



Veterinary
Medicines
Directorate

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.

Rheumocam 5 mg/ml Solution for Injection for Cattle and Pigs Vm 08749/5025

02 April 2025	Submission of a new Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
23 October 2024	Minor changes to an approved test procedure for finished product.
25 April 2024	Updated CEP submitted for the manufacture of an active substance.
09 June 2023	Change to comply with Ph. Eur. or with a national pharmacopoeia of a Member State, to reflect compliance with the Ph. Eur. by removing reference to the internal test method and test method number.
11 June 2021	Increase in the shelf-life of the finished product as packaged for sale, from 4 to 5 years.