Veterinary Medicines Directorate

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Post Authorisation Assessments

Zodon 88 mg Chewable Tablets for Dogs

Vm 15052/4128

•	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.
•	03 February 2022	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	06 October 2021	Change in the SPC, labelling or package leaflet due to new data.
•	26 July 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	08 July 2019	Renewal – UK CMS
•	23 May 2019	Replacement of a site where batch control/testing takes place
•	06 July 2018	Change in RMS from the UK to FR.
•	28 June 2018	Deletion of a manufacturer responsible for batch release
•	14 May 2018	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	25 March 2018	Change in product name in DK, FI, NO, LU and NL only.
•	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.
•	19 September 2017	Change in the name and/or address of the MAH in Spain only.
•	22 August 2017	Deletion of manufacturing site for an active substance Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer
•	20 October 2016	Change in shelf-life of the veterinary medicinal product as packaged for sale from 21 months to 3 years.
•	08 September 2016	Change in the name of a manufacturer of the finished product, also responsible for batch release. Change in the name of a manufacturer of the finished

		product, also responsible for batch release. Change in the name of the manufacturer of the finished product.
•	06 September 2016	Change in the name an address of the MAH in Italy only.
•	29 June 2016	Introduction of a new pharmacovigilance system which has been assessed by the relevant national competent authority/EMA for another product of the same MAH.
•	16 June 2016	Change of MAH, from Sogeval to Ceva Animal Health Ltd.
•	20 November 2014	Introduction of joint labelling with Ireland.