Post Authorisation Assessments

Tetramin 200 Powder Premix for medicated feed

•	13 July 2015	Submission of an updated certificate of suitability. Submission of a new certificate of suitability. Deletion of a batch release site.
•	26 March 2014	Extension of shelf life from 2 to 4 years.
•	18 May 2013	Grouped variation concerning a transfer of the Marketing Authorisation Holder as well as the distributor, the addition of an alternative site for batch release, a change of name of the site for batch testing, and the deletion of a manufacturer.
•	11 August 2010	Variation to register an alternative active substance manufacturer, and to delete an existing active substance manufacturer.
•	21 March 2009	Addition of a site responsible for QA testing of the finished product.
•	14 August 2007	Addition of a manufacturer and assembler of the dosage form. Replacement of a site for batch release in the EU.
•	08 March 2007	Variation to change the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
•	28 December 2006	Variation to change the Marketing Authorisation Holder.
•	20 October 2006	Renewal.
•	16 August 2006	Change in the test procedures for the finished product.
•	11 July 2005	Change of distributor.
•	28 August 2003	Addition of a distributor.
•	31 August 2001	Change of name of the Marketing Authorisation Holder.
•	28 September 1999	Change of name of the Marketing Authorisation Holder.
•	20 November 1998	Change in the finished product specification.