Veterinary Medicines Directorate

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

Post Authorisation Assessments

Zitac Vet 200 mg Tablets for Dogs

Vm 06376/5045

09 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
12 January 2026	Update to the information on batch control and quality testing sites for the finished product.
22 April 2025	Change of Marketing Authorisation Holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands.
03 March 2025	Alignment of the product information with version 9.0* of the QRD templates.
14 December 2024	Deletion of a non-significant specification parameter for an active substance. (NI)
01 December 2024	Addition of new specification parameters for the immediate packaging. Tightening of immediate packaging specification limits. Deletion of non-significant specification parameters for the immediate packaging. (NI)
01 December 2024	Addition of new specification parameters for the immediate packaging. Tightening of immediate packaging specification limits. Deletion of non-significant specification parameters for the immediate packaging. (GB)
26 November 2024	Deletion of non-significant specification parameter for active substance. (GB).
10 February 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
17 July 2020	Change of MAH from Intervet International BV represented by: Intervet UK Ltd., Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ to MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, Buckinghamshire, MK7 7AJ.
22 May 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance.
13 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS
01 December 2016	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
28 November 2014	Update to the DDPS.
15 January 2014	Deletion of a manufacturing site.
24 October 2012	Changes to the DDPS following assessment.
23 June 2011	Renewal Marketing Authorisation.

11 March 2009	Change to in-process test/limits during manufacture.