Veterinary Medicines Directorate

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Post Authorisation Assessments

Prazitel 230/20 mg Flavoured Film-coated Tablets for Cats

Vm 08749/3018

•	28 April 2024	Addition of a manufacturing site for the active substance.
•	06 March 2024	One-off alignment of the product information with version 9.0* of the QRD templates.
•	23 August 2023	Adverse Event added: hypersalivation.
•	16 February 2023	Updated certificate of suitability from an already approved manufacture.
•	27 January 2023	Addition of a manufacturing site for the active substance.
•	13 April 2022	Update to ASMF.
•	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.
•	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.
•	30 October 2018	Renewal – UK as CMS
•	02 July 2018	ASMF updated.
•	03 January 2018	Change in the invented name of the veterinary medicinal product from Prazitel 230/20mg flavoured film-coated tablets for cats to Exitel 230/20mg flavoured film-coated tablets for cats in Poland.
•	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.
•	17 October 2014	Change in the invented name of the medical product in the Czech Republic and Slovakia only, from ‘Prazitel 230/20 mg Flavoured Film-Coated Tablets for Cats’ to ‘Dehelmint 230/20 mg Film Coated tablets for cats’.