Veterinary Medicines Directorate

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Post Authorisation Assessments

AquaVac Vibrio Immersion and Injection

Vm 01708/4569

•	07 June 2023	Change in name and address details of a manufacturer of the finished product. Change in name and address details of a manufacturer of the active substance. (NI)
•	22 December 2022	Change in name and address details of a manufacturer of the finished product. Change in name and address details of a manufacturer of the active substance.
•	08 October 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. Addition of a batch release site of the finished product. Replacement of a secondary packaging site of the finished product.
•	25 April 2014	Change of Manufacturing Authorisation Holder.
•	18 January 2012	Change in immediate packaging of the finished product.
•	02 September 2011	To change the MAH in Portugal only.
•	13 July 2011	Renewal Marketing Authorisation – France as RMS.
•	09 February 2011	To change the MAH from Schering-Plough Ltd to Intervet UK Ltd.