



Post Authorisation Assessments

Kenocidin Chlorhexidine Digluconate 5mg/ml, Teat Dip Solution for Cattle (Dairy) Vm 22136/4002

•	18 April 2024	Changes in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). (NI)
•	09 September 2021	Change in the name of a supplier of active substance used in the manufacture of the active substance. Replacement of components (excipients) of the flavouring or colouring system of the finished product. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	05 May 2021	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	18 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. TSE certificate of suitability for a starting material from an already approved manufacturer.
•	04 October 2018	Change in RMS from UK to BE.
•	10 September 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	16 April 2015	Renewal – UK as RMS.
•	06 February 2015	Submission of an updated Ph. Eur. Certificate of Suitability.
•	20 April 2011	New MA - MRP