Veterinary Medicines Directorate

Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS
Telephone +44 (0)1932 336911

Post Authorisation Assessments

Vasotop 5 mg Tablets

Vm 01708/4400

•	09 September 2022	Submission of a new certificate of suitability for an already approved manufacturer.
•	11 February 2022	Change in storage conditions of the finished product from Do not store above 30°C. to Do not store above 25°C. Change in the specification parameters and/or limits of the finished product.
•	09 June 2021	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
•	23 August 2018	Reduction of the shelf life of the finished product as packaged for sale from 24 months to 18 months.
•	04 January 2013	Changes to the SPC and product literature following deletion of presentations.
•	17 March 2010	Changes to the SPC and product literature in order to maintain harmonisation across the product range.
•	08 July 2009	Change to shelf-life specifications.
•	20 June 2008	Renewal.
•	28 December 2006	Changes to the SPC and product literature to bring them into line with new legislation.
•	09 August 2006	Change in the qualitative and/or quantitative composition of the immediate packaging.
•	09 August 2006	Addition of a flavouring component and consequential replacement of an excipient.
•	09 August 2006	Change in the name of the product.
•	23 June 2006	Change in test procedure of an excipient.
•	25 May 2006	Change to the test procedure of the finished product.
•	21 July 2005	Changes to the package leaflet.
•	23 June 2005	Change of distributor.
•	17 March 2005	Change to design of packaging.
•	29 September 2004	Renewal.
•	11 April 2003	Additional manufacturer/assembler of dosage form.
•	03 July 2001	Addition of a distributor in Northern Ireland.
•	17 April 2000	Change in dosage schedule for standard dose.
•	18 February 2000	Change in name and address of MAH including name of manufacturer of the active substance and assembler of dosage form.
•	30 June 1999	Change in assembler of dosage form.
•	30 June 1999	Change in manufacturer of dosage form.