Veterinary Medicines Directorate

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Post Authorisation Assessments

Nobilis Rhino CV Lyophilisate for Oculonasal Suspension for Chickens

Vm 06376/3061

16 May 2026	The proposed variation is to add the use of animal-component-free recombinant trypsin in the manufacture of the hydrolysed gelatine Byco-C produced by Croda to ensure uninterrupted supply of this stabiliser component.
14 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
August 2025	Change in legal entity of MA 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.
04 July 2025	Align the product information with the current QRD v9 EU template.
26 February 2024	The variation is to mention the use of animal derived trypsin in the manufacture of the Byco-C hydrolysed gelatin provided by Croda and to declare a change in the hydrolysis step of the gelatin (excipient) by replacing porcine trypsin by the use of either porcine or bovine trypsin.
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.
10 April 2019	Addition of a manufacturer of a starting material used in the manufacturing process of the active.
24 February 2017	Changes to the branding of the mock ups.
02 July 2012	Variation to change the manufacturing process.
02 April 2012	Variation to change the name of the finished product Manufacturer.
07 July 2010	Renewal.
13 June 2008	Addition of a second site for finished product testing.
30 January 2008	Transfer of legal category from POM to POM-V in accordance with the Veterinary Regulations, 2005.
10 July 2007	Addition of a new TSE European Pharmacopoeia Certificate of Suitability.
05 May 2005	New Marketing Authorisation.