



Post Authorisation Assessments

ITCH WORMER for Cats & Kittens 230/20mg Flavoured Tablets

•	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.