1 September 2022
EMA/734182/2022
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: methohexital

Procedure no.: PSUSA/00010656/202201

Official address Domenico Scarlattilaan 6 ● 1083 HS Amsterdam ● The Netherlands
Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us
Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

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
Brietal® 500 mg - Trockenstechampulle	not available	15.983	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Brietal Hikma, poeder voor injectievloeistof 500 mg	not available	RVG 00544	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	NL
Brietal® 500 mg - Trockenstechampulle	not available	15.983	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	AT
Brevimytal® Hikma, 500 mg, Pulver zur Herstellung einer Injektions- bzw. Infusionslösung	not available	6128119.00.00	HIKMA FARMACÊUTICA (PORTUGAL), S.A.	DE