Veterinary Medicines Directorate

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Post Authorisation Assessments

Suvaxyn M. Hyo - Parasuis, Suspension for Injection for Pigs

•	25 October 2019	Change in the address of the marketing authorisation holder from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
•	25 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	09 March 2018	Change in RMS from UK to DE.
•	11 January 2017	Change in the specification limits of a starting material/intermediate used in the manufacturing process of the active substance.
•	08 December 2016	Change of MAH address in France, Czech Republic and Slovakia.
•	22 March 2016	Renewal – UK as RMS
•	12 February 2016	Change in the name of a manufacturer of the active substance. Change in the name of a manufacturer of the finished product.
•	28 October 2015	Change to in-process tests or limits applied during the manufacture of the finished product Changes in the manufacturing process of the active substance
•	05 May 2015	Change in the QPPV contact details.
•	10 April 2015	Update to the product dossier. Change of site for testing starting materials of biological origin. Change in test procedure for testing starting materials of biological origin.
•	30 January 2015	Change of test location for an in-process-control test.
•	11 October 2013	To change the name and/or address of the marketing authorisation holder from Pfizer to Zoetis in AT, BE, FR, and LU only.
•	10 October 2013	Change in the name of the manufacturer of the active substance. Changes in the name of the manufacturer of the finished product. Change of QPPV contact details.
•	8 August 2013	Change of MAH and distributor.
•	13 November 2012	Repeat Use procedure to add Estonia and Cyprus as CMS.
•	23 February 2012	Change in test procedure for the finished product.

•	28 December 2011	Renewal procedure. UK as RMS.
•	23 September 2011	To change the name of the manufacturer of the active substance, to change the name of the site for batch release, final product testing and labelling, and to change the site for filling , blending QC testing and assembly.
•	08 April 2011	To change the address of the MAH in Poland.
•	16 June 2010	To change the MAH and distributor from Fort Dodge Animal Health Ltd to Pfizer Limited.
•	04 June 2008	New MA (MRP)
•	05 December 2007	Additional batch release site
•	26 September 2007	Incorporation of new field study
•	26 September 2007	Change of the inactivation time for *M. hyopneumoniae* antigen
•	16 August 2007	Incorporation of new field study
•	16 May 2007	SPC/label changes (Vet Regs 2005)