Post Authorisation Assessments

Grisol V Powder 7.5% w/w Oral Powder

•	12 April 2011	Removal of test performed on the finished product
•	02 November 2010	Submission of a new Ph. Eur. Certificate of Suitability for an active substance from an already approved manufacturer Deletion of a manufacturer of the active substance
•	29 September 2009	Addition of a manufacturing site for batch release
•	27 November 2008	Harmonisation of the SPC and Product Literature
•	28 July 2008	Addition of a manufacturer of the active substance
•	14 February 2008	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
•	17 April 2007	Renewal
•	23 November 2005	Change of address of the manufacturing site of batch release
•	16 September 2004	Change of address of the MAH
•	24 August 2004	Removal of a manufacturing site for the finished product
•	29 July 2004	Change to the specification of the finished product Change of address of the manufacturing site for testing of the finished product
•	21 July 2004	Addition of a manufacturer of the dosage form
•	02 April 2004	Renewal
•	23 July 1998	Renewal Deletion of a target species
•	28 June 1996	Change of MAH