



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Amsterdam, 13 January 2022
EMA/35549/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: acipimox

Procedure no.: PSUSA/00000050/202105

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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
OLBETAM 250 mg capsule rigide	not available	025307024	PFIZER ITALIA S.R.L.	IT
Olbetam 250 mg kemény kapszula	not available	OGYI-T-1448/01	PFIZER KFT.	HU
Olbetam, capsules 250 mg	not available	RVG 13159	PFIZER B.V.	NL
Olbetam, kapsler, hårde	not available	12805	PFIZER APS	DK