



## Post Authorisation Assessments

### Eurican DAPPi Lyophilisate and Solvent for Suspension for Injection Vm 61700/3038

19 February 2026	To remove a supplier of a starting material and to delete the certificate of suitability.
13 February 2026	To delete a site for manufacturing of the diluent.
12 February 2026	To update the value of pH adjustment in the culture step of the manufacture of an active substance.
12 February 2026	To make editorial changes to the wording in the dossier (Part 2E) to clarify the frequency of finished product testing. To make editorial changes to include reference to a starting material (Phenol red) already authorised to adjust the pH of culture media in the dossier (Part 2B) and to include reference to previously omitted active substance (distemper virus AI) in the list of starting materials (Part 2C).
25 July 2025	Change of Marketing Authorisation Holder from Boehringer Ingelheim Animal Health UK Limited, Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS, United Kingdom to Boehringer Ingelheim Vetmedica GmbH, Binger Strasse 173, 55216 Ingelheim am Rhein, Germany (NI-only).
03 June 2025	Update to the description of starting materials of biological origin.
13 March 2025	This variation was aimed to change the name and address change of a sub-contractor for some of the approved release tests.
18 May 2024	Section 5.4 Nature and composition of immediate packaging Immediate container: type I glass vials with chlorobutyl rubber stoppers, sealed with aluminium caps.
18 May 2024	Addition of new pack sizes, to add the possibility to supply the lyophilisate and the solvent separately. Compatibility claim with the new Boehringer Ingelheim's Eurican L4 vaccine. Updating the product information according to the last QRD template in force.
22 September 2023	Extension of a storage period of the active substance supported by real time data.
11 April 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
15 March 2023	To reduce the minimum volume for the CPV active ingredient formulation in DAP and DAPPI lyophilisates.
10 February 2023	The variation is to introduce the use of recombinant trypsin as a substitute to porcine trypsin for the manufacture of the active substance.
31 January 2023	Reduction of the minimum volume criterion of the CPV active ingredient for the formulation of lyophilisate.

17 March 2022	Change of a test procedure for the active substance. Changes in the manufacturing process of the active substance.
21 January 2022	Deletion of a non-significant specification parameter of the finished product. Replacement to a test procedure for the finished product.
22 October 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
16 July 2021	Change in the batch size (including batch size range) of the active substance used in the manufacturing process of the active substance. Change in storage conditions of the active substance. Change in the manufacturing process of the active substance.
15 February 2021	Deletion of manufacturing site for the finished product.
04 February 2021	Renewal - UK as CMS.
26 November 2020	Change in the name of the manufacturer of the finished product.
22 July 2020	Change in the name of a manufacturer of the active substance.
18 June 2020	Change in the name of the manufacturer of the finished product.
27 May 2020	Change in the name of a manufacturer of the finished product, also responsible for batch release.
16 December 2019	Change of a test procedure for the active substance.
05 November 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
16 November 2018	Change in the name and address of the marketing authorisation holder from Merial Animal Health Limited, Sandringham House, Sandringham Avenue, Harlow Business Park, Harlow, Essex, CM19 5TG, United Kingdom to Boehringer Ingelheim Animal Health UK Limited, Ellesfield Avenue, Bracknell, Berkshire, RG12 8YS, United Kingdom.
12 April 2018	Change of a test procedure for the finished product.
31 October 2017	Replacement of a test procedure for the active substance.
22 December 2016	Tightening of specification limits of the finished product.