

Post Authorisation Assessments

Rispoval Pasteurella Lyophilisate and Solvent for Emulsion for Injection Vm 60021/3019

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| 16 December 2025 | To add an already authorised site for Physical/Chemical testing of the finished product to the manufacturing flow chart. |
| 12 December 2025 | One-off alignment of the product information with the national product information template v. 3. |
| 15 November 2024 | Change of Legal Entity for UK(NI) from Zoetis UK Limited to Zoetis Belgium S.A. |
| 01 May 2020 | Change in the address of the marketing authorisation holder from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP. |
| 23 May 2018 | Tightening of specification limits of the finished product. Tightening of specification limits of the finished product. Deletion of a non-significant specification parameter of the finished product. Submission of 4 updated Ph. Eur. TSE certificates of suitability for starting materials from an already approved manufacturer. Change in the specification limits of the finished product. |
| 26 January 2016 | Change in the sterility test and addition of an enhanced sterility test for the finished product (diluent). |
| 01 September 2014 | Change in a test procedure for a starting material used in the manufacturing process of the active substance. |
| 17 February 2010 | Addition of suppliers of starting material. |
| 22 January 2009 | Renewal |
| 18 July 2008 | Alignment of the SPC and product literature between the UK and Ireland and change of legal category from POM to POM-V. |
| 28 October 2005 | Renewal. |
| 25 July 2005 | Addition of a distributor. |
| 26 March 2004 | Change to ingredient specification. |
| 19 June 2002 | Change of shelf life of finished product. |