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Post Authorisation Assessments

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

Veraflox 60 mg Tablets for Dogs

Vm 04895/5011

16 June 2026	Minor editorial amendments.
15 April 2026	Introduction of a new active substance manufacturer for pradofloxacin for Veraflox Tablets and Oral Solution. Changes to the active substance test methods.
31 October 2025	Change in the name of the manufacture of the dosage form
22 October 2025	Update to MIC information in SPC. Alignment of the product information with version 9.0* of the QRD templates.
17 July 2025	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
17 August 2023	Change in the name and address of the marketing authorisation holder from Bayer Animal Health GmbH, 51368 Leverkusen, Germany to Elanco Animal Health GmbH, Alfred-Nobel-Str. 50, 40789 Monheim, Germany.
13 December 2022	Changes to labelling to include GB details in blue box.
24 March 2022	Change in the manufacturer of an intermediate used in the manufacturing process of the active Change in the manufacturer of an intermediate used in the manufacturing process of the active
10 February 2022	Change in the name of the manufacturer of the finished product.