Veterinary Medicines Directorate

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Post Authorisation Assessments

Nuflor 40 mg/g Premix for Medicated Feeding Stuff for Pigs

Vm 01708/5112

07 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.
27 October 2024	One-off alignment of the product information with version 3 template.
22 March 2024	Change in the specification parameters of an active substance.
February 2024	Change in the specification parameters of an active substance.
09 February 2023	Tightening of specification limits for the active substance.
27 January 2022	Addition of a manufacturer of the active substance or addition of a site of manufacture.
14 August 2020	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
07 June 2018	Updates to the SPC and QRD texts.
29 June 2016	Change in the manufacturer of a starting material used in the manufacturing process.
06 March 2015	Change in the batch size of the active substance. Tightening of specification limits of the active substance.
20 November 2014	Change in test procedure for the active substance. Change in the re-test period of the active substance. To introduce a second manufacturing process.
31 July 2014	Deletion of a manufacturing site for the active substance.
17 February 2012	To change the specification parameter and/or limits of the finished product.
25 January 2012	Variation to change the MAH and distributor from Schering Plough Ltd to Intervet UK Ltd.
23 January 2012	Deletion of an active substance manufacturer.
20 December 2011	Renewal procedure – France as RMS.
16 December 2011	To add an additional manufacturing site. To add an alternative site for primary and secondary packaging operations and batch control/release testing of the finished product. Minor changes to the manufacturing process.
02 September 2011	To update the specification limits of the active substance.
14 March 2011	To change the name of the MAH in Portugal only.
09 February 2011	To add a new manufacturer of the active substance.
18 October 2007	Change in part of primary packaging material not in contact with finished product. Change in composition of immediate packaging: all other pharm forms.