Veterinary Medicines Directorate

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Post Authorisation Assessments

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

Vm 36967/3001

•	14 December 2023	Deletion of an obsolete parameter in the specification parameters of the finished product. Deletion of an obsolete parameter in the specification parameters of the finished product. Deletion of an obsolete parameter in the specification parameters of the finished product.
•	10 November 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile. Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile.
•	26 June 2023	Change in the composition of a non-sterile finished product:– addition or replacement of a component or components of the flavouring or colouring system.
•	26 June 2023	Change in the batch size of the finished product.
•	26 June 2023	Change in the manufacturing process of the finished product, including an intermediate used in the manufacture of the finished product: - Minor change in the manufacturing process.
•	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.