

## Woodham Lane, New Haw, Addlestone, Surrey KT15 3LS Telephone +44 (0)1932 336911

## **Post Authorisation Assessments**

## Milaxyn 230/20 mg Flavoured Film-Coated Tablets for Cats Vm 40162/3000

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•	18 November 2023	SRP to add 9 CMS.
•	21 December 2022	Update to an approved Ph.Eur certificate of suitability for
		an active substance.
•	21 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.
•	20 December 2022	Submission of an updated Ph. Eur. CEP from an already
		approved manufacturer for a non-sterile: – active
		substance.
•	20 December 2022	Submission of a new Ph. Eur. CEP from a new
		manufacturer for a non-sterile: – active substance.
•	08 December 2022	One-off alignment of the product information with version
		9.0 of the QRD template.
•	11 March 2020	Renewal – UK as CMS
•	25 September 2019	Increase in the shelf-life of the finished product as
		packaged for sale, from 4 years to 5 years.
•	09 July 2019	Submission of an updated Ph. Eur. certificate of
		suitability for an active substance from an already
		approved manufacturer.
•	12 March 2019	Change in RMS from UK to ES.
•	12 July 2018	Update to the Active Substance Master File.
•	17 January 2018	Submission of a new Ph.Eur. Certificate of Suitability for
		an active substance from a new manufacturer.
•	19 August 2016	New mutual recognition procedure (MRP) – UK as RMS
•	07 January 2016	Deletion of manufacturing sites for an active substance.
•	12 August 2015	Submission of a new certificate of suitability from a new
		manufacturer.
•	08 December 2022  11 March 2020 25 September 2019  09 July 2019  12 March 2019  12 July 2018  17 January 2018  19 August 2016  07 January 2016	manufacturer for a non-sterile: – active substance.  One-off alignment of the product information with version 9.0 of the QRD template.  Renewal – UK as CMS  Increase in the shelf-life of the finished product as packaged for sale, from 4 years to 5 years.  Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer.  Change in RMS from UK to ES.  Update to the Active Substance Master File.  Submission of a new Ph.Eur. Certificate of Suitability for an active substance from a new manufacturer.  New mutual recognition procedure (MRP) – UK as RMS Deletion of manufacturing sites for an active substance.  Submission of a new certificate of suitability from a new