



Post Authorisation Assessments

Panacur Equine Guard 10% w/v Oral Suspension

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| • | 23 November 2022 | Extension or introduction of a re-test period/storage period supported by real time data. Minor change to the restricted part of an Active Substance Master File. |
| • | 30 December 2020 | Change in the name of the MAH from Intervet UK Limited to MSD Animal Health UK Limited. |
| • | 17 August 2020 | Addition of a new specification parameter with its corresponding test method of the active substance. Minor change to the restricted part of an Active Substance Master File. |
| • | 24 June 2020 | Change in the specification limits of the finished product. |
| • | 27 April 2020 | Addition of a manufacturer of the active substance or addition of a site of manufacture. |
| • | 10 January 2020 | Tightening of specification limits of the finished product. Reduction of the shelf life of the finished product as packaged for sale from 3 years to 2 years. |
| • | 25 April 2012 | Deletion of a dosing device. |
| • | 08 October 2008 | Changes to the SPC and product literature to bring them into line with new legislation. |
| • | 08 October 2008 | Change of legal category from PML to POM-VPS. |
| • | 06 November 2006 | Change in the batch size of the finished product. |
| • | 12 April 2006 | Renewal. |
| • | 05 October 2005 | Change in the name of the product from Panacur Equine Guard-Unflavoured to Panacur Equine Guard. |
| • | 12 May 2005 | Change to distributor in Northern Ireland. |
| • | 20 April 2005 | Change in the formulation of the finished product. |
| • | 07 November 2002 | Change of manufacturer and assembler of dosage form. |
| • | 03 July 2001 | Addition of a distributor for Northern Ireland. |