



Post Authorisation Assessments

Selectan 300 mg/ml Solution for Injection for Cattle and Swine Vm 17533/4006

•	03 October 2023	Change to an approved stability protocol of an active substance. Deletion of a manufacturing site for an active substance starting material. Minor changes to an approved test procedure for the active substance. Addition of a supplier of packaging components for the active substance.
•	03 October 2023	Minor updates to an ASMF.
•	03 October 2023	Change in the Summary of Product Characteristics, labelling or package leaflet intended to implement the outcome of a Union interest referral procedure according to Article 83 of Regulation 2019/6.
•	18 August 2021	Update to ASMF.
•	07 October 2019	Deletion of manufacturing site for an active substance. Addition of a manufacturer of the active substance.
•	15 July 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	12 June 2017	Addition to a test procedure for the finished product.
•	04 February 2016	Change in the name of the local UK representative.
•	31 July 2015	Change to an already approved manufacturer of the active substance.
•	27 February 2014	To add an additional manufacturer of the active substance.
•	14 December 2012	Renewal procedure.
•	27 July 2011	Change in immediate packaging of the finished product
•	18 May 2011	Repeat Use Comm – UK as CMS
•	9 December 2010	Change in batch size of the finished product.