



Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021,
please refer to the [EMA](#) website.

CircoMax Emulsion for Injection for Pigs Vm 42058/5003

18 December 2025	To add an already authorised site for Physical/Chemical testing of the finished product to the manufacturing flow chart.
21 February 2025	Removal of the EU Local Representative addresses.
19 January 2025	Extension of the shelf life of the finished product to 24 months.
10 July 2024	Approval of mock-ups.
08 May 2024	To add the option to administer CircoMax Myco and Circomax intramuscularly using needle-free devices.
03 July 2023	This variation is to remove from the product information the warning regarding the absence of safety data during pregnancy and lactation: "The safety of the veterinary medicinal product has not been established during pregnancy or lactation. Do not use during pregnancy and lactation" and substitute it with "Not applicable". One-off alignment of the product information text with version 1 of the GB and QRD template.
24 January 2023	Addition of a secondary packaging site of a finished product. Addition of a secondary packaging site of a finished product.