Veterinary Medicines Directorate

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Post Authorisation Assessments

Nobilis ND Clone 30 Live
Vm 01708/4276

•	12 March 2024	The registration dossiers of the concerned products are supplemented with (i) the information on the use of animal derived trypsin in the manufacture of the hydrolysed gelatin and with (ii) respective extraneous agents and TSE risk assessments.
•	15 November 2022	To include the non-mixed associated use claim with Innovax-ND-ILT in the product information of Nobilis ND Clone 30.
•	19 January 2022	Addition of new tests and limits applied during the manufacture of the finished product. Addition to a test procedure for the finished product. Addition to a test procedure for the finished product. Changes to a test procedure for the finished product. Addition of new tests and limits applied during the manufacture of the finished product. Addition of a new container for the finished product. Changes in the manufacturing process of the finished product.
•	04 August 2020	Change in the SPC, labelling or package leaflet due to new data.
•	03 July 2020	Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited.
•	14 September 2017	Change in the immediate packaging of the finished product Change in the SPC, labelling or package leaflet due to new data.
•	28 March 2012	Variation to change the manufacturer of the Active Substance.
•	22 March 2012	Variation to change the finished product manufacturer.
•	27 November 2009	Renewal.
•	30 May 2008	Addition of a site for sterility testing. Change of a sterility test method.
•	30 May 2008	Variation to allow additional QC testing site.
•	16 July 2007	Alignment of the SPC and product literature between the UK and Ireland.
•	01 June 2007	Submission of an updated Certificate of Suitability.
•	26 July 2006	Ammendment of packaging material. Change of legal category. Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005.
•	12 January 2006	Addition of a QC testing site.