Veterinary Medicines Directorate

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Post Authorisation Variations

Equibactin Vet. (333 mg/g + 67 mg/g) Oral Paste for Horses

•	11 May 2022	Deletion of Ph. Eur. certificates of suitability for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	11 August 2021	Minor change in the manufacturing process of the finished product. Change to the manufacturing process. Change to batch size of the finished product. Change to specification of the finished product. Increase in batch size (including batch size range) of the finished product. Deletion of a non-significant specification parameter of the finished product.
•	02 September 2019	Introduction of a new pharmacovigilance system.
•	22 March 2019	Change of distributor to Dechra Veterinary Products Limited, Sansaw Business Park, Hadnall, Shrewsbury, Shropshire SY4 4AS, United Kingdom
•	18 October 2018	Changes to the labelling.
•	17 January 2018	Addition of a manufacturer of the active substance.
•	07 October 2015	To add a new pack size of the finished product.
•	07 May 2014	Renewal procedure – Netherlands as RMS.
•	16 September 2011	To increase the shelf life of the finished product from 2 to 3 years.
•	15 September 2010	To change the address of the Marketing Authorisation Holder