Post Authorisation Assessments

| 10 December 2014 | To replace the text in section 4.11 of the SPC, and the corresponding text in the product literature. |
|-------------------|--|
| 10 November 2014 | Update to the DDPS. |
| 18 July 2013 | Submission of a new or updated Ph. Eur. Certificate of Suitability for an active substance manufacturer. |
| 20 June 2012 | Variation to change the distributor. |
| 28 January 2009 | Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. |
| 09 August 2007 | Replacement or addition of manufacturing site for part or all of the finished product. |
| 06 March 2007 | Transfer of the legal category from POM to POM-V. |
| 09 February 2006 | Renewal. |
| 13 December 2005 | New/updated Ph. Eur for component. |
| 10 November 2005 | Identical changes to a number of products. |
| 21 March 2001 | Renewal. |
| 12 September 1995 | Renewal. |
| | 10 November 2014 18 July 2013 20 June 2012 28 January 2009 09 August 2007 06 March 2007 09 February 2006 13 December 2005 10 November 2005 21 March 2001 |

Norofulvin 33.3% Equine Oral Paste