



Post Authorisation Assessments

AviPro Thymovac Lyophilisate for Use in Drinking Water. Vm 00879/5042

•	17 August 2021	Deletion of a non-significant parameter of an active substance used in the manufacturing process of the active substance Deletion of a specification parameter of the finished product.
•	16 December 2020	Change of MAH from Lohmann Animal Health GmbH, Heinz-Lohmann-Straße 4, D-27472 Cuxhaven, to Elanco Europe Ltd., Form 2, Bartley Way, Bartley Wood Business Park, Hook, RG27 9XA, United Kingdom.
•	13 June 2019	Update the Part 2 of the dossier in VNeS-format.
•	05 June 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	27 April 2017	Change in the manufacturing process of the active substance.
•	18 January 2017	Minor changes to an approved test procedure of the finished product. Changes to a test procedure for the finished product.
•	21 November 2014	Update to the DDPS.
•	07 February 2014	Changes to an existing pharmacovigilance system as described in the DDPS.
•	08 July 2013	Renewal.
•	16 April 2013	Change to batch release arrangements and quality control testing of the finished product. Change in test procedure for the finished product.
•	16 May 2012	Change in name of MAH. Change in name of Manufacturer of the finished product.
•	09 June 2011	Change to an existing Pharmacovigilance system as described in the DDPS.
•	30 November 2010	Change in container closure system of the finished product.
•	10 August 2010	To submit an updated part 2 to the current manufacturing and control.
•	03 February 2010	To change the described testing procedure for extraneous viruses
•	02 December 2009	To submit joint-labelling between the UK and Ireland after MRP.
•	14 September 2009	Change of QPPV vaccines.