Veterinary Medicines Directorate

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Post Authorisation Assessments

Mypravac Suis Suspension for Injection

Vm 17533/3004

21 January 2025	G.I.18 update of the product information to EU version 9.
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.