

Post Authorisation Assessments

Anthelmin 230 mg / 20 mg Film-coated Tablets for Cats

Vm 01656/3054

14 May 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
25 September 2025	Change in distributor details to KRKA UK Ltd, Thames House, Waterside Drive, Langley, SL3 6EZ, United Kingdom.
24 January 2025	Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance.
20 November 2024	Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex.
30 July 2024	Editorial changes to SPC, package leaflet or labelling if inclusion in an upcoming procedure is not possible.
04 April 2024	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer. Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer.
20 October 2023	One-off alignment of the product information with version 9.0 of the QRD templates.
06 June 2023	Submission of an updated certificate of suitability.
24 May 2023	Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.
23 January 2023	Change in the name or address of a batch testing site.
09 January 2023	Updated certificate of suitability from an already approved manufacturer.
20 December 2022	Change in the name or address of a batch testing site.
01 April 2022	Renewal – UK as CMS
28 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.
26 March 2020	Changes to the labelling and/or package leaflet.
19 February 2020	Increase in the shelf-life of the finished product as packaged for sale, from 2 years to 3 years.
26 April 2019	Addition of a manufacturer responsible for batch release including batch control/testing. Addition of a primary packaging site of the finished product. Addition of a secondary packaging site of the finished product. Addition of a manufacturing site of the finished product.
02 April 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

08 March 2019	Change in the contact details of the QPPV of an existing pharmacovigilance system as described in the DDPS.
24 May 2018	Change in the invented name of the veterinary medicinal product from Anthelmin to Dehinel in FR only
26 October 2017	Change in contact details for local representative.