Veterinary Medicines Directorate
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Post Authorisation Assessments

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

Credelio Plus 450 mg/16.88 mg Chewable Tablets for Dogs (>11 kg- 22 kg)

Vm 52127/5036

22 October 2026	Minor changes in the manufacturing process of an active substance.
23 July 2026	Addition of a site where batch control or testing of the active substance takes place. Minor changes to an approved test procedure for the active substance.
16 July 2026	Introduction of a new manufacturer of the active substance that is not supported by an ASMF and requires significant update to the relevant active substance section of the dossier. Change in the manufacturer (including where relevant quality control testing sites) of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier: - Other changes.
25 March 2026	One-off alignment of the product information with version 3 of the National QRD templates. Addition of indication for 'treatment of Sarcoptic mange'. Addition of indication 'reduction of the risk of infection with Babesia canis canis via transmission by Dermacentor reticulatus for one month'.
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	Minor changes to an approved test procedure for an active substance.
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. 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. 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 name of a manufacturer of a starting material used in the manufacturing process of the active substance. Change in the address of a manufacturer of a starting material used in the manufacturing process of the active substance. Minor change in the manufacturing process of the active substance. Minor change in the manufacturing process of the active substance. Deletion of manufacturing site for an active substance manufacturer responsible for the supply of a starting material.