Veterinary Medicines Directorate

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Post Authorisation Assessments

Rhemox Forte, 1000 mg/g Powder for Use in Drinking Water for Chickens, Ducks, Turkeys

Vm 43173/4001

•	27 June 2024	Addition of a site of batch control testing for the finished product. Replacement of a site of secondary manufacturing for the finished product.
•	24 January 2023	Updated certificate of suitability from an already approved manufacturer.
•	01 November 2022	Updated certificate of suitability from an already approved manufacturer.
•	02 July 2021	Renewal – UK as CMS.
•	14 October 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer Deletion of Ph. Eur. certificates of suitability for an active substance manufacturer
•	09 March 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	07 June 2018	Change in RMS from UK to ES.
•	06 November 2017	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Introduction of a re-test period of the active substance.
•	22 December 2016	Change in the name and address of the marketing authorisation holder from aniMedica Espana to LIVISTO Int'l, S.L.
•	02 August 2016	Submission of a new certificate of suitability for an active substance from an already approved manufacturer Submission of an updated certificate of suitability for an active substance from an already approved manufacturer
•	28 July 2016	Change in the invented name of the veterinary medicinal product in Spain and France.