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.

NexGard Combo Spot-On Solution for Cats <2.5 kg

Vm 04491/5000

•	14 February 2024	Addition of a site of batch control for the finished product.
•	01 June 2023	Change in the shelf-life or storage conditions of the finished product: - Change in storage conditions of the finished product or the diluted/reconstituted product.
•	21 April 2023	Change in the name or address or contact details of a qualified person for pharmacovigilance.
•	07 November 2022	Addition of a new in-process test applied during manufacture of the active substance.
•	22 July 2022	Deletion of a test procedure for the active substance.
•	13 July 2022	Minor changes to an approved test procedure for the active substance.
•	28 April 2022	Increase in batch size of the active substance used in the manufacturing process of the active substance.
•	15 February 2022	Change in the name of a manufacturer used in the manufacture of the active substance.
•	06 January 2022	Change in the name/address of a manufacturer used in the manufacture of the active substance.
•	22 December 2021	Change in the SPC, labelling or package leaflet due to new data. Addition of a new therapeutic indication. Addition of a new therapeutic indication.
•	30 September 2021	Change to an approved stability protocol.
•	06 July 2021	Extension of a re-test period of the active substance.