12 April 2018
EMA/221551/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): quetiapine

Procedure No.: PSUSA/00002589/201707

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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
SEROQUEL 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/001	ASTRAZENECA UK LIMITED	SI
SEROQUEL 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/007	ASTRAZENECA UK LIMITED	SI
SEROQUEL 100 mg filmom obložene tablete	NL/H/0156/002	HR-H-461808155	ASTRAZENECA D.O.O.	HR
SEROQUEL 200 mg filmom obložene tablete	NL/H/0156/003	HR-H-204008507	ASTRAZENECA D.O.O.	HR
SEROQUEL 25 mg filmom obložene tablete	NL/H/0156/001	HR-H-619090976	ASTRAZENECA D.O.O.	HR
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3073681	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3073582	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3073988	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL 25 mg, comprimés pelliculés	NL/H/0156/001	2010010629	ASTRAZENECA S.A. / N.V.	LU
Seroquel® 100 mg tabletti, kalvopäällysteinen	NL/H/0156/002	14758	ASTRAZENECA OY	FI
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3073780	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3074085	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg film-coated tablets	NL/H/0156/001	PL 17901/0038	ASTRAZENECA UK LIMITED	UK
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3539087	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel® 25 mg tabletti, kalvopäällysteinen	NL/H/0156/001	14757	ASTRAZENECA OY	FI
Seroquel, 100 mg õhukese polümeerikattegatabletid	NL/H/0156/002	257199	ASTRAZENECA UK LIMITED	EE
Seroquel 100 mg plėvele dengtos tabletės	NL/H/0156/002	LT/1/99/0294/002	ASTRAZENECA UK LIMITED	LT
Seroquel® 200 mg tabletti, kalvopäällysteinen	NL/H/0156/003	14759	ASTRAZENECA 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
Seroquel 100 mg film-coated tablets	NL/H/0156/002	PL 17901/0039	ASTRAZENECA UK LIMITED	UK
Seroquel, 200 mg õhukese polümeerikattega tabletid	NL/H/0156/003	257099	ASTRAZENECA UK LIMITED	EE
SEROQUEL 100 mg, comprimés pelliculés	NL/H/0156/002	2010010630	ASTRAZENECA S.A. / N.V.	LU
Seroquel 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/003	17718	ASTRAZENECA UK LIMITED	CY
Seroquel 100 mg Filmtabletten	NL/H/0156/002	1-23461	ASTRAZENECA OSTERREICH GMBH	AT
Seroquel® 150 mg tabletti, kalvopäällysteinen	NL/H/0156/006	16558	ASTRAZENECA OY	FI
SEROQUEL 150 mg, comprimés pelliculés	NL/H/0156/006	2010010632	ASTRAZENECA S.A. / N.V.	LU
Seroquel 150 mg Filmtabletten	NL/H/0156/006	1-24318	ASTRAZENECA OSTERREICH GMBH	AT
Seroquel® 200 mg Filmtabletten	NL/H/0156/003	47291.02.00	ASTRAZENECA GMBH	DE
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3074887	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075488	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3074986	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075181	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075389	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075082	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg Filmtabletten	NL/H/0156/001	1-23460	ASTRAZENECA OSTERREICH GMBH	AT
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075587	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/001	17716	ASTRAZENECA UK LIMITED	CY

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
Seroquel 200 mg comprimidos revestidos por película	NL/H/0156/003	3075280	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/002	17717	ASTRAZENECA UK LIMITED	CY
Seroquel® 25 mg Filmtabletten	NL/H/0156/001	47291.00.00	ASTRAZENECA GMBH	DE
Seroquel 25 mg plėvele dengtos tabletės	NL/H/0156/001	LT/1/99/0294/001	ASTRAZENECA UK LIMITED	LT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3734084	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733086	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733987	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733284	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733383	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733482	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL 200 mg, comprimés pelliculés	NL/H/0156/003	2010010633	ASTRAZENECA S.A. / N.V.	LU
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733789	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733185	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg film-coated tablets	NL/H/0156/003	PL 17901/0040	ASTRAZENECA UK LIMITED	UK
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733581	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg plėvele dengtos tabletės	NL/H/0156/003	LT/1/99/0294/004	ASTRAZENECA UK LIMITED	LT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733680	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 200 mg Filmtabletten	NL/H/0156/003	1-23463	ASTRAZENECA OSTERREICH 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
Seroquel® 100 mg Filmtabletten	NL/H/0156/002	47291.01.00	ASTRAZENECA GMBH	DE
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074382	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 150 mg comprimidos revestidos por película	NL/H/0156/006	3733888	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074580	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074481	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074283	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074788	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074184	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	3074689	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL 300 mg, comprimés pelliculés	NL/H/0156/007	2010010631	ASTRAZENECA S.A. / N.V.	LU
Seroquel 300 mg film-coated tablets	NL/H/0156/007	PL 17901/0088	ASTRAZENECA UK LIMITED	UK
Seroquel® 300 mg Filmtabletten	NL/H/0156/007	47291.04.00	ASTRAZENECA GMBH	DE
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734985	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg Filmtabletten	NL/H/0156/007	1-24319	ASTRAZENECA OSTERREICH GMBH	AT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3735081	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734688	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734480	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734381	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos	NL/H/0156/007	3735180	ASTRAZENECA PRODUTOS	PT

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revestidos por película			FARMACEUTICOS LDA	
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3735289	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734886	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734282	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734787	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 300 mg comprimidos revestidos por película	NL/H/0156/007	3734589	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel® 300 mg tabletti, kalvopäällysteinen	NL/H/0156/007	16559	ASTRAZENECA OY	FI
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C 32944112	ASTRAZENECA S.P.A.	IT
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C 32944011	ASTRAZENECA S.P.A.	IT
Seroquel 150 mg filmdragerade tabletter	NL/H/0156/006	18964	ASTRAZENECA AB	SE
Seroquel®-100, filmomhulde tabletten	NL/H/0156/002	RVG 20827	ASTRAZENECA BV	NL
SEROQUEL 200 mg, comprimés pelliculés	NL/H/0156/003	BE210375	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	32944086	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg, filmomhulde tabletten	NL/H/0156/003	BE210375	ASTRAZENECA S.A. / N.V.	BE
Seroquel 200 mg film-coated tablets	NL/H/0156/003	PA 970/18/3	ASTRAZENECA UK LIMITED	IE
SEROQUEL 100 mg, filmomhulde tabletten	NL/H/0156/002	BE210366	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	32944074	ASTRAZENECA S.P.A.	IT
Seroquel®-25, filmomhulde tabletten	NL/H/0156/001	RVG 20826	ASTRAZENECA BV	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
SEROQUEL 100 mg, comprimés pelliculés	NL/H/0156/002	BE210366	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C 32944035	ASTRAZENECA S.P.A.	IT
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C 32944023	ASTRAZENECA S.P.A.	IT
Seroquel 100 mg film-coated tablets	NL/H/0156/002	PA 970/18/2	ASTRAZENECA UK LIMITED	IE
Seroquel 200 mg film-coated tablets	NL/H/0156/003	MA044/00103	ASTRAZENECA UK LIMITED	MT
SEROQUEL 25 mg, comprimés pelliculés	NL/H/0156/001	BE210357	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 150 mg, filmomhulde tabletten	NL/H/0156/006	BE228891	ASTRAZENECA S.A. / N.V.	BE
Seroquel 100 mg filmdragerade tabletter	NL/H/0156/002	18959	ASTRAZENECA AB	SE
SEROQUEL 150 mg, comprimés pelliculés	NL/H/0156/006	BE228891	ASTRAZENECA S.A. / N.V.	BE
Seroquel 25 mg film-coated tablets	NL/H/0156/001	PA 970/18/1	ASTRAZENECA UK LIMITED	IE
Seroquel®-200, filmomhulde tabletten	NL/H/0156/003	RVG 20828	ASTRAZENECA BV	NL
Seroquel®-150, filmomhulde tabletten	NL/H/0156/006	RVG 25602	ASTRAZENECA BV	NL
Seroquel 25 mg film-coated tablets	NL/H/0156/001	MA044/00101	ASTRAZENECA UK LIMITED	MT
Seroquel 25 mg filmdragerade tabletter	NL/H/0156/001	18958	ASTRAZENECA AB	SE
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C 32944050	ASTRAZENECA S.P.A.	IT
Seroquel 100 mg film-coated tablets	NL/H/0156/002	MA044/00102	ASTRAZENECA UK LIMITED	MT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C 32944047	ASTRAZENECA S.P.A.	IT
Seroquel®-300, filmomhulde tabletten	NL/H/0156/007	RVG 25603	ASTRAZENECA BV	NL
SEROQUEL 25 mg, filmomhulde	NL/H/0156/001	BE210357	ASTRAZENECA S.A. / N.V.	BE

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tabletten				
SEROQUEL 300 mg, filmomhulde tabletten	NL/H/0156/007	BE228907	ASTRAZENECA S.A. / N.V.	BE
Seroquel 300 mg film-coated tablets	NL/H/0156/007	PA 970/18/7	ASTRAZENECA UK LIMITED	IE
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C 032944100	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C 032944098	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg filmdragerade tabletter	NL/H/0156/003	18960	ASTRAZENECA AB	SE
SEROQUEL 300 mg, comprimés pelliculés	NL/H/0156/007	BE228907	ASTRAZENECA S.A. / N.V.	BE
Seroquel 300 mg filmdragerade tabletter	NL/H/0156/007	18961	ASTRAZENECA AB	SE
Seroquel 25 mg comprimidos recubiertos con película	NL/H/0156/001	63.054	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel 100 mg comprimidos recubiertos con película	NL/H/0156/002	63.055	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel 200 mg comprimidos recubiertos con película	NL/H/0156/003	63.056	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel, filmovertrukne tabletter	NL/H/0156/003	32336	ASTRAZENECA A/S	DK
Seroquel, filmovertrukne tabletter	NL/H/0156/002	32335	ASTRAZENECA A/S	DK
Seroquel 300 mg comprimidos recubiertos con película	NL/H/0156/007	64.436	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel, filmovertrukne tabletter	NL/H/0156/001	32334	ASTRAZENECA A/S	DK
Seroquel, filmovertrukne tabletter	NL/H/0156/007	32600	ASTRAZENECA A/S	DK
Seroquel, filmovertrukne tabletter	NL/H/0156/006	32599	ASTRAZENECA A/S	DK
Seroquel 100 mg filmdrasjerte tabletter	NL/H/0156/002	02-1436	ASTRAZENECA AS	NO
Seroquel 200 mg filmdrasjerte tabletter	NL/H/0156/003	02-1438	ASTRAZENECA AS	NO
Seroquel 300 mg filmdrasjerte tabletter	NL/H/0156/007	02-1439	ASTRAZENECA AS	NO
Seroquel 150 mg filmdrasjerte	NL/H/0156/006	02-1437	ASTRAZENECA AS	NO

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tabletter				
Seroquel 25 mg filmdrasjerte tabletter	NL/H/0156/001	02-1435	ASTRAZENECA AS	NO
Seroquel 100 mg filmuhúðaðar töflur	NL/H/0156/002	IS/1/03/002/02	ASTRAZENECA AB	IS
Seroquel 150 mg filmuhúðaðar töflur	NL/H/0156/006	IS/1/03/002/03	ASTRAZENECA AB	IS
Seroquel 200 mg filmuhúðaðar töflur	NL/H/0156/003	IS/1/03/002/04	ASTRAZENECA AB	IS
Seroquel 25 mg filmuhúðaðar töflur	NL/H/0156/001	IS/1/03/002/01	ASTRAZENECA AB	IS
Seroquel 300 mg filmuhúðaðar töflur	NL/H/0156/007	IS/1/03/002/05	ASTRAZENECA AB	IS
Seroquel 200 mg plėvele dengtos tabletės	NL/H/0156/003	LT/1/99/0294/005	ASTRAZENECA UK LIMITED	LT
Seroquel 100 mg plevele dengtos tabletes	NL/H/0156/002	LT/1/99/0294/003	ASTRAZENECA UK LIMITED	LT
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/010	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/009	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/058	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/011	ASTRAZENECA UK LIMITED	SI
Seroquel 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/002	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/012	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/008	ASTRAZENECA UK LIMITED	SI
Seroquel 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/005	ASTRAZENECA UK LIMITED	SI
Seroquel 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/003	ASTRAZENECA UK LIMITED	SI
Seroquel 25 mg filmsko obložene	NL/H/0156/001	H/99/01407/004	ASTRAZENECA UK LIMITED	SI

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tablete				
Seroquel 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/057	ASTRAZENECA UK LIMITED	SI
Seroquel 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/056	ASTRAZENECA UK LIMITED	SI
Seroquel 100 mg comprimidos revestidos por película	NL/H/0156/002	5575980	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel 25 mg comprimidos revestidos por película	NL/H/0156/001	3073889	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroque1 100 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/002	54573/15-06-2017	ASTRAZENECA S.A	GR
Seroque1 25 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/001	54572/15-06-2017	ASTRAZENECA S.A	GR
Seroque1 300 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/007	54578/15-06-2017	ASTRAZENECA S.A	GR
Seroquel 200 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/003	54574/15-06-2017	ASTRAZENECA S.A	GR
Seroque1150 mg επικαλυμμένα με λεπτό υμένιο δισκία	NL/H/0156/006	54577/15-06-2017	ASTRAZENECA S.A	GR
SEROQUEL Starter Pack, filmomhulde tabletten (combinatieverpakking)	NL/H/0156/004	BE210341	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL Starter Pack, comprimés pelliculés (emballage combiné)	NL/H/0156/004	BE210341	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C. 032944213	ASTRAZENECA S.P.A.	IT
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C. 032944249	ASTRAZENECA S.P.A.	IT
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C. 032944237	ASTRAZENECA S.P.A.	IT
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C. 032944252	ASTRAZENECA S.P.A.	IT
SEROQUEL 100 mg compresse rivestite con film	NL/H/0156/002	A.I.C. 032944225	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse	NL/H/0156/006	A.I.C. 032944314	ASTRAZENECA S.P.A.	IT

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rivestite con film				
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944338	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944290	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944326	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944288	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944302	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944264	ASTRAZENECA S.P.A.	IT
SEROQUEL 150 mg compresse rivestite con film	NL/H/0156/006	A.I.C. 032944276	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C. 032944389	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C. 032944340	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C. 032944365	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C. 032944353	ASTRAZENECA S.P.A.	IT
SEROQUEL 200 mg compresse rivestite con film	NL/H/0156/003	A.I.C. 032944377	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944427	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944415	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944403	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944391	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944441	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944454	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944466	ASTRAZENECA S.P.A.	IT
SEROQUEL 300 mg compresse rivestite con film	NL/H/0156/007	A.I.C. 032944439	ASTRAZENECA S.P.A.	IT
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C. 032944201	ASTRAZENECA S.P.A.	IT
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C. 032944175	ASTRAZENECA S.P.A.	IT
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C. 032944187	ASTRAZENECA S.P.A.	IT
SEROQUEL 25 mg compresse rivestite con film	NL/H/0156/001	A.I.C. 032944199	ASTRAZENECA S.P.A.	IT
Seroquel Embalagem de início de 3 Dias (25 mg) + (100 mg) comprimidos revestidos por película	NL/H/0156/004	3075686	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL 4-day Starter Pack, comprimés pelliculés	NL/H/0156/005	BE217305	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL 4-day Starter Pack, filmomhulde tabletten	NL/H/0156/005	BE217305	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL Starter Pack, comprimés pelliculés	NL/H/0156/004	2006088994	ASTRAZENECA S.A. / N.V.	LU
Seroquel, filmovertrukne tabletter	NL/H/0156/005	32337	ASTRAZENECA A/S	DK
Seroquel Confezione Starter 4-Giorni	NL/H/0156/005	032944062	ASTRAZENECA S.P.A.	IT
SEROQUEL 4-day Starter Pack, comprimés pelliculés	NL/H/0156/005	2006088995	ASTRAZENECA S.A. / N.V.	LU
Seroquel Embalagem de início de 4 Dias (25 mg) + (100 mg) + (200 mg) comprimidos revestidos por película	NL/H/0156/005	3233681	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel Confezione Starter 3-Giorni (confezione combinata)	NL/H/0156/004	A.I.C. 032944478	ASTRAZENECA BV	IT
SEROQUEL 300 mg compresse	NL/H/0156/007	A.I.C. 032944439	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rivestite con film				
SEROQUEL 100 mg filmsko obložene tablete	NL/H/0156/002	H/99/01407/013	ASTRAZENECA UK LIMITED	SI
SEROQUEL μικτή συσκευασία έναρξης θεραπείας 3 ημερών (3 Day Starterpack)	NL/H/0156/004	54575/15-06-2017	ASTRAZENECA S.A	GR
SEROQUEL μικτή συσκευασία έναρξης θεραπείας 4 ημερών (4 Day Starterpack)	NL/H/0156/005	54576/15-06-2017	ASTRAZENECA S.A	GR
SEROQUEL 25 mg filmsko obložene tablete	NL/H/0156/001	H/99/01407/006	ASTRAZENECA UK LIMITED	SI
Q Mind 150 mg comprimate filmate	DE/H/2610/004/DC	8352/2015/01-04	TORRENT PHARMA SRL	RO
Quetiapin Heumann 50 mg Filmtabletten	DE/H/2610/002	79951.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Quetiapin Heumann 150 mg Filmtabletten	DE/H/2610/004/DC	79953.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Quetiapin Heumann 400 mg Filmtabletten	DE/H/2610/007/DC	79956.00.00	HEUMANN PHARMA GMBH & CO. GENERICA KG	DE
Alzen SR 150 mg comprimidos de libertação prolongada	PT/H/1439/002	5176953	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 150 mg comprimidos de libertação prolongada	PT/H/1439/002	5176938	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 150 mg comprimidos de libertação prolongada	PT/H/1439/002	5176961	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 150 mg comprimidos de libertação prolongada	PT/H/1439/002	5176979	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 150 mg comprimidos de libertação prolongada	PT/H/1439/002	5176946	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 200 mg comprimidos de libertação prolongada	PT/H/1439/003	5090832	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 200 mg comprimidos de libertação prolongada	PT/H/1439/003	5090857	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 400 mg comprimidos de libertação prolongada	PT/H/1439/005	5090964	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 300 mg comprimidos de	PT/H/1439/004	5090915	ASTRAZENECA PRODUTOS	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
libertação prolongada			FARMACEUTICOS LDA	
Alzen SR 200 mg comprimidos de libertação prolongada	PT/H/1439/003	5090865	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 200 mg comprimidos de libertação prolongada	PT/H/1439/003	5090873	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 300 mg comprimidos de libertação prolongada	PT/H/1439/004	5090949	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 300 mg comprimidos de libertação prolongada	PT/H/1439/004	5090907	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 300 mg comprimidos de libertação prolongada	PT/H/1439/004	5090923	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 200 mg comprimidos de libertação prolongada	PT/H/1439/003	5090840	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 400 mg comprimidos de libertação prolongada	PT/H/1439/005	5090972	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 400 mg comprimidos de libertação prolongada	PT/H/1439/005	5091012	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 50 mg comprimidos de libertação prolongada	PT/H/1439/001	5090808	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 50 mg comprimidos de libertação prolongada	PT/H/1439/001	5090824	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 50 mg comprimidos de libertação prolongada	PT/H/1439/001	5090774	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 50 mg comprimidos de libertação prolongada	PT/H/1439/001	5090816	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 400 mg comprimidos de libertação prolongada	PT/H/1439/005	5090956	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 50 mg comprimidos de libertação prolongada	PT/H/1439/001	5090766	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 400 mg comprimidos de libertação prolongada	PT/H/1439/005	5091004	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Alzen SR 300 mg comprimidos de libertação prolongada	PT/H/1439/004	5090931	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
XeroquelÒ LP 300 mg, comprimé à libération prolongée	NL/H/1983/004	NL 39 380	ASTRAZENECA S.A.S.	FR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
XeroquelÒ LP 400 mg, comprimé à libération prolongée	NL/H/1983/005	NL 39 381	ASTRAZENECA S.A.S.	FR
XeroquelÒ LP 50 mg, comprimé à libération prolongée	NL/H/1983/001	NL 39 377	ASTRAZENECA S.A.S.	FR
Quetiapine XR AstraZeneca 150 mg, tabletten met verlengde afgifte	NL/H/1983/002	RVG 106978	ASTRAZENECA BV	NL
Quetiapine XR AstraZeneca 400 mg, tabletten met verlengde afgifte	NL/H/1983/005	RVG 106981	ASTRAZENECA BV	NL
Quetiapine XR AstraZeneca 50 mg, tabletten met verlengde afgifte	NL/H/1983/001	RVG 106971	ASTRAZENECA BV	NL
Quetiapine XR AstraZeneca 200 mg, tabletten met verlengde afgifte	NL/H/1983/003	RVG 106979	ASTRAZENECA BV	NL
Quetiapine XR AstraZeneca 300 mg, tabletten met verlengde afgifte	NL/H/1983/004	RVG 106980	ASTRAZENECA BV	NL
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/051	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/031	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/046	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/047	ASTRAZENECA UK LIMITED	SI
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/037	ASTRAZENECA UK LIMITED	SI
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/036	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/041	ASTRAZENECA UK LIMITED	SI
Seroquel XR 200 mg tablety s predĺženým uvoľňovaním	NL/H/0156/009	68/0231/07-S	ASTRAZENECA UK LIMITED	SK
Seroquel XR 50 mg tablete s produljenim oslobađanjem	NL/H/0156/008	HR-H-955743091	ASTRAZENECA D.O.O.	HR
Seroquel XR 200 mg tablete s produljenim oslobađanjem	NL/H/0156/009	HR-H-607016913	ASTRAZENECA D.O.O.	HR
Seroquel XR 300 mg tablete s	NL/H/0156/010	HR-H-408184728	ASTRAZENECA BV	HR

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
produljenim oslobađanjem				
Seroquel XR 400 mg tablete s produljenim oslobađanjem	NL/H/0156/011	HR-H-789589749	ASTRAZENECA D.O.O.	HR
Seroquel XR 150 mg tablete s produljenim oslobađanjem	NL/H/0156/012	HR-H-202528018	ASTRAZENECA D.O.O.	HR
SEROQUEL XR 150 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/012	20699	ASTRAZENECA UK LIMITED	CY
Seroquel Prolong 150 mg depottabletti	NL/H/0156/012	24948	ASTRAZENECA OY	FI
Seroquel Prolong 400 mg depottabletti	NL/H/0156/011	24067	ASTRAZENECA OY	FI
Seroquel XR, 300 mg toimeainet prolongeeritult vabastavad tabletid	NL/H/0156/010	565007	ASTRAZENECA UK LIMITED	EE
Seroquel SR 200 mg comprimidos de libertação prolongada	NL/H/0156/009	5085238	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel SR 200 mg comprimidos de libertação prolongada	NL/H/0156/009	5085220	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel Prolong 300 mg depottabletti	NL/H/0156/010	24066	ASTRAZENECA OY	FI
SEROQUEL XR 300 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/010	20358	ASTRAZENECA UK LIMITED	CY
Seroquel XR 300 mg tablety s predĺženým uvoľňovaním	NL/H/0156/010	68/0232/07-S	ASTRAZENECA UK LIMITED	SK
Seroquel SR 150 mg comprimidos de libertação prolongada	NL/H/0156/012	5168455	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel SR 150 mg comprimidos de libertação prolongada	NL/H/0156/012	5168463	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL XR 400 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/011	20359	ASTRAZENECA UK LIMITED	CY
Seroquel SR 300 mg comprimidos de libertação prolongada	NL/H/0156/010	5085246	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel SR 300 mg comprimidos de libertação prolongada	NL/H/0156/010	5085253	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel XR, 400 mg toimeainet prolongeeritult vabastavad tabletid	NL/H/0156/011	565207	ASTRAZENECA UK LIMITED	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
Seroquel Prolong 50 mg depottabletti	NL/H/0156/008	24064	ASTRAZENECA OY	FI
Seroquel SR 400 mg comprimidos de libertação prolongada	NL/H/0156/011	5085261	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel SR 50 mg comprimidos de libertação prolongada	NL/H/0156/008	5178165	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel XR 50 mg tablety s predĺženým uvoľňovaním	NL/H/0156/008	68/0230/07-S	ASTRAZENECA UK LIMITED	SK
Seroquel SR 50 mg comprimidos de libertação prolongada	NL/H/0156/008	5085212	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
SEROQUEL XR 50 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/008	20356	ASTRAZENECA UK LIMITED	CY
SEROQUEL XR 150 mg, tabletten met verlengde afgifte	NL/H/0156/012	BE331947	ASTRAZENECA S.A. / N.V.	BE
Seroquel XR 150 mg Retardtabletten	NL/H/0156/012	1-28377	ASTRAZENECA OSTERREICH GMBH	AT
Seroquel Prolong® 150 mg Retardtabletten	NL/H/0156/012	73290.00.00	ASTRAZENECA GMBH	DE
SEROQUEL XR 150 mg, comprimé à libération prolongée	NL/H/0156/012	BE331947	ASTRAZENECA S.A. / N.V.	BE
Seroquel XR 300 mg Retardtabletten	NL/H/0156/010	1-27422	ASTRAZENECA OSTERREICH GMBH	AT
SEROQUEL XR 300 mg, comprimé à libération prolongée	NL/H/0156/010	BE314264	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL XR 400 mg, tabletten met verlengde afgifte	NL/H/0156/011	BE314282	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL XR 400 mg, comprimé à libération prolongée	NL/H/0156/011	BE314282	ASTRAZENECA S.A. / N.V.	BE
Seroquel XR 400 mg Retardtabletten	NL/H/0156/011	1-27419	ASTRAZENECA OSTERREICH GMBH	AT
SEROQUEL XR 300 mg, tabletten met verlengde afgifte	NL/H/0156/010	BE314264	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL XR 50 mg, comprimé à libération prolongée	NL/H/0156/008	BE314221	ASTRAZENECA S.A. / N.V.	BE
Seroquel XR 50 mg Retardtabletten	NL/H/0156/008	1-27420	ASTRAZENECA OSTERREICH	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
			GMBH	
SEROQUEL XR 50 mg, tabletten met verlengde afgifte	NL/H/0156/008	BE314221	ASTRAZENECA S.A. / N.V.	BE
Seroquel Prolong® 50 mg Retardtabletten	NL/H/0156/008	70561.00.00	ASTRAZENECA GMBH	DE
Seroquel Prolong® 300 mg Retardtabletten	NL/H/0156/010	70563.00.00	ASTRAZENECA GMBH	DE
Seroquel Prolong® 400 mg Retardtabletten	NL/H/0156/011	70564.00.00	ASTRAZENECA GMBH	DE
Seroquel XR 50 mg pailginto atpalaidavimo tabletės	NL/H/0156/008	LT/1/08/0964/001	ASTRAZENECA UK LIMITED	LT
Seroquel 50 mg compresse a rilascio prolungato	not available	32944124	ASTRAZENECA S.P.A.	IT
Seroquel XR 50 mg, comprimés à libération prolongée	NL/H/0156/008	2008040019	ASTRAZENECA S.A. / N.V.	LU
Seroquel XR 50 mg prolonged-release tablets	NL/H/0156/008	PA 970/18/8	ASTRAZENECA UK LIMITED	IE
Seroquel XR 150 mg, comprimés à libération prolongée	NL/H/0156/012	2009030009	ASTRAZENECA S.A. / N.V.	LU
Seroquel XR 300 mg, comprimés à libération prolongée	NL/H/0156/010	2008040021	ASTRAZENECA S.A. / N.V.	LU
Seroquel XR 400 mg pailginto atpalaidavimo tabletės	NL/H/0156/011	LT/1/08/0964/004	ASTRAZENECA UK LIMITED	LT
Seroquel XR 150 mg pailginto atpalaidavimo tabletės	NL/H/0156/012	LT/1/08/0964/006	ASTRAZENECA UK LIMITED	LT
Seroquel XR 300 mg pailginto atpalaidavimo tabletės	NL/H/0156/010	LT/1/08/0964/003	ASTRAZENECA UK LIMITED	LT
Seroquel XR 400 mg, comprimés à libération prolongée	NL/H/0156/011	2008040022	ASTRAZENECA S.A. / N.V.	LU
Seroquel 150 mg compresse a rilascio prolungato	NL/H/0156/012	A.I.C. 032944163	ASTRAZENECA S.P.A.	IT
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C 32944151	ASTRAZENECA S.P.A.	IT
Seroquel XR 300 mg prolonged-release tablets	NL/H/0156/010	PA 970/18/10	ASTRAZENECA UK LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C 32944148	ASTRAZENECA S.P.A.	IT
Seroquel XR 400 mg prolonged-release tablets	NL/H/0156/011	PA 970/18/11	ASTRAZENECA UK LIMITED	IE
Seroquel XR 150 mg prolonged-release tablets	NL/H/0156/012	PA 970/18/12	ASTRAZENECA UK LIMITED	IE
Seroquel XR® 400 mg, tabletten met verlengde afgifte	NL/H/0156/011	RVG 34628	ASTRAZENECA BV	NL
Seroquel Depot 50 mg depottabletter	NL/H/0156/008	25986	ASTRAZENECA AB	SE
Seroquel Depot 400 mg depottabletter	NL/H/0156/011	25989	ASTRAZENECA AB	SE
Seroquel XR® 50 mg, tabletten met verlengde afgifte	NL/H/0156/008	RVG 34625	ASTRAZENECA BV	NL
Seroquel XL 400 mg prolonged-release tablets	NL/H/0156/011	PL 17901/0252	ASTRAZENECA UK LIMITED	UK
Seroquel XL 50 mg prolonged-released tablets	NL/H/0156/008	PL 17901/0249	ASTRAZENECA UK LIMITED	UK
Seroquel XR® 300 mg, tabletten met verlengde afgifte	NL/H/0156/010	RVG 34627	ASTRAZENECA BV	NL
Seroquel Depot 150 mg depottabletter	NL/H/0156/012	26908	ASTRAZENECA AB	SE
Seroquel XL 300 mg prolonged-release tablets	NL/H/0156/010	PL 17901/0251	ASTRAZENECA UK LIMITED	UK
Seroquel XL 150 mg prolonged-release tablets	NL/H/0156/012	17901/0259	ASTRAZENECA UK LIMITED	UK
Seroquel XR® 150 mg, tabletten met verlengde afgifte	NL/H/0156/012	RVG 102408	ASTRAZENECA BV	NL
Seroquel XR® 200 mg, tabletten met verlengde afgifte	NL/H/0156/009	RVG 34626	ASTRAZENECA BV	NL
Seroquel Depot 300 mg depottabletter	NL/H/0156/010	25988	ASTRAZENECA AB	SE
Seroquel Prolong 400 mg comprimidos de liberación prolongada	NL/H/0156/011	69.648	ASTRAZENECA FARMACÉUTICA SPAIN, 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
Seroquel XR 50 mg retard tabletta	NL/H/0156/008	OGYI-T-5863/02	ASTRAZENECA KFT.	HU
Seroquel Prolong 50 mg comprimidos de liberación prolongada	NL/H/0156/008	69.645	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel XR 300 mg retard tabletta	NL/H/0156/010	OGYI-T-5863/04	ASTRAZENECA KFT.	HU
Seroquel Prolong 150 mg comprimidos de liberación prolongada	NL/H/0156/012	70.859	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel Prolong 300 mg comprimidos de liberación prolongada	NL/H/0156/010	69.647	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel XR 400 mg retard tabletta	NL/H/0156/011	OGYI-T-5863/05	ASTRAZENECA KFT.	HU
Seroquel XR 150 mg retard tabletta	NL/H/0156/012	OGYI-T-5863/08	ASTRAZENECA KFT.	HU
Seroquel XR 150 mg retard tabletta	NL/H/0156/012	OGYI-T-5863/09	ASTRAZENECA KFT.	HU
Seroquel Prolong, depottabletter	NL/H/0156/010	42033	ASTRAZENECA A/S	DK
Seroquel Prolong, depottabletter	NL/H/0156/011	42034	ASTRAZENECA A/S	DK
Seroquel Prolong, depottabletter	NL/H/0156/008	42031	ASTRAZENECA A/S	DK
Seroquel Prolong, depottabletter	NL/H/0156/012	43173	ASTRAZENECA A/S	DK
Seroquel Depot 150 mg depottabletter	NL/H/0156/012	08-5880	ASTRAZENECA AS	NO
Seroquel Depot 200 mg depottabletter	NL/H/0156/009	07-5215	ASTRAZENECA AS	NO
Seroquel Depot 300 mg depottabletter	NL/H/0156/010	07-5216	ASTRAZENECA AS	NO
Seroquel Depot 400 mg depottabletter	NL/H/0156/011	07-5217	ASTRAZENECA AS	NO
Seroquel Depot 50 mg depottabletter	NL/H/0156/008	07-5214	ASTRAZENECA AS	NO
Seroquel Prolong 150 mg forðatöflur	NL/H/0156/012	IS/1/08/081/01	ASTRAZENECA A/S	IS
Seroquel Prolong 200 mg forðatöflur	NL/H/0156/009	IS/1/08/003/02	ASTRAZENECA A/S	IS
Seroquel Prolong 300 mg forðatöflur	NL/H/0156/010	IS/1/08/003/03	ASTRAZENECA A/S	IS
Seroquel Prolong 400 mg	NL/H/0156/011	IS/1/08/003/04	ASTRAZENECA A/S	IS

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
forðatöflur				
Seroquel Prolong 50 mg forðatöflur	NL/H/0156/008	IS/1/08/003/01	ASTRAZENECA A/S	IS
Seroquel XR 50 mg retard tabletta	NL/H/0156/008	OGYI-T-5863/10	ASTRAZENECA KFT.	HU
Seroquel XR 400 mg retard tabletta	NL/H/0156/011	OGYI-T-5863/16	ASTRAZENECA KFT.	HU
Seroquel XR 200 mg retard tabletta	NL/H/0156/009	OGYI-T-5863/11	ASTRAZENECA KFT.	HU
Seroquel XR 300 mg retard tabletta	NL/H/0156/010	OGYI-T-5863/15	ASTRAZENECA KFT.	HU
Seroquel XR 400 mg retard tabletta	NL/H/0156/011	OGYI-T-5863/13	ASTRAZENECA KFT.	HU
Seroquel XR 200 mg retard tabletta	NL/H/0156/009	OGYI-T-5863/14	ASTRAZENECA KFT.	HU
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/01	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/03	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/02	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/04	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/05	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/06	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg comprimate cu eliberare prelungita	NL/H/0156/012	9529/2016/07	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/03	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/02	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/01	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/07	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/04	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/05	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu	NL/H/0156/010	9531/2016/04	ASTRAZENECA UK 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
eliberare prelungita				
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/05	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/01	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/06	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/02	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/07	ASTRAZENECA UK LIMITED	RO
Seroquel XR 300 mg comprimate cu eliberare prelungita	NL/H/0156/010	9531/2016/03	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/04	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/01	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/03	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/06	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/05	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/02	ASTRAZENECA UK LIMITED	RO
Seroquel XR 400 mg comprimate cu eliberare prelungita	NL/H/0156/011	9532/2016/07	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/07	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/05	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/04	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/01	ASTRAZENECA UK 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
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/06	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/03	ASTRAZENECA UK LIMITED	RO
Seroquel XR 50 mg comprimate cu eliberare prelungita	NL/H/0156/008	9528/2016/02	ASTRAZENECA UK LIMITED	RO
Seroquel XR 200 mg comprimate cu eliberare prelungita	NL/H/0156/009	9530/2016/06	ASTRAZENECA UK LIMITED	RO
Seroquel XR 150 mg pailginto atpalaidavimo tabletės	NL/H/0156/012	LT/1/08/0964/005	ASTRAZENECA UK LIMITED	LT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	32944136	ASTRAZENECA S.P.A.	IT
Seroquel XR 200 mg, comprimés à libération prolongée	NL/H/0156/009	2008040020	ASTRAZENECA S.A. / N.V.	LU
Seroquel XR 300 mg retard tabletta	NL/H/0156/010	OGYI-T-5863/12	ASTRAZENECA KFT.	HU
SEROQUEL XR 200 mg, comprimé à libération prolongée	NL/H/0156/009	BE314246	ASTRAZENECA S.A. / N.V.	BE
SEROQUEL XR 200 mg, tabletten met verlengde afgifte	NL/H/0156/009	BE314246	ASTRAZENECA S.A. / N.V.	BE
Seroquel Prolong, depottabletter	NL/H/0156/009	42032	ASTRAZENECA A/S	DK
Seroquel Prolong 200 mg depottabletti	NL/H/0156/009	24065	ASTRAZENECA OY	FI
Seroquel Prolong® 200 mg Retardtabletten Wirkstoff: Quetiapin	NL/H/0156/009	70562.00.00	ASTRAZENECA GMBH	DE
Seroquel XR 200 mg prolonged-release tablets	NL/H/0156/009	PA 970/18/9	ASTRAZENECA UK LIMITED	IE
Seroquel Depot 200 mg depottabletter	NL/H/0156/009	25987	ASTRAZENECA AB	SE
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/038	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/043	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/045	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s	NL/H/0156/010	H/99/01407/066	ASTRAZENECA UK 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
podaljšanim sproščanjem				
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/062	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/064	ASTRAZENECA UK LIMITED	SI
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/039	ASTRAZENECA UK LIMITED	SI
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/040	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/063	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/044	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/048	ASTRAZENECA UK LIMITED	SI
Seroquel SR 150 mg tablete s podaljšanim sproščanjem	NL/H/0156/012	H/99/01407/061	ASTRAZENECA UK LIMITED	SI
Seroquel SR 200 mg tablete s podaljšanim sproščanjem	NL/H/0156/009	H/99/01407/042	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/049	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/050	ASTRAZENECA UK LIMITED	SI
Seroquel SR 300 mg tablete s podaljšanim sproščanjem	NL/H/0156/010	H/99/01407/065	ASTRAZENECA UK LIMITED	SI
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/052	ASTRAZENECA UK LIMITED	SI
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/068	ASTRAZENECA UK LIMITED	SI
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/067	ASTRAZENECA UK LIMITED	SI
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/054	ASTRAZENECA UK LIMITED	SI
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/055	ASTRAZENECA UK 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
Seroquel SR 400 mg tablete s podaljšanim sproščanjem	NL/H/0156/011	H/99/01407/053	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/035	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/032	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/060	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/034	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/059	ASTRAZENECA UK LIMITED	SI
Seroquel SR 50 mg tablete s podaljšanim sproščanjem	NL/H/0156/008	H/99/01407/033	ASTRAZENECA UK LIMITED	SI
Seroquel XR 200 mg retard tabletta	NL/H/0156/009	OGYI-T-5863/03	ASTRAZENECA KFT.	HU
Seroquel SR 150 mg comprimidos de libertação prolongada	NL/H/0156/012	5456033	ASTRAZENECA PRODUTOS FARMACEUTICOS LDA	PT
Seroquel XR 200 mg pailginto atpalaidavimo tabletės	NL/H/0156/009	LT/1/08/0964/002	ASTRAZENECA UK LIMITED	LT
SEROQUEL XR 200 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/009	54581/15.06.2017	ASTRAZENECA S.A	GR
SEROQUEL XR 150 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/012	54580/15.06.2017	ASTRAZENECA S.A	GR
SEROQUEL XR 400 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/011	54583/15.06.2017	ASTRAZENECA S.A	GR
SEROQUEL XR 300 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/010	54582/15.06.2017	ASTRAZENECA S.A	GR
SEROQUEL XR 50 mg δισκία παρατεταμένης αποδέσμευσης	NL/H/0156/008	54579/15.06.2017	ASTRAZENECA S.A	GR
Seroquel Prolong 200 mg comprimidos de liberación prolongada	NL/H/0156/009	69.646	ASTRAZENECA FARMACÉUTICA SPAIN, S.A.	ES
Seroquel 150 mg compresse a rilascio prolungato	NL/H/0156/012	A.I.C. 032944530	ASTRAZENECA S.P.A.	IT
Seroquel 150 mg compresse a	NL/H/0156/012	A.I.C. 032944542	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
rilascio prolungato				
Seroquel 150 mg compresse a rilascio prolungato	NL/H/0156/012	A.I.C. 032944579	ASTRAZENECA S.P.A.	IT
Seroquel 150 mg compresse a rilascio prolungato	NL/H/0156/012	A.I.C. 032944567	ASTRAZENECA S.P.A.	IT
Seroquel 150 mg compresse a rilascio prolungato	NL/H/0156/012	A.I.C. 032944555	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	A.I.C. 032944593	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	A.I.C. 032944581	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	A.I.C. 032944617	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	A.I.C. 032944629	ASTRAZENECA S.P.A.	IT
Seroquel 200 mg compresse a rilascio prolungato	NL/H/0156/009	A.I.C. 032944605	ASTRAZENECA S.P.A.	IT
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C. 032944670	ASTRAZENECA S.P.A.	IT
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C. 032944643	ASTRAZENECA S.P.A.	IT
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C. 032944668	ASTRAZENECA S.P.A.	IT
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C. 032944631	ASTRAZENECA S.P.A.	IT
Seroquel 300 mg compresse a rilascio prolungato	NL/H/0156/010	A.I.C. 032944656	ASTRAZENECA S.P.A.	IT
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C. 032944718	ASTRAZENECA S.P.A.	IT
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C. 032944720	ASTRAZENECA S.P.A.	IT
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C. 032944694	ASTRAZENECA S.P.A.	IT
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C. 032944682	ASTRAZENECA S.P.A.	IT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Seroquel 400 mg compresse a rilascio prolungato	NL/H/0156/011	A.I.C. 032944706	ASTRAZENECA S.P.A.	IT
Seroquel 50 mg compresse a rilascio prolungato	NL/H/0156/008	A.I.C. 032944480	ASTRAZENECA S.P.A.	IT
Seroquel 50 mg compresse a rilascio prolungato	NL/H/0156/008	A.I.C. 032944528	ASTRAZENECA S.P.A.	IT
Seroquel 50 mg compresse a rilascio prolungato	NL/H/0156/008	A.I.C. 032944492	ASTRAZENECA S.P.A.	IT
Seroquel 50 mg compresse a rilascio prolungato	NL/H/0156/008	A.I.C. 032944516	ASTRAZENECA S.P.A.	IT
Seroquel 50 mg compresse a rilascio prolungato	NL/H/0156/008	A.I.C. 032944504	ASTRAZENECA S.P.A.	IT
Seroquel XL 200 mg prolonged-release tablets	NL/H/0156/009	PL 17901/0250	ASTRAZENECA UK LIMITED	UK
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/030	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/031	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/032	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/033	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/034	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/035	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/036	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/037	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/038	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/039	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene	DK/H/1527/004	H/09/01997/040	LEK PHARMACEUTICALS D.D.	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
tablete			LJUBLJANA	
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/041	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/042	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/043	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 300 mg filmsko obložene tablete	DK/H/1527/004	H/09/01997/044	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/004	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/007	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/014	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/015	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/016	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/018	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/025	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/026	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/027	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/028	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 200 mg filmsko obložene tablete	DK/H/1527/003	H/09/01997/029	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/001	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/002	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/003	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/006	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/005	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/008	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/009	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/010	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/011	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/012	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 100 mg filmsko obložene tablete	DK/H/1527/001	H/09/01997/013	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/017	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene	DK/H/1430/002	H/09/01933/018	LEK PHARMACEUTICALS D.D.	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
tablete			LJUBLJANA	
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/019	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/020	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/021	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/022	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/023	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/024	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/025	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/026	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/027	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/028	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/029	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/030	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/031	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/032	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/033	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/034	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/036	LEK PHARMACEUTICALS D.D. LJUBLJANA	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
Kvelux 50 mg filmsko obložene tablete	DK/H/1430/002	H/09/01933/035	LEK PHARMACEUTICALS D.D. LJUBLJANA	SI
Quetiapin 1A Pharma 100 mg - Filmtabletten	DK/H/1529/001	1-29169	1A PHARMA GMBH	AT
Quetiapin Sandoz 50 mg – Filmtabletten	DK/H/1430/002	1-28698	SANDOZ GMBH	AT
Quetiapin 1A Pharma 50 mg - Filmtabletten	DK/H/1432/002	1-28720	1A PHARMA GMBH	AT
Quetiapin Sandoz 150 mg – Filmtabletten	DK/H/1527/002	1-29175	SANDOZ GMBH	AT
Quetiapin 1A Pharma 150 mg - Filmtabletten	DK/H/1529/002	1-29170	1A PHARMA GMBH	AT
Quetiapin Sandoz 200 mg – Filmtabletten	DK/H/1527/003	1-29176	SANDOZ GMBH	AT
Quetiapin 1A Pharma 200 mg - Filmtabletten	DK/H/1529/003	1-29171	1A PHARMA GMBH	AT
Quetiapin Sandoz 300 mg – Filmtabletten	DK/H/1527/004	1-29177	SANDOZ GMBH	AT
Quetiapin 1A Pharma 300 mg - Filmtabletten	DK/H/1529/004	1-29172	1A PHARMA GMBH	AT
Quetiapin 1A Pharma 400 mg - Filmtabletten	DK/H/1529/005	1-29173	1A PHARMA GMBH	AT
Quetiapin HEXAL 100 mg Filmtabletten	DK/H/1528/001	73558.00.00	HEXAL AG	DE
Quetiapin - 1 A Pharma 100 mg Filmtabletten	DK/H/1529/001	73497.00.00	1 A PHARMA GMBH	DE
Quetiapin HEXAL 150 mg Filmtabletten	DK/H/1528/002	73559.00.00	HEXAL AG	DE
Quetiapin HEXAL 200 mg Filmtabletten	DK/H/1528/003	73560.00.00	HEXAL AG	DE
Quetiapin - 1 A Pharma 200 mg Filmtabletten	DK/H/1529/003	73499.00.00	1 A PHARMA GMBH	DE
Quetiapin HEXAL 300 mg Filmtabletten	DK/H/1528/004	73561.00.00	HEXAL AG	DE
Quetiapin - 1 A Pharma 300 mg	DK/H/1529/004	73500.00.00	1 A PHARMA GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Filmtabletten				
Quetiapin - 1 A Pharma 400 mg Filmtabletten	DK/H/1529/005	73501.00.00	1 A PHARMA GMBH	DE
Quetiapin HEXAL 50 mg Filmtabletten	DK/H/1431/002	71239.00.00	HEXAL AG	DE
Quetiapin - 1 A Pharma 150 mg Filmtabletten	DK/H/1529/002	73498.00.00	1 A PHARMA GMBH	DE
Quetiapin HEXAL 400 mg Filmtabletten	DK/H/1528/005	73562.00.00	HEXAL AG	DE
Quetiapine 200 mg Film-coated Tablets	DK/H/1527/003	PL 04416/0964	SANDOZ LTD	UK
Quetiapine 100 mg Film-coated Tablets	DK/H/1527/001	PL 04416/0962	SANDOZ LTD	UK
Quetiapine 150 mg Film-coated Tablets	DK/H/1527/002	PL 04416/0963	SANDOZ LTD	UK
Quetiapine 300 mg Film-coated Tablets	DK/H/1527/004	PL 04416/0965	SANDOZ LTD	UK
QUETIAPINE SANDOZ 300 MG, FILMOMHULDE TABLETTEN	DK/H/1527/004	RVG 102993	SANDOZ B.V.	NL
Quetiapine Sandoz 200 mg, filmomhulde tabletten	DK/H/1527/003	RVG 102992	SANDOZ B.V.	NL
Quetiapin "Sandoz"	DK/H/1430/002	42919	SANDOZ A/S	DK
Quetiapin "1A Farma"	DK/H/1432/002	42923	1A FARMA A/S	DK
Seresano	DK/H/1431/002	42921	HEXAL A/S	DK
Quetiapin "Sandoz"	DK/H/1527/001	43568	SANDOZ A/S	DK
Quetiapin "Sandoz"	DK/H/1527/002	43569	SANDOZ A/S	DK
Quetiapin "Sandoz"	DK/H/1527/003	43570	SANDOZ A/S	DK
Quetiapin "Sandoz"	DK/H/1527/004	43571	SANDOZ A/S	DK
Seresano, filmovertrukne tabletter 100 mg	DK/H/1528/001	43573	HEXAL A/S	DK
Seresano, filmovertrukne tabletter 150 mg	DK/H/1528/002	43574	HEXAL A/S	DK
Seresano, filmovertrukne tabletter 200 mg	DK/H/1528/003	43575	HEXAL A/S	DK
Seresano, filmovertrukne tabletter	DK/H/1528/004	43576	HEXAL A/S	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
300 mg				
Quetiapin ”1A Farma”, filmovertrukne tabletter 100 mg	DK/H/1529/001	43582	1A FARMA A/S	DK
Quetiapin "1A Farma", filmovertrukne tabletter 150 mg	DK/H/1529/002	43583	1A FARMA A/S	DK
Quetiapin "1A Farma", filmovertrukne tabletter 200 mg	DK/H/1529/003	43584	1A FARMA A/S	DK
Quetiapin "1A Farma", filmovertrukne tabletter 300 mg	DK/H/1529/004	43585	1A FARMA A/S	DK
Seresano, filmovertrukne tabletter 400 mg	DK/H/1528/005	43577	HEXAL A/S	DK
Quetiapin "1A Farma", filmovertrukne tabletter 400 mg	DK/H/1529/005	43586	1A FARMA A/S	DK
Quetiapin Sandoz 100 mg kalvopäällysteinen tabletti	DK/H/1527/001	25268	SANDOZ A/S	FI
Quetiapin Sandoz 300 mg kalvopäällysteinen tabletti	DK/H/1527/004	25271	SANDOZ A/S	FI
Quetiapin Sandoz 200 mg kalvopäällysteinen tabletti	DK/H/1527/003	25270	SANDOZ A/S	FI
Quetiapin Sandoz 100 mg filmdragerade tabletter	DK/H/1527/001	27752	SANDOZ A/S	SE
Quetiapin Sandoz 200 mg filmdragerade tabletter	DK/H/1527/003	27754	SANDOZ A/S	SE
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968012	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968048	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968075	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968099	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968101	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968152	SANDOZ GMBH	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
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968176	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968190	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968226	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968240	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968253	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968315	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968327	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968339	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968378	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968024	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968051	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968063	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968087	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968113	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968125	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968137	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968149	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg	DK/H/1527/003	040968164	SANDOZ GMBH	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
compresse rivestite con film				
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968188	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968238	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968202	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968214	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 200 mg compresse rivestite con film	DK/H/1527/003	040968265	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968277	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968291	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968303	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968341	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968354	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968366	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968380	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968392	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 100 mg compresse rivestite con film	DK/H/1527/001	040968036	SANDOZ GMBH	IT
Quetiapina Sandoz GmbH 300 mg compresse rivestite con film	DK/H/1527/004	040968289	SANDOZ GMBH	IT
QUETIAPINE SANDOZ 100 MG, FILMOMHULDE TABLETTEN	DK/H/1527/001	RVG 102974	SANDOZ B.V.	NL
Quetex 100 mg Film-Coated Tablets	DK/H/1528/001	PA 711/156/3	ROWEX LTD	IE
Quetex 200 mg Film-Coated Tablets	DK/H/1528/003	PA 711/156/4	ROWEX LTD	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Quetex 300 mg Film-Coated Tablets	DK/H/1528/004	PA 711/156/5	ROWEX LTD	IE
Quetiapin Sandoz 100 mg – Filmtabletten	DK/H/1527/001	1-29174	SANDOZ GMBH	AT
Quetiapina Qualigen 50 mg comprimidos recubiertos con película	not available	80989	QUALIGEN, S.L.	ES
Tomel 150 mg Film-coated tablets	not available	21146	DELORBIS PHARMACEUTICALS LTD	CY
Tomel 300 mg Film-coated tablets	not available	21148	DELORBIS PHARMACEUTICALS LTD	CY
Ketilept 300 mg apvalkotās tabletes	HU/H/0126/001-005/MR	07-0335	EGIS PHARMACEUTICALS PLC	LV
Ketilept 150 mg filmtabletta	HU/H/0126/003/MR	OGYI-T-20056/10, 12	EGIS PHARMACEUTICALS PLC	HU
Ketilept 150 mg comprimate filmate	HU/H/0126/003/MR/E001	4214/2012/03-04	EGIS PHARMACEUTICALS PLC	RO
Ketilept 300 mg potahované tablety	HU/H/0126/001-005/MR	68/255/07-C	EGIS PHARMACEUTICALS PLC	CZ
Ketilept 150 mg filmom obalené tablety	HU/H/0126/003/MR	68/0143/07-S	EGIS PHARMACEUTICALS PLC	SK
Ketilept 150 mg apvalkotās tabletes	HU/H/0126/001-005/MR	07-0333	EGIS PHARMACEUTICALS PLC	LV
Ketileppt 150 mg filmtabletta	HU/H/0126/003/MR	OGYI-T-20056/09, 11	EGIS PHARMACEUTICALS PLC	HU
Psicotric 50 mg comprimidos recubiertos con película	not available	80459	QUALIGEN, S.L.	ES
Quetiapine Neuraxpharm 50 mg filmomhulde tabletten	NL/H/1616/006	RVG 110369	NEURAXPHARM ARZNEIMITTEL GMBH	NL
Quetiapin-neuraxpharm 50 mg Filmtabletten	NL/H/1616/006	85926.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Quetiapin NeuroPharma, 50 mg, tabletki powlekane	NL/H/1616/006	20577	NEURAXPHARM ARZNEIMITTEL GMBH	PL
Quetiapin Lannacher 50 mg-Retardtabletten	not available	1-31091	G.L. PHARMA GMBH	AT
Quetiapin Lannacher 200 mg-Retardtabletten	not available	1-31092	G.L. PHARMA GMBH	AT
Quetiapin Lannacher 300 mg-Retardtabletten	not available	1-31096	G.L. PHARMA GMBH	AT
Quetiapin Lannacher 400 mg-Retardtabletten	not available	1-31097	G.L. PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Quetiapin Rosemont 20 mg/ml mikstur, suspensjon	UK/H/5869/01/DC	14-10296	ROSEMONT PHARMACEUTICALS LIMITED	NO
Quetiapin Rosemont, oral suspension	UK/H/5869/01/DC	MTNR. 54693	ROSEMONT PHARMACEUTICALS LIMITED	DK
Quetiapine Rosemont 20mg/ml Oral Suspension	UK/H/5869/01/DC	PL 00427/0240	ROSEMONT PHARMACEUTICALS LIMITED	UK
Desiquet 20mg/ml Suspension zum Einnehmen	UK/H/5869/01/DC	93617.00.00	ROSEMONT PHARMACEUTICALS LIMITED	DE
Quetiapine Rosemont 20mg/ml Oral Suspension	UK/H/5869/01/DC	PA 312/028/005	ROSEMONT PHARMACEUTICALS LIMITED	IE
Quetiapine/Rosemont 20mg/ml πόσιμο εναιώρημα	UK/H/5869/01/DC	8988 / 02-02/2017	ROSEMONT PHARMACEUTICALS LIMITED	GR
Quetiapin Gerot 50 mg-Retardtabletten	not available	1-31085	G.L. PHARMA GMBH	AT
Quetiapin Gerot 200 mg-Retardtabletten	not available	1-31088	G.L. PHARMA GMBH	AT
Quetiapin Gerot 300 mg-Retardtabletten	not available	1-31089	G.L. PHARMA GMBH	AT
Quetiapin Gerot 400 mg-Retardtabletten	not available	1-31090	G.L. PHARMA GMBH	AT
Quetiapine 150 mg film-coated tablets	PT/H/0604/003	MA807/03008	AUROBINDO PHARMA (MALTA) LIMITED	MT
Quetiapine 300 mg film-coated tablets	PT/H/0604/005	MA807/03010	AUROBINDO PHARMA (MALTA) LIMITED	MT
Quetiapin Rosemont 20 mg/ml mikstur, suspensjon	UK/H/5869/01/DC	14-10296	ROSEMONT PHARMACEUTICALS LIMITED	NO
Quetiapin Rosemont, oral suspension	UK/H/5869/01/DC	MTNR. 54693	ROSEMONT PHARMACEUTICALS LIMITED	DK
Quetiapine Rosemont 20mg/ml Oral Suspension	UK/H/5869/01/DC	PL 00427/0240	ROSEMONT PHARMACEUTICALS LIMITED	UK
Desiquet 20mg/ml Suspension zum Einnehmen	UK/H/5869/01/DC	93617.00.00	ROSEMONT PHARMACEUTICALS LIMITED	DE
Quetiapine Rosemont 20mg/ml Oral Suspension	UK/H/5869/01/DC	PA 312/028/005	ROSEMONT PHARMACEUTICALS LIMITED	IE
Quetiapine/Rosemont 20mg/ml	UK/H/5869/01/DC	8988 / 02-02/2017	ROSEMONT	GR

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
πόσιμο εναιώρημα			PHARMACEUTICALS LIMITED	
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/11	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/13	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg apvalkotās tabletes	DK/H/1059/001	07-0229	KRKA, D.D., NOVO MESTO	LV
Kventiax 200 mg apvalkotās tabletes	DK/H/1059/004	07-0232	KRKA, D.D., NOVO MESTO	LV
Kventiax 300 mg apvalkotās tabletes	DK/H/1059/005	07-0233	KRKA, D.D., NOVO MESTO	LV
Kventiax 300 mg tabletki powlekane	DK/H/1059/005	14113	KRKA, D.D., NOVO MESTO	PL
Квентиакс 150 mg филмирани таблетки	DK/H/1059/003	20100176	KRKA, D.D., NOVO MESTO	BG
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/01	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/01	KRKA, D.D., NOVO MESTO	RO
Kventiax, 300 mg õhukese polümeerikattega tabletid	DK/H/1059/005	558007	KRKA, D.D., NOVO MESTO	EE
Kventiax, 25 mg õhukese polümeerikattega tabletid	DK/H/1059/001	558307	KRKA, D.D., NOVO MESTO	EE
Kventiax 150 mg filmom obalené tablety	DK/H/1059/003	68/0266/07-S	KRKA, D.D., NOVO MESTO	SK
Kventiax 300 mg potahované tablety	DK/H/1059/005	68/476/07-C	KRKA, D.D., NOVO MESTO	CZ
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/02	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/03	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/04	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/05	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/06	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/07	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/08	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/09	KRKA, D.D., NOVO MESTO	RO
Kventiax 25 mg comprimate filmate	DK/H/1059/001	5522/2013/10	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate	DK/H/1059/003	5524/2013/02	KRKA, D.D., NOVO MESTO	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
filmate				
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/03	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/04	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/05	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/06	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/07	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/08	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/09	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/10	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/11	KRKA, D.D., NOVO MESTO	RO
Kventiax 150 mg comprimate filmate	DK/H/1059/003	5524/2013/12	KRKA, D.D., NOVO MESTO	RO
Quetiapin-ratiopharm® 200 mg Filmtabletten	UK/H/1228/004	2009020018	RATIOPHARM GMBH	LU
Quetiapin-ratiopharm® 300 mg Filmtabletten	UK/H/1228/005	2009020019	RATIOPHARM GMBH	LU
Quetiapin-ratiopharm® 300 mg Filmtabletten	UK/H/1228/005	70576.00.00	RATIOPHARM GMBH	DE
Quetiapin-ratiopharm® 100 mg Filmtabletten	UK/H/1228/002	2009020017	RATIOPHARM GMBH	LU
Quetiapine Teva 200 mg apvalkotās tabletes	UK/H/1228/004	08-0340	TEVA PHARMA B.V.	LV
Quetiapin-ratiopharm® 25 mg Filmtabletten	UK/H/1228/001	2009020016	RATIOPHARM GMBH	LU
Quetiapin-ratiopharm® 100 mg Filmtabletten	UK/H/1228/002	70573.00.00	RATIOPHARM GMBH	DE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Quetiapin-ratiopharm® 200 mg Filmtabletten	UK/H/1228/004	70575.00.00	RATIOPHARM GMBH	DE
Quetiapine Teva 25 mg apvalkotās tabletes	UK/H/1228/001	08-0342	TEVA NEDERLAND B.V.	LV
Quetiapin-ratiopharm® 25 mg Filmtabletten	UK/H/1228/001	70572.00.00	RATIOPHARM GMBH	DE
Biquelle XL 50 mg prolonged-release tablets	not available	PL 35533/0051	ASPIRE PHARMA LIMITED	UK
Biquelle XL 150 mg prolonged-release tablets	not available	PL 35533/0052	ASPIRE PHARMA LIMITED	UK
Biquelle XL 200 mg prolonged-release tablets	not available	PL 35533/0053	ASPIRE PHARMA LIMITED	UK
Biquelle XL 300 mg prolonged-release tablets	not available	PL 35533/0054	ASPIRE PHARMA LIMITED	UK
Biquelle XL 400 mg prolonged-release tablets	not available	PL 35533/0055	ASPIRE PHARMA LIMITED	UK
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/030	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/031	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/032	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/033	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/034	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/035	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/036	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/037	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/038	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele	UK/H/3525/003	LT/1/11/2387/039	ACCORD HEALTHCARE	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
dengtos tabletės			LIMITED	
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/040	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/041	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/042	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/058	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/059	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/060	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/061	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/062	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/063	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/064	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/065	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/066	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/067	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/068	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/069	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/070	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/030	ACCORD HEALTHCARE 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
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/031	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/032	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/033	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/034	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/035	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/036	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/037	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/038	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/039	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/040	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/041	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/042	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/058	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/059	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/060	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/061	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/062	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele	UK/H/3525/005	LT/1/11/2387/063	ACCORD HEALTHCARE	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
dengtos tabletes			LIMITED	
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/064	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/065	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/066	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/067	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/068	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/069	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele dengtos tabletes	UK/H/3525/005	LT/1/11/2387/070	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/02	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/03	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/04	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/05	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/06	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/07	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/08	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/09	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/10	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/11	ACCORD HEALTHCARE 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
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/12	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/13	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/14	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/02	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/03	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/04	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/05	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/06	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/07	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/08	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/09	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/10	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/11	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/12	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/14	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/13	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg plevele dengtos tabletes	UK/H/3525/003	LT/1/11/2387/029	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plevele	UK/H/3525/005	LT/1/11/2387/057	ACCORD HEALTHCARE	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
dengtos tabletes			LIMITED	
Alcreno , 150 mg, tabletki powlekane	UK/H/3525/003	18456	ACCORD HEALTHCARE LIMITED	PL
Alcreno , 300 mg, tabletki powlekane	UK/H/3525/005	18458	ACCORD HEALTHCARE LIMITED	PL
Quetiapine Accord 25 mg comprimate filmate	UK/H/3525/001	3590/2011/01	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg comprimate filmate	UK/H/3525/003	3592/2011/01	ACCORD HEALTHCARE LIMITED	RO
Quetiapine Accord 150 mg filmom obalené tablety	UK/H/3525/003	68/0051/11-S	ACCORD HEALTHCARE LIMITED	SK
Quetiapine Accord 150 mg plėvele dengtos tabletės	UK/H/3525/003	LT/1/11/2387/029	ACCORD HEALTHCARE LIMITED	LT
Quetiapine Accord 300 mg plėvele dengtos tabletės	UK/H/3525/005	LT/1/11/2387/057	ACCORD HEALTHCARE LIMITED	LT
Setinin 150 mg filmsko obložene tablete	DK/H/1390/003	H/10/01411/008	ACTAVIS GROUP PTC EHF.	SI
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/025	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/027	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/030	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/031	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/029	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/036	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/026	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/032	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plėvele dengtos tabletės	DK/H/1390/003	LT/1/09/1555/028	ACTAVIS GROUP PTC EHF.	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
Setinin 150 mg filmsko obložene tablete	DK/H/1390/003	H/10/01411/007	ACTAVIS GROUP PTC EHF.	SI
Quetiapine Actavis 150 mg plévele dengtos tabletés	DK/H/1390/003	LT/1/09/1555/033	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plévele dengtos tabletés	DK/H/1390/003	LT/1/09/1555/034	ACTAVIS GROUP PTC EHF.	LT
Quetiapine Actavis 150 mg plévele dengtos tabletés	DK/H/1390/003	LT/1/09/1555/035	ACTAVIS GROUP PTC EHF.	LT
QUETIAPINE ACTAVIS 25 mg apvalkotās tabletes	DK/H/1390/001	09-0108	ACTAVIS GROUP PTC EHF.	LV
Setinin 150 mg film-coated tablets	DK/H/1390/003	MA628/02303	ACTAVIS GROUP PTC EHF.	MT