IS/1/07/164/03	GLAXOSMITHKLINE PHARMA A/S	IS
Requip Depot 8 mg depottablett	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

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 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

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 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	UP/I-530-09/13-02/392	GLAXOSMITHKLINE D.O.O.	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	UP/I-530-09/13-02/393	GLAXOSMITHKLINE D.O.O.	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	UP/I-530-09/13-02/394	GLAXOSMITHKLINE D.O.O.	HR
Requip Modutab 8 mg tablety s prodlouženým uvolňováním	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip XL 2 mg prolonged-release tablets.	not available	PL 10592/0293	SMITHKLINE BEECHAM 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
Requip XL 2 mg δισκία παρατεταμένης αποδέσμευσης.	FR/H/0111/006	2330606	GLAXOSMITHKLINE AEBE	GR
Requip XL 4 mg prolonged-release tablets.	not available	PL 10592/0295	SMITHKLINE BEECHAM LTD	UK
Requip XL 4 mg δισκία παρατεταμένης αποδέσμευσης.	FR/H/0111/008	2330608	GLAXOSMITHKLINE AEBE	GR
Requip XL 8 mg prolonged-release tablets.	not available	PL 10592/0296	SMITHKLINE BEECHAM LTD	UK
Requip XL 8 mg δισκία παρατεταμένης αποδέσμευσης.	FR/H/0111/009	2330609	GLAXOSMITHKLINE AEBE	GR
Requip-Modutab 2 mg - Retardtabletten	FR/H/0111/006	1-27360	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 2 mg comprimate cu eliberare prelungita	not available	842/2008/05	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/06	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/07	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
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/08	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	842/2008/04	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 ilgstošās darbības tabletes	not available	06-0261	GLAXOSMITHKLINE LATVIA SIA	LV
Requip-Modutab 2 mg prolonged-release tablets	FR/H/0111/006	PA 1077/037/006	GLAXOSMITHKLINE (IRELAND) 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
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 KFT.	HU
Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/04	GLAXOSMITHKLINE KFT.	HU
Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/20	GLAXOSMITHKLINE KFT.	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	2008060006	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 2 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/007	GLAXOSMITHKLINE D.O.O.	SI
REQUIP-Modutab 2 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/010	GLAXOSMITHKLINE 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
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 SLOVAKIA S.R.O.	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 prelungita	not available	844/2008/01	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	844/2008/03	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	844/2008/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	844/2008/02	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
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 ilgstošās darbības tabletes	not available	06-0263	GLAXOSMITHKLINE LATVIA SIA	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 KFT.	HU
Requip-Modutab 4 mg retard filmtabletta	not available	OGYI-T-6200/21	GLAXOSMITHKLINE KFT.	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

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 Retardtabletten	FR/H/0111/008	2008060008	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/011	GLAXOSMITHKLINE D.O.O.	SI
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/008	GLAXOSMITHKLINE D.O.O.	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 predĺženým uvoľňovaním	not available	27/0313/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	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	845/2008/03	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
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	845/2008/04	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	845/2008/02	GLAXOSMITHKLINE (IRELAND) LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	845/2008/01	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 prolongée	FR/H/0111/009	2008 06 0004	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
Requip-Modutab 8 mg ilgstošās darbības tabletes	not available	06-0264	GLAXOSMITHKLINE LATVIA SIA	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 KFT.	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
Requip-Modutab 8 mg retard filmtabletta	not available	OGYI-T-6200/06	GLAXOSMITHKLINE KFT.	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	2008060004	GLAXOSMITHKLINE PHARMACEUTICALS SA	LU
REQUIP-Modutab 8 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/012	GLAXOSMITHKLINE D.O.O.	SI
REQUIP-Modutab 8 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/009	GLAXOSMITHKLINE D.O.O.	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 predĺženým uvoľňovaním	not available	27/0314/06-S	GLAXOSMITHKLINE SLOVAKIA S.R.O.	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 przedłużonym 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
Ropilynz XL 2 mg prolonged-release tablets	not available	PL 35507/0153	LUPIN HEALTHCARE (UK) LIMITED	UK
Ropilynz XL 4 mg prolonged-release tablets	not available	PL 35507/0154	LUPIN HEALTHCARE (UK) LIMITED	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
Ropilynz XL 8 mg prolonged-release tablets	not available	PL 35507/0155	LUPIN HEALTHCARE (UK) LIMITED	UK
Ropinict	DK/H/1031/006	45562	SANDOZ A/S	DK
Ropinict	DK/H/1031/007	45563	SANDOZ A/S	DK
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", depottabletter	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099304	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099437	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol Genedec 3 mg comprimidos revestidos por película	not available	5099270	DECOMED FARMACÊUTICA, S.A.	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
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099320	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099452	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099460	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol Genedec 4 mg comprimidos revestidos por película	not available	5099312	DECOMED FARMACÊUTICA, S.A.	PT
Ropinirol STADA 3 mg Filmtabletten	DE/H/1052/005	68829.00.00	STADAPHARM GMBH	DE
Ropinirol STADA 4 mg Filmtabletten	DE/H/1052/006	68830.00.00	STADAPHARM GMBH	DE
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

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
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	NL34109	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 5 mg, comprimé pelliculé	DE/H/1052/007	NL34110	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
Sindranol3 mg tablety s predĺženým uvol'ňovaním	DE/H/2132/002	27/0020/12-S	EGIS PHARMACEUTICALS PLC	SK