Veterinary Medicines Directorate

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Post Authorisation Assessments

Selehold 240 mg Spot-on Solution for Dogs 20.1–40.0 kg

Vm 01656/5061

25 August 2026	Change in the specification parameters and/or limits of an excipient.
13 April 2026	Change(s) in the name or address or contact details of a qualified person for pharmacovigilance.
07 July 2025	Change in the address of a manufacturer of the finished product.
24 March 2025	Change in the manufacturer of a starting material used in the manufacturing process of the active substance, where no Ph. Eur. Certificate of Suitability is part of the approved dossier:- Other changes.
14 January 2025	Deletion of a manufacturing site for an active substance.
03 May 2024	Unlimited renewal
13 February 2024	Alignment with version 9 of the QRD templates.
21 March 2023	Minor changes to an approved test procedure for the finished product.
16 January 2023	Changes to comply with the Ph. Eur. monograph of the active substance - Removal of references to internal test methods and test method numbers. Change in the address of the manufacturer of the starting material.
01 September 2022	Extension of the re-test period of the active substance.
22 March 2022	Minor changes to an approved test procedure of the finished product. Minor change of an analytical procedure for an in-process control applied during the manufacture of the finished product.
27 August 2021	Increase in the shelf-life of the finished product as packaged for sale, from 2 to 3 years.
01 August 2019	Changes to the labelling.