Veterinary Medicines Directorate

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Post Authorisation Assessments

AquaVac ERM Concentrate for Dip Suspension for Rainbow Trout

Vm 01708/3047

•	28 April 2024	Change in name and address details of the manufacturer of the finished product. Change in name and address details of the manufacturer of the active substance.
•	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.