



Post Authorisation Assessments

Veteglan 0.075 mg/ml Solution for Injection for Cows, Sows and Mares Vm 20634/5004

28 April 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV.)
17 April 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
27 March 2025	Extension of the shelf-life of the finished product as packaged for sale. Change in storage condition of the finished product. Addition of a new pack size of the finished product.
4 June 2024	Change in distributor details from LABORATORIOS CALIER S.A to EGG TECHNOLOGIES INTERNATIONAL LTD, 18 Springfield Park, Tisbury, Salisbury, Wiltshire SP3 6QN, United Kingdom. Changes to the Product Literature.
15 April 2024	Name of active substance manufacturer changed.
15 February 2024	Changes to the package leaflet not connected to the SPC.
15 August 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
07 March 2023	Change in the name and address of a manufacturer of the finished product responsible for sterility testing.
02 February 2023	Updates of the QRD/SPC information in line with V1 of the GB national QRD template.
05 May 2022	Updates to the ASMF.
25 November 2021	Renewal – UK as CMS.
28 July 2021	Addition of a manufacturer of the active substance or addition of a site of manufacture.
19 October 2020	Changes to the SPC and QRD text following a Repeat Use Procedure.
05 February 2020	Repeat Use application to add 1 new member state.
06 March 2019	Increase in batch size of the finished product. Addition of a manufacturing site of the finished product.
07 September 2017	Addition of a site where batch control/testing takes place. Addition of a site where batch control/testing takes place.
13 April 2017	To update the SPC and QRD post Repeat-Use Mutual Recognition Procedure.