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

12 March 2026	Submission of a Ph. Eur. CEP for an active substance. Submission of a Ph. Eur. CEP for an active substance.
06 January 2026	Change in the dimensions of the closure of the immediate packaging for the finished product.
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.