11 April 2024
EMA/160401/2024
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): zolpidem

Procedure No. PSUSA-00003151-202308

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
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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
Bikalm 10 mg Filmtabletten	not available	25938.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Bikalm 10 mg Filmtabletten	not available	25938.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Bikalm 10 mg Filmtabletten	not available	25938.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Bikalm 10 mg Filmtabletten	not available	25938.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Dalparan 10 mg comprimidos recubiertos con película	not available	59264	SANOFI-AVENTIS, S.A.	ES
Dalparan 10 mg comprimidos recubiertos con película	not available	59264	SANOFI-AVENTIS, S.A.	ES
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/01	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/02	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/03	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/04	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/05	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/06	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/07	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/08	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/01	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/02	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/03	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate	SE/H/1046/002	10616/2018/04	VIATRIS HEALTHCARE LTD	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
sublinguale				
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/05	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/06	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/07	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimate sublinguale	SE/H/1046/002	10616/2018/08	VIATRIS HEALTHCARE LTD	RO
Edluar 10 mg comprimido sublingual	SE/H/1046/002	5481031	VIATRIS HEALTHCARE, LDA	PT
Edluar 10 mg comprimido sublingual	SE/H/1046/002	5481023	VIATRIS HEALTHCARE, LDA	PT
Edluar 10 mg comprimido sublingual	SE/H/1046/002	5481031	VIATRIS HEALTHCARE, LDA	PT
Edluar 10 mg comprimido sublingual	SE/H/1046/002	5481023	VIATRIS HEALTHCARE, LDA	PT
Edluar 10 mg comprimidos sublinguales	SE/H/1046/002	76723	MYLAN IRE HEALTHCARE LIMITED	ES
Edluar 10 mg comprimidos sublinguales	SE/H/1046/002	76723	MYLAN IRE HEALTHCARE LIMITED	ES
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/08	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/09	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/10	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/11	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/12	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/13	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/14	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti	SE/H/1046/002	OGYI-T-22265/16	VIATRIS HEALTHCARE LTD	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletta				
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/08	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/09	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/10	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/11	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/12	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/13	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/14	VIATRIS HEALTHCARE LTD	HU
EDLUAR 10 mg nyelvalatti tabletta	SE/H/1046/002	OGYI-T-22265/16	VIATRIS HEALTHCARE LTD	HU
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/010	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/011	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/012	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/013	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/014	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/015	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/009	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/016	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/010	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične	SE/H/1046/002	H/13/00526/011	MEDA PHARMA GMBH & CO.	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			KG	
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/012	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/013	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/014	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/015	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/009	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg podjezične tablete	SE/H/1046/002	H/13/00526/016	MEDA PHARMA GMBH & CO. KG	SI
Edluar 10 mg resoriblett, sublingual	SE/H/1046/002	45050	MEDA AB	SE
Edluar 10 mg resoriblett, sublingual	SE/H/1046/001	29165	VIATRIS OY	FI
Edluar 10 mg resoriblett, sublingual	SE/H/1046/002	45050	MEDA AB	SE
Edluar 10 mg resoriblett, sublingual	SE/H/1046/001	29165	VIATRIS OY	FI
Edluar 10 mg resoribletti	SE/H/1046/002	29165	VIATRIS OY	FI
Edluar 10 mg resoribletti	SE/H/1046/002	29165	VIATRIS OY	FI
Edluar 10 mg sublingual tablets	SE/H/1046/002	PL 46302/0204	MYLAN PRODUCTS LIMITED	XI
Edluar 10 mg sublingual tablets	SE/H/1046/002	PA2010/050/002	MYLAN IRE HEALTHCARE LIMITED	IE
Edluar 10 mg sublingual tablets	SE/H/1046/002	PL 46302/0204	MYLAN PRODUCTS LIMITED	XI
Edluar 10 mg sublingual tablets	SE/H/1046/002	PA2010/050/002	MYLAN IRE HEALTHCARE LIMITED	IE
Edluar 10 mg Sublingualtabletten	SE/H/1046/002	1-31494	MYLAN ÖSTERREICH GMBH	AT
Edluar 10 mg Sublingualtabletten	SE/H/1046/002	1-31494	MYLAN ÖSTERREICH GMBH	AT
Edluar 10 mg tungurótartöflur	SE/H/1046/002	IS/1/12/076/02	VIATRIS APS	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
Edluar 10 mg tungurótartöflur	SE/H/1046/002	IS/1/12/076/02	VIATRIS APS	IS
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 583 283 4 3	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 583 282 8 2	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 269 404 7 8	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 127 2 6	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 126 6 5	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 124 3 6	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 123 7 5	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 122 0 7	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 583 283 4 3	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 583 282 8 2	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 269 404 7 8	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 127 2 6	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 126 6 5	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 124 3 6	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 123 7 5	MEDA PHARMA SAS	FR
EDLUAR 10 mg, comprimé sublingual	SE/H/1046/002	34009 266 122 0 7	MEDA PHARMA SAS	FR
Edluar 10 mg, tablet voor sublinguaal gebruik	SE/H/1046/002	RVG 108439	MYLAN HEALTHCARE B.V.	NL
Edluar 10 mg, tablet voor	SE/H/1046/002	RVG 108439	MYLAN HEALTHCARE B.V.	NL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
sublinguaal gebruik				
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/01	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/02	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/03	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/04	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/05	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/06	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/07	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/08	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/01	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/02	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/03	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/04	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/05	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/06	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/07	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimate sublinguale	SE/H/1046/001	10615/2018/08	VIATRIS HEALTHCARE LTD	RO
Edluar 5 mg comprimido sublingual	SE/H/1046/001	5481015	VIATRIS HEALTHCARE, LDA	PT
Edluar 5 mg comprimido	SE/H/1046/001	5481007	VIATRIS HEALTHCARE, LDA	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
sublingual				
Edluar 5 mg comprimido sublingual	SE/H/1046/001	5481015	VIATRIS HEALTHCARE, LDA	PT
Edluar 5 mg comprimido sublingual	SE/H/1046/001	5481007	VIATRIS HEALTHCARE, LDA	PT
Edluar 5 mg comprimidos sublinguales	SE/H/1046/001	76724	MYLAN IRE HEALTHCARE LIMITED	ES
Edluar 5 mg comprimidos sublinguales	SE/H/1046/001	76724	MYLAN IRE HEALTHCARE LIMITED	ES
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/01	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/02	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/03	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/04	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/05	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/06	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/07	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/15	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/01	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/02	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/03	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/04	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/05	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti	SE/H/1046/001	OGYI-T-22265/06	VIATRIS HEALTHCARE LTD	HU

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
tabletta				
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/07	VIATRIS HEALTHCARE LTD	HU
EDLUAR 5 mg nyelvalatti tabletta	SE/H/1046/001	OGYI-T-22265/15	VIATRIS HEALTHCARE LTD	HU
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/001	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/006	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/005	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/004	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/003	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/002	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/007	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/008	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/001	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/006	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/005	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/004	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/003	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/002	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične tablete	SE/H/1046/001	H/13/00526/007	MEDA PHARMA GMBH & CO. KG	SI
Edluar 5 mg podjezične	SE/H/1046/001	H/13/00526/008	MEDA PHARMA GMBH & CO.	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			KG	
Edluar 5 mg resoriblett, sublingual	SE/H/1046/001	45049	MEDA AB	SE
Edluar 5 mg resoriblett, sublingual	SE/H/1046/002	29164	VIATRIS OY	FI
Edluar 5 mg resoriblett, sublingual	SE/H/1046/001	45049	MEDA AB	SE
Edluar 5 mg resoriblett, sublingual	SE/H/1046/002	29164	VIATRIS OY	FI
Edluar 5 mg resoribletti	SE/H/1046/001	29164	VIATRIS OY	FI
Edluar 5 mg resoribletti	SE/H/1046/001	29164	VIATRIS OY	FI
Edluar 5 mg sublingual tablets	SE/H/1046/001	PL 46302/0203	MYLAN PRODUCTS LIMITED	XI
Edluar 5 mg sublingual tablets	SE/H/1046/001	PA2010/050/001	MYLAN IRE HEALTHCARE LIMITED	IE
Edluar 5 mg sublingual tablets	SE/H/1046/001	PL 46302/0203	MYLAN PRODUCTS LIMITED	XI
Edluar 5 mg sublingual tablets	SE/H/1046/001	PA2010/050/001	MYLAN IRE HEALTHCARE LIMITED	IE
Edluar 5 mg Sublingualtabletten	SE/H/1046/001	83439.00.00	MEDA PHARMA GMBH & CO. KG	DE
Edluar 5 mg Sublingualtabletten	SE/H/1046/001	1-31493	MYLAN ÖSTERREICH GMBH	AT
Edluar 5 mg Sublingualtabletten	SE/H/1046/001	83439.00.00	MEDA PHARMA GMBH & CO. KG	DE
Edluar 5 mg Sublingualtabletten	SE/H/1046/001	1-31493	MYLAN ÖSTERREICH GMBH	AT
Edluar 5 mg tungurótartöflur	SE/H/1046/001	IS/1/12/076/01	VIATRIS APS	IS
Edluar 5 mg tungurótartöflur	SE/H/1046/001	IS/1/12/076/01	VIATRIS APS	IS
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 072 3 4	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 074 6 3	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 076 9 2	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé	SE/H/1046/001	34009 266 077 5 3	MEDA PHARMA SAS	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
sublingual				
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 269 403 0 0	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 583 236 6 9	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 583 237 2 0	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 075 2 4	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 072 3 4	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 074 6 3	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 076 9 2	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 077 5 3	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 269 403 0 0	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 583 236 6 9	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 583 237 2 0	MEDA PHARMA SAS	FR
EDLUAR 5 mg, comprimé sublingual	SE/H/1046/001	34009 266 075 2 4	MEDA PHARMA SAS	FR
Edluar 5 mg, tablet voor sublinguaal gebruik	SE/H/1046/001	RVG 108438	MYLAN HEALTHCARE B.V.	NL
Edluar 5 mg, tablet voor sublinguaal gebruik	SE/H/1046/001	RVG 108438	MYLAN HEALTHCARE B.V.	NL
Edluar, sublinguale resoribletter	SE/H/1046/001	47607	VIATRIS APS	DK
Edluar, sublinguale resoribletter	SE/H/1046/002	47608	VIATRIS APS	DK
Edluar, sublinguale resoribletter	SE/H/1046/001	47607	VIATRIS APS	DK
Edluar, sublinguale	SE/H/1046/002	47608	VIATRIS APS	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
resoribletter				
Ivadal Filmtabletten	not available	1-20472	SANOFI-AVENTIS GMBH OSTERREICH	AT
Ivadal Filmtabletten	not available	1-20472	SANOFI-AVENTIS GMBH OSTERREICH	AT
Ivadal Filmtabletten	not available	1-20472	SANOFI-AVENTIS GMBH OSTERREICH	AT
Ivadal Filmtabletten	not available	1-20472	SANOFI-AVENTIS GMBH OSTERREICH	AT
NOTTEM 10 mg compresse rivestite con film	not available	028445017	SANOFI S.R.L.	IT
NOTTEM 10 mg compresse rivestite con film	not available	028445031	SANOFI S.R.L.	IT
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Oniria 5 mg δισκία διασπειρόμενα στο στόμα	IT/H/0187/003	80459/26-07-2022	ITF HELLAS S.A.	GR
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés	not available	BE146492	SANOFI BELGIUM	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
pelliculés				
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés pelliculés	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg comprimés pelliculés	not available	0323502	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0149217	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0149203	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0883864	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0883851	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0897941	SANOFI BELGIUM	LU
Stilnoct 10 mg comprimés pelliculés	not available	0897923	SANOFI BELGIUM	LU
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerad tablett	not available	11697	SANOFI OY	FI
Stilnoct 10 mg filmdragerade tabletter	not available	12020	SANOFI AB	SE
Stilnoct 10 mg filmdragerade	not available	12020	SANOFI 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
tabletter				
Stilnoct 10 mg filmdragerade tabletter	not available	12020	SANOFI AB	SE
Stilnoct 10 mg filmdragerade tabletter	not available	12020	SANOFI AB	SE
Stilnoct 10 mg filmdragerade tabletter	not available	12020	SANOFI AB	SE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmomhulde tabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
STILNOCT 10 mg Filmtabletten	not available	BE146492	SANOFI BELGIUM	BE
Stilnoct 10 mg filmuhúðaðar	not available	880059	SANOFI-AVENTIS NORGE AS	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
töflur.				
Stilnoct 10 mg filmuhúðaðar töflur.	not available	880059	SANOFI-AVENTIS NORGE AS	IS
Stilnoct 10 mg filmuhúðaðar töflur.	not available	880059	SANOFI-AVENTIS NORGE AS	IS
Stilnoct 10 mg tablett, filmdrasjert	not available	8202	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 10 mg tablett, filmdrasjert	not available	8202	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 10 mg tablett, filmdrasjert	not available	8202	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10 mg tabletti, kalvopäällysteinen	not available	11697	SANOFI OY	FI
Stilnoct 10mg Film-coated Tablets	not available	PA 540/160/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg filmdragerade	not available	12019	SANOFI 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
tabletter				
Stilnoct 5 mg filmdragerade tabletter	not available	12019	SANOFI AB	SE
Stilnoct 5 mg tablett, filmdrasjert	not available	8201	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 5 mg tablett,filmdrasjert	not available	8201	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 5 mg tablett,filmdrasjert	not available	8201	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 5 mg tabletter, filmdrasjert	not available	8201	SANOFI-AVENTIS NORGE AS	NO
Stilnoct 5mg Film Coated Tablets	not available	PA 540/160/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Stilnoct, filmovertrukne tabletter	not available	13198	SANOFI A/S	DK
Stilnoct, filmovertrukne tabletter	not available	13198	SANOFI A/S	DK
Stilnoct, filmovertrukne tabletter	not available	13198	SANOFI A/S	DK
Stilnoct, filmovertrukne tabletter	not available	13198	SANOFI A/S	DK
Stilnox 10 mg apvalkotās tabletes	not available	99-1021	SANOFI WINTHROP INDUSTRIE	LV
Stilnox 10 mg apvalkotās tabletes	not available	99-1021	SANOFI WINTHROP INDUSTRIE	LV
STILNOX 10 mg compresse rivestite con film	not available	026695054	SANOFI S.R.L.	IT
STILNOX 10 mg compresse rivestite con film	not available	026695015	SANOFI S.R.L.	IT
STILNOX 10 mg compresse rivestite con film	not available	026695027	SANOFI S.R.L.	IT
STILNOX 10 mg compresse rivestite con film	not available	026695041	SANOFI S.R.L.	IT
Stilnox 10 mg comprimate filmate	not available	1344/2009/03	SANOFI ROMANIA SRL	RO
Stilnox 10 mg comprimate	not available	1344/2009/02	SANOFI ROMANIA SRL	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				
Stilnox 10 mg comprimate filmate	not available	1344/2009/01	SANOFI ROMANIA SRL	RO
Stilnox 10 mg comprimate filmate	not available	1344/2009/04	SANOFI ROMANIA SRL	RO
STILNOX 10 mg comprimidos recubiertos con película	not available	58470	SANOFI-AVENTIS, S.A.	ES
STILNOX 10 mg comprimidos recubiertos con película	not available	58470	SANOFI-AVENTIS, S.A.	ES
Stilnox 10 mg comprimidos revestidos por película	not available	4508495	SANOFI - PRODUTOS FARMACEUTICOS LDA	PT
Stilnox 10 mg film-coated tablet	not available	MA1359/03401	SANOFI S.R.L.	MT
Stilnox 10 mg filmom obalené tablety	not available	57/0887/92-CS	SANOFI WINTHROP INDUSTRIE	SK
Stilnox 10 mg filmom obalené tablety	not available	57/0887/92-CS	SANOFI WINTHROP INDUSTRIE	SK
Stilnox 10 mg filmom obalené tablety	not available	57/0887/92-CS	SANOFI WINTHROP INDUSTRIE	SK
Stilnox 10 mg filmom obalené tablety	not available	57/0887/92-CS	SANOFI WINTHROP INDUSTRIE	SK
Stilnox 10 mg filmom obalené tablety	not available	57/0887/92-CS	SANOFI WINTHROP INDUSTRIE	SK
Stilnox 10 mg filmtabletta	not available	OGYI-T-2244/03	SANOFI-AVENTIS ZRT	HU
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	DE
Stilnox 10 mg Filmtabletten	not available	16345.00.00	SANOFI-AVENTIS DEUTSCHLAND 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**
Stilnox 10 mg plėvele dengtos tabletės	not available	LT/1/95/2002/001	SANOFI WINTHROP INDUSTRIE	LT
Stilnox 10 mg plėvele dengtos tabletės	not available	LT/1/95/2002/002	SANOFI WINTHROP INDUSTRIE	LT
Stilnox 10 mg plėvele dengtos tabletės	not available	LT/1/95/2002/003	SANOFI WINTHROP INDUSTRIE	LT
Stilnox 10 mg potahované tablety	not available	57/887/92-C	SANOFI S.R.O.	CZ
Stilnox 10 mg potahované tablety	not available	57/887/92-C	SANOFI S.R.O.	CZ
Stilnox 10 mg potahované tablety	not available	57/887/92-C	SANOFI S.R.O.	CZ
Stilnox 10 mg potahované tablety	not available	57/887/92-C	SANOFI S.R.O.	CZ
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45258/21-10-2009	SANOFI-AVENTIS MONOPROSOPI A.E.B.E	GR
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45258/21-10-2009	SANOFI-AVENTIS MONOPROSOPI A.E.B.E	GR
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45258/21-10-2009	SANOFI-AVENTIS MONOPROSOPI A.E.B.E	GR
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45258/21-10-2009	SANOFI-AVENTIS MONOPROSOPI A.E.B.E	GR
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	45258/21-10-2009	SANOFI-AVENTIS MONOPROSOPI A.E.B.E	GR
Stilnox 10 mg επικαλυμμένα με λεπτό υμένιο δισκία	not available	19612	SANOFI WINTHROP INDUSTRIE	CY
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 555 845 1 3	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 339 036 1 9	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 329 611 3 9	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 346 586 3 1	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 346 588 6 0	SANOFI WINTHROP INDUSTRIE	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
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 563 132 0 4	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 557 763 2 1	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 329 610 7 8	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 555 846 8 1	SANOFI WINTHROP INDUSTRIE	FR
STILNOX 10 mg, comprimé pelliculé sécable	not available	34009 346 585 7 0	SANOFI WINTHROP INDUSTRIE	FR
Stilnox, 10 mg õhukese polümeerikattega tabletid	not available	327500	SANOFI WINTHROP INDUSTRIE	EE
Stilnox, 10 mg õhukese polümeerikattega tabletid	not available	327500	SANOFI WINTHROP INDUSTRIE	EE
Stilnox, 10 mg õhukese polümeerikattega tabletid	not available	327500	SANOFI WINTHROP INDUSTRIE	EE
STILNOX, 10 mg, tabletki powlekane	not available	R/0375	SANOFI WINTHROP INDUSTRIE	PL
STILNOX, 10 mg, tabletki powlekane	not available	R/0375	SANOFI WINTHROP INDUSTRIE	PL
STILNOX, 10 mg, tabletki powlekane	not available	R/0375	SANOFI WINTHROP INDUSTRIE	PL
STILNOX, 10 mg, tabletki powlekane	not available	R/0375	SANOFI WINTHROP INDUSTRIE	PL
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540080	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540092	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540104	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540128	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540130	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540116	VIATRIS ITALIA S.R.L.	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
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540142	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540167	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540080	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540092	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540104	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540128	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540130	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540116	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540142	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg compresse sublinguali	SE/H/1046/002	040540167	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 10 mg comprimés sublinguaux	SE/H/1046/002	2012120234	VIATRIS HEALTHCARE	LU
Zolpeduar 10 mg comprimés sublinguaux	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 10 mg comprimés sublinguaux	SE/H/1046/002	2012120234	VIATRIS HEALTHCARE	LU
Zolpeduar 10 mg comprimés sublinguaux	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 10 mg Sublingualtabletten	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 10 mg Sublingualtabletten	SE/H/1046/002	2012120234	VIATRIS HEALTHCARE	LU
Zolpeduar 10 mg Sublingualtabletten	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 10 mg Sublingualtabletten	SE/H/1046/002	2012120234	VIATRIS HEALTHCARE	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
Zolpeduar 10 mg tabletten voor sublinguaal gebruik	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 10 mg tabletten voor sublinguaal gebruik	SE/H/1046/002	BE 424295	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540015	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540027	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540039	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540041	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540054	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540066	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540078	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540155	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540015	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540027	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540039	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540041	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540054	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540066	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540078	VIATRIS ITALIA S.R.L.	IT
Zolpeduar 5 mg compresse sublinguali	SE/H/1046/001	040540155	VIATRIS ITALIA S.R.L.	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
Zolpeduar 5 mg comprimés sublinguaux	SE/H/1046/001	2012120233	VIATRIS HEALTHCARE	LU
Zolpeduar 5 mg comprimés sublinguaux	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg comprimés sublinguaux	SE/H/1046/001	2012120233	VIATRIS HEALTHCARE	LU
Zolpeduar 5 mg comprimés sublinguaux	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg Sublingualtabletten	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg Sublingualtabletten	SE/H/1046/001	2012120233	VIATRIS HEALTHCARE	LU
Zolpeduar 5 mg Sublingualtabletten	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg Sublingualtabletten	SE/H/1046/001	2012120233	VIATRIS HEALTHCARE	LU
Zolpeduar 5 mg tabletten voor sublinguaal gebruik	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpeduar 5 mg tabletten voor sublinguaal gebruik	SE/H/1046/001	BE 424286	VIATRIS HEALTHCARE	BE
Zolpidem 10 mg Film-coated Tablets	not available	PL 20416/0599	CRESCENT PHARMA LIMITED	XI
Zolpidem Aristo 5 mg comprimidos	not available	65310	ARISTO PHARMA IBERIA, S.L.	ES
Zolpidem Aurobindo 5 mg film-coated tablets	PT/H/0866/001	MA807/07801	AUROBINDO PHARMA (MALTA) LIMITED	MT
Zolpidem Dune, filmovertrukne tabletter	DK/H/2309/001	52454	DUNE MEDICARE APS	DK
Zolpidem Sandoz 5 mg filmdragerade tabletter	NL/H/0253/001	17289	SANDOZ A/S	SE
Zolpidem Tartrate 10 mg Tablets	not available	PL 17780/0018	ZENTIVA PHARMA UK LIMITED	XI
Zolpidem Tartrate 5 mg Tablets.	not available	PL 17780/0017	ZENTIVA PHARMA UK LIMITED	XI
Zolpidem Tevagen 5 mg comprimidos recubiertos con	NL/H/1585/001	75795	TEVA PHARMA S.L.U.,	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
película				
ZOLPIDEM ZENTIVA 10 mg compresse rivestite con film	not available	031850023	ZENTIVA ITALIA S.R.L.	IT
ZOLPIDEM ZENTIVA 10 mg compresse rivestite con film	not available	031850011	ZENTIVA ITALIA S.R.L.	IT
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 347 972 4 8	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 347 974 7 7	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 347 978 2 8	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 347 977 6 7	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 564 534 5 0	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 333 547 4 9	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 362 195 5 7	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 367 493 4 4	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 564 533 9 9	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 367 492 8 3	ZENTIVA FRANCE	FR
ZOLPIDEM ZENTIVA 10 mg, comprimé pelliculé sécable	not available	34009 347 975 3 8	ZENTIVA FRANCE	FR
Zolpidem-neuraxpharm 5 mg Filmtabletten	not available	49458.00.00	NEURAXPHARM ARZNEIMITTEL GMBH	DE
Zolpidemtartraat Aurobindo 5 mg, filmomhulde tabletten	NL/H/4052/001	RVG 121164	AUROBINDO PHARMA B.V.	NL
ZOLPIDEMTARTRAAT SANDOZ 10 MG, FILMOMHULDE TABLETTEN	NL/H/0262/001	RVG 25030	SANDOZ B.V.	NL
Zolpidemtartraat Teva 5 mg, filmomhulde tabletten	NL/H/1585/001	RVG 103724	TEVA B.V	NL

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Zolpi-Lich 10 mg Filmtabletten	not available	48888.00.00	WINTHROP ARZNEIMITTEL GMBH	DE