



Post Authorisation Assessments

Panacur Small Animal 100 mg/ml Oral Suspension Vm 06376/4077

20 April 2026	Replacement of a measuring or administration device which is not an integrated part of the primary packaging.
08 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
17 October 2025	One-off alignment of the product information with version 3 of the GB QRD templates.
16 October 2025	Deletion of a non-significant specification parameter of an active substance.
15 July 2025	Change in the specification parameters of the immediate packaging of the finished product.
22 December 2024	Replacement of a test procedure for the finished product.
22 December 2024	Change in immediate packaging of 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.
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.
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.
23 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.
27 March 2020	Tightening of specification limits of the finished product. Reduction of the shelf life of the finished product as packaged for sale from 3 years to 2 years.
21 December 2016	Minor change in the manufacturing process of the active substance.

04 November 2015	Updates to the SPC, labelling and package leaflet.
01 September 2015	Changes to the labelling and package leaflet.
11 January 2008	Renewal
06 November 2006	Change in the batch size of the finished product.
18 July 2006	Change to product name as a result of new guidelines.
18 July 2006	Changes to the SPC and product literature to bring them into line with new legislation.
18 July 2006	Change in legal category from PML to NFA-VPS.
20 April 2005	Change in formulation of the finished product.
20 January 2003	Renewal
07 November 2002	Change in the manufacturer and assembler of the finished product.
17 March 2000	Change to name and address of MAH.
23 August 1999	Change in packaging presentation.
18 August 1999	Renewal
17 August 1998	Change in the product name.
04 November 1996	Change in name of MAH.
27 August 1996	Change in therapeutic indications.