Veterinary Medicines Directorate

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Post Authorisation Assessments

Easimax Plus Tablets for Dogs

Vm 08749/3004

24 February 2026	Updates to adverse events section accepted and update to the contact details.
19 January 2025	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
16 January 2025	Submission of an updated Ph. Eur. CEP from an already approved manufacturer. Submission of an updated Ph. Eur. CEP from an already approved manufacturer. Submission of an updated Ph. Eur. CEP from an already approved manufacturer.
01 September 2023	SRP to add 3 new CMS.
05 July 2023	Deletion of a manufacturing site of the active substance.
20 June 2023	One-off alignment of the product information with version 9.0* of the QRD templates i.e., major update of the QRD templates in accordance with Regulation (EU) 2019/6, for veterinary medicinal products placed on the market in accordance with Directive 2001/82/EC or Regulation (EC) No 726/2004.
12 May 2023	Substantial changes in the updated version of the ASMF or the active substance part of the dossier.
20 April 2022	Update to ASMF.
22 April 2021	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. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
07 October 2019	Increase in the shelf-life of the finished product as packaged for sale, from 3 years to 5 years.
15 July 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
10 January 2019	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
02 July 2018	ASMF updated.
22 December 2017	Addition of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
26 October 2016	Change in the (invented) name of the veterinary medicinal product in Hungary.
15 May 2015	Submission of a new certificate of suitability.
18 July 2014	Renewal procedure – Ireland as RMS.
• 10 June 2013	To add the additional active substance manufacturer.

01 February 2013	Submission of an updated Ph. Eur. Certificate of Suitability for an already approved manufacturer.
02 May 2012	Submission of a new or updated Ph. Eur. Certificate of Suitability.
02 May 2012	Submission of a new or updated Ph. Eur. Certificate of Suitability.
22 December 2010	To submit packaging for approval in the UK.
02 November 2010	Change of name of the Veterinary Medicinal Product from Excitel Plus Tablets for Dogs to Easimax Plus Tablets for Dogs.
02 July 2010	Variation to add a pork flavour to the tablets.