Veterinary Medicines Directorate
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Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021, please refer to the EMA website.

Forceris 30 mg/ml + 133.4 mg/ml Suspension for Injection for Piglets

Vm 15052/5007

•	22 March 2024	Addition of a site of batch control for the finished product.
•	25 May 2023	Change in the manufacturer of a starting material/reagent/intermediate used in the manufacturing process of the active substance. Changes in the manufacturing process of the active substance: - Minor change to the restricted part of an Active Substance Master File.
•	14 February 2023	Approval of mock ups.
•	11 October 2022	Change in the address of the MAH from Unit 3 Anglo Office Park, White Lion Road Amersham, Buckinghamshire HP7 9FB to Explorer House, Mercury Park, Wycombe Lane, Wooburn Green, High Wycombe, Buckinghamshire, HP10 0HH, United Kingdom.
•	17 May 2022	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product. Minor change in the manufacturing process of the finished product. Introduction of an additional sterilisation site of the packaging for the finished product. Introduction of an additional sterilisation site of the packaging for the finished product.
•	17 May 2022	Addition of a manufacturing site of the finished product. Addition of a secondary packaging site of the finished product. Addition of a site where batch control/testing takes place.
•	17 January 2022	Additional supplier of starting material for the manufacture of the active substance.
•	24 November 2021	Extension of a re-test period of the active substance.