



Post Authorisation Assessments

Nobilis ND Clone 30 Live Lyophilisate for Oculonasal Suspension/Use in Drinking Water for Chickens and Turkeys

Vm 06376/5047

05 March 2026	One-off alignment of the product information with the national product information template v. 3 for GB-National MA and with the QRD template 9.1 for NI-National MA.
26 February 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
09 September 2025	Change in primary packaging material not in contact with the finished product formulation.
06 May 2025	Change of Marketing Authorisation Holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands.
10 January 2025	Change in test procedure for starting material used in the manufacturing process of the active substance. Change in test procedure for the finished product. Change in the storage conditions of the finished product. Change to in-process tests applied during the manufacture of the active substance. Changes in the manufacturing process of the active substance. Change in the manufacturing process of the active substance.
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	Amendment 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.