



Post Authorisation Assessments

Carprieve 50 mg Flavoured Tablets for Dogs

•	13 January 2020	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	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.
•	26 July 2019	Addition of a manufacturer responsible for batch release of the finished product.
•	21 May 2019	Change in the manufacturer of a starting material used in the manufacturing process of the active substance.
•	28 January 2019	Change in RMS from UK to IE.
•	07 August 2018	Tightening of specification limits of an excipient. Change in the specification parameters and/or limits of an excipient.
•	20 May 2016	Renewal – UK as RMS
•	25 September 2014	Change of QPPV and update to the DDPS.
•	26 October 2012	Variation to change the product name in France and Germany only.
•	18 July 2012	To change the distributor address.
•	22 February 2012	Reduction of text on the single carton to incorporate three languages.
•	07 July 2011	Change in invented name of the veterinary medicinal product in Hungary and Sweden only.