Post Authorisation Assessments

Grisol V Granules 7.5%

•	14 August 2014	Deletion of a site of secondary assembly.
•	12 August 2014	Change to the text in section 4.11 of the SPC and in the product literature.
•	09 December 2013	Submission of an updated Ph. Eur. Certificate of Suitability for an already approved manufacturer.
•	01 August 2009	Addition of a manufacturing site of assembly.
•	27 July 2009	Addition of a manufacturing site of assembly.
•	17 September 2008	Change to batch release arrangements and quality control testing of the finished product.
•	18 June 2008	Change to withdrawal period for horses – addition of statements regarding treated horses not being used for human consumption.
•	28 November 2007	Change of legal category from POM to POM-V. Changes to the SPC and Product Literature to bring in line with new legislation.
•	13 September 2006	Renewal.
•	29 March 2006	Addition of a manufacturer of the dosage form.
•	25 January 2006	Submission of a new Ph. Eur. Certificate of Suitability for an active substance from a new manufacturer. Change of manufacturing site of batch release.
•	24 August 2004	Change of manufacturer and assembler of the dosage form.
•	19 August 2004	Change of address of the MAH.
•	02 December 2003	Addition of a manufacturer for assembly of the dosage form.
•	25 May 2000	Renewal.
•	29 May 1998	Deletion of a target species.
•	10 October 1996	Addition of 40g, 70g and 100g sachets.
•	08 July 1996	Change of MAH.