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Post Authorisation Assessments

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

Halocur 0.5 mg/ml Oral Solution for Calves

Vm 01708/5037

14 March 2025	Deletion of a manufacturing site for an active substance.
02 October 2024	One-off alignment of the product information with version 9.0* of the QRD templates.
02 June 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 of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier: -Introduction of a manufacturer of the active substance supported by an ASMF.
07 July 2022	Changes to labelling to include GB details in blue box.