



Post Authorisation Assessments

Animec 18.7 mg/g Oral Paste for Horses Vm 13277/4019

•	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.

