



Post Authorisation Assessments

Panacur 10% Oral Suspension

Vm 06376/4076

26 March 2026	Change in the shelf-life of the finished product. Change in the immediate packaging of the finished product.
01 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
16 October 2025	Deletion of a non-significant specification parameter of an active substance.
25 July 2025	Update to dosing instructions.
25 July 2025	One-off alignment of the product information with version 3.0* of the QRD templates.
22 December 2024	Replacement of a test procedure for the finished product.
19 December 2024	Approval of mock ups.
28 June 2024	Change in legal entity of the MAH from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International BV, Wim de Korverstraat 35, 5831 AN, Boxmeer, Netherlands.
14 May 2024	Updates to an ASMF and active substance specification in order to comply with an update of the relevant monograph of the Ph. Eur.
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.