Veterinary Medicines Directorate

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Post Authorisation Assessments

Canigen Rabies
Vm 01708/4521

•	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.
•	02 December 2020	Change in the name of the MAH from Intervet UK Limited 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	Corrections to the SPC and Product Literature
•	23 March 2011	Change to starting materials for the production of the active substance
•	05 July 2010	Renewal
•	05 November 2008	Addition of a manufacturer of the active substance Minor changes to the manufacturing process
•	11 June 2008	Change of packaging style
•	26 July 2006	Change of legal category from POM to POM-V Changes to the SPC and Product Literature to bring in line with new legislation
•	30 November 2005	Change of formulation
•	25 October 2005	Review