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

04 June 2025	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
16 April 2025	Change in a manufacturer for an active substance. Change in a manufacturer for an excipient.
27 March 2025	One-off alignment of the product information with version 3 of the QRD template.
13 March 2025	Other changes to the active substance: - Substantial changes in the updated version of the ASMF or the active substance part of the dossier.
11 February 2025	Minor changes to an approved test procedure for an excipient.
14 August 2024	Changes to the production process or the storage of active substance where no Ph. Eur. CEP is part of the approved dossier of an active substance.
20 June 2024	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
13 May 2024	Unlimited Renewal
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.