Veterinary Medicines Directorate

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Post Authorisation Assessments

Methoxasol 20/100 mg/ml Solution for Use in Drinking Water for Pigs and Chickens

Vm 16849/3007

•	20 November 2024	Minor change in immediate packaging of the finished product.
•	15 December 2023	Inclusion of SPC and product literature changes required following the Article 83 referral for products containing NMP. Alignment with version 9.0 of the QRD template.
•	07 April 2022	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product
•	01 February 2022	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.
•	November 2021	Deletion of manufacturing site for a finished product.
•	20 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	24 January 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	22 May 2018	Replacement of a manufacturer responsible for batch release including batch control/testing. Addition of a secondary packaging site of the finished product. Addition of a primary packaging site of the finished product. Addition of a manufacturing site for the manufacturing process of the finished product.
•	02 June 2017	Renewal – UK as CMS.
•	08 December 2016	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	23 April 2014	Change to the shape of the immediate packaging.
•	24 October 2013	Change to the test procedure for the finished product.
•	6 March 2013	Change of QPPV and QPPV contact details for an existing pharmacovigilance system.