



## Post Authorisation Assessments

### ITCH WORMER for Cats & Kittens 230/20mg Flavoured Tablets Vm 08749/4068

•	11 May 2024	Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible.
•	04 May 2024	Change in pack size of the finished product outside the range of the currently approved pack sizes.
•	13 December 2022	Other changes to the active substance: - Substantial changes in the updated version of the ASMF or the active substance part of the dossier.
•	13 May 2022	Changes to the labelling and/or package leaflet.
•	12 May 2022	Deletion of Ph. Eur. certificates of suitability for an active substance. Submission of a new 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 an already approved manufacturer. Submission of a new Ph. Eur. certificate of suitability for an active substance.
•	18 June 2021	Renewal
•	04 July 2019	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.
•	01 May 2019	Change in the invented name of the veterinary medicinal product from EziWormer Cats & Kittens 230/20mg Flavoured Tablets to ITCH WORMER for Cats & Kittens 230/20mg Flavoured Tablets.
•	24 July 2018	Update to the active substance master file.
•	04 January 2018	Submission of a new Ph. Eur. certificate of suitability for an active substance.
•	04 September 2017	Change of legal distribution category from NFA-VPS to AVM-GSL and associated changes to the SPC and product literature.
•	09 August 2017	Change in the invented name of the veterinary medicinal product from Extrontel 230/20 mg Flavoured Film-coated Tablets for Cats to EziWormer Cats & Kittens 230/20mg Flavoured Tablets.

