



Post Authorisation Assessments

Amlodipine 1.25 mg Ceva Chewable Tablets for Cats

•	15 March 2019	The RMS has changed from UK to FR
•	07 August 2018	Change in the invented name of the veterinary medicinal product from Amodip 1.25 mg chewable tablets for cats to Amlodipine 1.25 mg Ceva chewable tablets for cats, in DE/FR/NL and to Amlodipina Ceva 1.25 mg, compresse masticabili per gatti in IT.
•	24 May 2018	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	11 October 2017	Change in the invented name of the veterinary medicinal product in the United Kingdom only from Amodip 1.25 mg Chewable Tablets for Cats to Amlodipine 1.25 mg Ceva Chewable Tablets for Cats.
•	19 September 2017	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 March 2017	Change in shelf-life of the finished product from 30 months to 3 years.
•	29 November 2016	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	15 September 2016	Change in the name of a manufacturer of the finished product including manufacturer responsible for batch release. Change in the name of a manufacturer of the finished product. Deletion of a manufacturing site for an active substance.
•	06 September 2016	Change in the name and address of the MAH in Italy only.
•	29 June 2016	Introduction of a new pharmacovigilance system which has been assessed by the relevant national competent authority/EMA for another product of the same MAH.
•	16 June 2016	Change of MAH, from Sogeval to Ceva Animal Health Ltd.