



Post Authorisation Assessments

Nobivac Parvo-C Lyophilisate for Suspension for Injection for Dogs Vm 06376/4105

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| 01 March 2026 | Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. |
| 26 February 2026 | One-off alignment of the product information with the national product information template v.3. |
| 09 February 2026 | To delete sites for the manufacture of the active substance and the solvent. |
| 19 December 2024 | Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands |
| 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. |
| 05 January 2023 | To replace the tissue culture medium used during finished product formulation (blending) with a basal medium. |
| 18 November 2022 | Change(s) in the SPC, labelling or package leaflet to section 4.6 and 6. Changes to the SPC and / or product literature to sections 4.3, 4.4, 4.5; 4.8; 4.9 and 4.10. |
| 20 January 2022 | Change in the SPC, labelling or package leaflet due to new data. |
| 23 April 2021 | Change in the address of a manufacturer used in the manufacture of the active substance. |
| 05 November 2020 | Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited. |
| 27 July 2017 | Update of the pharmaceutical form on the SPC to align with the EDQM Standard Term and to update the full product name in line with QRD requirements. |
| 05 September 2013 | Change of name of a manufacturer if the finished product |
| 15 December 2011 | Change in batch size of the finished product |
| 08 July 2010 | Renewal |
| 18 March 2009 | Addition of a manufacturing site for part of the manufacturing process of the finished product |
| 19 January 2009 | Change to batch safety test on the finished product |
| 07 December 2007 | Introduction of new packaging material |
| 01 June 2007 | Submission of an updated Ph. Eur. Certificate of Suitability for an excipient |
| 14 September 2006 | Change of shelf life of the active substance from 24 to 36 months |
| 26 July 2006 | Change of legal category from POM to POM-V |

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| | Changes to the SPC and Product Literature to bring in line with new legislation |
| 24 October 2005 | Review |
| 13 June 2005 | Renewal |
| 10 June 2005 | Change of distributor |
| 09 December 2003 | Change of re-vaccination interval from every 2 years to every 3 years |
| 04 April 2003 | Submission of a TSE Certificate |
| 17 February 2003 | Change of batch size of the finished product |
| 13 August 2002 | Change of re-vaccination interval from every 1 year to every 2 years |
| 04 February 2002 | Change of manufacturing site of the active substance |
| 08 November 2001 | Addition of a distributor in Northern Ireland |
| 22 October 2001 | Change of formulation |
| 30 November 2000 | Change of manufacturing site of the dosage form |
| 25 July 2000 | Renewal Change of MAH address |
| 21 September 1999 | Change of dosage particulars |
| 25 February 1998 | Change of specification for the finished product |
| 26 March 1997 | Change of product name from 'Nobivac Parvo c' to 'Nobivac Parvo-C' |
| 19 September 1996 | Additional presentation |
| 20 November 1995 | Change of product name from 'Nobi-vac Parvo-c' to 'Nobivac Parvo c' |
| 07 August 1995 | Renewal Change of manufacturer of the active substance |