



Post Authorisation Assessments

Pentobarbital for Euthanasia 20% w/v Solution for Injection Vm 16431/5001

•	25 November 2022	Conversion of UK licence number into separate Great Britain (GB) and Northern Ireland (NI) Licence numbers.
•	25 July 2017	Change to in-process tests or limits applied during the manufacture of the finished product. Change in specification parameters or limits of the finished product.
•	26 September 2014	Minor change to a test procedure for the finished product.
•	23 September 2014	Reduction in shelf-life from 3 years to 2 years.
•	11 November 2009	Variation to change a test procedure for the finished product.
•	29 July 2009	Variation to change the Marketing Authorisation Holder.
•	02 April 2009	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
•	19 September 2008	Variation to replace a Manufacturer, Assembler, and Distributor for the finished product.
•	04 September 2008	Variation to delete an Assembler and Manufacturer.
•	04 September 2008	Variation to change batch release arrangements and quality control testing of the finished product.
•	29 March 2006	Renewal.
•	17 May 2001	Change to the Active Substance Manufacturer.
•	25 February 2000	Renewal.
•	04 May 1999	Change of Distributor.