Veterinary Medicines Directorate

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Post Authorisation Assessments

Mycoflor, 300 mg/ml Solution for Injection for Cattle and Pigs
Vm 36967/4000

•	08 April 2024	Incorporation of the changes required following the outcome of the referral for products including NMP as an excipient. (NI)
•	08 April 2024	Incorporation of the changes required following the outcome of the referral for products including NMP as an excipient. (GB)
•	13 July 2023	Change in the manufacturer of a starting material/reagent/intermediate used in the manufacturing process of the active substance or change in the manufacturer of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier: - Introduction of a manufacturer of the active substance supported by an ASMF.
•	31 March 2020	Updated Active Substance Master File.
•	26 June 2018	Change in the RMS from UK to IE.
•	09 May 2017	Addition of a manufacturer of the active substance or addition of a site of manufacture
•	08 December 2016	Renewal – UK as RMS
•	14 April 2015	Addition of a target animal species.