Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Equine Guard 10% w/v Oral Suspension

Vm 06376/4093

07 April 2025	Change in immediate packaging of the finished product: - addition of a new container.
22 November 2024	Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to Intervet International B.V., Wim de Körverstraat, 35, 5831 AN Boxmeer, The Netherlands.
11 June 2024	Updates to an ASMF and active substance specification in order to comply with an update of the relevant monograph of the Ph. Eur.
24 November 2023	Change in the specification limits of the finished product.
07 July 2023	Delete odour testing specification parameter of the active substance.
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.