



Post Authorisation Assessments

Clinacin 300mg Tablets for Dogs

Vm 11990/4053

•	28 January 2022	Changes in the qualitative and quantitative composition of the immediate packaging of the finished product for solid pharmaceutical forms.
•	15 October 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer. Deletion of Ph. Eur. certificates of suitability for an active substance. Deletion of Ph. Eur. certificates of suitability for an active substance.
•	30 September 2020	Addition of components (excipients) of the flavouring or colouring system of the finished product. Increase in the shelf-life of the finished product after first opening, to 72 hours. Changes in break lines intended to divide into equal doses.
•	17 February 2020	Deletion of a non-significant specification parameter of the finished product.
•	22 March 2019	Change in the shelf-life of the finished product stored in HDPE containers to 5 years
•	20 October 2016	Submission of an updated certificate of suitability.
•	26 February 2015	Submission of a new Ph. Eur. Certificate of Suitability for an active substance, from an already approved manufacturer.
•	31 October 2014	Renewal.
•	06 May 2014	Extension – submission of a new or updated certificate of suitability.