



Post Authorisation Assessments

Benefortin Flavour 20 mg Tablets for Dogs Vm 32823/4008

•	12 July 2023	Minor changes to an approved test procedure for the finished product.
•	18 June 2020	Change in the address of the marketing authorisation holder from Lavet Pharmaceuticals Ltd, H-1161 Budapest, Ottó u. 14, Hungary to Lavet Pharmaceuticals Ltd., 2143 Kistarcsa, Batthyány u. 6., Hungary.
•	24 October 2019	Deletion of manufacturing site for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	25 September 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	11 June 2019	Change of distributor from Boehringer Ingelheim Limited to Boehringer Ingelheim Animal Health UK Limited.
•	05 March 2019	Changes to the labelling, or the package leaflet, which are not connected with the SPC.
•	17 May 2016	Renewal – UK CMS
•	11 December 2015	Change in the (invented) name of the medicinal product in Poland
•	25 September 2014	Joint-labelling variation.
•	14 February 2014	Change to the SPC and product literature.
•	07 February 2014	Submission of a new and an updated Ph. Eur. Certificate of Suitability.