Veterinary Medicines Directorate

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Post Authorisation Assessments

Strantel Plus XL Tablets for Dogs

•	20 April 2022	Update to ASMF.
•	20 January 2022	Deletion of manufacturing site for an active substance.
•	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.
•	12 November 2019	Increase in the shelf-life of the finished product as packaged for sale, from 3 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
•	13 June 2019	Change in the invented name of the Italian veterinary medicinal product from Exitel Plus XL tablets for dogs to Panadron Plus XL compresse per cani
•	10 January 2019	Submission of a new Ph. Eur. certificate of suitability for an active substance from a new manufacturer.
•	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.
•	12 April 2017	Renewal – UK as CMS
•	15 May 2015	Submission of a new certificate of suitability.
•	06 November 2011	Change to the invented name of the veterinary medicinal product in the UK from Exitel Plus XL Tablets for Dogs to Strantel Plus XL Tablets for Dogs. Submission of two updated Certificates of suitability.
•	10 October 2013	Change in test procedure for the finished product.