Veterinary Medicines Directorate

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Post Authorisation Assessments

Nobivac Rabies
Vm 01708/4325

•	23 December 2021	Change in the SPC, Labelling o/Package Leaflet intended to implement the outcome of a procedure concerning a PSUR. Editorial and QRD template updates to the SPC with associated product literature updates.
•	25 June 2021	Replacement of a test procedure for the finished product.
•	05 November 2020	Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited.
•	03 August 2017	Change in name of manufacturer. Change of specification of a former non Pharmacopoeial excipient starting material to comply with the Ph. Eur. or with a national pharmacopoeia of a Member State.
•	06 June 2017	Change of indications for minimum vaccination age
•	20 October 2016	Changes to section 4.8 of SPC for clarification of compatibility, as requested by VMD.
•	24 May 2016	Replacement of a test procedure for the active substance. Replacement of a test procedure for the finished product.
•	14 September 2011	Variation to change the SPC/Package Leaflet.
•	23 March 2011	Variation to change formulation of product.
•	05 July 2010	Renewal.
•	06 November 2008	Addition of a Manufacturer.
•	06 November 2008	Variation concerning the introduction of minor changes to the production procedure.
•	12 December 2007	Introduction of a new packaging style.
•	17 May 2006	Variation to bring the SPC/Product Literature in line with the Veterinary Regulations, 2005.
•	30 November 2005	Inclusion of a preservative and change of excipient.
•	25 October 2005	Reviewed Marketing Authorisation.