Veterinary Medicines Directorate

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Post Authorisation Assessments

Nobivac FeLV, Suspension for Injection for Cats

Vm 05653/4059

•	20 July 2024	To modify the in-process controls during manufacture of the antigen including: add kinetic chromogenic technique for endotoxin detection; add a new pre-cast gel system and reducing agent for SDS-PAGE and Western Blot assays for identity/purity testing; an update to Ph. Eur. compliant starting materials for antigen manufacture and other editorial changes.
•	29 August 2023	Change in the SPC/QRD to implement the recommendation of a competent authority concerning pharmacovigilance data.
•	16 October 2020	Change in distributor details from Intervet UK Limited to MSD Animal Health UK Limited.
•	07 January 2020	Modification of an approved indication.
•	26 March 2019	Modification of an approved indication.
•	25 July 2018	Change in test procedure to reflect compliance with the Ph. Eur. and remove reference to outdated internal test methods and test method numbers.
•	06 July 2018	Changes to the labelling and/or package leaflet.
•	24 March 2017	Tightening of specification limits of the finished product. Minor change in the manufacturing process of an immediate release solid oral dosage form or oral solutions.
•	07 September 2011	Variation to seek approval for revised packaging.
•	05 October 2010	Renewal.
•	08 September 2010	Changes to an existing pharmacovigilance system.
•	08 July 2009	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category from POM to POM-V.
•	19 December 2008	Alterations to the SPC and Product Literature regarding compatibility of combined usage.
•	23 January 2008	Simple changes to packaging materials.
•	22 February 2007	Variation to change the concurrent use with another product.
•	11 November 2005	New Marketing Authorisation.