



Post Authorisation Assessments

For details of post authorisation assessments prior to 1st January 2021,
please refer to the [EMA](#) website.

Credelio Plus 225 mg/8.44 mg Chewable Tablets for Dogs (>5.5 kg - 11 kg) Vm 52127/5035

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| 18 December 2025 | Deletion of a non-significant specification parameter of an active substance. |
| 18 December 2025 | Deletion of an active substance manufacturing site where batch control takes place. Deletion of an active substance manufacturing site where batch control takes place. |
| 28 October 2025 | Changes to the manufacturing process of the active substance. Changes to the specification parameters of an intermediate during the manufacturing process of the active substance. |
| 08 October 2025 | Minor change to an approved test procedure for an active substance. |
| 11 June 2025 | Change(s) in the name or address or contact details of a qualified person for pharmacovigilance (QPPV). |
| 07 May 2025 | Minor changes to an approved test procedure for the finished product. |
| 04 April 2025 | Change in the manufacturer of an intermediate used in the manufacturing process of the active substance. |
| 04 April 2025 | Addition of a new specification parameter to the specification with its corresponding test method. Change in the specification parameters or limits of reagent used in the manufacturing process of the active substance. Change in the specification parameters of reagents used in the manufacturing process of the active substance. Minor changes:– in the manufacturing process of an active substance. |
| 21 January 2025 | Change in the name or address or contact details of: a manufacturer of starting material. Changes to the quality part of the dossier: Deletion of a supplier of a starting material for an active substance. |
| 30 July 2024 | One-off alignment of the product information with version 9.0* of the QRD template. |
| 23 February 2023 | Addition of a new indication: For the treatment of demodicosis. |
| 31 May 2022 | Change(s) in the manufacturing process of the active substance. |
| 19 November 2021 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 21 October 2021 | Extension of the shelf-life of an excipient. |
| 28 June 2021 | Addition of a manufacturer of a starting material. Changes in the specification parameters of a starting material. Change in the manufacturer of a starting material. Change in the manufacturer of a starting material used in the manufacturing process of the active where no Ph. Eur. Certificate of Suitability is part of the approved dossier. Addition of a manufacturer of a starting material. Changes in the specification parameters of a starting material. Change in the manufacturer of a starting material. |

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