Veterinary Medicines Directorate

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Post Authorisation Assessments

Suiseng Suspension for Injection for Pigs

Vm 17533/4009

•	13 February 2024	Change in the name and address details of the manufacturer of the active substance. (NI)
•	14 April 2023	Change in the name and address details of the manufacturer of the active substance.
•	15 July 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	21 June 2019	Deletion of a test procedure for the intermediate used in the manufacturing process of the active substance if an alternative test procedure is already authorised. Deletion of a non-significant in-process test applied during the manufacture of the active substance. Changes to a test procedure (including replacement) for the active substance. Change to in-process tests applied during the manufacture of the active substance. Change in the batch size (including batch size range) of the active substance used in the manufacturing process of the active substance. Qualitative composition changes to the immediate packaging of the active substance. Changes in the manufacturing process of the active substance.
•	21 June 2019	Minor changes to an approved test procedure of the finished product. Minor changes to an approved test procedure of the finished product. Minor change in the manufacturing process of an immediate release solid oral dosage form or oral solutions. Change of a test procedure for the finished product.
•	07 February 2018	Change in RMS from UK to IE.
•	04 February 2016	Change in the name of the local UK representative.
•	21 October 2015	Repeat use application to add one new member state.
•	06 February 2015	Renewal.
•	06 November 2014	Increase in the storage period of the *clostridium* antigens to 24 months.
•	16 December 2010	Variation to extend the shelf-life of the finished

		product from fifteen to eighteen months.
•	16 December 2010	Variation to extend the shelf-life of the *E.coli* antigens.