Veterinary Medicines Directorate

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Post Authorisation Assessments

Readycef 50 mg/ml Suspension for Injection for Swine and Cattle

Vm 20634/4003

•	13 February 2024	Change in the address of the manufacturer of the finished product. (NI)
•	15 August 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
•	20 July 2023	Change in the address of the manufacturer of the finished product.
•	16 October 2019	Changes in the manufacturing process of the finished product.
•	07 September 2017	Addition of a site where batch control/testing takes place.
•	14 January 2016	Change in the (invented) name of the medicinal product in AT, DE and EL only
•	25 June 2014	To change the QPPV.
•	22 May 2014	Renewal procedure – Portugal as RMS.
•	27 March 2013	The addition of a 250 ml glass vial presentation.
•	26 March 2012	Changes in Summary of Product Characteristics (SPC), labelling or package leaflet following a referral procedure.
•	04 February 2011	To add a manufacturer of the active substance.
•	24 June 2010	Change in the (invented) name of the veterinary medicinal product. Previous name Ceftiomax 50 mg/ml Suspension for Injection for Swine and Cattle.