Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Easimax Plus Tablets for Dogs

Vm 08749/5040

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| 12 December 2024 | Submission of an updated Ph. Eur. CEP from an already approved manufacturer.<br>Submission of an updated Ph. Eur. CEP from an already approved manufacturer.<br>Submission of an updated Ph. Eur. CEP from an already approved manufacturer. |
| 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. |
| 28 March 2023 | Deletion of an active substance manufacturer. |
| 16 January 2023 | Additional manufacturing site for the active substance pryantel embonate. |
| 23 September 2022 | Updated certificate of suitability from an already approved manufacturer. |
| 20 April 2022 | Update to ASMF. |
| 22 April 2021 | Deletion of manufacturing site for an active substance.<br>Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.<br>Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.<br>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. |

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| 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. |