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Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021, please refer to the EMA website.

Zolvix 25 mg/ml Oral Solution for Sheep

Vm 52127/5029

•	29 August 2023	One-off alignment of the product information with version 9.0* of the QRD templates.
•	28 July 2023	Extension in the re-test period of the active substance where no Ph. Eur. Certificate of Suitability covering the retest period is part of the approved dossier.
•	10 May 2023	Change in manufacturer responsible for batch control of the active substance.
•	18 April 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.