10 June 2022
EMA/586078/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: olmesartan

Procedure no.: PSUSA/00002207/202110

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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
Тенсар 10 mg филмирани таблетки	not available	20050545	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Тенсар 20 mg филмирани таблетки	not available	20050546	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Тенсар 40 mg филмирани таблетки	not available	20050547	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BG
Mesar 10 mg plėvele dengtos tabletės	not available	LT/1/04/3171/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 10 mg plėvele dengtos tabletės	not available	LT/1/04/3171/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 10 mg plėvele dengtos tabletės	not available	LT/1/04/3171/001	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 10 mg plėvele dengtos tabletės	not available	LT/1/04/3171/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 40 mg plėvele dengtos tabletės	not available	LT/1/04/3171/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 20 mg plėvele dengtos tabletės	not available	LT/1/04/3171/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 20 mg plėvele dengtos tabletės	not available	LT/1/04/3171/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 40 mg plėvele dengtos tabletės	not available	LT/1/04/3171/011	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Mesar 20 mg plėvele dengtos tabletės	not available	LT/1/04/3171/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 40 mg plėvele dengtos tabletės	not available	LT/1/04/3171/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 40 mg plėvele dengtos tabletės	not available	LT/1/04/3171/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 20 mg plėvele dengtos tabletės	not available	LT/1/04/3171/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LT
Mesar 10 mg apvalkotās tabletes	not available	04-0062	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Mesar 20 mg apvalkotās tabletes	not available	04-0063	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Mesar 40 mg apvalkotās tabletes	not available	04-0064	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LV
Olmesartan G.L. 5 mg-Filmtabletten	not available	139893	G.L. PHARMA GMBH	AT
MESAR 20 mg õhukese polümeerikattega tabletid	not available	424703	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
MESAR 40 mg õhukese polümeerikattega tabletid	not available	424803	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
MESAR 10 mg õhukese polümeerikattega tabletid	not available	424603	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	EE
Olmeblo 5 mg-Filmtabletten	EE/H/0716/001	137112	G.L. PHARMA GMBH	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Olmetec 10 mg, Filmtabletten	DE/H/0384/001	50202.00.00	DAIICHI SANKYO EUROPE GMBH	DE
Olmetec 20 mg, Filmtabletten	DE/H/0384/002	50202.01.00	DAIICHI SANKYO EUROPE GMBH	DE
Olmetec 40 mg, Filmtabletten	DE/H/0384/003	50202.02.00	DAIICHI SANKYO EUROPE GMBH	DE
Olmetec 10 mg comprimés pelliculés	DE/H/0384/001	BE254852	DAIICHI SANKYO BELGIUM S.A	BE
Olmetec 20 mg comprimés pelliculés	DE/H/0384/002	BE254861	DAIICHI SANKYO BELGIUM S.A	BE
Olmetec 40 mg comprimés pelliculés	DE/H/0384/003	BE254877	DAIICHI SANKYO BELGIUM S.A	BE
Olmetec 40 mg Filmtabletten	DE/H/0384/003	1-24928	DAIICHI SANKYO AUSTRIA GMBH	AT
Olmetec 10 mg Filmtabletten	DE/H/0384/001	1-24926	DAIICHI SANKYO AUSTRIA GMBH	AT
Olmetec 20 mg Filmtabletten	DE/H/0384/002	1-24927	DAIICHI SANKYO AUSTRIA GMBH	AT
OLMETEC 20 mg compresse rivestite con film	DE/H/0384/002	036027062	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 20 mg compresse rivestite con film	DE/H/0384/002	036027074	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 20 mg compresse rivestite con film	DE/H/0384/002	036027086	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 20 mg compresse rivestite con film	DE/H/0384/002	036027098	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 20 mg compresse rivestite con film	DE/H/0384/002	036027100	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
OLMETEC 40 mg compresse rivestite con film	DE/H/0384/003	036027112	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 40 mg compresse rivestite con film	DE/H/0384/003	036027124	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 40 mg compresse rivestite con film	DE/H/0384/003	036027136	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 40 mg compresse rivestite con film	DE/H/0384/003	036027148	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 40 mg compresse rivestite con film	DE/H/0384/003	036027151	DAIICHI SANKYO ITALIA S.P.A	IT
Olmetec 20 mg tabletti, kalvopäällysteinen	DE/H/0384/002	17890	DAIICHI SANKYO EUROPE GMBH	FI
Olmetec, filmovertrukne tabletter	DE/H/0384/001	34529	DAIICHI SANKYO EUROPE GMBH	DK
Olmetec, filmovertrukne tabletter	DE/H/0384/002	34530	DAIICHI SANKYO EUROPE GMBH	DK
Olmetec, filmovertrukne tabletter	DE/H/0384/003	34531	DAIICHI SANKYO EUROPE GMBH	DK
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4729083	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4729182	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4729281	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
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4729380	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4822888	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4841185	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4841284	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 20 mg comprimidos revestidos por película	DE/H/0384/002	4841383	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4729489	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4729588	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4729687	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4729786	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4822987	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4841482	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
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4841581	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 40 mg comprimidos revestidos por película	DE/H/0384/003	4841680	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg, filmomhulde tabletten	DE/H/0384/001	RVG 28782	DAIICHI SANKYO NEDERLAND B.V.	NL
Olmetec 20 mg, filmomhulde tabletten	DE/H/0384/002	RVG 28783	DAIICHI SANKYO NEDERLAND B.V.	NL
Olmetec 40 mg, filmomhulde tabletten	DE/H/0384/003	RVG 28784	DAIICHI SANKYO NEDERLAND B.V.	NL
Olmetec 10 mg comprimidos recubiertos	DE/H/0384/001	65.492	DAIICHI SANKYO ESPAÑA, S.A.	ES
Olmetec 20 mg comprimidos recubiertos	DE/H/0384/002	65.496	DAIICHI SANKYO ESPAÑA, S.A.	ES
Olmetec 40 mg comprimidos recubiertos	DE/H/0384/003	65.497	DAIICHI SANKYO ESPAÑA, S.A.	ES
Olmetec 10 mg film-coated tablets	DE/H/0384/001	PL 08265/0015	DAIICHI SANKYO UK LTD	XI
Olmetec 20 mg film-coated tablets	DE/H/0384/002	PL 08265/0016	DAIICHI SANKYO UK LTD	XI
Olmetec 40 mg film-coated tablets	DE/H/0384/003	PL 08265/0017	DAIICHI SANKYO UK LTD	XI
Olmetec 10 mg filmuhúðaðar töflur	DE/H/0384/001	IS/1/03/011/01	DAIICHI SANKYO EUROPE GMBH	IS
Olmetec 20 mg filmuhúðaðar töflur	DE/H/0384/002	IS/1/03/011/02	DAIICHI SANKYO EUROPE GMBH	IS
Olmetec 40 mg filmuhúðaðar töflur	DE/H/0384/003	IS/1/03/011/03	DAIICHI SANKYO EUROPE GMBH	IS
Olmetec 10 mg comprimés pelliculés	DE/H/0384/001	1413/03/11/0020	DAIICHI SANKYO BELGIUM S.A	LU
Olmetec 20 mg comprimés pelliculés	DE/H/0384/002	1413/03/11/0021	DAIICHI SANKYO BELGIUM S.A	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
Olmetec 40 mg comprimés pelliculés	DE/H/0384/003	1413/03/11/0022	DAIICHI SANKYO BELGIUM S.A	LU
Olmetec 10 mg tabletter, filmdrasjerte	DE/H/0384/001	02-1565	DAIICHI SANKYO EUROPE GMBH	NO
Olmetec 20 mg tabletter, filmdrasjerte	DE/H/0384/002	02-1567	DAIICHI SANKYO EUROPE GMBH	NO
Olmetec 40 mg tabletter, filmdrasjerte	DE/H/0384/003	02-1568	DAIICHI SANKYO EUROPE GMBH	NO
Olmetec 40 mg tabletti, kalvopäällysteinen	DE/H/0384/003	17891	DAIICHI SANKYO EUROPE GMBH	FI
Olmetec 10 mg tabletti, kalvopäällysteinen	DE/H/0384/001	17889	DAIICHI SANKYO EUROPE GMBH	FI
OLMETEC 10 mg compresse rivestite con film	DE/H/0384/001	036027011	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 10 mg compresse rivestite con film	DE/H/0384/001	036027023	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 10 mg compresse rivestite con film	DE/H/0384/001	036027035	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 10 mg compresse rivestite con film	DE/H/0384/001	036027047	DAIICHI SANKYO ITALIA S.P.A	IT
OLMETEC 10 mg compresse rivestite con film	DE/H/0384/001	036027050	DAIICHI SANKYO ITALIA S.P.A	IT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4728689	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4728788	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
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4728887	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4728986	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4822789	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4840880	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4840989	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Olmetec 10 mg comprimidos revestidos por película	DE/H/0384/001	4841086	DAIICHI SANKYO PORTUGAL, UNIP. LDA	PT
Benetor 10 mg film-coated tablets	DE/H/0384/001	PA 1595/1/1	DAIICHI SANKYO IRELAND LIMITED	IE
Benetor 20 mg film-coated tablets	DE/H/0384/002	PA 1595/1/2	DAIICHI SANKYO IRELAND LIMITED	IE
Benetor 40 mg film-coated tablets	DE/H/0384/003	PA 1595/1/3	DAIICHI SANKYO IRELAND LIMITED	IE
Olmetec 10 mg filmomhulde tabletten	DE/H/0384/001	BE254852	DAIICHI SANKYO BELGIUM S.A	BE
Olmetec 20 mg filmomhulde tabletten	DE/H/0384/002	BE254861	DAIICHI SANKYO BELGIUM S.A	BE
Olmetec 40 mg filmomhulde tabletten	DE/H/0384/003	BE254877	DAIICHI SANKYO BELGIUM S.A	BE
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 565 329 6 4	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 362 323 3 4	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
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 374 016 3 0	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 362 322 7 3	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 374 014 0 1	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 362 321 0 5	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 362 320 4 4	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 10 mg, comprimé pelliculé	DE/H/0384/001	34009 374 015 7 9	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 565 330 4 6	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 362 328 5 3	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 374 020 0 2	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 374 019 2 0	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 362 327 9 2	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 374 018 6 9	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 362 326 2 4	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 20 mg, comprimé pelliculé	DE/H/0384/002	34009 362 325 6 3	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 362 329 1 4	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 565 331 0 7	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 362 333 9 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
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 374 024 6 0	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 374 022 3 1	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 362 332 2 5	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 374 021 7 0	DAIICHI SANKYO FRANCE SAS	FR
OLMETEC 40 mg, comprimé pelliculé	DE/H/0384/003	34009 362 331 6 4	DAIICHI SANKYO FRANCE SAS	FR
Olmetec 20 mg filmdragerade tabletter	DE/H/0384/002	17890	DAIICHI SANKYO EUROPE GMBH	FI
Olmetec 40 mg filmdragerade tabletter	DE/H/0384/003	17891	DAIICHI SANKYO EUROPE GMBH	FI
Olmetec 10 mg filmdragerade tabletter	DE/H/0384/001	17889	DAIICHI SANKYO EUROPE GMBH	FI
Belsar 40 mg filmomhulde tabletten	DE/H/0386/003	BE256995	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 10 mg Filmtabletten	DE/H/0386/001	BE256977	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 20 mg filmomhulde tabletten	DE/H/0386/002	BE256986	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 40 mg Filmtabletten	DE/H/0386/003	BE256995	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 20 mg Filmtabletten	DE/H/0386/002	BE256986	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 10 mg filmomhulde tabletten	DE/H/0386/001	BE256977	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
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/027	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/021	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/004	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/010	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/033	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/022	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/007	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/016	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/032	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/025	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/011	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
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/020	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/029	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/008	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/026	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/003	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/031	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/028	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/006	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/009	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/018	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/030	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
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/015	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/019	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/005	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/014	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Mencord® 40 mg Filmtabletten	DE/H/0386/003	1-24932	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Mencord® 20 mg Filmtabletten	DE/H/0386/002	1-24931	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT
Belsar 40 mg comprimés pelliculés	DE/H/0386/003	BE256995	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 10 mg comprimés pelliculés	DE/H/0386/001	BE256977	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Belsar 20 mg comprimés pelliculés	DE/H/0386/002	BE256986	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	BE
Mencord® 10 mg Filmtabletten	DE/H/0386/001	1-24930	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	AT

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Olartan 10 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/001	19688	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Olartan 20 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/002	19689	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Olartan 40 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/003	19690	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CY
Sarten 10 mg potahované tablety	DE/H/0386/001	58/074/05-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sarten 40 mg potahované tablety	DE/H/0386/003	58/076/05-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
Sarten 20 mg potahované tablety	DE/H/0386/002	58/075/05-C	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	CZ
OLPRESS 10 mg compresse rivestite con film	DE/H/0386/001	036026019	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Belsar 40 mg comprimés pelliculés	DE/H/0386/003	0951/09010080	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
Omesar 10 mg film-coated tablets	DE/H/0386/001	MA204/00301	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Omesar 20 mg film-coated tablets	DE/H/0386/002	MA204/00302	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
Belsar 20 mg comprimés pelliculés	DE/H/0386/002	0951/09010079	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	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
Omesar 40 mg film-coated tablets	DE/H/0386/003	MA204/00303	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	MT
OLPRESS 10 mg compresse rivestite con film	DE/H/0386/001	036026033	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 10 mg compresse rivestite con film	DE/H/0386/001	036026045	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 10 mg compresse rivestite con film	DE/H/0386/001	036026021	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 10 mg compresse rivestite con film	DE/H/0386/001	036026058	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 20 mg compresse rivestite con film	DE/H/0386/002	036026060	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 20 mg compresse rivestite con film	DE/H/0386/002	036026072	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 20 mg compresse rivestite con film	DE/H/0386/002	036026084	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 20 mg compresse rivestite con film	DE/H/0386/002	036026096	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 20 mg compresse rivestite con film	DE/H/0386/002	036026108	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 362 947 7 6	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
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 362 945 4 7	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 362 946 0 8	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 372 068 6 0	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 372 066 3 1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 372 069 2 1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 362 944 8 6	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 10 mg, comprimé pelliculé	DE/H/0386/001	34009 565 394 2 0	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 362 953 7 7	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Olartan 10 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/001	42802	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Olartan 40 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/003	42804	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
OLPRESS 40 mg compresse rivestite con film	DE/H/0386/003	036026110	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
OLPRESS 40 mg compresse rivestite con film	DE/H/0386/003	036026134	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 40 mg compresse rivestite con film	DE/H/0386/003	036026146	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 40 mg compresse rivestite con film	DE/H/0386/003	036026159	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
OLPRESS 40 mg compresse rivestite con film	DE/H/0386/003	036026122	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
Belsar 10 mg comprimés pelliculés	DE/H/0386/001	0951/09010078	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	LU
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 362 948 3 7	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 372 070 0 3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 362 952 0 9	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 362 950 8 7	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 372 071 7 1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 372 072 3 2	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
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 362 951 4 8	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 20 mg, comprimé pelliculé	DE/H/0386/002	34009 565 395 9 8	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 362 954 3 8	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Votum® 40 mg, Filmtabletten	DE/H/0386/003	50208.02.00	BERLIN-CHEMIE AG	DE
Votum® 20 mg, Filmtabletten	DE/H/0386/002	50208.01.00	BERLIN-CHEMIE AG	DE
Olartan 20 mg επικαλυμμένο με λεπτό υμένιο δισκίο	DE/H/0386/002	42803	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	GR
Olsar 20 mg comprimidos revestidos por película	DE/H/0386/002	4730586	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/024	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 40 mg filmsko obložene tablete	DE/H/0386/003	H/05/01511/023	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Olsar 20 mg comprimidos revestidos por película	DE/H/0386/002	4730289	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Olsar 20 mg comprimidos revestidos por película	DE/H/0386/002	4730487	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 372 074 6 1	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
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 362 956 6 7	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 372 075 2 2	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 362 957 2 8	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 372 076 9 0	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Votum® 10 mg, Filmtabletten	DE/H/0386/001	50208.00.00	BERLIN-CHEMIE AG	DE
ALTEIS 40 mg, comprimé pelliculé	DE/H/0386/003	34009 565 396 5 9	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	FR
Olsar 40 mg comprimidos revestidos por película	DE/H/0386/003	4730982	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Ixia 20 mg comprimidos recubiertos	DE/H/0386/002	65.499	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olsar 40 mg comprimidos revestidos por película	DE/H/0386/003	4730685	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Ixia 10 mg comprimidos recubiertos	DE/H/0386/001	65.498	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Olsar 40 mg comprimidos revestidos por película	DE/H/0386/003	4730784	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
Olsar 40 mg comprimidos revestidos por película	DE/H/0386/003	4730883	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Omesar 40 mg film-coated tablets	DE/H/0386/003	PA 865/11/3	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Omesar 10 mg film-coated tablets	DE/H/0386/001	PA 865/11/1	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Olsar 20 mg comprimidos revestidos por película	DE/H/0386/002	4730388	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Olsar 10 mg comprimidos revestidos por película	DE/H/0386/001	4730081	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Olsar 10 mg comprimidos revestidos por película	DE/H/0386/001	4729984	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Olsar 10 mg comprimidos revestidos por película	DE/H/0386/001	4729885	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Olsar 10 mg comprimidos revestidos por película	DE/H/0386/001	4730180	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PT
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/013	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 20 mg filmsko obložene tablete	DE/H/0386/002	H/05/01511/012	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/001	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
Ixia 40 mg comprimidos recubiertos	DE/H/0386/003	65.500	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	ES
Tensiol 10 mg filmsko obložene tablete	DE/H/0386/001	H/05/01511/002	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SI
Omesar 20 mg film-coated tablets	DE/H/0386/002	PA 865/11/2	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IE
Revival, 20 mg, tabletki powlekane	DE/H/0386/002	11403	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Revival, 40 mg, tabletki powlekane	DE/H/0386/003	11402	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Revival, 10 mg, tabletki powlekane	DE/H/0386/001	11404	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	PL
Olmesartan medoxomil 10mg Film coated-tablets	not available	PL 20416/0540	CRESCENT PHARMA LIMITED	XI
Olmesartan medoxomil 20mg Film coated-tablets	not available	PL 20416/0541	CRESCENT PHARMA LIMITED	XI
Olmesartan medoxomil 40mg Film coated-tablets	not available	PL 20416/0542	CRESCENT PHARMA LIMITED	XI
Santini 20 mg comprimate filmate	not available	11447/2019/02	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/04	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/04	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/01	BERLIN-CHEMIE AG	RO
Santini 20 mg comprimate filmate	not available	11447/2019/01	BERLIN-CHEMIE AG	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
Santini 10 mg comprimate filmate	not available	11446/2019/03	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/01	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/02	BERLIN-CHEMIE AG	RO
Santini 20 mg comprimate filmate	not available	11447/2019/03	BERLIN-CHEMIE AG	RO
Santini 20 mg comprimate filmate	not available	11447/2019/04	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/03	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/02	BERLIN-CHEMIE AG	RO
Olmetec 10 mg comprimate filmate	not available	12684/2019/02	LABORMED PHARMA S.A.	RO
Olmetec 10 mg comprimate filmate	not available	12684/2019/01	LABORMED PHARMA S.A.	RO
Olmetec 10 mg comprimate filmate	not available	12684/2019/03	LABORMED PHARMA S.A.	RO
Olmetec 10 mg comprimate filmate	not available	12684/2019/04	LABORMED PHARMA S.A.	RO
Olmetec 20 mg comprimate filmate	not available	12685/2019/01	LABORMED PHARMA S.A.	RO
Olmetec 20 mg comprimate filmate	not available	12685/2019/02	LABORMED PHARMA S.A.	RO
Olmetec 20 mg comprimate filmate	not available	12685/2019/03	LABORMED PHARMA S.A.	RO
Olmetec 20 mg comprimate filmate	not available	12685/2019/04	LABORMED PHARMA S.A.	RO
Olmetec 40 mg comprimate filmate	not available	12686/2019/01	LABORMED PHARMA S.A.	RO
Olmetec 40 mg comprimate filmate	not available	12686/2019/02	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
Olmetec 40 mg comprimate filmate	not available	12686/2019/03	LABORMED PHARMA S.A.	RO
Olmetec 40 mg comprimate filmate	not available	12686/2019/04	LABORMED PHARMA S.A.	RO
Laresin 40 mg filmtabletta	not available	OGYI-T-9810/14	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 40 mg filmtabletta	not available	OGYI-T-9810/12	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 40 mg filmtabletta	not available	OGYI-T-9810/13	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 40 mg filmtabletta	not available	OGYI-T-9810/11	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 40 mg filmtabletta	not available	OGYI-T-9810/15	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 20 mg filmtabletta	not available	OGYI-T-9810/08	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 20 mg filmtabletta	not available	OGYI-T-9810/10	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 20 mg filmtabletta	not available	OGYI-T-9810/06	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 20 mg filmtabletta	not available	OGYI-T-9810/07	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 20 mg filmtabletta	not available	OGYI-T-9810/09	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
Laresin 10 mg filmtabletta	not available	OGYI-T-9810/02	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 10 mg filmtabletta	not available	OGYI-T-9810/01	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 10 mg filmtabletta	not available	OGYI-T-9810/04	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 10 mg filmtabletta	not available	OGYI-T-9810/05	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Laresin 10 mg filmtabletta	not available	OGYI-T-9810/03	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	HU
Santini 20 mg comprimate filmate	not available	11447/2019/02	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/04	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/04	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/01	BERLIN-CHEMIE AG	RO
Santini 20 mg comprimate filmate	not available	11447/2019/01	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/03	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/01	BERLIN-CHEMIE AG	RO
Santini 10 mg comprimate filmate	not available	11446/2019/02	BERLIN-CHEMIE AG	RO
Santini 20 mg comprimate filmate	not available	11447/2019/03	BERLIN-CHEMIE AG	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
Santini 20 mg comprimate filmate	not available	11447/2019/04	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/03	BERLIN-CHEMIE AG	RO
Santini 40 mg comprimate filmate	not available	11448/2019/02	BERLIN-CHEMIE AG	RO
Tenzar 10 mg filmom obalené tablety	not available	58/0118/04-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Tenzar 40 mg filmom obalené tablety	not available	58/0120/04-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Tenzar 20 mg filmom obalené tablety	not available	58/0119/04-S	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	SK
Olmes 10 mg, Filmtabletten	DE/H/0385/001	50205.00.00	DAIICHI SANKYO EUROPE GMBH	DE
Olmes 20 mg, Filmtabletten	DE/H/0385/002	50205.01.00	DAIICHI SANKYO EUROPE GMBH	DE
Olmes 40 mg, Filmtabletten	DE/H/0385/003	50205.02.00	DAIICHI SANKYO EUROPE GMBH	DE
Openvas 10 mg comprimidos recubiertos	DE/H/0385/001	65.493	DAIICHI SANKYO ESPAÑA, S.A.	ES
Openvas 20 mg comprimidos recubiertos	DE/H/0385/002	65.494	DAIICHI SANKYO ESPAÑA, S.A.	ES
Openvas 40 mg comprimidos recubiertos	DE/H/0385/003	65.495	DAIICHI SANKYO ESPAÑA, S.A.	ES
PLAUNAC 10 mg compresse rivestite con film	DE/H/0385/001	036025017	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 10 mg compresse rivestite con film	DE/H/0385/001	036025043	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
PLAUNAC 10 mg compresse rivestite con film	DE/H/0385/001	036025029	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 10 mg compresse rivestite con film	DE/H/0385/001	036025031	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 10 mg compresse rivestite con film	DE/H/0385/001	036025056	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 20 mg compresse rivestite con film	DE/H/0385/002	036025068	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 20 mg compresse rivestite con film	DE/H/0385/002	036025094	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 20 mg compresse rivestite con film	DE/H/0385/002	036025082	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 20 mg compresse rivestite con film	DE/H/0385/002	036025070	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 20 mg compresse rivestite con film	DE/H/0385/002	036025106	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 40 mg compresse rivestite con film	DE/H/0385/003	036025118	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 40 mg compresse rivestite con film	DE/H/0385/003	036025144	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 40 mg compresse rivestite con film	DE/H/0385/003	036025132	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
PLAUNAC 40 mg compresse rivestite con film	DE/H/0385/003	036025157	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT
PLAUNAC 40 mg compresse rivestite con film	DE/H/0385/003	036025120	MENARINI INTERNATIONAL OPERATIONS LUXEMBOURG S.A.	IT