Veterinary Medicines Directorate

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Post Authorisation Assessments

Bravoxin Suspension for Injection for Cattle and Sheep

Vm 06376/5030

14 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
11 June 2025	To remove the finished product test for residual toxicity to comply with the updates to Ph. Eur.
25 February 2025	Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to Intervet International B.V., Wim de Körverstraat, 35, 5831 AN Boxmeer, Netherlands.
01 December 2024	To change the potency release limits of the Clostridium spp. components that are tested by rabbit serology.
08 May 2024	Update to product information G.I.18.
18 March 2024	Removal of the bottle of 20 mL presentation.
29 March 2023	Number of changes introduced were to the production process of antigens.
08 February 2022	Change in the invented name of the veterinary medicinal product in IE only from Tribovax suspension for injection for cattle and sheep to Tribovax 10 suspension for injection for cattle and sheep.
28 April 2021	Deletion of manufacturer responsible for batch release.
11 March 2021	Change in the invented name of the veterinary medicinal product from Bovilis Bravoxin 10 Suspension for Injection for Cattle and Sheep to Bravoxin Suspension for Injection for Cattle and Sheep in Germany.