



Post Authorisation Assessments

Ovipast Plus Suspension for Injection for Sheep Vm 06376/5064

30 October 2025	<p>Change in shape and dimensions of the closure and container.</p> <p>Addition of a test procedure for active substances.</p> <p>Addition of a test procedure for active substances.</p> <p>Minor change to the in-process control test used in the manufacturing process of the active substances.</p> <p>Minor change to the in-process control test used in the manufacturing process of the active substances.</p> <p>Deletion of in-process test applied during the manufacture of the active substances</p> <p>Deletion of in-process test applied during the manufacture of the active substances.</p> <p>Changes in the manufacturing process of the active substances.</p> <p>Changes in the manufacturing process of the active substances</p> <p>Addition of a manufacturing site for manufacturing of active substances.</p> <p>Addition of a manufacturing site for manufacturing of active substances.</p>
04 August 2025	<p>One-off alignment of the product information with version 9.0 of the QRD templates.</p>
18 November 2024	<p>Change in legal entity of MA holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands.</p>
17 April 2023	<p>Addition of an in vitro potency ELISA for <i>C. perfringens</i> type C beta toxoid.</p> <p>Optimisation of the statistical analysis method for the Fab ELISA to confirm <i>M. haemolytica</i> identity and <i>P. trehalosi</i> identity/potency.</p> <p>Addition of Intervet International B.V., Boxmeer, NL, as finished product Quality Control test site for the <i>C. perfringens</i> type C beta toxoid potency test, <i>M. haemolytica</i> identity test and <i>P. trehalosi</i> identity/potency test.</p>
19 October 2022	<p>Endotoxin test: replacement turbidimetric kinetic method with chromogenic kinetic method at QC testing site in Boxmeer (NL).</p>
12 August 2022	<p>Replacement of test method for the finished product.</p>
16 February 2022	<p>Changes to a test procedure (including replacement or addition) for the active substance.</p> <p>Change(s) in the manufacturing process of the active substance.</p> <p>Change(s) in the manufacturing process of the active substance.</p> <p>Change(s) in the manufacturing process of the active substance.</p> <p>Change(s) in the manufacturing process of the active substance.</p> <p>Change(s) in the manufacturing process of the active substance.</p>

	Change(s) in the manufacturing process of the active substance.
15 July 2021	Minor changes to an approved test procedure of the finished product. Minor changes to an approved test procedure of the finished product. Update of the test procedure to comply with the updated general Ph. Eur monograph. Addition to a test procedure for the finished product. Addition of a site where batch control/testing takes place.
15 June 2021	Change in the manufacturing process of the finished product.
01 October 2020	Change in the name of a manufacturer of the finished product, also responsible for batch release.
11 September 2020	Change in the name 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.
24 April 2019	Addition of a manufacturer responsible for batch release of the finished product.
18 May 2018	Change in the manufacturing process of the active substance. Change in the manufacturing process of the active substance.
01 May 2018	Change in RMS from UK to ES.
30 March 2012	Change in the name of the manufacturer of the finished product.
08 March 2012	Updates to the Part II dossier including; change in the control of the active substance, change in the control of the finished product, change in the manufacturer of the active substance, change in the manufacturer of the finished product.
28 October 2009	Addition of a site of filling of the finished product.
01 June 2006	Renewal
20 June 2005	Change of distributor.
30 August 2002	Addition of sites for finished product labelling and packaging and change in site of batch release.
04 January 2002	Change of a supplier of an ingredient.
22 October 2001	Change of distributor.