



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

5 September 2024
EMA/475205/2024
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: pentoxyverine

Procedure no.: PSUSA/00002345/202312

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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
Silotoc 50 mg filmdragerad tablett	not available	9606	OPELLA HEALTHCARE FRANCE SAS	FI
Silotoc 50 mg tabletti, kalvopäällysteinen	not available	9606	OPELLA HEALTHCARE FRANCE SAS	FI
PENTOXYVERINE CLARIX 0,15 % ADULTES, solution buvable édulcorée au maltitol liquide	not available	34009 318 316 5 5	COOPER	FR
PENTOXYVERINE CLARIX 0,15 % ADULTES, solution buvable édulcorée au maltitol liquide	not available	34009 344 276 7 1	COOPER	FR
PENTOXYVERINE CLARIX 0,15 % ADULTES, solution buvable édulcorée au maltitol liquide	not available	34009 382 669 2 4	COOPER	FR
PENTOXYVERINE CLARIX 0,15 % ENFANTS, sirop	not available	34009 382 670 0 6	COOPER	FR
VICKS SIROP PECTORAL 0,15 %, sirop	not available	34009 353 577 6 2	P&G HEALTH FRANCE	FR