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

## **Post Authorisation Assessments**

## Moxiclear 40 mg + 4 mg Spot-on Solution for Small Cats and Ferrets Vm 02000/4435

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•	18 April 2024	Additional site of batch control for the finished product. (NI)
		Minor changes to an approved test procedure for the finished product. (NI)
	22 December 2023	Introduction of a summary of the PSMF or changes to
•	22 December 2023	the summary of the PSMF not already covered
		elsewhere in this Annex. (NI)
_	03 August 2023	Change to batch control arrangements for the finished
•	03 August 2023	1
		product. (GB)
		Minor change to an approved test procedure of the
	06 luna 0000	finished product.(GB)
•	26 June 2023	Submission of a new or updated Ph. Eur. certificate of
_	28 October 2022	suitability. (GB) Change in distributor details from Norbrook Laboratories
•	20 October 2022	
		(GB) Ltd, 1 Saxon Way East, Oakley Hay Industrial
		Estate, Corby, Northamptonshire, NN18 9EX, United Kingdom to Norbrook Laboratories Limited, Carnbane
		,
		Industrial Estate, Newry, Co Down, BT35 6QQ, Northern Ireland.
	29 July 2022	Unlimited renewal.
•	•	-
•	13 May 2021	Submission of an updated Ph. Eur. certificate of
		suitability for an active substance from an already approved manufacturer.
_	27 May 2020	
•	27 May 2020	Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already
		1
	25 September 2019	approved manufacturer.  Minor change in the manufacturing process of the
•	20 Gepterriber 2019	finished product.
•	22 August 2019	Addition of a manufacturer responsible for batch release
		of the finished product.
•	30 July 2019	Change in the QPPV of an existing pharmacovigilance
	-	system as described in the DDPS.
		Change of the back-up procedure of the QPPV of an
		existing pharmacovigilance system as described in the
		DDPS.
•	27 June 2019	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.
		approved manufacturer. Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already

22 August 2018	Minor changes to SPC/QRD
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