Veterinary Medicines Directorate

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 14966/3028

01 June 2026	Addition of a new specification parameter to the immediate packaging of the finished product. Change to comply with Ph. Eur. Change to comply with Ph. Eur. Deletion of a non-significant specification parameter of the finished product.
12 May 2026	Change in the pharmacovigilance system master file (PSMF) location. Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
12 May 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance. Change in the pharmacovigilance system master file (PSMF) location.
26 September 2025	Change of legal entity of the Marketing Authorisation Holder from Ceva Animal Health Ltd, Explorer House, Mercury Park, Wycombe Lane, Wooburn Green, High Wycombe, Buckinghamshire, HP10 0HH United Kingdom to Ceva Sante Animale, 8 rue de Logrono, 33500 Libourne, France.
17 January 2025	One-off alignment of the product information with version 9.0* of the QRD templates
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, 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 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 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. Submission of a Ph. Eur. Certificate of Suitability.
28 November 2013	Renewal procedure.
24 July 2012	Change to legal entity.
01 February 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 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.