Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Bolus 12 g, Continuous Release Intraruminal Device

Vm 01708/4447

•	7 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.
•	30 August 2023	Change in the specification parameters 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.
•	25 March 2022	Minor changes to an approved test procedure for the finished product.
•	30 December 2020	Change in the name of the MAH from Intervet UK Limited to MSD Animal Health UK Limited.
•	15 October 2020	Change in the name of the manufacturer of the finished product.
•	01 October 2020	Change in the name of a manufacturer of the finished product, also responsible for batch release.
•	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.
•	21 December 2016	Minor change in the manufacturing process of the active substance.
•	11 December 2015	Changes to SPC and package leaflet
•	12 September 2012	Update to package leaflet.
•	02 May 2012	Change of batch control/testing site.
•	15 December 2009	Change in quantitative composition of the immediate packaging.
•	18 September 2009	Changes to the SPC and product literature to bring them into line with new legislation.
•	18 September 2009	Change of legal category from PML to POM-VPS.
•	07 February 2006	Change in the composition of the immediate packaging.
•	07 February 2006	Renewal.
•	08 August 2005	Change in distributor for Northern Ireland.
•	23 September 2004	Change in wording on the packaging (carton).
•	27 February 2004	Change to wording of indications on the SPC.

•	03 July 2001	Addition of a distributor for Northern Ireland.
•	06 March 2001	Renewal.
•	28 March 2000	Change of assembler of dosage form.
•	17 March 2000	Change in name of MAH including manufacturer of the active substance and manufacturer and assembler of dosage form.
•	23 February 1998	Change in manufacturer of the active substance.
•	30 April 1997	Change to finished product specifications.
•	13 March 1997	Change of MAH.