Veterinary Medicines Directorate

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Post Authorisation Assessments

Clavamox LC Intramammary Suspension for Lactating Cattle

•	28 October 2021	Submission of a new certificate of suitability for an active substance (used in manufacturing process of active).
•	21 October 2020	Deletion of manufacturing site for an active substance. Deletion of Ph. Eur. certificates of suitability for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	13 November 2019	Change in the address of the marketing authorisation holder from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
•	09 May 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer
•	23 October 2018	Addition of a manufacturing site of the finished product.
•	15 October 2018	Renewal – UK as RMS.
•	09 October 2018	Change in RMS from UK to DE.
•	25 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	28 March 2018	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	08 March 2018	Change in the specification limits of the finished product.
•	18 October 2017	Addition of an alternative sterilisation site for the active substance.
•	28 June 2017	Repeat Use application to add 2 new member states
•	13 July 2016	Addition of an alternative sterilisation site for the active substance. Addition of an alternative sterilisation site for the active substance. Introduction of a new site of manufacture Submission of a new certificate of suitability for an active substance
•	17 May 2016	Submission of an updated Ph. Eur. certificate of suitability. Submission of an updated Ph. Eur. certificate of suitability.
•	02 June 2015	Addition of quantitative specification limits in finished product specification.
•	20 April 2015	Addition of a specification limit for the active substance.