Veterinary Medicines Directorate

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Post Authorisation Assessments

Ecomectin 6mg/g Premix for Medicated Feeding Stuff for Pigs

Vm 13277/5008

24 July 2026	Change to importer, batch control arrangements and quality testing (replacement or addition of a site) for a finished product.
16 May 2026	Change in the specification parameters and/or limits of an excipient: - Change outside the approved specifications limits range.
23 March 2026	Minor change to an approved test procedure for an excipient.
14 January 2026	Change to comply with Ph. Eur. or with a national pharmacopoeia of a Member State.
15 September 2025	Change to comply with a pharmacopoeial monograph. (NI).
11 August 2025	Change to comply with Ph. Eur. (GB)
14 May 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI).
02 January 2025	Addition of a site of batch control.
21 December 2023	Change in the holding time of an intermediate or bulk product.
01 June 2023	Change of MAH address from Eco Animal Health Ltd, 78 Coombe Road, New Malden, Surrey, KT3 4QS, United Kingdom to Eco Animal Health Ltd, The Grange, 100 High Street, London, N14 6BN, United Kingdom.
19 August 2021	Changes to the labelling and package leaflet.
15 April 2021	Replacement to a test procedure for the finished product.
15 February 2021	Addition of a site where batch control/testing takes place.
18 December 2020	Addition of a manufacturer responsible for batch release of the finished product.
06 November 2020	Change in shape or dimensions of the container or closure (immediate packaging). Increase in batch size of the finished product. Deletion of manufacturing site for an active substance.
09 March 2020	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
17 May 2017	Change in the name of a manufacturer of the finished product, also responsible for batch release. Change in the specification parameters of an excipient. Change in the specification parameters of an excipient. Change in the specification parameters of an excipient. Change in the specification parameters of an excipient. Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State.

	Change in the manufacturing process of the finished product. Change in the specification parameters of an excipient. Change in the specification parameters of an excipient.
25 October 2016	Change in the QPPV and/or QPPV contact details and/or back-up procedure. Change in the QPPV and/or QPPV contact details and/or back-up procedure.
25 August 2013	Repeat Use Comm.
12 June 2013	Renewal