



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

30 November 2023
EMA/549530/2023
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): methylphenobarbital

Procedure No. PSUSA/00002025/202303



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
PHEMITON 200 mg tablete	not available	HR-H-034063606	PLIVA HRVATSKA D.O.O.	HR
PHEMITON 200 mg tablete	not available	H/94/01253/001	PLIVA LJUBLJANA D.O.O.;	SI