

## Post Authorisation Assessments

### Curofen 50 mg/g Oral Powder for Pigs Vm 05150/3004

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| 17 February 2026 | Submission of a Ph. Eur. CEP for an active substance.   |
| 08 April 2025    | Submission of a new or updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile active substance.   |
| June 2024        | Introduction of a re-test period/storage period supported by real time data.  |
| 14 May 2024      | Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.  |
| 14 May 2024      | Submission of a new Ph. Eur. certificate of suitability for a manufacturer of the active substance.   |
| 20 February 2024 | Submission of a new Ph. Eur. certificate of suitability for a manufacturer of the active substance.   |
| 08 December 2023 | Submission of an updated Ph. Eur. CEP from an already approved manufacturer for a non-sterile: – active substance.  |
| 22 April 2021    | Deletion of manufacturing site for an active substance.<br>Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.<br>Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.<br>Update to the ASMF. |
| 18 March 2021    | Minor change in the manufacturing process of the finished product.<br>Increase in batch size (816kg to 816kg-1502kg) of the finished product.   |
| 09 March 2021    | Change in the invented name of the veterinary medicinal product from Fendoral 50 mg/g oral powder for pigs to Curazole 50 mg/g oral powder for pigs in Belgium.   |
| 27 October 2020  | Renewal – UK as CMS   |
| 07 February 2019 | Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.  |