



## Post Authorisation Assessments

### Strantel 230/20 mg Flavoured Film-coated Tablets for Cats Vm 08749/5053

•	06 March 2024	One-off alignment of the product information with version 9.0* of the QRD templates.
•	07 November 2023	Added SPC text section 4.6 and leaflet text section 6: In very rare cases, gastrointestinal disorders such as hypersalivation and /or vomiting & neurological signs such as ataxia and muscle tremors have been observed.
•	17 February 2023	Updated certificate of suitability from an already approved manufacturer.
•	27 January 2023	Addition of a manufacturing site for the active substance.
•	21 December 2022	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV).
•	13 April 2022	Update to ASMF.
•	25 March 2022	Change in the invented name of the veterinary medicinal product in IE only from Strantel 230/20 mg Flavoured Film-Coated Tablets for Cats to Wormaway 230/20 mg Flavoured Film-coated Tablets for Cats.
•	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.
•	16 October 2018	Renewal - UK as CMS
•	02 July 2018	ASMF updated.
•	22 December 2017	Addition of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	04 October 2016	Change in the name of the Product in the Netherlands only.
•	15 May 2015	Submission of a new certificate of suitability

