

Post Authorisation Assessments

Avishield ND, Lyophilisate for Oculonasal Suspension/Use in Drinking Water, for Chickens and Turkeys Vm 61812/5000

12 March 2026	Changing the legal entity of the MAH from Genera d.d., Svetonedeljska cesta 2, Kalinovica, 10436 Rakov Potok, Croatia to IZO S.r.l. a socio unico Via San Zeno 99/A, 25124 Brescia, Italy.
12 August 2025	Change in the manufacturing process of the finished product. Deletion of an obsolete in-process parameter.
20 May 2025	Addition of Dechra Veterinary Products Limited, Sansaw Business Park, Hadnall, Shrewsbury, Shropshire, SY4 4AS, UK as a local representative (NI).
30 April 2025	Change of named distributor details to Genera d.d., Svetonedeljska cesta 2, Kalinovica, 10436 Rakov Potok, Croatia.
23 April 2025	Addition of details of the local representative on the outer package and the package leaflet.
12 February 2025	Add claim for use in egg laying period. Deletion of the safety warning not to use the vaccine during lay. G.I.18 update of the product information to version 9.
18 April 2023	Changes in the manufacturing process of the active substance. Changes in the manufacturing process of the active substance. Changes in the manufacturing process of the active substance. Changes in the manufacturing process of the active substance.
09 February 2023	Tightening of specification limits for finished products subject to Official Control Authority Batch Release.
20 October 2022	Tightening of specification limits for finished products subject to Official Control Authority Batch Release.
08 March 2022	Changes to a test procedure for a starting material.
12 January 2022	Deletion of a non-significant parameter of an active substance in the manufacturing process of the active substance. Changes to a test procedure for the active substance. Changes to a test procedure for the active substance. Deletion of a specification parameter of the finished product.
25 June 2021	Changes to a test procedure for the finished product.
25 February 2021	Renewal – UK as CMS.
10 March 2020	Addition of a supplier of packaging components or devices. Deletion of a non-significant parameter of an active substance used in the manufacturing process of the active substance.
07 August 2019	Increase in the shelf-life of the finished product, from 12 months to 24 months
17 May 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS

02 April 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS
05 December 2018	Change in RMS from UK to NL.
17 August 2018	Change in the local representative and distributor in Germany
23 May 2018	Change in the address of a manufacturer used in the manufacture of the active substance. Change in the address of a manufacturer of the finished product, also responsible for batch release. Change in the address of the Marketing Authorisation Holder, to add 'Cesta'.
15 May 2018	Addition of a supplier of packaging components or devices. Addition of a supplier of packaging components or devices. Deletion of a non-significant specification parameter of an excipient. Deletion of a non-significant specification parameter of the immediate packaging of the finished product. Deletion of a non-significant specification parameter of the immediate packaging of the finished product.
01 March 2018	Change in the fill volume of the finished product.
26 July 2017	Changes in the manufacturing process of the finished product.
21 March 2017	Change to the QPPV. Change to the database system outlined in the DDPS.