Veterinary Medicines Directorate

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Post Authorisation Assessments

Alomec 18.7 mg/g Oral Paste for Horses

Vm 50146/4034

10 December 2024	Minor change in the manufacturing process of the finished product.
20 October 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
10 April 2024	Deletion of a Ph. Eur. CEP for an active substance manufacturer. Deletion of a Ph. Eur. CEP for an active substance manufacturer.
25 January 2024	Submission of a new Ph. Eur. CEP from a new manufacturer for a non-sterile.
18 March 2021	Replacement of a secondary packaging site of the finished product.
26 January 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
17 February 2020	Change in distributor details. From Farm and Stable Supplies LLP, Bridgelands, Ingrams Green, Midhurst, GU29 OLJ to GBA International LLP, Prospect House, Neston, CH64 3RU.
23 July 2019	Change in the name of a manufacturer used in the manufacture of the active substance. Change in the name and address of a manufacturer of the finished product, also responsible for batch release.
26 October 2018	Changes to an existing pharmacovigilance system as described in the DDPS. Change of MAH, from Cross Vetpharm Group Ltd, Broomhill Road, Tallaght, Dublin 24, Ireland to Bimeda Animal Health Limited, 2 / 3 / 4 Airton Close, Tallaght, Dublin 24, Ireland.
06 January 2016	Submission of an updated certificate of suitability.
09 September 2014	Change in the name of the medicinal product, from “Equipaste 1.87% w/w oral paste" to "Alomec, 18.7 mg/g oral paste".
09 September 2014	Renewal procedure.
7th February 2012	Submission of a new or updated Ph. Eur Certificate of Suitability.