Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

Noroclav 500 mg Chewable Flavoured Tablets for Dogs

•	30 July 2019	Addition of a manufacturer responsible for batch release of the finished product.
•	30 July 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	01 August 2017	Renewal – UK CMS
•	12 July 2017	Submission of an updated Ph. Eur. certificate of suitability for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance.
•	11 May 2016	Increase in batch size (including batch size range*) of the finished product.
•	19 January 2016	Submission of a new certificate of suitability.
•	10 April 2015	Submission of a new and an updated Ph. Eur. Certificate of Suitability.
•	28 November 2014	Update to the DDPS.
•	11 February 2014	Change in test procedure for the finished product.
•	30 May 2013	Change in name of medicinal product in Denmark only from ‘Veclavam Flavour vet’ to ‘Veclavam vet’.
•	13 February 2013	Alteration to carton text size.
•	30 November 2012	To change the name of the veterinary medicinal product in Belgium, Denmark, France, Luxembourg, Netherlands, Norway, Portugal, Spain and Sweden only.