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

Post Authorisation Assessments

Vetergesic Multidose, 0.3 mg/ml, Solution for Injection for Dogs and Cats

Vm 15052/5055

•	18 October 2022	Change in the address of the MAH from Unit 3 Anglo Office
		Park, White Lion Road, Amersham, Buckinghamshire, HP7
		9FB to Explorer House, Mercury Park, Wycombe Lane,
		Wooburn Green, High Wycombe, Buckinghamshire,
	20. A	HP10 0HH, United Kingdom.
•	26 August 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved
		manufacturer.
	13 February 2019	Deletion of manufacturing site for an active substance
	101 obradily 2010	Deletion of manufacturing site for finished product and
		packaging site
		Submission of an updated Ph. Eur. certificate of suitability
		for an active substance from an already approved
		manufacturer
•	07 August 2018	Increase in the shelf-life of the finished product as packaged
	00 Falamiam, 2010	for sale, from 2 years to 3 years.
•	06 February 2018	Change in storage conditions of the finished product.
•	19 September 2017	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.
•	22 June 2015	Change in legal entity.
•	09 April 2015	Introduction of a new pharmacovigilance system.
•	07 April 2015	Change of distributor and mock-up approval.
•	05 December 2014	Change in MAH name from Alstoe Limited to Sogeval UK Limited.
•	09 January 2014	Addition of a manufacturer for the finished product, including
		secondary packaging and batch release.
•	09 January 2014	Change in a specification parameter of an active substance.
	00 Novembre - 0040	Submission of a Ph. Eur. Certificate of Suitability.
•	28 November 2013	Renewal procedure.
•	24 July 2012 01 February 2012	Change to legal entity. Change in the specification parameters/ and or limits of the
•	UT Febluary 2012	Change in the specification parameters/ and or limits of the finished product.
•	04 August 2010	Change in any part of the (primary) packaging material not
	017 (agaot 2010	in contact with the finished formulation.
•	04 August 2010	Change in the batch size (including batch size ranges) of
		the finished product.
•	04 August 2010	Change to batch release arrangements and quality control

		testing of the final product.
•	04 August 2010	Replacement or addition of a manufacturing site.
•	04 August 2010	Changes in the specification parameters and /or limits of the
	-	finished product.