



Post Authorisation Assessments

Cazitel 230/20 mg Flavoured Film-coated Tablets for Cats Vm 08749/5044

•	01 December 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.
•	22 June 2023	Updates to SPC text section 4.6 and leaflet text section 6.
•	22 June 2023	One-off alignment of the product information with version 9.0* of the QRD templates.
•	17 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.
•	06 May 2021	Change in distributor from Zoetis UK Limited, 5th Floor, 6 St. Andrew Street, London, EC4A 3AE to Zoetis UK Limited, 1st Floor, Birchwood Building, Springfield Drive, Leatherhead, Surrey, KT22 7LP.
•	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.
•	23 September 2020	Change in the invented name of the veterinary medicinal product from Cazitel 230/20 mg Flavoured Film-Coated Tablets for Cats to Vermicat 230/20 mg Flavoured Film-Coated Tablets for Cats in Romania only.
•	15 October 2019	Increase in the shelf-life of the finished product as packaged for sale, from 4 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.
•	16 October 2018	Renewal - UK as CMS
•	25 September 2018	Change in the invented name of the veterinary medicinal product from Cazitel 230/20 mg to Strantel kat 230/20 mg in NL only.
•	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.
•	15 May 2015	Submission of a new certificate of suitability