Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Equine Oral Paste 18.75% w/w

Vm 01708/4439

•	07 July 2023	Delete odour testing specification parameter of the active substance.
•	03 July 2023	Deletion of a non-significant specification parameter in the specification limits of the finished product.
•	03 July 2023	Deletion of a non-significant specification parameter for an excipient.
•	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.
•	21 January 2022	Addition of a site where batch control/testing takes place. Addition of a site where batch control/testing takes place.
•	03 September 2021	Addition of a new specification parameter to the specification with its corresponding test method of the finished product
•	01 April 2021	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.
•	27 April 2020	Addition of a manufacturer of the active substance or addition of a site of manufacture.
•	19 September 2018	Changes to the labelling.
•	24 October 2013	Change in the finished product specifications.
•	20 June 2013	Change to logo on product literature and update to QRD format.
•	07 December 2011	Change in shelf-life of finished product.
•	12 April 2011	Minor change to the manufacturing process of the finished product.
•	16 December 2008	Harmonisation of the finished product specifications in countries where the product is registered.
•	08 October 2008	Changes to the SPC and product literature to bring them into line with new legislation.
•	24 August 2007	Renewal.
•	03 July 2007	Change in batch size of the finished product.
•	05 August 2005	Change in primary packaging.
•	12 May 2005	Change to distributor for Northern Ireland.
•	04 September 2003	Renewal
•	03 July 2001	Addition of a distributor for Northern Ireland.

•	17 March 2000	Change in the name/address of the MAH.
•	04 May 1999	Change in the formulation of the finished product.
•	02 April 1998	Renewal.
•	02 April 1998	Change in target species.
•	23 February 1998	Change in the manufacturer of the active substance.