Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Equine Oral Paste 18.75% w/w

Vm 06376/4094

•	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.
•	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.