



Post Authorisation Assessments

Proactive 1.52 mg/ml Teat Dip/Spray Solution Vm 17140/5001

10 December 2025	Updates to Part II of the dossier following an SRP application.
11 August 2025	Amendments to product literature subsequent to an SRP.
11 December 2024	One-off alignment of the product information with version 9.0.
25 April 2024	Change in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
19 March 2024	Change in the name or address or contact details of a qualified person for pharmacovigilance.
22 June 2018	Change of RMS from UK to IE.
03 March 2017	Harmonisation of the SPC between the first wave and repeat use member states.
16 May 2016	Variation to change the name of the MAH from DeLaval International AB to DeLaval NV
05 September 2011	Grouped variation concerning the addition of a batch size for the finished product, and a change to the manufacturing process.
27 July 2011	Extension of the finished product shelf life as packaged for sale.
14 October 2009	Variation to change the SPC as requested by CMS.
26 February 2009	MRP – UK as RMS.
11 January 2007	Renewal.
11 February 2005	Replacement of an excipient with a comparable excipient.
10 January 2001	Change of name of the Marketing Authorisation Holder.