16 May 2025
EMA/CVMP/82901/2025
Committee for Veterinary Medicinal Products (CVMP)

Summary of opinion[1] (post-authorisation)

NexGard Combo

International non-proprietary name (INN): esafoxolaner / eprinomectin / praziquantel

On 15 May 2025, the Committee for Veterinary Medicinal Products (CVMP) adopted a positive opinion[2], recommending the granting of a group of variations to the terms of the marketing authorisation for the veterinary medicinal product NexGard Combo. The marketing authorisation holder for this veterinary medicinal product is Boehringer Ingelheim Vetmedica GmbH.

NexGard Combo is currently authorised as spot-on solution for use in cats. The grouped variation concerns change(s) to therapeutic indication(s) - addition of a new therapeutic indication or modification of an approved one: treatment of infections with eye worms (*Thelazia callipaeda*) and immediate tick killing activity against *Ixodes hexagonus*.

Detailed conditions for the use of this product are described in the summary of product characteristics (SPC), for which an updated version reflecting the changes will be published in the Union Product Database (UPD) and will be available in all official European Union languages after the variation to the marketing authorisation has been granted by the European Commission.

[1] Summaries of opinion are published without prejudice to the Commission Decision.
[2] Applicants may appeal any CVMP opinion, provided they notify the European Medicines Agency in writing of their intention to appeal within 15 days of receipt of the opinion.

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