



Post Authorisation Assessments

AquaVac FNM Plus Emulsion for Injection for Fish

•	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.
•	12 December 2018	Addition of a manufacturer responsible for batch release of the finished product. Replacement of a manufacturer responsible for batch release of the finished product. Replacement of a secondary packaging site of the finished product
•	09 February 2018	Change in the RMS from UK to ES.
•	15 May 2014	Change in the name and address of the manufacturing authorisation holder.
•	27 January 2012	Change of immediate packaging for the veterinary medicinal product.
•	04 August 2011	To change the name of the MAH in Portugal.
•	14 February 2011	Renewal procedure – UK as RMS.
•	15 September 2010	To change the Marketing Authorisation Holder and distributor.
•	07 July 2010	To change the legal category from POM-V to POM-VPS.
•	14 August 2009	To change the name of the Marketing Authorisation Holder in France.
•	22 May 2008	Change of constituent.
•	07 March 2007	New Marketing Authorisation – MRP. UK as RMS.
•	19 December 2006	Change in in-process controls applied during manufacture of the product.
•	07 December 2006	Change of Marketing Authorisation Holder address (Schering-Plough AS) in Denmark.
•	30 June 2005	Variation to bring the data in line with the updated dossier.
•	22 December 2004	Replacement of an in-process test.
•	22 December 2004	Addition of QP release.
•	26 November 2004	Change of Marketing Authorisation Holder.
•	18 November 2004	Addition of a site of assembly.
•	03 August 2004	Addition of a batch size.
•	18 November 2003	Change to in-process controls.
•	23 October 2002	Change in the storage time of the active ingredient.
•	20 September 2002	Change to the in-process controls.

•	30 November 2001	Deletion of an in-process test.
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