29 November 2018
EMA/856936/2018
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance(s): methylphenobarbital

Procedure No.: PSUSA/00002025/201803

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
Telephone +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
Send a question via our website www.ema.europa.eu/contact

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	UP/I-530-09/12-02/500	PLIVA HRVATSKA D.O.O.	HR
PHEMITON 200 mg tablete	not available	H/94/01253/001	PLIVA LJUBLJANA D.O.O.;	SI