Veterinary Medicines Directorate

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Post Authorisation Assessments

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

Vm 01656/5054

•	22 February 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.
•	17 March 2023	Updated certificate of suitability from an already approved manufacturer
•	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.