Veterinary Medicines Directorate

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Post Authorisation Assessments

Doxipulvis 500 mg/g Powder for Use in Drinking Water / Milk Replacer

Vm 36967/5001

•	February 2024	Addition of a flavouring agent to the formulation of the finished product.
•	23 February 2024	Minor changes in the manufacturing process.
•	14 July 2023	Deletion of a non-significant specification parameter in the specification parameters or limits of the finished product.
•	08 March 2023	Updated certificate of suitability from an already approved manufacturer. Updated certificate of suitability from an already approved manufacturer.
•	14 February 2023	Change in batch size of the finished product.
•	01 February 2023	Change in name of a manufacturer of the finished product.
•	28 September 2022	Change in name of a manufacturer of the finished product.
•	09 December 2021	Renewal – UK as CMS.
•	25 August 2020	Introduction of a re-test period of the active substance.
•	06 July 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	17 September 2019	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.