



Post Authorisation Assessments

Pentobarbital for Euthanasia 20% w/v Solution for Injection

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|---|-------------------|---|
| • | 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. |