



Post Authorisation Assessments

Eficur 50 mg/ml Suspension for Injection for Pigs and Cattle Vm 17533/3009

11 November 2025	Alignment of the product information with version 9.0* of the QRD templates.
08 May 2025	Change to the manufacturing process of the finished product - other changes. Change to in-process tests applied during finished product manufacture.
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.