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# Post Authorisation Assessments

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

## Apoquel 5.4 mg Chewable Tablets for Dogs

Vm 42058/5001

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| 1 July 2026 | Minor changes to an approved test procedure for the finished product. |
| 13 May 2026 | Clarifications to dosing table regarding animal weights and numbers of tablets. |
| 16 February 2026 | Change in the test procedure for an excipient. |
| 22 December 2025 | Change to batch control arrangements and quality testing for a finished product. |
| 22 December 2025 | Removal of references to the local representative. |
| 04 September 2025 | Minor change in the manufacturing process of the finished product. Change in the holding time of the bulk product. |
| 04 July 2025 | Change to comply with Ph. Eur. |
| 12 March 2024 | Change in the specification parameters and/or limits of the finished product: - Change outside the approved specifications limits range. |
| 12 March 2024 | List of EU Local Representatives deleted from GB QRD documents. |
| 15 September 2023 | Change in the shelf-life or storage conditions of the finished product: - Extension of the shelf life of the finished product - As packaged for sale. |
| 25 August 2023 | Added to Safety Sections the adverse event ' Convulsion' with the frequency of Very rare: <1 animal / 10,000 animals treated, including isolated reports. |
| 15 June 2023 | Minor changes to an approved test procedure for active substance. |
| 14 June 2023 | Change in test procedure for an excipient: - Other changes to a test procedure (including replacement or addition). |
| 01 June 2023 | One-off alignment of the product information with version 9.0* of the QRD template. |
| 23 December 2022 | Minor changes to an approved test procedure, for a starting material, reagent or intermediate used in the manufacturing process of the active substance. |
| 23 December 2022 | Minor changes to an approved test procedure for active substance. |
| 01 November 2022 | Change in name and address details of a manufacturer of the active substance. |
| 28 June 2022 | Change in the manufacturer of a starting material/reagent/intermediate used in the manufacturing process of the active substance or change in the manufacture. |
| 20 May 2022 | Change in the number of units (blister) in a pack within the range of the currently approved pack sizes of the finished product. |
| 18 May 2022 | Change in test procedure for an excipient. |