



Post Authorisation Assessments

Doxycycline Hyclate VMD 500 mg/g Powder for Use in Drinking Water for Chickens Vm 19968/4001

•	22 July 2022	Deletion of manufacturing site for an active substance. Submission of an updated certificate of suitability. Submission of an updated certificate of suitability.
•	28 February 2018	Submission of an updated Ph. Eur. Certificate of Suitability for an active substance from an already approved manufacturer. Addition of a new Ph. Eur. Certificate of Suitability for an active substance from a new manufacturer.
•	19 February 2018	Change in the invented name of the veterinary medicinal product from Doxyveto 500 mg/g powder for use in drinking water for chickens to Doxycycline Hyclate VMD 500 mg/g Powder for Use in Drinking Water for Chickens
•	17 November 2016	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
•	16 November 2016	Renewal.
•	20 July 2016	Submission of an updated certificates of suitability.
•	21 November 2014	Submission of a new and an updated Ph. Eur Certificate of Suitability.
•	23 May 2012	Minor change in the manufacturing process of the finished product. Replacement of batch control/testing site. Addition of a secondary packaging site. Replacement of primary and secondary packaging site. Replacement of finished product manufacturing site.