



## Post Authorisation Assessments

### AquaVac ERM Concentrate for Dip Suspension for Rainbow Trout Vm 01708/4564

• 27 June 2023	Change in the name and address of the manufacturer of the finished product. (GB) Change in the name and address of the manufacturer of the active substance. (GB)
• 27 September 2021	Addition to a test procedure for the finished product. Addition of a site where batch control/testing takes place.
• 01 October 2020	Change in the name of a manufacturer of the finished product, also responsible for batch release.
• 14 August 2020	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
• 20 December 2018	Replacement of a manufacturer responsible for batch release of the finished product. Replacement of a manufacturer responsible for batch release of the finished product. Deletion of a packaging site. Replacement of a secondary packaging site of the finished product.
• 09 February 2018	Change in the RMS from UK to IE.
• 08 July 2015	Change to the design and technical specifications of the product labels.
• 15 May 2014	Change in the name and address of the manufacturing authorisation holder.
• 27 January 2012	Change in the immediate packaging for the finished product.
• 04 August 2011	To change the name of the MAH in Portugal only, from Schering-Plough II to Intervet Portugal.
• 15 October 2010	Renewal.
• 08 September 2009	Change of Marketing Authorisation Holder.
• 14 August 2009	Change of Marketing Authorisation Holder.
• 10 January 2007	Variation to increase the size of maximum blending volume of final product.
• 11 December 2006	Variation to change Marketing Authorisation Holder in Denmark only.
• 21 March 2006	MRP procedure – UK as RMS.