



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

10 April 2025
EMA/129679/2025
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: dexamfetamine

Procedure no.: PSUSA/00000986/202409



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
Adhesa 5 mg tabletten	not available	RVG 128087	ACE PHARMACEUTICALS BV	NL
Amfexa 10 mg Tablets	SE/H/1719/002	PL 11243/0023	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI
Amfexa 20 mg Tablets	SE/H/1719/003/DC	PL 11243/0024	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI
Amfexa 5 mg Tablets	SE/H/1719/001/DC	PL 11243/0021	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	XI
Attadex 10 mg tabletit	SE/H/2484/002	42950	ABBOXIA AB	FI
Attadex 10 mg tablett	SE/H/2484/002	42950	ABBOXIA AB	FI
Attadex 20 mg tabletit	SE/H/2484/003	42951	ABBOXIA AB	FI
Attadex 20 mg tablett	SE/H/2484/003	42951	ABBOXIA AB	FI
Attadex 5 mg tabletit	SE/H/2484/001	42949	ABBOXIA AB	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
Attadex 5 mg tablett	SE/H/2484/001	42949	ABBOXIA AB	FI
Attenfarm 10 mg tabletten	SE/H/2344/002	RVG 131135	ORIFARM HEALTHCARE A/S	NL
Attenfarm 10 mg tabletten	SE/H/2344/002	RVG 131135	ORIFARM HEALTHCARE A/S	NL
Attenfarm 10 mg tabletten	SE/H/2344/002	RVG 131135	ORIFARM HEALTHCARE A/S	NL
Attenfarm 10 mg tabletten	SE/H/2344/002	RVG 131135	ORIFARM HEALTHCARE A/S	NL
Attenfarm 20 mg tabletten	SE/H/2344/003	RVG 131136	ORIFARM HEALTHCARE A/S	NL
Attenfarm 20 mg tabletten	SE/H/2344/003	RVG 131136	ORIFARM HEALTHCARE A/S	NL
Attenfarm 20 mg tabletten	SE/H/2344/003	RVG 131136	ORIFARM HEALTHCARE A/S	NL
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	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
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	NL
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	NL
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	NL
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	NL
Attenfarm 5 mg tabletten	SE/H/2344/001	RVG 131133	ORIFARM HEALTHCARE A/S	NL
Attentin 10 mg tabletit	SE/H/1719/002	33554	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Attentin 10 mg Tablette	not available	94042.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Attentin 10 mg Tabletten	SE/H/1719/002/DC	2017040061	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Attentin 10 mg tabletter	SE/H/1719/002/DC	15-10771	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO

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
Attentin 10 mg tabletter	SE/H/1719/002/DC	53180	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Attentin 10 mg töflur	SE/H/1719/002	IS/1/17/018/02	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Attentin 20 mg tabletit	SE/H/1719/003/DC	33555	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FI
Attentin 20 mg Tablette	not available	94043.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Attentin 20 mg Tabletten	SE/H/1719/003/DC	2017040062	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Attentin 20 mg tabletter	SE/H/1719/003/DC	15-10772	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
Attentin 20 mg tabletter	SE/H/1719/003	53181	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Attentin 20 mg töflur	SE/H/1719/003/DC	IS/1/17/018/03	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Attentin 5 mg tabletit	SE/H/1719/001/DC	30066	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
Attentin 5 mg Tabletten	SE/H/1719/001	2014120350	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	LU
Attentin 5 mg tabletter	SE/H/1719/001/DC	11-8529	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NO
Attentin 5 mg tabletter	SE/H/1719/001/DC	46616	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	SE
Attentin 5 mg töflur.	SE/H/1719/001	IS/1/17/018/01	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	IS
Attentin 5 mg, Tablette	not available	74643.00.00	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DE
Attentin, tabletter 10 mg	SE/H/1719/002	56271	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Attentin, tabletter 20 mg	SE/H/1719/003/DC	56272	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Attentin, tabletter 5 mg	SE/H/1719/001	49372	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	DK
Deksamfetamin Waymade B.V. 5 mg tabletter	MT/H/0389/001/E/001	22-14964	WAYMADE B.V.	NO

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
Dexamfetamin Sea Pharma 10 mg tabletter	SE/H/2509/002	16116	SEA PHARMA AB	NO
Dexamfetamin Sea Pharma 10 mg tabletter.	SE/H/2509/002	61077	SEA PHARMA AB	SE
Dexamfetamin Sea Pharma 20 mg tabletter	SE/H/2509/003	16117	SEA PHARMA AB	NO
Dexamfetamin Sea Pharma 20 mg tabletter.	SE/H/2509/003	61078	SEA PHARMA AB	SE
Dexamfetamin Sea Pharma 5 mg tabletter	SE/H/2509/001	16115	SEA PHARMA AB	NO
Dexamfetamin Sea Pharma 5 mg tabletter.	SE/H/2509/001	61076	SEA PHARMA AB	SE
Dexamfetamin Waymade	MT/H/0389/001/E/001	63798	WAYMADE B.V.	SE
Dexamfetamine Sea Pharma 10 mg tabletten	SE/H/2509/002	RVG 133096	SEA PHARMA AB	NL
Dexamfetamine Sea Pharma 20 mg tabletten	SE/H/2509/003	RVG 133097	SEA PHARMA AB	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
Dexamfetamine Sea Pharma 5 mg tabletten	SE/H/2509/001	RVG 133093	SEA PHARMA AB	NL
Dexamfetamine Sulfate 1 mg/ ml Oral Solution	NO/H/0246/001	PL 00427/0282	ROSEMONT PHARMACEUTICALS LIMITED	XI
Dexamfetamine Sulfate 5 mg Tablets	not available	PL 25298/0152	BROWN & BURK UK LIMITED	XI
Dexamfetamine Sulfate 5 mg Tablets	not available	PL 25298/0152	BROWN & BURK UK LIMITED	XI
Dexamfetamine Sulfate 5 mg Tablets	not available	PL 00289/2278	TEVA UK LIMITED	XI
Dexamfetamine Sulfate 5 mg Tablets	MT/H/0389/001	PL06464/3115	WAYMADE PLC	XI
Dexamfetamine Sulfate ATNAHS 5 mg Tablets.	MT/H/0389/001	MA1409/00101	WAYMADE B.V.	MT
Dexamfetamine Taw Pharma 1 mg/ml mixtúra lausn	NO/H/0246/001	IS/1/15/060/01	TAW PHARMA (IRELAND) LTD	IS
Dexamfetamine Taw Pharma 1 mg/ml mikstur, oppløsning	NO/H/0246/001	13-9893	TAW PHARMA (IRELAND) LTD	NO

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
Dexamfetaminesulfaat DMB 5 mg, tabletten	not available	RVG 115446	TIOFARMA BV	NL
Dexamfetaminesulfaat Waymade 5 mg tabletten	MT/H/0389/001/E/001	RVG 130466	WAYMADE B.V.	NL
Dexamfetaminhemisulfat 5 mg Tabletten	MT/H/0389/001	7010987.00.00	WAYMADE B.V.	DE
Dexamfetaminsulfat "Waymade"	MT/H/0389/001/E/001	68351	WAYMADE B.V.	DK
Dexatin 10 mg tablett	SE/H/2484/002	63409	ABBOXIA AB	SE
Dexatin 10 mg tabletter	SE/H/2484/002	23-15744	ABBOXIA AB	NO
Dexatin 20 mg tablett	SE/H/2484/003	63410	ABBOXIA AB	SE
Dexatin 20 mg tabletter	SE/H/2484/003	23-15745	ABBOXIA AB	NO
Dexatin 5 mg tablett	SE/H/2484/001	63408	ABBOXIA AB	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dexatin 5 mg tabletter	SE/H/2484/001	23-15743	ABBOXIA AB	NO
Dexatin, tabletter	SE/H/2484/001	70285	ABBOXIA AB	DK
Dexatin, tabletter	SE/H/2484/002	70287	ABBOXIA AB	DK
Dexatin, tabletter	SE/H/2484/003	70288	ABBOXIA AB	DK
Dexfarm 10 mg tabletit	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletit	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletit	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletit	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletter	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	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
Dexfarm 10 mg tabletter	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletter	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletter	SE/H/2344/002	42172	ORIFARM HEALTHCARE A/S	FI
Dexfarm 10 mg tabletter	SE/H/2344/002	23-15314	ORIFARM HEALTHCARE A/S	NO
Dexfarm 10 mg tabletter	SE/H/2344/002	23-15314	ORIFARM HEALTHCARE A/S	NO
Dexfarm 10 mg tabletter	SE/H/2344/002	23-15314	ORIFARM HEALTHCARE A/S	NO
Dexfarm 10 mg tabletter	SE/H/2344/002	23-15314	ORIFARM HEALTHCARE A/S	NO
Dexfarm 10 mg tabletter	SE/H/2344/002	64416	ORIFARM HEALTHCARE A/S	SE
Dexfarm 10 mg tabletter	SE/H/2344/002	64416	ORIFARM HEALTHCARE A/S	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dexfarm 10 mg tabletter	SE/H/2344/002	64416	ORIFARM HEALTHCARE A/S	SE
Dexfarm 10 mg tabletter	SE/H/2344/002	64416	ORIFARM HEALTHCARE A/S	SE
Dexfarm 20 mg tabletit	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletit	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletit	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletter	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletter	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletter	SE/H/2344/003	42173	ORIFARM HEALTHCARE A/S	FI
Dexfarm 20 mg tabletter	SE/H/2344/003	23-15315	ORIFARM HEALTHCARE A/S	NO

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
Dexfarm 20 mg tabletter	SE/H/2344/003	23-15315	ORIFARM HEALTHCARE A/S	NO
Dexfarm 20 mg tabletter	SE/H/2344/003	23-15315	ORIFARM HEALTHCARE A/S	NO
Dexfarm 20 mg tabletter	SE/H/2344/003	64417	ORIFARM HEALTHCARE A/S	SE
Dexfarm 20 mg tabletter	SE/H/2344/003	64417	ORIFARM HEALTHCARE A/S	SE
Dexfarm 20 mg tabletter	SE/H/2344/003	64417	ORIFARM HEALTHCARE A/S	SE
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	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
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletit	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	42171	ORIFARM HEALTHCARE A/S	FI
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO

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
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO
Dexfarm 5 mg tabletter	SE/H/2344/001	23-15313	ORIFARM HEALTHCARE A/S	NO
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE
Dexfarm 5 mg tabletter	SE/H/2344/001	64415	ORIFARM HEALTHCARE A/S	SE
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/002	69170	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/003	69171	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Dexfarm, tabletter	SE/H/2344/001	69168	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/002	69170	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/002	69170	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/002	69170	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/003	69171	ORIFARM HEALTHCARE A/S	DK
Dexfarm, tabletter	SE/H/2344/003	69171	ORIFARM HEALTHCARE A/S	DK
Philla 10 mg Tabletten	SE/H/1719/002	142166	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Philla 20 mg Tabletten	SE/H/1719/003	142167	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	AT
Philla 5 mg Tabletten	SE/H/1719/001	142165	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
Tentin 10 mg comprimidos	SE/H/1719/002	89596	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Tentin 10 mg comprimidos	SE/H/1719/002	5880422	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 10 mg comprimidos	SE/H/1719/002	5880430	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 10 mg comprimidos	SE/H/1719/002	5880455	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 10 mg comprimidos	SE/H/1719/002	5880448	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 10 mg tabletid	SE/H/1719/002	1154224	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Tentin 10 mg tabletten	not available	RVG 126113	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
TENTIN 10 mg, comprimé	SE/H/1719/002	CIS: 6 174 671 2	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Tentin 20 mg comprimidos	SE/H/1719/003	89597	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tentin 20 mg comprimidos	SE/H/1719/003	5880463	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 20 mg comprimidos	SE/H/1719/003	5880505	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 20 mg comprimidos	SE/H/1719/003	5880471	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 20 mg tabletid	SE/H/1719/003	1154324	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Tentin 20 mg tabletten	not available	RVG 126116	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
TENTIN 20 mg, comprimé	SE/H/1719/003	CIS: 6 905 420 3	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Tentin 5 mg comprimidos	SE/H/1719/001	89601	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	ES
Tentin 5 mg comprimidos	SE/H/1719/001	5880414	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 5 mg comprimidos	SE/H/1719/001	5880356	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	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
Tentin 5 mg comprimidos	SE/H/1719/001	5880349	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 5 mg comprimidos	SE/H/1719/001	5880372	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 5 mg comprimidos	SE/H/1719/001	5880364	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 5 mg comprimidos	SE/H/1719/001	5880406	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PT
Tentin 5 mg tabletid	SE/H/1719/001	1155524	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	EE
Tentin 5 mg tabletten	not available	RVG 124800	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	NL
TENTIN 5 mg, comprimé	SE/H/1719/001	CIS: 6 189 992 5	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	FR
Tentin, 10 mg, tabletki	SE/H/1719/002	28609	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL
Tentin, 20 mg, tabletki	SE/H/1719/003	28610	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Tentin, 5 mg, tabletki	SE/H/1719/001	28608	MEDICE ARZNEIMITTEL PÜTTER GMBH & CO. KG	PL