



Post Authorisation Assessments

CIDR OVIS 0.35 g Vaginal Delivery System for Sheep Vm 42058/5118

•	04 May 2024	Update to the latest version of the QRD.
•	11 January 2024	Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance.
•	27 July 2022	Deletion of a manufacturer of an active substance.
•	21 June 2022	Deletion of a manufacturer of an active substance.
•	18 May 2022	Renewal.
•	24 November 2021	Submission of a new Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Deletion of Ph. Eur. certificates of suitability for an active substance.
•	05 October 2020	Replacement to a test procedure for the finished product. Addition to a test procedure for the finished product.
•	06 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active from an already approved manufacturer
•	17 January 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. certificate of suitability for an active substance from an already approved manufacturer.
•	12 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.
•	23 January 2019	Change(s) in the SPC, labelling or package leaflet further to a veterinary PSUR.
•	25 September 2018	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	17 April 2018	Change of RMS from UK to ES.
•	13 February 2018	Repeat use MRP to add 1 new member state