Veterinary Medicines Directorate

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Post Authorisation Assessments

Vetivex 3 (Sodium Chloride 0.9 % w/v and Glucose 5 % w/v Intravenous Infusion BP)
Vm 10434/4055

•	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.
•	17 September 2015	Change in the address of the MAH.
•	08 September 2015	Deletion of non-significant specification parameters of the finished product.
•	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.
•	25 September 2008	Variation to change the Marketing Authorisation Holder.
•	09 May 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.
•	01 August 2005	Variation to change the name and address of the Marketing Authorisation Holder.
•	22 December 2004	Change of the name of a manufacturer/assembler of the finished product.
•	30 November 2004	Renewal.
•	22 November 2004	Addition of a distributor.
•	28 November 2003	Addition of an active substance manufacturer.