

Post Authorisation Assessments

Itch Wormer for Cats & Kittens 230/20mg Flavoured Tablets Vm 08749/4068

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| 22 December 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. |
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| 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. |

