Veterinary Medicines Directorate
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Post Authorisation Assessments

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

Meloxoral 1.5 mg/ml Oral Suspension for Dogs

Vm 50406/5001

30 April 2026	Submission of a Ph. Eur. CEP for an active substance.
13 January 2026	Change of a finished product administration device.
16 April 2025	Submission of new mock ups.
06 July 2024	Addition of a manufacturer responsible for batch release including batch control or testing of a non-sterile finished product.
13 May 2024	One-off alignment of the product information with version 9.0* of the QRD templates.
15 February 2022	Deletion of a non-significant specification parameter of the finished product. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
07 October 2021	Deletion of Local Representative information from the package leaflet.
04 August 2021	Change of MAH, from Le Vet B.V., Wilgenweg 7, 3421 TV Oudewater, The Netherlands to Dechra Regulatory B.V., Handelsweg 25, 5531 AE Bladel, The Netherlands.
04 February 2021	Change in the specification limits of the finished product.