

Post Authorisation Assessments

Animec 18.7 mg/g Oral Paste for Horses

Vm 61301/5000

10 December 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
09 December 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
29 August 2025	Change of legal entity of the Marketing Authorisation Holder from Eco Animal Health Ltd to ACME DRUGS s.r.l.
25 July 2025	Change in the address of the marketing authorisation holder from The Grange, 100 High Street, London, UK, N14 6BN to 78 Coombe Road, New Malden Surrey, KT3 4QS, United Kingdom. (GB+NI)
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).
11 December 2024	Addition of a supplier of packaging components or devices. (GB).
07 March 2024	Removal of a site responsible for the manufacture of the active substance. (NI)
07 December 2023	Removal of a manufacturing site responsible for batch release. (GB)
07 December 2023	Removal of a site responsible for the manufacture of the finished product. (NI)
06 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)
1 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.
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.
4 September 2018	RU MRP to add two new CMS
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.
29 May 2013	Repeat use, authorising product for use in Poland.
13 December 2012	Change in the name of the veterinary medicinal product in Greece only.
28 June 2012	Change in test procedure for the finished product.
30 May 2012	Renewal – Ireland as RMS
16 March 2012	Change in the shape or dimensions of the container/closure

16 March 2012	Change in batch size of the finished product
16 March 2012	Change in pack size of the finished product
29 February 2012	To change the name of the product from Vectimax 18.7 mg/g pasta orale per cavella to Avatar 18.7 mg/g pasta orale per cavalli.
17 January 2011	Change to batch release arrangements and quality control testing of the finished product.
17 January 2011	Replacement/addition of a manufacturing site.