



Post Authorisation Assessments

Nobilis Reo + IB + G + ND Vm 01708/4332

•	28 May 2021	Minor change to an approved test procedure for the active substance used in the manufacturing process of the active substance.
•	23 April 2021	Change in the address of a manufacturer used in the manufacture of the active substance.
•	03 July 2020	Change in the name of the marketing authorisation holder from Intervet UK Ltd to MSD Animal Health UK Limited.
•	11 November 2013	Changes in manufacturing process of the Active Substance.
•	28 March 2012	Variation to change the name/address of an Active Substance Manufacturer.
•	22 March 2012	Variation to change the Manufacturer of the finished product.
•	19 May 2011	Variation concerning a change to various test methods.
•	19 March 2010	Renewal.
•	18 October 2007	Addition of a site for finished product QC Testing.
•	31 May 2007	Addition of an Active Substance Manufacturer.
•	04 January 2007	Variation to bring the SPC/Labelling in lien with the Veterinary Regulations, 2005. Transfer of legal category from PML to POM-V.
•	23 September 2005	Renewal.
•	23 September 2005	Renewal.
•	23 September 2005	Renewal.
•	13 September 2005	Marketing Authorisation Conversion.
•	11 August 2005	Variation to update test methods to comply with the Ph. Eur.
•	27 May 2005	Change of Distributor.
•	07 November 2001	Change of Manufacturer/Assembler of Dosage Form.
•	26 October 2001	Variation to change a container type.
•	31 August 2001	Addition of a Distributor.
•	02 August 2000	Renewal.
•	13 July 2000	Change in the address of the Marketing Authorisation Holder.
•	13 July 2000	Renewal.
•	05 August 1998	Variation concerning the product formulation.
•	16 July 1998	Renewal.
•	16 July 1997	Renewal.
•	24 April 1997	Change of product name.

•	24 April 1997	Change in control test particulars.
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