



Post Authorisation Assessments

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

17 March 2026	Submission of a Ph. Eur. CEP for an active substance. Submission of a Ph. Eur CEP for an active substance.
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.