25 February 2026
EMADOC-1700519818-3052528
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): ropinirole

EURD list No. PSUSA/00002661/202507

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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/0258/001	67.919	GLAXOSMITHKLINE, S.A.	ES
Adartrel 0,25 mg comprimidos revestidos por película	FR/H/0258/001	5825583	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 0,25 mg filmdragerade tabletter.	FR/H/0258/001	21478	GLAXOSMITHKLINE AB	SE
ADARTREL 0,25 mg filmom obalené tablety	FR/H/0258/001	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/0258/001	6 942 205 7	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel 0,5 mg comprimidos revestidos por película	FR/H/0258/002	5825682	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 0,5 mg filmdragerade tabletter.	FR/H/0258/002	21479	GLAXOSMITHKLINE AB	SE
ADARTREL 0,5 mg filmdragerade tabletter.	FR/H/0258/002	21479	GLAXOSMITHKLINE AB	SE
ADARTREL 0,5 mg filmom obalené tablety	FR/H/0258/002	27/0219/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
ADARTREL 0,5 mg filmom obalené tablety	FR/H/0258/002	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/0258/002	67.921	GLAXOSMITHKLINE, S.A.	ES

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ADARTREL 0,50 mg comprimidos recubiertos con película	FR/H/0258/002	67.921	GLAXOSMITHKLINE, S.A.	ES
ADARTREL 0,50 mg, comprimé pelliculé	FR/H/0258/002	6 998 292 8	LABORATOIRE GLAXOSMITHKLINE	FR
ADARTREL 0,50 mg, comprimé pelliculé	FR/H/0258/002	6 998 292 8	LABORATOIRE GLAXOSMITHKLINE	FR
ADARTREL 2 mg comprimidos recubiertos con película	FR/H/0258/004	67.922	GLAXOSMITHKLINE, S.A.	ES
ADARTREL 2 mg comprimidos recubiertos con película	FR/H/0258/004	67.922	GLAXOSMITHKLINE, S.A.	ES
Adartrel 2 mg comprimidos revestidos por película	FR/H/0258/004	5825880	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
ADARTREL 2 mg filmdragerade tabletter.	FR/H/0258/004	21481	GLAXOSMITHKLINE AB	SE
ADARTREL 2 mg filmdragerade tabletter.	FR/H/0258/004	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	6 727 699 5	LABORATOIRE GLAXOSMITHKLINE	FR
ADARTREL 2 mg, comprimé pelliculé	FR/H/0258/004	6 727 699 5	LABORATOIRE GLAXOSMITHKLINE	FR
Adartrel, 0,5 mg, tabletki powlekane	FR/H/0258/002	12424	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Adartrel, 0,5 mg, tabletki powlekane	FR/H/0258/002	12424	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Ipinnia XL 2 mg prolonged-release tablets	not available	PL 01883/0326	MACARTHYS LABORATORIES 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
Ipinnia XL 2 mg prolonged-release tablets	not available	PL 01883/0326	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 3 mg prolonged-release tablets	not available	PL 01883/0327	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 3 mg prolonged-release tablets	not available	PL 01883/0327	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 4 mg prolonged-release tablets	not available	PL 01883/0328	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 4 mg prolonged-release tablets	not available	PL 01883/0328	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 6 mg prolonged-release tablets	not available	PL 01883/0329	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 6 mg prolonged-release tablets	not available	PL 01883/0329	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 8 mg prolonged-release tablets	not available	PL 01883/0330	MACARTHYS LABORATORIES LTD	XI
Ipinnia XL 8 mg prolonged-release tablets	not available	PL 01883/0330	MACARTHYS LABORATORIES LTD	XI
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
Raponer, prolonged-	DK/H/2043/004	47992	TEVA B.V	DK

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release tablets				
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
Raponer, prolonged-release tablets	DK/H/2043/004	47992	TEVA B.V	DK
REQUIP - MODUTAB 2 mg pailginto atpalaidavimo tabletės	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/1/08/1077/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletės	FR/H/0255/008	LT/1/08/1077/007	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 4 mg pailginto atpalaidavimo tabletės	FR/H/0255/008	LT/1/08/1077/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletės	FR/H/0255/009	LT/1/08/1077/009	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LT
REQUIP - MODUTAB 8 mg pailginto atpalaidavimo tabletės	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/0111/001	61.464	GLAXOSMITHKLINE, S.A.	ES
Requip 0,25 mg filmdragerade tabletter.	FR/H/0111/001	13222	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 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 Filmtabletten	FR/H/0111/001	38707.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP 0,25 mg tabletki powlekane	FR/H/0111/001	11115	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 0,25 mg, comprimé pelliculé	FR/H/0111/001	6 149 795 1	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 0,25, filmomhulde tabletten 0,25 mg	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/0111/002	61.465	GLAXOSMITHKLINE, S.A.	ES
Requip 0,5 mg filmtabletta	not available	OGYI-T-6200/12	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
REQUIP 0,5 mg Filmtabletten	FR/H/0111/002	38707.01.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP 0,5 mg tabletki powlekane	FR/H/0111/002	11116	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 0,50 mg, comprimé pelliculé	FR/H/0111/002	6 669 563 8	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

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	MA 192/00101	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
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 EN ABREGE GLAXOSMITHKLINE	BE
Requip 1 mg comprimés pelliculés	FR/H/0111/003	2009080539	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 1 mg comprimidos recubiertos con película	FR/H/0111/003	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 film-coated tablets.	FR/H/0111/003	MA 192/00103	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 1 mg filmdragerade tabletter.	FR/H/0111/003	13224	GLAXOSMITHKLINE AB	SE
Requip 1 mg filmomhulde tabletten.	FR/H/0111/003	BE182314	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
REQUIP 1 mg filmsko obložene tablete	not available	H/02/01344/004	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
Requip 1 mg Filmtabletten	FR/H/0111/003	BE182314	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
REQUIP 1 mg Filmtabletten	FR/H/0111/003	38707.02.00	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 1 mg Filmtabletten	FR/H/0111/003	2009080539	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 1 mg tabletki powlekane	FR/H/0111/003	11122	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 1 mg tabletki powlekane	FR/H/0111/003	11122	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 1 mg, comprimé pelliculé	FR/H/0111/003	6 664 952 3	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 1, filmomhulde tabletten 1 mg	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	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 EN ABREGE GLAXOSMITHKLINE	BE
Requip 2 mg comprimés pelliculés	FR/H/0111/004	2009080540	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 2 mg comprimidos recubiertos con película	FR/H/0111/004	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 film-coated tablets.	FR/H/0111/004	MA 192/00104	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 2 mg filmdragerade tabletter.	FR/H/0111/004	13225	GLAXOSMITHKLINE AB	SE
Requip 2 mg filmomhulde tabletten.	FR/H/0111/004	BE182305	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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 EN ABREGE GLAXOSMITHKLINE	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	2009080540	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 2 mg tabletki powlekane	FR/H/0111/004	11113	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 2 mg tabletki powlekane	FR/H/0111/004	11113	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 2 mg tabletki powlekane	FR/H/0111/004	11113	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 2 mg tabletki powlekane	FR/H/0111/004	11113	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 2 mg, comprimé pelliculé	FR/H/0111/004	6 820 907 0	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 2, filmomhulde tabletten 2 mg	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

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 - Filmtabletten	FR/H/0111/005	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/0111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Requip 5 mg comprimés pelliculés	FR/H/0111/005	2009080541	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 5 mg comprimidos recubiertos con película	FR/H/0111/005	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 film-coated tablets.	FR/H/0111/005	MA 192/00105	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 5 mg film-coated tablets.	FR/H/0111/005	MA 192/00105	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip 5 mg filmdragerade tabletter.	FR/H/0111/005	13226	GLAXOSMITHKLINE AB	SE
Requip 5 mg filmomhulde tabletten.	FR/H/0111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
Requip 5 mg filmtabletta	not available	OGYI-T-6200/19	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
Requip 5 mg Filmtabletten	FR/H/0111/005	BE184502	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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	2009080541	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP 5 mg tabletki	FR/H/0111/005	11114	GLAXOSMITHKLINE	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
powlekane			TRADING SERVICES LIMITED	
REQUIP 5 mg tabletki powlekane	FR/H/0111/005	11114	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 5 mg tabletki powlekane	FR/H/0111/005	11114	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 5 mg tabletki powlekane	FR/H/0111/005	11114	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP 5 mg, comprimé pelliculé	FR/H/0111/005	6 754 812 1	LABORATOIRE GLAXOSMITHKLINE	FR
Requip 5, filmomhulde tabletten 5 mg	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 depottabletter	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/0111/006	25843	GLAXOSMITHKLINE AB	SE
Requip Depot 2 mg depottabletter.	FR/H/0111/006	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	FR/H/0111/008	25845	GLAXOSMITHKLINE AB	SE

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depottabletter.				
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 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 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/0111/006	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	6 155 384 1	LABORATOIRE GLAXOSMITHKLINE	FR
REQUIP LP 2 mg, comprimé à libération prolongée	FR/H/0111/006	6 155 384 1	LABORATOIRE GLAXOSMITHKLINE	FR
Requip LP 4 mg comprimidos de libertação prolongada	FR/H/0111/008	5080809	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E	PT

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			QUÍMICOS, LDA	
Requip LP 4 mg comprimidos de libertação prolongada	FR/H/0111/008	5073929	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
REQUIP LP 4 mg, comprimé à libération prolongée	FR/H/0111/008	6 249 478 7	LABORATOIRE GLAXOSMITHKLINE	FR
REQUIP LP 4 mg, comprimé à libération prolongée	FR/H/0111/008	6 249 478 7	LABORATOIRE GLAXOSMITHKLINE	FR
Requip LP 8 mg comprimidos de libertação prolongada	FR/H/0111/009	5080817	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
Requip LP 8 mg comprimidos de libertação prolongada	FR/H/0111/009	5073937	BEECHAM PORTUGUESA, PRODUTOS FARMACÊUTICOS E QUÍMICOS, LDA	PT
REQUIP LP 8 mg, comprimé à libération prolongée	FR/H/0111/009	6 901 692 5	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 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 2 mg tablety s prodlouženým uvolňováním	not available	27/461/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 2 mg tablety s prodlouženým	not available	27/461/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
uvolňováním				
Requip Modutab 2 mg tablety s prodlouženým uvolňováním	not available	27/461/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 2 mg tablety s prodlouženým uvolňováním	not available	27/461/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
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 4 mg tablety s prodlouženým uvolňováním	not available	27/463/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 4 mg tablety s prodlouženým uvolňováním	not available	27/463/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
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 uvolnováním	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 8 mg tablety s prodlouženým uvolnováním	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip Modutab 8 mg tablety s prodlouženým	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
uvolňováním				
Requip Modutab 8 mg tablety s prodlouženým uvolňováním	not available	27/464/07-C	GLAXOSMITHKLINE (IRELAND) LIMITED	CZ
Requip XL 2 mg δισκία παρατεταμένης αποδέσμευσης.	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 - Retardtabletten	FR/H/0111/006	1-27360	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 2 mg comprimate cu eliberare prelungita	not available	13169/2020/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungita	not available	13169/2020/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/02	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/05	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/06	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 2 mg	not available	13169/2020/07	GLAXOSMITHKLINE	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
comprimate cu eliberare prelungită			TRADING SERVICES LIMITED	
Requip-Modutab 2 mg comprimate cu eliberare prelungită	not available	13169/2020/08	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
REQUIP-MODUTAB 2 mg comprimés à libération prolongée.	FR/H/0111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
REQUIP-MODUTAB 2 mg comprimés à libération prolongée.	FR/H/0111/006	2008060006	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg ilgstošas darbibas tabletes	not available	06-0261	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 2 mg	FR/H/0111/006	PA 1077/037/006	GLAXOSMITHKLINE	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
prolonged-release tablets.			(IRELAND) LIMITED	
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/0255/006	MA192/00106	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
REQUIP-MODUTAB 2 mg prolonged-release tablets.	FR/H/0255/006	MA192/00106	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
Requip-Modutab 2 mg retard filmtabletta	not available	OGYI-T-6200/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	HU
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/0111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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 EN ABREGE GLAXOSMITHKLINE	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	FR/H/0111/006	BE316531	GLAXOSMITHKLINE PHARMACEUTICALS EN	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
afgifte			ABREGE GLAXOSMITHKLINE	
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 TRADING SERVICES LIMITED	EE
Requip-Modutab 2 mg, tabletten met verlengde afgifte	FR/H/0111/006	RVG 100571	GLAXOSMITHKLINE B.V.	NL
Requip-Modutab 2 mg, tabletten met verlengde afgifte	FR/H/0111/006	RVG 100571	GLAXOSMITHKLINE B.V.	NL
Requip-Modutab 4 mg - Retardtabletten	FR/H/0111/008	1-27362	GLAXOSMITHKLINE PHARMA GMBH.	AT
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	13170/2020/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/02	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 4 mg comprimate cu eliberare prelungită	not available	13170/2020/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
REQUIP-MODUTAB 4 mg comprimés à libération prolongée.	FR/H/0111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
REQUIP-MODUTAB 4 mg comprimés à libération prolongée.	FR/H/0111/008	2008060008	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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-MODUTAB 4 mg comprimés à libération prolongée.	FR/H/0111/008	2008060008	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Requip-Modutab 4 mg ilgstošās darbības tabletes	not available	06-0263	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 4 mg ilgstošās darbības tabletes	not available	06-0263	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 4 mg ilgstošās darbības tabletes	not available	06-0263	GLAXOSMITHKLINE TRADING SERVICES LIMITED	LV
Requip-Modutab 4 mg ilgstošās darbības 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/0111/008	PA 1077/037/008	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REQUIP-MODUTAB 4 mg prolonged-release tablets.	FR/H/0255/008	MA192/00108	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
REQUIP-MODUTAB 4 mg prolonged-release tablets.	FR/H/0255/008	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/0111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/0111/008	69869.00.00	GLAXOSMITHKLINE GMBH & CO. KG	DE
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/0111/008	2008060008	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP-MODUTAB 4 mg Retardtabletten	FR/H/0111/008	2008060008	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/008	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-Modutab 4 mg tablete s podaljšanim sproščanjem	not available	H/02/01344/011	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SI
REQUIP-MODUTAB 4 mg tabletten met verlengde afgifte	FR/H/0111/008	BE316556	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
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 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 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	not available	532106	GLAXOSMITHKLINE	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
toimeainet prolongeeritult vabastavad tabletid.			TRADING SERVICES LIMITED	
Requip-Modutab 4 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532106	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 4 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532106	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 4 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532106	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 4 mg, tabletten met verlengde afgifte	FR/H/0111/008	RVG 100574	GLAXOSMITHKLINE B.V.	NL
Requip-Modutab 8 mg - Retardtabletten	FR/H/0111/009	1-27364	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 8 mg - Retardtabletten	FR/H/0111/009	1-27364	GLAXOSMITHKLINE PHARMA GMBH.	AT
Requip-Modutab 8 mg comprimate cu eliberare prelungita	not available	13171/2020/01	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungita	not available	13171/2020/04	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungita	not available	13171/2020/03	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
Requip-Modutab 8 mg comprimate cu eliberare prelungită	not available	13171/2020/02	GLAXOSMITHKLINE TRADING SERVICES LIMITED	RO
REQUIP-MODUTAB 8 mg comprimés à libération	FR/H/0111/009	BE316565	GLAXOSMITHKLINE PHARMACEUTICALS EN	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
prolongée.			ABREGE GLAXOSMITHKLINE	
REQUIP-MODUTAB 8 mg comprimés à libération prolongée.	FR/H/0111/009	2008060004	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
REQUIP-MODUTAB 8 mg comprimés à libération prolongée.	FR/H/0111/009	2008060004	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	LU
Requip-Modutab 8 mg ilgstošās darbības 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/0111/009	PA 1077/037/009	GLAXOSMITHKLINE (IRELAND) LIMITED	IE
REQUIP-MODUTAB 8 mg prolonged-release tablets.	FR/H/0255/009	MA192/00109	GLAXOSMITHKLINE (IRELAND) LIMITED	MT
REQUIP-MODUTAB 8 mg prolonged-release tablets.	FR/H/0255/009	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/0111/009	BE316565	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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 EN ABREGE GLAXOSMITHKLINE	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-MODUTAB 8 mg Retardtabletten	FR/H/0111/009	2008060004	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	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/0111/009	BE316565	GLAXOSMITHKLINE PHARMACEUTICALS EN ABREGE GLAXOSMITHKLINE	BE
REQUIP-MODUTAB 8 mg tablety s predĺženým uvoľňovaním	not available	27/0314/06-S	GLAXOSMITHKLINE TRADING SERVICES LIMITED	SK
Requip-Modutab 8 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 8 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 8 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 8 mg toimeainet prolongeeritult vabastavad tabletid.	not available	532006	GLAXOSMITHKLINE TRADING SERVICES LIMITED	EE
Requip-Modutab 8 mg, tabletten met verlengde afgifte	FR/H/0111/009	RVG 100576	GLAXOSMITHKLINE B.V.	NL
Requip-Modutab, 2 mg, tabletki o przedłużonym	FR/H/0255/006	14577	GLAXOSMITHKLINE TRADING SERVICES	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
uwalnianiu			LIMITED	
Requip-Modutab, 2 mg, tabletki o przedłużonym uwalnianiu	FR/H/0255/006	14577	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Requip-Modutab, 4 mg, tabletki o przedluzonym uwalnianiu	FR/H/0255/008	14579	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Requip-Modutab, 4 mg, tabletki o przedłużonym uwalnianiu	FR/H/0255/008	14579	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Requip-Modutab, 8 mg, tabletki o przedłużonym uwalnianiu	FR/H/0255/009	14580	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
Requip-Modutab, 8 mg, tabletki o przedłużonym uwalnianiu	FR/H/0255/009	14580	GLAXOSMITHKLINE TRADING SERVICES LIMITED	PL
REQUIP-PROLIB 2 mg comprimidos de liberación prolongada.	FR/H/0111/006	69.700	GLAXOSMITHKLINE, S.A.	ES
REQUIP-PROLIB 4 mg comprimidos de liberación prolongada.	FR/H/0111/008	69.701	GLAXOSMITHKLINE, S.A.	ES
REQUIP-PROLIB 4 mg comprimidos de liberación prolongada.	FR/H/0111/008	69.701	GLAXOSMITHKLINE, S.A.	ES
REQUIP-PROLIB 8 mg comprimidos de liberación prolongada.	FR/H/0111/009	69.702	GLAXOSMITHKLINE, S.A.	ES
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	XI
Ropilynz XL 4 mg prolonged-release	not available	PL 35507/0154	LUPIN HEALTHCARE (UK) LIMITED	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
tablets				
Ropilynz XL 8 mg prolonged-release tablets	not available	PL 35507/0155	LUPIN HEALTHCARE (UK) LIMITED	XI
Ropinirol - 1 A Pharma 3 mg Filmtabletten	not available	71161.00.00	1 A PHARMA GMBH	DE
Ropinirol - 1 A Pharma 3 mg Filmtabletten	not available	71161.00.00	1 A PHARMA GMBH	DE
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 - 1 A Pharma 4 mg Filmtabletten	not available	71162.00.00	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 "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol "Actavis", prolonged-release tablets	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK
Ropinirol "Actavis", prolonged-release	DK/H/2042/004	47982	ACTAVIS GROUP PTC EHF.	DK

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
tablets				
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 946 8 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 516 0 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 936 2 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 932 7 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 509 4 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 927 3 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 512 5 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 929 6 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 514 8 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 937 9 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 508 8 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 941 6 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 939 1 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 945 1 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 513 1 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 944 5 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 930 4 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25	DE/H/1052/001	34009 388 948 0 6	EG LABO LABORATOIRES	FR

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mg, comprimé pelliculé			EUROGENERICS	
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 938 5 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 507 1 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 931 0 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 947 4 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 942 2 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 935 6 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 943 9 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 388 933 3 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 510 2 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 515 4 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	34009 273 511 9 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,25 mg, comprimé pelliculé	DE/H/1052/001	3400938892673	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 520 8 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 951 1 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 958 6 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 950 5 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 526 6 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg,	DE/H/1052/002	34009 388 954 0 7	EG LABO LABORATOIRES	FR

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comprimé pelliculé			EUROGENERICS	
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 952 8 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 953 4 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 962 3 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 524 3 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 949 7 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 961 7 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 955 7 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 964 6 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 523 7 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 518 3 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 522 0 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 960 0 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 388 959 2 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	34009 273 521 4 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 0,5 mg, comprimé pelliculé	DE/H/1052/002	3400938895636	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 538 4 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 971 2 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg,	DE/H/1052/003	34009 417 729 6 2	EG LABO LABORATOIRES	FR

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comprimé pelliculé			EUROGENERICS	
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 724 4 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 969 8 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 967 5 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 537 8 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 968 1 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 727 3 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 966 9 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 535 5 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 539 0 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 534 9 9	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 388 970 6 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 731 0 5	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 536 1 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 732 7 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 273 540 9 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 735 6 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg, comprimé pelliculé	DE/H/1052/003	34009 417 736 2 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 1 mg,	DE/H/1052/003	3400938896527	EG LABO LABORATOIRES	FR

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comprimé pelliculé			EUROGENERICS	
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 273 545 0 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 273 542 1 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 978 7 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 972 9 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 975 8 6	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 714 9 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 720 9 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 722 1 4	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 716 1 3	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 718 4 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 977 0 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 976 4 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 712 6 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 417 723 8 2	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 273 546 7 0	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 974 1 8	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 388 973 5 7	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg,	DE/H/1052/004	34009 273 543 8 0	EG LABO LABORATOIRES	FR

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comprimé pelliculé			EUROGENERICS	
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 273 544 4 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	34009 273 547 3 1	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE EG 2 mg, comprimé pelliculé	DE/H/1052/004	3400927354151	EG LABO LABORATOIRES EUROGENERICS	FR
ROPINIROLE GSK LP 2 mg, comprimé à libération prolongée	FR/H/0255/006	6 907 203 0	LABORATOIRE GLAXOSMITHKLINE	FR
ROPINIROLE GSK LP 2 mg, comprimé à libération prolongée	FR/H/0255/006	6 907 203 0	LABORATOIRE GLAXOSMITHKLINE	FR
ROPINIROLE GSK LP 4 mg, comprimé à libération prolongée	FR/H/0255/008	6 241 363 2	LABORATOIRE GLAXOSMITHKLINE	FR
ROPINIROLE GSK LP 4 mg, comprimé à libération prolongée	FR/H/0255/008	6 241 363 2	LABORATOIRE GLAXOSMITHKLINE	FR
ROPINIROLE GSK LP 8 mg, comprimé à libération prolongée	FR/H/0255/009	6 677 737 3	LABORATOIRE GLAXOSMITHKLINE	FR
ROPINIROLE GSK LP 8 mg, comprimé à libération prolongée	FR/H/0255/009	6 677 737 3	LABORATOIRE GLAXOSMITHKLINE	FR
Ropinirol-neuraxpharm 3 mg Filmtabletten	not available	68843.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Ropinirol-neuraxpharm 3 mg Filmtabletten	not available	68843.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
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
Ropinirol-neuraxpharm 4 mg Filmtabletten	not available	68844.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Ropinirol-neuraxpharm 4	not available	68844.00.00	NEURAXPHARM	DE

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mg Filmtabletten			ARZNEIMITTEL GMBH	