Veterinary Medicines Directorate

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Post Authorisation Assessments

Amproline 400 mg/mL Solution for Use in Drinking Water for Chickens and Turkeys

Vm 41623/5000

•	28 April 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
•	April 2024	One-off alignment of the product information with version 9.0* of the QRD templates.
•	13 January 2022	Change of specification(s) of a former non Pharmacopoeial active substance to comply with the Ph. Eur. or with a national pharmacopoeia of a Member State.
•	02 December 2021	Change in the specification limits of the finished product.
•	08 April 2021	Renewal - UK as CMS.
•	08 October 2020	Change in batch size range of the finished product. Addition of a manufacturer responsible for batch release including batch control/testing. Addition of a primary packaging site of the finished product. Addition of a manufacturing site of the finished product.
•	19 June 2020	Minor change in the manufacturing process of the finished product.
•	11 June 2020	Change in storage conditions of the finished product.
•	17 April 2020	Changes (Safety/Efficacy) to a Veterinary Medicinal Product following a Repeat Use Procedure.
•	26 July 2019	Repeat Use application to add 11 new member states
•	08 July 2019	Changes to the labelling and package leaflet. Change of distributor from Vetsonic (UK) LTD, Riccal Drive, Malton, Y017 6YE to MAH address.
•	04 April 2019	Change in the name and address of a manufacturer
•	04 April 2019	Change of MAH name and address to: HUVEPHARMA SA 34 RUE JEAN MONNET ZI D'ETRICHE SEGRE 49500 SEGRE-EN-ANJOU BLEU FRANCE
•	14 February 2019	Change in storage conditions of the finished product.
•	July 2018	Change in shape or dimensions of the container or closure (immediate packaging). Changes in the qualitative and quantitative composition of the immediate packaging of the finished product.