Veterinary Medicines Directorate

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Post Authorisation Assessments

Panacur Bolus 12 g, Continuous Release Intraruminal Device

Vm 06376/4087

10 December 2024	Change in legal entity from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes MK7 7AJ to Intervet International B.V., Wim de Korverstraat 35, 5831 AN, Boxmeer, Netherlands.
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.