

## Post Authorisation Assessments

### Oxytobel 10 IU/ml Solution for Injection for Horses, Cattle, Pigs, Sheep, Goats, Dogs and Cats Vm 41816/4000

02 April 2026	Changes to the quality part of the dossier: Deletion of a manufacturing site for an active substance. (GB + NI).
19 January 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
16 June 2023	Submission of a new or updated Ph. Eur. certificate of suitability or deletion of Ph. Eur. certificate of suitability.
06 October 2022	Change in the specification parameters and/or limits of the finished product: - Change outside the approved specifications limits range.
07 July 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
16 March 2021	Minor changes to an approved test procedure of the finished product.
11 July 2019	Change in distributor details.
31 January 2019	Renewal – UK as CMS
23 February 2017	Change in test procedure for the finished product. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
25 June 2015	Change of distributor details. Changes to the labelling and product literature.
06 March 2015	Submission of updated Ph. Eur. Certificates of Suitability.
21 October 2014	Repeat Use Comms.
06 August 2014	Change in the invented name of the veterinary medicinal product from 'Oxytocin 10 bela-pharm IU/ml Solution for Injection for Horses, Cattle, Pigs, Sheep, Goats, Dogs and Cats' to 'Oxytobel 10 IU/ml solution for injection for Horses, Cattle, Pigs, Sheep, Goats, Dogs and Cats'.