Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur 10% Oral Suspension

Vm 01708/4435

•	16 January 2024	Change in any part of the primary packaging material not in contact with the finished product formulation.
•	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.
•	17 May 2022	Addition of a new container for the finished product. Addition of a new container for the finished product. Change in the fill weight / fill volume of the finished product. Changes to the labelling and/or package leaflet.
•	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.
•	22 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.
•	22 October 2015	Approval of Mock ups and update of section 4.10 of the SPC.
•	25 July 2014	Tightening of specification limits for the finished product.
•	15 June 2010	Addition of the statement ‘Do not mix with other products’ to the SPC and product literature.
•	30 September 2009	Addition of warnings regarding anthelmintics to the SPC and product literature.
•	06 November 2006	Change in batch size of the finished product.
•	26 July 2006	Changes to the SPC and product literature to bring them into line with new legislation.
•	02 February 2006	Change of distributor.
•	18 November 2005	Renewal
•	20 April 2005	Change in formulation of the finished product.
•	06 August 2004	Change in design of product literature.
•	02 January 2003	Change of withdrawal period for sheep.
•	08 November 2002	Change in site of manufacturer and assembler of dosage form.

•	03 July 2001	Additional distributor in Northern Ireland.
•	17 March 2000	Change of name and address of MAH.
•	19 January 2000	Renewal.
•	19 October 1998	Deletion of a presentation.
•	09 September 1998	Variation to manufacturer of active substance.
•	04 November 1996	Company name change.
•	15 August 1996	Additional Indication.
•	12 July 1995	Change in Quality procedures.