



## Post Authorisation Assessments

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### Myorelax 100 mg/ml Solution for Infusion for Horses

Vm 16849/5010

•	13 October 2023	Change in the manufacturing process of the finished product. Change in the manufacturing process of the finished product. Submission of an updated Ph. Eur. certificate of suitability for an active substance. Submission of an updated Ph. Eur. certificate of suitability for an active substance.
•	02 October 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	24 January 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	05 December 2017	Submission of an updated certificate of suitability.
•	30 August 2017	Renewal – UK as CMS.
•	21 November 2016	Submission of an updated Ph. Eur. certificate of suitability for an active substance. Deletion of Ph. Eur. certificate of suitability for an active substance.
•	08 March 2013	Mock-ups reviewed and approved.
•	06 march 2013	Change of QPPV and change of contact details for QPPV for an existing pharmacovigilance system..