Veterinary Medicines Directorate

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Post Authorisation Assessments

Aquatet 100% w/w Premix for Medicated Feeding Stuff

Vm 11003/4002

•	13 July 2021	Deletion of manufacturing site for an active substance.
•	27 October 2020	Deletion of a non-significant parameter of an active substance. Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State.
•	18 September 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance excipient from a new manufacturer.
•	21 March 2018	Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State.
•	23 November 2012	Submission of a new Ph. Eur. Certificate of Suitability for a new manufacturer of the active substance.
•	24 May 2011	Increase of shelf life of the finished product from 1 year to 18 months.
•	12 January 2011	Addition of a 25kg presentation.
•	12 January 2011	Change in specification of the finished product.
•	20 May 2009	Removal of manufacturing site for the active substance and change of address of a manufacturer of an active substance.
•	24 February 2009	Corrections/simple text layout changes to the SPC and/or Product Literature.
•	09 July 2008	Changes to bring the SPC and Product Literature in line with new legislation and change in legal category from MFS to POM-V.
•	22 June 2007	Renewal.
•	22 October 2004	Change of MAH from Alpharma Animal Health Ltd to Pharmaq Limited.
•	22 July 2004	Renewal.
•	22 July 2004	Increase of withdrawal period from 400 to 720 degree days.
•	30 May 2001	Change of MA holder name from Vetrepharm Ltd to Alpharma Animal Health Ltd.
•	13 March 1997	Addition of a manufacturer of the active substance.
•	22 February 1996	Addition of a manufacturer of the active substance.