



Post Authorisation Assessments

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

Draxxin 25 mg/ml Solution for Injection for Pigs Vm 42058/5022

24 September 2025	Addition of a test procedure for the finished product.
14 May 2025	Removed mention of local representatives from adverse event sections in both SPC & QRD as there are no local representative details provided. Deletion of a test procedure for the finished product. Deletion of a manufacturer responsible for batch release of the finished product. Editorial Changes to the SPC.
11 December 2024	Deletion of a non significant specification parameter for an active substance.
14 August 2024	One-off alignment of the product information with version 9.0.
18 April 2024	Deletion of a manufacturing site of the finished product.
11 August 2023	Approval of mock ups.
24 August 2021	Updates to Section 4.5 of the Summary of Product Characteristics and to Section 12 of the product literature with regard to adverse reactions.
23 April 2021	Alignment of SPC and QRD texts.