



## Post Authorisation Assessments

### Leventia 1 mg/ml Oral Solution for Dogs Vm 06376/3058

14 March 2026	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
02 September 2025	Change of legal entity of the 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.
24 July 2025	One-off alignment of the product information with version 9.0* of the QRD templates.
14 December 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI)
14 December 2024	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB)
26 April 2024	Change in the dimensions of the immediate packaging for the finished product. (NI)
25 January 2024	Alternative primary packaging container registered for the finished product. (GB)
25 August 2021	Addition of a site where batch control/testing takes place.
27 April 2021	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
14 August 2020	Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited.
13 November 2018	Change in the safety database of an existing pharmacovigilance system as described in the DDPS
16 January 2018	Change in the specification limits of the finished product.
01 December 2016	Change in the safety database of an existing pharmacovigilance system as described in the DDPS.
27 April 2016	Submission of an updated Ph. Eur. certificate of suitability.
02 April 2014	Reduction in the shelf-life of the finished product, from 3 years to 2 years.
27 November 2014	Update of the pharmacovigilance system as described in the DDPS.
06 October 2014	Change in specification limit of the finished product.
22 October 2012	Changes to the DDPS following assessment of the same DDPS in relation to another medicinal product of the same MAH.
13 June 2012	Renewal procedure – Ireland as RMS.

18 March 2009	Change shelf life of finished product (as packaged for sale)
19 November 2008	Simple dosage instruction changes
22 May 2008	Change shelf life of finished product (after first opening)
22 May 2008	Formulation
04 July 2007	Deletion of any manufacturing site