



Post Authorisation Assessments

Nobivac Tricat Trio, Lyophilisate and Solvent for Suspension for Injection for Cats Vm 01708/3004

•	28 May 2021	Change in the address of a manufacturer used in the manufacture of the active substance.
•	14 August 2020	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
•	13 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS
•	01 June 2018	Change in the specification limits of the finished product.
•	05 January 2017	Changes to a DDPS following the assessment of the same DDPS in relation to another medicinal product of the same MAH.
•	28 April 2016	Change in test procedure for the finished product.
•	24 August 2015	Repeat Use Comms.
•	28 November 2014	Update to the DDPS.
•	30 October 2014	Approval of mock-ups.
•	31 July 2014	Addition of a distributor.
•	20 January 2014	Introduction of a new type of secondary packaging.
•	02 October 2013	Change in the name of manufacturer of the finished product.
•	12 October 2011	Renewal – UK as CMS.
•	29 June 2010	Changes to an existing pharmacovigilance system as described in the DDPS.
•	21 October 2009	To add an additional manufacturer of the active.
•	29 July 2009	To amend the SPC section 4.6 on adverse reactions, and update to section 4.8 to come in line with the latest QRD template.
•	14 January 2009	Correction/simple text layout changes to SPC and/or product literature.
•	09 July 2008	Variation to change reagent.
•	09 July 2008	Variation to change fill volume.
•	26 July 2007	New/Updates TSE Ph. Eur. Certificate of Suitability for excipient. TSE susceptible species.