Veterinary Medicines Directorate

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

Post Authorisation Assessments

Vasotop 1.25 mg Tablets

Vm 01708/4403

•	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 SPC and product literature due to cancellation of presentations.
•	17 March 2010	Changes to the SPC and product literature 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 material.
•	09 August 2006	Modification to the formulation.
•	09 August 2006	Change to the product name.
•	23 June 2006	Change to test procedure of an excipient.
•	25 May 2006	Change in the test procedure of the finished product.
•	20 July 2005	Amendments to the package leaflet.
•	24 June 2005	Change if distributor.
•	17 March 2005	Update of packaging design.
•	29 September 2004	Renewal.
•	11 April 2003	Additional manufacturer/assembler of dosage form.
•	22 June 2001	Additional distributor in Northern Ireland.
•	17 April 2000	Change in dosage schedule for standard dose.
•	29 February 2000	Change in anme and address of MAH including name change of manufacturer of active ingredientand assembler of dosage form.
•	30 June 1999	Change to site of manufacturer of dosage form (assembly).
•	30 June 1999	Change to site of manufacturer of dosage form.