



Post Authorisation Assessments

Trimacare Tablets Bolus

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| • | 30 July 2019 | Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| • | 24 July 2013 | Change of distributor details. |
| • | 16 July 2013 | Grouped variation to update European Pharmacopoeia Certificates of Suitability from already approved active substance manufacturers. |
| • | 20 January 2009 | Submission of an updated European Pharmacopoeia Certificate of Suitability from an already approved active substance manufacturer. |
| • | 20 November 2008 | Variation to update a European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer. |
| • | 20 January 2009 | Submission of an updated European Pharmacopoeia Certificate of Suitability for an already approved active substance manufacturer. |
| • | 05 November 2008 | Variation to update the product literature and outer packaging. |
| • | 08 October 2008 | Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. |
| • | 21 November 2007 | Renewal. |
| • | 06 March 2007 | Change of legal category from POM to POM-V. |
| • | 10 November 2005 | Addition of a site of assembly (labelling). |
| • | 05 July 2004 | Renewal. |
| • | 09 October 2003 | Change of withdrawal period for calves. |
| • | 15 May 1998 | Copycat. |