

Post Authorisation Assessments

Anthelmin Plus Flavour Tablets for Dogs

Vm 01656/4015

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| 04 March 2026 | Correction of the target species weight on product literature. |
| 17 February 2025 | Alignment of the product information with version 9.0* of the QRD templates. |
| 15 January 2025 | Change in the manufacturers of the active substance where no Ph. Eur. CEP is part of the approved dossier. Minor changes to an approved test procedure for the active substance. (GB and NI). |
| 15 January 2025 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI) |
| 01 December 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB). |
| 23 February 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB) |
| 23 February 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI) |
| 19 January 2024 | Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. (NI) |
| January 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB) |
| 04 December 2023 | Introduction of a summary of the PSMF. (GB) |
| 05 October 2023 | Substantial changes in the updated version of the ASMF. |
| 07 June 2023 | Submission of a new certificate of suitability. |
| 16 March 2023 | New certificate of suitability from a new manufacturer. |
| 06 January 2023 | Updated certificate of suitability from an already approved manufacturer. |
| 11 October 2022 | Updated certificate of suitability from an already approved manufacturer. |
| 07 April 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. |
| 13 April 2021 | Minor changes to an approved test procedure of the finished product. |
| 16 March 2021 | Changes to the labelling and/or package leaflet. |
| 26 January 2021 | 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. |
| 25 November 2020 | Submission of a new Ph. Eur. certificate of suitability for an |

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| | active substance from a new manufacturer. |
| 14 August 2020 | Changes to the labelling. |
| 09 July 2020 | Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Minor change in the manufacturing process of the finished product. Addition of a manufacturer responsible for batch release of the finished product. Change in the shape or dimensions of the pharmaceutical form. Addition of a manufacturing site of the finished product. |
| 16 August 2019 | Addition of a site where batch control/testing takes place. |
| 27 June 2019 | 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. |
| 27 June 2019 | Addition of a site where batch control/testing takes place. Deletion of manufacturing site for finished product. Replacement of a secondary packaging site of the finished product. Replacement of a primary packaging site of the finished product. |
| 11 June 2019 | Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS. |
| 26 October 2018 | Change in RMS from UK to IE. |
| 23 October 2018 | Update to the Local Representative details. |
| 25 July 2017 | Change to comply with an update of the relevant monograph of the Ph. Eur. or national pharmacopoeia of a Member State. |
| 12 April 2017 | Deletion of manufacturing site for an active substance |
| 01 December 2016 | Mock-ups approved. |
| 07 October 2016 | Addition of a new test method. Addition of an active substance manufacturer. |
| 05 September 2016 | Change in the (invented) name of the medicinal product in Spain and Portugal. |
| 05 August 2016 | Increase in the shelf-life of the finished product as packaged for sale, from 2 to 3 years. |
| 02 June 2016 | Renewal – UK as RMS |
| 25 November 2015 | To update the administration advice in section 4.9 of the SPC |
| 22 April 2015 | Addition of UK local representative information to package leaflet. |
| 23 March 2015 | Change in distributor details. |
| 09 May 2014 | Receipt of an updated Certificate of Suitability for an active substance. |
| 19 December 2013 | To update the pharmacovigilance system. |
| 19 December 2013 | Deletion of a manufacturing site for primary and secondary packaging. |

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| 01 November 2013 | Submission of an updated Ph. Eur. certificate of suitability. |
| 21 March 2013 | Changes to the labelling or the package leaflet which are not connected with summary of product characteristics |
| 01 March 2013 | Deletion of a secondary packaging site and deletion of a manufacturing site responsible for batch release. |
| 12 October 2012 | Deletion of a manufacturing site for the active substance febantel; deletion of a manufacturing site for the active substance praziquantel; addition of a primary packaging site for the finished product; addition of a manufacturing site (not responsible for batch release or batch control) for the finished product. |
| 10 August 2012 | Change of name of veterinary medicinal product. |
| 06 January 2012 | Submission of a new or updated Ph. Eur. certificate of suitability. |