



Post Authorisation Assessments

Aquatet 100% w/w Premix for Medicated Feeding Stuff Vm 11003/5001

05 March 2025	Update to the finished product specification based on the latest Ph.Eur Oxytetracycline hydrochloride monograph (0198) as a 100% active product.
14 November 2024	Change to comply with Ph. Eur.: change to comply with an update of the relevant monograph of the Ph. Eur.
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.

