



Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021,
please refer to the [EMA](#) website.

Metacam 0.5 mg/ml Oral Suspension for Cats and Guinea Pigs Vm 61700/5053

14 August 2025	Deletion of a secondary packaging site.
06 March 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
18 June 2024	Variation to harmonise the indications and dosing instructions for Metacam 0.5 mg/ml oral suspension for cats and guinea pigs, Metacam 2 mg/ml solution for injection for cats and Metacam 5 mg/ml solution for injection for cats and dogs. Variation to harmonise the indications and dosing instructions for Metacam 0.5 mg/ml oral suspension for cats and guinea pigs, Metacam 2 mg/ml solution for injection for cats and Metacam 5 mg/ml solution for injection for cats and dogs.
17 May 2023	Changes to the quality part of the dossier: Deletion of - a non-significant specification parameter (e.g. deletion of an obsolete parameter) of a measuring or administration device.
17 May 2023	Change of a measuring or administration device:
14 April 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
13 December 2022	Deletion of a non-significant specification parameter for an excipient.
15 February 2022	Addition of two secondary packaging sites of the finished product.
20 December 2021	Minor changes to an approved test procedure of the finished product.
01 December 2021	Changes to the labelling and/or package leaflet.
24 June 2021	Change in the test parameter.
17 May 2021	Change to part of the (primary) packaging material not in contact with the finished product formulation.
25 March 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.