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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

•	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.