Veterinary Medicines Directorate
Woodham Lane, New Haw, Addlestone, Surrey, KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

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

Suvaxyn Circo + MH RTU Emulsion for Injection for Pigs

Vm 42058/5072

18 December 2025	To add an already authorised site for Physical/Chemical testing of the finished product to the manufacturing flow chart.
15 December 2025	To remove the spike control and initial 1:50 dilution for the positive control from the in-process ELISA test for porcine IgG content in the active substance.
21 January 2025	Increase in batch size of active substance without process change.
22 June 2024	To align the product information with GB version 2 of the SPC/QRD templates.
26 April 2023	Change in the specification limits of an active substance.
25 January 2023	Addition of a secondary packaging site of a finished product. Addition of a secondary packaging site of a finished product.
24 January 2023	To widen their approved bovine viral diarrhoea virus specifications for the Foetal Bovine Serum starting material used in the manufacture of porcine circovirus antigen.
14 June 2022	Changes to the labelling.
05 November 2021	Increase in the shelf-life of the finished product as packaged for sale, from 18 to 24 months.
22 April 2021	Change in source of active substance.