



Post Authorisation Assessments

AviPro Salmonella Duo Lyophilisate for Use in Drinking Water Vm 52127/3005

07 January 2026	To add a new withdrawal period after the 4th dose of vaccine. To include the possibility of administration of an additional dose of vaccine to chickens during lay.
18 March 2025	Change in legal entity from Elanco Europe Ltd., Form 2, Bartley Way, Bartley Wood, Business Park, Hook, RG27 9XA, United Kingdom to Elanco GmbH, Heinz-Lohmann Strasse 4, Groden, D-27472, Cuxhaven, Germany.
19 January 2025	Change(s) 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. (NI).
13 December 2024	Add a new pack size of 4000 doses. Add type I bromobutyl rubber stoppers. Update of the cooling temperature of the fermentation culture from "20 – 25°C" to "2 – 25°C". Inclusion of a real-time polymerase chain reaction method to differentiate Salmonella vaccine strains from wild-type field isolates. Removal of gelatin and peptone from the finished product. Update of the product information to version 9.0 of the QRD-template.
04 May 2024	Changes to a non-significant specification parameter of a starting material used in the manufacturing process of the active substance.
19 March 2024	Changes to a non-significant specification parameter of a starting material used in the manufacturing process of the active substance.
31 May 2022	Repeat Use to add one CMS
23 June 2021	Changes in the manufacturing process of the finished product. Change in the specification limits 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.
16 September 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Update of format of Part 2 of the dossier.
05 June 2019	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
29 November 2017	Minor change in the manufacturing process of the active substance.

12 July 2017	Increase in batch size of active substance or intermediate used in the manufacturing process of the active substance.
27 March 2017	Repeat Use application to add one new Member State.
28 July 2016	Renewal – UK CMS
17 December 2014	Minor change to a test procedure for an excipient.
21 November 2014	Update to the DDPS.
19 March 2014	Addition of turkey as a target species.
07 February 2014	Changes to an existing pharmacovigilance system.
16 May 2012	Change to the name of the MA holder, change to the name of the active substance manufacturer and change to the name of the finished product manufacturers responsible for packaging, quality control and batch release.