



## Post Authorisation Assessments

### AquaVac PD3 Emulsion for Injection for Atlantic Salmon

Vm 06376/5055

09 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
14 August 2025	Changing the legal entity of the Marketing Authorisation Holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands.
02 July 2025	One-off alignment of the product information with the national product information template v. 3.
04 April 2025	To add an additional method for the sterilisation of simethicone (anti-foaming agent) using irradiation $\geq 25$ kGy.
19 September 2023	Addition of alternative sterilisation method of the immediate packaging of the finished product.
20 December 2022	Increase of the maximum production volume of the IPNV antigen from 300 l to 1000 l. IPNV bulk antigen shelf-life extension from 20 to 22 months. Addition of a PE storage container for IPNV antigen. Addition of a site for the manufacture of the active substance infectious pancreatic necrosis virus (IPNV) antigen and related in-process controls. Addition of a site for finished product Quality Control testing.
14 August 2020	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
18 March 2020	Renewal – UK as CMS.
13 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS
01 December 2016	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
12 November 2015	Change in the shelf life of the finished product up to 19 months.