Veterinary Medicines Directorate

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Post Authorisation Assessments

Eficur 50 mg/ml Suspension for Injection for Pigs and Cattle

Vm 17533/4010

•	28 April 2024	Introduction of a manufacturer of the active substance supported by an ASMF.
•	07 January 2020	Minor changes to an approved test procedure of the finished product. Replacement to a test procedure for the finished product. Replacement of a specification parameter with its corresponding test method of the finished product.
•	15 July 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	10 April 2017	Addition of a manufacturing site for part of the manufacturing process of the finished product
•	04 February 2016	Change in the name of the UK local representative.
•	08 May 2015	Introduction of three new presentations of the finished product, and the addition of a sterilisation site as a consequence of this change.
•	29 May 2013	Deletion of manufacturing site responsible for terminal sterilisation.
•	13 July 2012	A new site to perform the finished product final terminal sterilisation was added.
•	02 May 2012	Variation to change the distributor.
•	22/03/2012	Renewal procedure – Ireland as RMS.
•	08/12/2010	Addition of a manufacturer of the active substance.
•	09/08/2010	To add a new pack size of the finished product.