Veterinary Medicines Directorate

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Post Authorisation Assessments

Effectimec 18.7 mg/g Oral Paste for Horses

Vm 61301/3002

15 May 2026	One-off alignment of the product information with version 9.0 of the QRD templates.
16 February 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
29 August 2025	Change of legal entity of the Marketing Authorisation Holder from Eco Animal Health Ltd to ACME DRUGS s.r.l.
18 June 2025	Change in shape or dimensions of the container or closure (GB)
14 March 2025	Addition of a supplier of packaging components or devices. (NI).
14 December 2024	Addition of a supplier of packaging components or devices. (GB)
19 March 2024	Removal of a site responsible for the manufacture of the active substance. (NI)
29 November 2023	Removal of a manufacturing site responsible for batch release. (GB)
29 November 2023	Removal of a site responsible for the manufacture of the finished product. (NI)
29 November 2023	Removal of a site responsible for the manufacture of the finished product. (GB)
26 October 2023	Removal of a site responsible for the manufacture of the active substance. (GB)
02 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.
18 May 2020	Change in the invented name of the veterinary medicinal product from Ecomectin Ivermectine 18,7 mg/g Pate Orale pour Chevaux to Nexmectin 18,7 mg/g Pate Orale pour Chevaux in France only.
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.
26 September 2018	Repeat Use application to add 3 new member states
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.
14 November 2014	Update of Certificates of Suitability. Minor change to product specifications.

28 June 2012	Removal of a redundant reference.
30 May 2012	Renewal procedure – Ireland as CMS.
16 march 2012	Change in the pack size of the finished product.
16 march 2012	Change in the shape or dimensions of the immediate package
16 march 2012	Change in the batch size of the finished product.
29 February 2012	To change the name of the medicinal product in Ireland and France.