Veterinary Medicines Directorate

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# Post Authorisation Assessments

## Moxiclear 100 mg + 25 mg Spot-on Solution for Medium Dogs

Vm 02000/4438

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| 04 March 2026 | Submission of an updated Ph. Eur. certificate of suitability for an active substance. (GB + NI). |
| 11 February 2026 | Submission of a Ph. Eur. CEP for an active substance. (NI). |
| 15 December 2025 | Submission of a Ph. Eur. CEP for an active substance. (GB). |
| 19 June 2025 | Alignment of the product information with version 9.0* of the EU QRD templates. |
| 18 April 2024 | Additional site of batch control for the finished product. (NI)<br>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 the summary of the PSMF not already covered elsewhere in this Annex. (NI) |
| 03 August 2023 | Change to batch control arrangements for the finished product. (GB)<br>Minor change to an approved test procedure of the finished product. (GB) |
| 26 June 2023 | Submission of a new or updated Ph. Eur. certificate of suitability. (GB) |
| 28 October 2022 | Change in distributor details from Norbrook Laboratories (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 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 25 September 2019 | Minor change in the manufacturing process of the 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.<br>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.<br>Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 22 August 2018 | Minor changes to SPC/QRD |