

Post Authorisation Assessments

Panacur Equine 222 mg/g Granules Vm 06376/4092

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.
22 February 2026	Removal of the prescriptive dosing programme.
22 February 2026	Alignment of the product information with version 9.0* of the QRD templates.
04 February 2026	Addition of a new specification parameter with test method.
16 October 2025	Deletion of a non-significant specification parameter of an active substance.
22 November 2024	Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to Intervet International B.V., Wim de Körverstraat, 35, 5831 AN Boxmeer, The Netherlands.
11 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.
13 September 2023	Change in the specification limits of the finished product. Change to the method description of an in-process control test. Change to an in-process control test limit applied during the manufacture 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.
02 August 2021	Increase in batch size (from 400 - 500 kg to 400 - 600 kg) of the finished product.
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.
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.
08 June 2016	Reduction of the shelf life of the finished product as packaged for sale from 5 years to 36 months.

18 January 2012	Change of logo and formatting on the product literature.
18 September 2008	Changes to the SPC and product literature to bring them into line with new legislation.
18 September 2008	Change of legal category from PML to POM-VPS.
03 June 2008	Deletion of a 1kg pack size.
03 June 2008	Replacement of a site of manufacturer of finished product and batch release.
19 March 2008	Change of equipment used for an in-process control.
21 February 2008	Change in batch size of the finished product.
24 August 2007	Renewal
28 March 2007	Change in the test procedure of the finished product.
12 May 2005	Change of distributor for Northern Ireland.
04 September 2003	Renewal.
03 July 2001	Addition of a distributor for Northern Ireland.
30 November 2000	Change in manufacturer and assembler of dosage form.
17 March 2000	Change in name and address of the MAH.
18 June 1998	Renewal.
23 February 1998	Addition of a manufacture of the active substance.
19 September 1996	Change of MAH.
19 September 1996	Addition of a manufacturer and assembler of the finished product.
05 September 1996	Change in therapeutic indications.
15 August 1996	Change in dosage and administration.