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# Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021, please refer to the EMA website.

## Bravecto 112.5 mg Spot-on Solution for Very Small Dogs (2-4.5 kg)

Vm 01708/5018

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|---|---|
| 08 March 2026 | Introduction of a summary of the PSMF or changes to the summary of the PSMF not already covered elsewhere in this Annex. |
| 08 August 2025 | Change to in-process tests or limits applied during the manufacture of the finished product.<br>Minor change in the manufacturing process of the finished product. |
| 07 August 2025 | Terms removed from adverse events table as outcome of signal management. |
| 23 July 2025 | Deletion of a non-significant specification parameter of an active substance. |
| 10 December 2024 | Editorial changes to part 2 of the dossier if inclusion in an upcoming procedure concerning part 2 is not possible. |
| 24 January 2024 | Alignment of product information with the national SPC/QRD template. |
| 09 June 2023 | Unlimited renewal |
| 22 June 2022 | Change in name of manufacturer of the active substance. |
| 08 April 2022 | Changes to the labelling and/or package leaflet. |
| 30 July 2021 | Addition of a pair of gloves per pipette in the carton box. |
| 06 May 2021 | Addition of a primary packaging site.<br>Addition of a secondary packaging site.<br>Change in the manufacturing process of the finished product.<br>Change in the manufacturing process of the finished product.<br>Addition of a bulk manufacturing site. |
| 04 March 2021 | Deletion of site where batch control takes place |