Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Equine Guard 10% w/v Oral Suspension

Vm 01708/4466

•	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.