1 December 2022
EMA/933404/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: amlodipine / olmesartan

Procedure no.: PSUSA/00002208/202204

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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
Capenon 20 mg/5 mg, filmomhulde tabletten	NL/H/1114/001	RVG 100989	DAIICHI SANKYO EUROPE GMBH	NL
Capenon 40 mg/5 mg, filmomhulde tabletten	NL/H/1114/002	RVG 100990	DAIICHI SANKYO EUROPE GMBH	NL
Capenon 40 mg/10 mg, filmomhulde tabletten	NL/H/1114/003	RVG 100991	DAIICHI SANKYO EUROPE GMBH	NL
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946176	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946137	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946152	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946188	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946101	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946125	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946087	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946099	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946051	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946048	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946113	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946149	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946164	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Capenon 20 mg /5 mg comprimidos recubiertos con película	NL/H/1114/001	70.067	DAIICHI SANKYO ESPAÑA, S.A.	ES
Capenon 40 mg /5 mg comprimidos recubiertos con película	NL/H/1114/002	70.070	DAIICHI SANKYO ESPAÑA, S.A.	ES
Capenon 40 mg /10 mg comprimidos recubiertos con película	NL/H/1114/003	70.071	DAIICHI SANKYO ESPAÑA, S.A.	ES
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/01	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/09	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/10	LABORMED PHARMA S.A.	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
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/11	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/02	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/03	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/04	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/05	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/06	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/07	LABORMED PHARMA S.A.	RO
Sevikar 20 mg/5 mg comprimate filmate	NL/H/1113/001	5953/2013/08	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/04	LABORMED PHARMA S.A.	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
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/05	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/06	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/07	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/08	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/09	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/10	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/01	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/02	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/03	LABORMED PHARMA S.A.	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
Sevikar 40 mg/10 mg comprimate filmate	NL/H/1113/003	5955/2013/11	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/01	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/02	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/10	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/11	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/03	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/04	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/05	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/06	LABORMED PHARMA S.A.	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
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/07	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/08	LABORMED PHARMA S.A.	RO
Sevikar 40 mg/5 mg comprimate filmate	NL/H/1113/002	5954/2013/09	LABORMED PHARMA S.A.	RO
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725023	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725031	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725049	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725056	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725064	TAD PHARMA GMBH	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
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725072	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725106	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725114	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725205	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725213	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725221	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725239	TAD PHARMA GMBH	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
Olssa 20 mg/5 mg potahované tablety	CZ/H/0715/001	58/704/16-C	KRKA, D.D., NOVO MESTO	CZ
Olssa 40 mg/5 mg potahované tablety	CZ/H/0715/002	58/705/16-C	KRKA, D.D., NOVO MESTO	CZ
Olssa 40 mg/10 mg potahované tablety	CZ/H/0715/003	58/706/16-C	KRKA, D.D., NOVO MESTO	CZ
Olssa 20 mg/5 mg õhukese polümeerikattega tabletid	CZ/H/0715/001	949617	KRKA, D.D., NOVO MESTO	EE
Olssa 40 mg/5 mg õhukese polümeerikattega tabletid	CZ/H/0715/002	949717	KRKA, D.D., NOVO MESTO	EE
Olssa 40 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0715/003	949817	KRKA, D.D., NOVO MESTO	EE
Olssa 20 mg/5 mg apvalkotās tabletes	CZ/H/0715/001	17-0199	KRKA, D.D., NOVO MESTO	LV
Olssa 40 mg/5 mg apvalkotās tabletes	CZ/H/0715/002	17-0200	KRKA, D.D., NOVO MESTO	LV
Olssa 40 mg/10 mg apvalkotās tabletes	CZ/H/0715/003	17-0201	KRKA, D.D., NOVO MESTO	LV

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
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725023	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725031	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725049	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 20 mg comprimidos revestidos por película	CZ/H/0715/001	5725056	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725064	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725072	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725106	TAD PHARMA GMBH	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
Amlodipina + olmesartan medoxomilo TAD 5 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/002	5725114	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725205	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725213	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725221	TAD PHARMA GMBH	PT
Amlodipina + olmesartan medoxomilo TAD 10 mg + 40 mg comprimidos revestidos por película	CZ/H/0715/003	5725239	TAD PHARMA GMBH	PT
Olssa 20 mg/5 mg potahované tablety	CZ/H/0715/001	58/704/16-C	KRKA, D.D., NOVO MESTO	CZ
Olssa 40 mg/5 mg potahované tablety	CZ/H/0715/002	58/705/16-C	KRKA, D.D., NOVO MESTO	CZ
Olssa 40 mg/10 mg potahované tablety	CZ/H/0715/003	58/706/16-C	KRKA, D.D., NOVO MESTO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Olssa 20 mg/5 mg õhukese polümeerikattega tabletid	CZ/H/0715/001	949617	KRKA, D.D., NOVO MESTO	EE
Olssa 40 mg/5 mg õhukese polümeerikattega tabletid	CZ/H/0715/002	949717	KRKA, D.D., NOVO MESTO	EE
Olssa 40 mg/10 mg õhukese polümeerikattega tabletid	CZ/H/0715/003	949817	KRKA, D.D., NOVO MESTO	EE
Olssa 20 mg/5 mg apvalkotās tabletes	CZ/H/0715/001	17-0199	KRKA, D.D., NOVO MESTO	LV
Olssa 40 mg/5 mg apvalkotās tabletes	CZ/H/0715/002	17-0200	KRKA, D.D., NOVO MESTO	LV
Olssa 40 mg/10 mg apvalkotās tabletes	CZ/H/0715/003	17-0201	KRKA, D.D., NOVO MESTO	LV
Forzaten 40 mg/5 mg filmomhulde tabletten	NL/H/1115/002	BE325464	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten® 20 mg / 5 mg Filmtabletten	NL/H/1115/001	BE325455	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten® 40 mg / 10 mg Filmtabletten	NL/H/1115/003	BE325473	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Forzaten® 40 mg / 5 mg Filmtabletten	NL/H/1115/002	BE325464	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten 40 mg/10 mg filmomhulde tabletten	NL/H/1115/003	BE325473	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Forzaten 20 mg/5 mg filmomhulde tabletten	NL/H/1115/001	BE325455	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/001	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/023	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/013	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/10 mg comprimate filmate	NL/H/1115/003	7208/2014/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 40 mg/5 mg comprimate filmate	NL/H/1115/002	7207/2014/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/028	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/031	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/030	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/029	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/015	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/019	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/025	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/033	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/027	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/032	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 20 mg/5 mg filmsko obložene tablete	NL/H/1115/001	H/09/01160/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/021	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/016	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/10 mg filmsko obložene tablete	NL/H/1115/003	H/09/01160/026	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/020	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olectan 40 mg/5 mg filmsko obložene tablete	NL/H/1115/002	H/09/01160/022	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Inovum 20 mg/5 mg comprimate filmate	NL/H/1115/001	7206/2014/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	RO
Balzak 20 mg /5 mg comprimidos recubiertos con película	NL/H/1115/001	70.091	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
AXELER 40 mg/10 mg, comprimé pelliculé	NL/H/1115/003	34009 388 541 8 3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
AXELER 40 mg/10 mg, comprimé pelliculé	NL/H/1115/003	34009 388 543 0 5	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947230	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Vocado® 40 mg/5 mg Filmtabletten	NL/H/1115/002	70064.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 40 mg/10 mg	NL/H/1115/003	13406	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/20	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sintonyn 20 mg/5 mg potahované tablety	NL/H/1115/001	58/526/08-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Balzak 40 mg /10 mg comprimidos recubiertos con película	NL/H/1115/003	70.096	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Folgan 40 mg/5 mg filmom obalené tablety	NL/H/1115/002	58/0420/08-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947103	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
AXELER 40 mg/10 mg, comprimé pelliculé	NL/H/1115/003	34009 573 838 3 1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/17	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Sanoral, 40 mg/10 mg õhukese polümeerikattega tabletid	NL/H/1115/003	606308	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Vocado® 40 mg/10 mg Filmtabletten	NL/H/1115/003	70065.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 40 mg/5 mg	NL/H/1115/002	13405	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Sanoral, 40 mg/5 mg õhukese polümeerikattega tabletid	NL/H/1115/002	606408	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
AXELER 40 mg/5 mg, comprimé pelliculé	NL/H/1115/002	34009 388 544 7 3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
AXELER 20 mg/5 mg, comprimé pelliculé	NL/H/1115/001	34009 388 540 1 5	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
AXELER 20 mg/5 mg, comprimé pelliculé	NL/H/1115/001	34009 388 538 7 2	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
AXELER 20 mg/5 mg, comprimé pelliculé	NL/H/1115/001	34009 573 840 8 1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
AXELER 40 mg/5 mg, comprimé pelliculé	NL/H/1115/002	34009 388 547 6 3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Vocado® 20 mg/5 mg Filmtabletten	NL/H/1115/001	70063.00.00	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	DE
AXELER 40 mg/5 mg, comprimé pelliculé	NL/H/1115/002	34009 573 837 7 0	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Konverge 20 mg/5 mg film-coated tablets	NL/H/1115/001	PA 865/17/1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947040	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947077	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947115	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947127	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947091	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947141	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947192	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947139	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947065	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/22	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947038	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947204	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/21	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Sintonyn 40 mg/10 mg potahované tablety	NL/H/1115/003	58/528/08-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 40 mg/10 mg	NL/H/1115/003	20614	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Amelior® 40 mg/10 mg Filmtabletten	NL/H/1115/003	1-27896	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947216	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947228	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947178	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947166	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947154	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Sanoral 40 mg/10 mg apvalkotās tabletes	NL/H/1115/003	08-0272	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/16	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Konverge 40 mg/10 mg film-coated tablets	NL/H/1115/003	PA 865/17/3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947026	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947089	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Теспадан 20 mg/5 mg филмирани таблетки	NL/H/1115/001	20080196	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Sanoral 40 mg/5 mg apvalkotās tabletes	NL/H/1115/002	08-0271	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Amelior® 20 mg/5 mg Filmtabletten	NL/H/1115/001	1-27894	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Forzaten 20 mg/5 mg comprimés pelliculés	NL/H/1115/001	BE325455	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Amelior® 40 mg/5 mg Filmtabletten	NL/H/1115/002	1-27895	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 40 mg/5 mg	NL/H/1115/002	20613	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/18	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 20 mg/5 mg	NL/H/1115/001	13403	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947317	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Konverge 40 mg/5 mg film-coated tablets	NL/H/1115/002	PA 865/17/2	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Sanoral, 20 mg/5 mg õhukese polümeerikattega tabletid	NL/H/1115/001	606208	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Sintonyn 40 mg/5 mg potahované tablety	NL/H/1115/002	58/527/08-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 20 mg/5 mg filmtabletta	NL/H/1115/001	OGYI-T-20884/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/09	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/19	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Forzaten 40mg/5mg, comprimés pelliculés	NL/H/1115/002	BE325464	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Επικαλυμμένα με λεπτό υμένιο δισκία Orizal 20 mg/5 mg	NL/H/1115/001	20612	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Sanoral 40 mg/5 mg plėvele dengtos tabletės	NL/H/1115/002	LT/1/08/1363/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
BIVIS 20 mg/5 mg compresse rivestite con film	NL/H/1115/001	038947053	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Sanoral 20 mg/5 mg apvalkotās tabletes	NL/H/1115/001	08-0270	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV

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
Теспадан 40 mg/10 mg филмирани таблетки	NL/H/1115/003	20080198	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947242	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947293	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947329	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947255	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947267	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947331	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947281	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
BIVIS 40 mg/5 mg compresse rivestite con film	NL/H/1115/002	038947180	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947279	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/23	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Duactan 40 mg/10 mg filmtabletta	NL/H/1115/003	OGYI-T-20884/24	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Теспадан 40 mg/5 mg филмирани таблетки	NL/H/1115/002	20080197	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Forzaten 40 mg/10 mg, comprimés pelliculés	NL/H/1115/003	BE325473	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Balzak 40 mg /5 mg comprimidos recubiertos con película	NL/H/1115/002	70.094	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Zolnor 40 mg + 5 mg comprimidos revestidos por película	NL/H/1115/002	5148333	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor 40 mg + 10 mg comprimidos revestidos por película	NL/H/1115/003	5148374	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
BIVIS 40 mg/10 mg compresse rivestite con film	NL/H/1115/003	038947305	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
Sanoral 20 mg/5 mg plėvele dengtos tabletės	NL/H/1115/001	LT/1/08/1363/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 20 mg/5 mg plėvele dengtos tabletės	NL/H/1115/001	LT/1/08/1363/001	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 20 mg/5 mg plėvele dengtos tabletės	NL/H/1115/001	LT/1/08/1363/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 20 mg/5 mg plėvele dengtos tabletės	NL/H/1115/001	LT/1/08/1363/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 40 mg/10 mg plėvele dengtos tabletės	NL/H/1115/003	LT/1/08/1363/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 40 mg/10 mg plėvele dengtos tabletės	NL/H/1115/003	LT/1/08/1363/011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 40 mg/10 mg plėvele dengtos tabletės	NL/H/1115/003	LT/1/08/1363/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 40 mg/10 mg plėvele dengtos tabletės	NL/H/1115/003	LT/1/08/1363/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Zolnor 20 mg + 5 mg comprimidos revestidos por película	NL/H/1115/001	5148275	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Forzaten 40 mg/5 mg comprimés pelliculés	NL/H/1115/002	2008120008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Konverge 40 mg/5 mg film-coated tablets	NL/H/1115/002	MA204/00402	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Zolnor 20 mg + 5 mg comprimidos revestidos por película	NL/H/1115/001	5148317	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor 20 mg + 5 mg comprimidos revestidos por película	NL/H/1115/001	5148309	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Sanoral 40 mg/5 mg plėvele dengtos tabletės	NL/H/1115/002	LT/1/08/1363/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Belfor 20mg/5mg, filmomhulde tabletten	NL/H/1115/001	RVG 100993	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Sanoral 40 mg/5 mg plėvele dengtos tabletės	NL/H/1115/002	LT/1/08/1363/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Sanoral 40 mg/5 mg plėvele dengtos tabletės	NL/H/1115/002	LT/1/08/1363/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Folgan 40 mg/10 mg filmom obalené tablety	NL/H/1115/003	58/0421/08-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Konverge 40 mg/10 mg film-coated tablets	NL/H/1115/003	MA204/00403	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Belfor 40mg/10mg, filmomhulde tabletten	NL/H/1115/003	RVG 100995	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Elestar, 40 mg + 5 mg, tabletki powlekane	NL/H/1115/002	16194	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Elestar, 20 mg + 5 mg, tabletki powlekane	NL/H/1115/001	16195	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Belfor 40mg/5mg, filmomhulde tabletten	NL/H/1115/002	RVG 100994	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	NL
Zolnor 40 mg + 10 mg comprimidos revestidos por película	NL/H/1115/003	5148366	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor 40 mg + 5 mg comprimidos revestidos por película	NL/H/1115/002	5148325	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor 40 mg + 10 mg comprimidos revestidos por película	NL/H/1115/003	5148358	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Zolnor 40 mg + 5 mg comprimidos revestidos por película	NL/H/1115/002	5148341	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Folgan 20 mg/5 mg filmom obalené tablety	NL/H/1115/001	58/0419/08-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Forzaten 40 mg/10 mg, comprimés pelliculés	NL/H/1115/003	2008120009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Konverge 20 mg/5 mg film-coated tablets	NL/H/1115/001	MA204/00401	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Forzaten 20 mg/5 mg comprimés pelliculés	NL/H/1115/001	2008120007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Elestar, 40 mg + 10 mg, tabletki powlekane	NL/H/1115/003	16193	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Duactan 40 mg/5 mg filmtabletta	NL/H/1115/002	OGYI-T-20884/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Olmesartan/Amlodipine Krka 20 mg/5 mg potahované tablety	CZ/H/0717/001	58/707/16-C	KRKA, D.D., NOVO MESTO	CZ
Olmesartan/Amlodipine Krka 40 mg/5 mg potahované tablety	CZ/H/0717/002	58/708/16-C	KRKA, D.D., NOVO MESTO	CZ
Olmesartan/Amlodipine Krka 40 mg/10 mg potahované tablety	CZ/H/0717/003	58/709/16-C	KRKA, D.D., NOVO MESTO	CZ

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Olmesartan/Amlodipine Krka d.d. Novo mesto 20 mg/5 mg filmomhulde tabletten	CZ/H/0717/001	BE514880	KRKA, D.D., NOVO MESTO	BE
Olmesartan/Amlodipine Krka d.d. Novo mesto 40 mg/5 mg filmomhulde tabletten	CZ/H/0717/002	BE514897	KRKA, D.D., NOVO MESTO	BE
Olmesartan/Amlodipine Krka d.d. Novo mesto 40 mg/10 mg filmomhulde tabletten	CZ/H/0717/003	BE514906	KRKA, D.D., NOVO MESTO	BE
Olmesartán/Amlodipino Krka 20 mg /5 mg comprimidos recubiertos con película	CZ/H/0717/001	82464	KRKA, D.D., NOVO MESTO	ES
Olmesartán/Amlodipino Krka 40 mg /5 mg comprimidos recubiertos con película	CZ/H/0717/002	82465	KRKA, D.D., NOVO MESTO	ES
Olmesartán/Amlodipino Krka 40 mg /10 mg comprimidos recubiertos con película	CZ/H/0717/003	82466	KRKA, D.D., NOVO MESTO	ES
Capenon 20 mg/5 mg, filmomhulde tabletten	NL/H/1114/001	RVG 100989	DAIICHI SANKYO EUROPE GMBH	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
Capenon 40 mg/5 mg, filmomhulde tabletten	NL/H/1114/002	RVG 100990	DAIICHI SANKYO EUROPE GMBH	NL
Capenon 40 mg/10 mg, filmomhulde tabletten	NL/H/1114/003	RVG 100991	DAIICHI SANKYO EUROPE GMBH	NL
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946075	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946176	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946137	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946152	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946188	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946101	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946125	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946087	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946099	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946051	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946036	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946048	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/5 mg compresse rivestite con film	NL/H/1114/002	038946113	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946149	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.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
GIANT 20 mg/5 mg compresse rivestite con film	NL/H/1114/001	038946012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
GIANT 40 mg/10 mg compresse rivestite con film	NL/H/1114/003	038946164	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Capenon 20 mg /5 mg comprimidos recubiertos con película	NL/H/1114/001	70.067	DAIICHI SANKYO ESPAÑA, S.A.	ES
Capenon 40 mg /5 mg comprimidos recubiertos con película	NL/H/1114/002	70.070	DAIICHI SANKYO ESPAÑA, S.A.	ES
Capenon 40 mg /10 mg comprimidos recubiertos con película	NL/H/1114/003	70.071	DAIICHI SANKYO ESPAÑA, S.A.	ES
Olmedipin 40 mg/5 mg Filmtabletten	DE/H/5342/002	2200820.00.00	RATIOPHARM GMBH	DE
Olmedipin 40 mg/10 mg Filmtabletten	DE/H/5342/003	2200821.00.00	RATIOPHARM GMBH	DE
Olmedipin 20 mg/5 mg Filmtabletten	DE/H/5342/001	2200819.00.00	RATIOPHARM GMBH	DE
Sevikar 20mg/5mg, comprimés pelliculés	NL/H/1113/001	BE325482	DAIICHI SANKYO BELGIUM S.A	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
Sevikar 40mg/5mg, comprimés pelliculés	NL/H/1113/002	BE325491	DAIICHI SANKYO BELGIUM S.A	BE
Sevikar 40mg/10mg, comprimés pelliculés	NL/H/1113/003	BE325507	DAIICHI SANKYO BELGIUM S.A	BE
SEVIKAR 20 mg/5 mg Filmtabletten	NL/H/1113/001	1-27891	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR 40 mg/5 mg Filmtabletten	NL/H/1113/002	1-27892	DAIICHI SANKYO AUSTRIA GMBH	AT
SEVIKAR 40 mg/10 mg Filmtabletten	NL/H/1113/003	1-27893	DAIICHI SANKYO AUSTRIA GMBH	AT
Sevikar 20 mg/5 mg, filmomhulde tabletten	NL/H/1113/001	BE325482	DAIICHI SANKYO BELGIUM S.A	BE
Sevikar 40 mg/5 mg, filmomhulde tabletten	NL/H/1113/002	BE325491	DAIICHI SANKYO BELGIUM S.A	BE
Sevikar 40 mg/10 mg, filmomhulde tabletten	NL/H/1113/003	BE325507	DAIICHI SANKYO BELGIUM S.A	BE
Sevikar 20 mg/5 mg tabletti, kalvopäällysteinen	NL/H/1113/001	24129	DAIICHI SANKYO EUROPE GMBH	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
Sevikar 40 mg/5 mg tabletti, kalvopäällysteinen	NL/H/1113/002	24130	DAIICHI SANKYO EUROPE GMBH	FI
Sevikar 40 mg/10 mg tabletti, kalvopäällysteinen	NL/H/1113/003	24131	DAIICHI SANKYO EUROPE GMBH	FI
Sevikar 20 mg/5 mg tablett, filmdrasjert	NL/H/1113/001	07-5221	DAIICHI SANKYO EUROPE GMBH	NO
Sevikar 40 mg/5 mg tablett, filmdrasjert	NL/H/1113/002	07-5222	DAIICHI SANKYO EUROPE GMBH	NO
Sevikar 40 mg/10 mg tablett, filmdrasjert	NL/H/1113/003	07-5223	DAIICHI SANKYO EUROPE GMBH	NO
SEVIKAR 20 mg/5 mg Filmtabletten	NL/H/1113/001	70001.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR 40 mg/10 mg Filmtabletten	NL/H/1113/003	70003.00.00	DAIICHI SANKYO EUROPE GMBH	DE
SEVIKAR 40 mg/5 mg Filmtabletten	NL/H/1113/002	70002.00.00	DAIICHI SANKYO EUROPE GMBH	DE
Sevikar, filmovertrukne tabletter	NL/H/1113/001	42061	DAIICHI SANKYO EUROPE GMBH	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
Sevikar, filmovertrukne tabletter	NL/H/1113/002	42062	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar, filmovertrukne tabletter	NL/H/1113/003	42063	DAIICHI SANKYO EUROPE GMBH	DK
Sevikar 20mg/5mg, comprimés pelliculés	NL/H/1113/001	1851/08 11 0049	DAIICHI SANKYO BELGIUM S.A	LU
Sevikar 40mg/5mg, comprimés pelliculés	NL/H/1113/002	1851/08 11 0050	DAIICHI SANKYO BELGIUM S.A	LU
Sevikar 40mg/10mg, comprimés pelliculés	NL/H/1113/003	1851/08 11 0051	DAIICHI SANKYO BELGIUM S.A	LU
Sevikar 20 mg/5 mg filmuhúðaðar töflur	NL/H/1113/001	IS/1/08/005/01	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar 40 mg/5 mg filmuhúðaðar töflur	NL/H/1113/002	IS/1/08/005/02	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar 40 mg/10 mg filmuhúðaðar töflur	NL/H/1113/003	IS/1/08/005/03	DAIICHI SANKYO EUROPE GMBH	IS
Sevikar 20 mg/5 mg film-coated tablets	NL/H/1113/001	PA 1595/005/001	DAIICHI SANKYO IRELAND LIMITED	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Sevikar 40 mg/5 mg film-coated tablets	NL/H/1113/002	PA 1595/005/002	DAIICHI SANKYO IRELAND LIMITED	IE
Sevikar 40 mg/10 mg film-coated tablets	NL/H/1113/003	PA 1595/005/003	DAIICHI SANKYO IRELAND LIMITED	IE
Sevikar 20 mg/5 mg, filmomhulde tabletten	NL/H/1113/001	RVG 100984	DAIICHI SANKYO NEDERLAND B.V.	NL
Sevikar 40 mg/5 mg, filmomhulde tabletten	NL/H/1113/002	RVG 100986	DAIICHI SANKYO NEDERLAND B.V.	NL
Sevikar 40 mg/10 mg, filmomhulde tabletten	NL/H/1113/003	RVG 100987	DAIICHI SANKYO NEDERLAND B.V.	NL
SEVIKAR 20 mg/5 mg, comprimé pelliculé	NL/H/1113/001	388 582-6	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 20 mg/5 mg, comprimé pelliculé	NL/H/1113/001	573 867-3	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 20 mg/5 mg, comprimé pelliculé	NL/H/1113/001	388 584-9	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 40 mg/10 mg, comprimé pelliculé	NL/H/1113/003	388 574-3	DAIICHI SANKYO FRANCE 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
SEVIKAR 40 mg/10 mg, comprimé pelliculé	NL/H/1113/003	573 871-0	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 40 mg/10 mg, comprimé pelliculé	NL/H/1113/003	388 577-2	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 40 mg/5 mg, comprimé pelliculé	NL/H/1113/002	388 578-9	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 40 mg/5 mg, comprimé pelliculé	NL/H/1113/002	573 870-4	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 40 mg/5 mg, comprimé pelliculé	NL/H/1113/002	388 580-3	DAIICHI SANKYO FRANCE SAS	FR
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983019	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983021	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983033	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983045	DAIICHI SANKYO ITALIA 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
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983058	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983060	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983072	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983084	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983096	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983108	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 20 mg/5 mg compresse rivestite con film	NL/H/1113/001	038983110	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983159	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983161	DAIICHI SANKYO ITALIA 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
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983173	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983185	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983197	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983209	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983211	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983223	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983146	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983134	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/5 mg compresse rivestite con film	NL/H/1113/002	038983122	DAIICHI SANKYO ITALIA 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
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983235	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983247	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983250	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983262	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983274	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983286	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983298	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983300	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983312	DAIICHI SANKYO ITALIA 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
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983324	DAIICHI SANKYO ITALIA S.P.A	IT
SEVIKAR 40 mg/10 mg compresse rivestite con film	NL/H/1113/003	038983336	DAIICHI SANKYO ITALIA S.P.A	IT
Sevikar 20 mg + 5 mg comprimidos revestidos por película	NL/H/1113/001	5148168	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 20 mg + 5 mg comprimidos revestidos por película	NL/H/1113/001	5148176	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 20 mg + 5 mg comprimidos revestidos por película	NL/H/1113/001	5148200	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 40 mg + 5 mg comprimidos revestidos por película	NL/H/1113/002	5148218	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 40 mg + 5 mg comprimidos revestidos por película	NL/H/1113/002	5148226	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 40 mg + 5 mg comprimidos revestidos por película	NL/H/1113/002	5148234	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 40 mg + 10 mg comprimidos revestidos por película	NL/H/1113/003	5148242	DAIICHI SANKYO PORTUGAL, UNIP. 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
Sevikar 40 mg + 10 mg comprimidos revestidos por película	NL/H/1113/003	5148259	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 40 mg + 10 mg comprimidos revestidos por película	NL/H/1113/003	5148267	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Sevikar 20 mg /5 mg comprimidos recubiertos con película	NL/H/1113/001	70.079	DAIICHI SANKYO ESPAÑA, S.A.	ES
Sevikar 40 mg /5 mg comprimidos recubiertos con película	NL/H/1113/002	70.072	DAIICHI SANKYO ESPAÑA, S.A.	ES
Sevikar 40 mg /10 mg comprimidos recubiertos con película	NL/H/1113/003	70.069	DAIICHI SANKYO ESPAÑA, S.A.	ES
Sevikar 20 mg/5 mg film-coated tablets	NL/H/1113/001	PL 08265/0026	DAIICHI SANKYO UK LTD	XI
Sevikar 40 mg/5 mg film-coated tablets	NL/H/1113/002	PL 08265/0027	DAIICHI SANKYO UK LTD	XI
Sevikar 40 mg/10 mg film-coated tablets	NL/H/1113/003	PL 08265/0028	DAIICHI SANKYO UK LTD	XI
Sevikar 40 mg/5 mg filmdragerade tabletter	NL/H/1113/002	24130	DAIICHI SANKYO EUROPE GMBH	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
Sevikar 40 mg/10 mg filmdragerade tabletter	NL/H/1113/003	24131	DAIICHI SANKYO EUROPE GMBH	FI
Sevikar 20 mg/5 mg filmdragerade tabletter	NL/H/1113/001	24129	DAIICHI SANKYO EUROPE GMBH	FI
Olmesartan/Amlodipine Krka 20 mg/5 mg potahované tablety	CZ/H/0717/001	58/707/16-C	KRKA, D.D., NOVO MESTO	CZ
Olmesartan/Amlodipine Krka 40 mg/5 mg potahované tablety	CZ/H/0717/002	58/708/16-C	KRKA, D.D., NOVO MESTO	CZ
Olmesartan/Amlodipine Krka 40 mg/10 mg potahované tablety	CZ/H/0717/003	58/709/16-C	KRKA, D.D., NOVO MESTO	CZ
Olmesartan/Amlodipine Krka d.d. Novo mesto 20 mg/5 mg filmomhulde tabletten	CZ/H/0717/001	BE514880	KRKA, D.D., NOVO MESTO	BE
Olmesartan/Amlodipine Krka d.d. Novo mesto 40 mg/5 mg filmomhulde tabletten	CZ/H/0717/002	BE514897	KRKA, D.D., NOVO MESTO	BE
Olmesartan/Amlodipine Krka d.d. Novo mesto 40 mg/10 mg filmomhulde tabletten	CZ/H/0717/003	BE514906	KRKA, D.D., NOVO MESTO	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
Olmesartán/Amlodipino Krka 20 mg /5 mg comprimidos recubiertos con película	CZ/H/0717/001	82464	KRKA, D.D., NOVO MESTO	ES
Olmesartán/Amlodipino Krka 40 mg /5 mg comprimidos recubiertos con película	CZ/H/0717/002	82465	KRKA, D.D., NOVO MESTO	ES
Olmesartán/Amlodipino Krka 40 mg /10 mg comprimidos recubiertos con película	CZ/H/0717/003	82466	KRKA, D.D., NOVO MESTO	ES