



Post Authorisation Assessments

Mypravac Suis Suspension for Injection

Vm 17533/4001

•	15 July 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	18 May 2018	Change in RMS from UK to ES.
•	18 June 2009	Change to packaging material not in contact with final formulation.
•	12 June 2005	Renewal procedure.
•	24 January 2007	Pack type.
•	05 October 2005	Repeat Use Procedure (UK as RMS).
•	30 March 2005	Inclusion of a bio burden control test.
•	19 January 2005	Inclusion of a new laboratory trial concerning the efficacy of the vaccine.
•	20 March 2003	MRP (UK as RMS).
•	31 July 2002	Change of batch size of the active substance.
•	08 July 2002	Change of the finished product shelf life.
•	19 June 2002	Addition of an active substance manufacturer.