Veterinary Medicines Directorate

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Post Authorisation Assessments

Neoprinil Pour-on 5 mg/ml Pour-on Solution for Cattle

Vm 05653/3047

18 February 2026	Change in batch size of the active substance.
05 August 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
21 July 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
30 January 2025	Changes in the manufacturing process of the active substance - Minor change to the restricted part.
30 January 2025	Change in the batch size of an active substance. Deletion of a non-significant specification parameter of an active substance.
19 November 2024	Change in dimensions of the container or closure of a non-sterile finished product. Deletion of a non-significant specification parameter in the specification parameters or limits of the immediate packaging of the finished product.
19 November 2024	Change in immediate packaging of the finished product.
24 May 2023	Change in batch size (including batch size ranges) of active substance or intermediate used in the manufacturing process of the active substance:– up to 10-fold increase compared to the originally approved batch size.
24 May 2023	Change in the re-test period/storage period of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier: - Extension or introduction of a re-test period/storage period supported by real time data.
27 March 2020	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
14 February 2019	Renewal UK as CMS
14 February 2018	Addition of a manufacturer of the active substance.
20 September 2017	Change in shape or dimensions of the container or closure (immediate packaging)
18 December 2015	Change in the shelf-life from 2 years to 3 years Minor changes to images on SPC and labelling
27 October 2015	Tightening of the test limits of the contents of the active substance.
14 October 2015	Change in the manufacturing process of the finished product. Changes in the manufacturing process of the active substance.

17 April 2015	Change to the in-use shelf-life of the pouches from 1 year to 2 years.
13 February 2015	Change in dimensions of the container. Addition of a site for batch control/testing.