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Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021, please refer to the EMA website.

Innovax-ND-IBD Concentrate and Solvent for Suspension for Injection for Chickens

Vm 01708/5040

08 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
28 November 2025	To delete a site for antigen production, in process control testing, blending, primary packaging and secondary packaging.
07 February 2024	Update to the national template.
14 June 2022	Increase in the shelf-life of the finished product, from 2 years to 3 years.
06 April 2022	Changes to the labelling.
15 February 2022	Renewal.
27 January 2022	Change in the number of units in a pack within the range of the currently approved pack sizes of the finished product. Deletion of manufacturing site. Deletion of manufacturing site. Change in the number of units in a pack outside the range of the currently approved pack sizes of the finished product. Increase in the shelf-life of the finished product as packaged for sale, from 2 to 3 years. Change in storage conditions of the finished product. Deletion of the solvent container from the pack. Changes in the composition (excipients) of the finished product.
26 October 2021	Change in the manufacturer of a starting material used in the manufacturing process of the active.
24 March 2021	Change in the address of manufacture of the active substance.
17 March 2021	Change in the address of the manufacturer of the finished product.