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Post Authorisation Assessments

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

Recocam 20 mg/ml Solution for Injection for Cattle, Pigs and Horses

Vm 50146/5000

17 April 2026	One-off alignment of the product information with version 3 of the QRD templates.
13 April 2026	Align PL with reference product - updated indication and warning.
06 February 2026	Update to Adverse event section. Frequency of Injection site swelling from 'Undetermined frequency' to 'Very Rare' for cattle and horses.
01 August 2025	Submission of a Ph. Eur. CEP for an intermediate used in the manufacturing process of the active substance. Submission of a Ph. Eur. CEP for an active substance Submission of a Ph. Eur. CEP for an active substance
22 July 2025	Replacement of a manufacturer for batch release. Replacement of a secondary packaging site of a finished product.
24 May 2022	Change in the pack size of the finished product.
08 March 2022	Increase in batch size (including batch size range) of the finished product Minor change in the manufacturing process of an immediate release solid oral dosage form.