



Post Authorisation Assessments

Vetivex 9 (Ringers Solution for Injection) Vm 10434/4054

•	12 February 2019	Changes to an existing pharmacovigilance system as described in the DDPS.
•	22 October 2015	Addition of a manufacturer responsible for batch release. Addition of a secondary packaging site. Changes in test procedure for the finished product. Minor changes in the manufacturing process. Addition of a manufacturing site of the finished product.
•	14 August 2015	Deletion of non-significant specification parameters and/or limits of the finished product.
•	13 May 2014	Change of active substance manufacturing site.
•	08 August 2012	Variation to change the composition of the primary packaging.
•	24 April 2012	Variation to change the name of a manufacturer.
•	19 January 2011	Variation to change the distributor.
•	22 December 2010	Variation to change the name of a manufacturer.
•	30 September 2008	Variation to change the Marketing Authorisation Holder.
•	23 April 2008	Renewal.
•	20 December 2006	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005. Transfer of the legal category.
•	07 September 2005	Variation to change the active substance manufacturer.
•	05 August 2005	Variation to change the name and address of the Marketing Authorisation Holder.
•	13 January 2005	Renewal.
•	22 December 2004	Change of the name of a manufacturer/assembler of the finished product.
•	22 November 2004	Addition of a distributor.
•	28 November 2003	Variation concerning a change to the active substance manufacturer.
•	12 July 2003	Change in a finished product test procedure.