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Post Authorisation Assessments

Norodine Bolus Tablet

Vm 02000/4079

•	30 July 2019	Change in the QPPV of an existing pharmacovigilance system as described in the DDPS. Change of the back-up procedure of the QPPV of an existing pharmacovigilance system as described in the DDPS.
•	16 July 2013	Variation to update Ph. Eur. Certificates of Suitability from already approved manufacturers.
•	20 June 2012	Variation to change a Distributor address.
•	16 December 2008	Variation to bring the SPC/Labelling in line with the Veterinary Regulations, 2005.
•	07 October 2008	Deletion of an Active Substance Manufacturer.
•	08 May 2008	Variation to update a Ph. Eur. Certificate of Suitability.
•	16 January 2008	Addition of an Active Substance manufacturer.
•	16 January 2008	Addition of an Active Substance manufacturer.
•	20 February 2007	Transfer of legal category from POM to POM-V.
•	24 May 2006	Renewal.
•	23 November 2005	Addition of an Assembler.
•	28 November 2003	Renewal.
•	17 July 2003	Variation to change a withdrawal period.
•	27 September 2000	Addition of an Active Substance Manufacturer.
•	19 October 1999	Addition of an Active Substance Manufacturer.
•	12 May 1998	Renewal.
•	23 April 1998	Variation concerning and alternative Active Substance Manufacturer.
•	10 October 1996	Addition of an Active Substance Manufacturer.