12 September 2025
EMADOC-1700519818-2407766
Committee for Veterinary Medicinal Products (CVMP)

Summary of opinion[1] (post-authorisation)

Bravecto TriUNO

International non-proprietary name (INN): Fluralaner / Moxidectin / Pyrantel

On 10 September 2025, the Committee for Veterinary Medicinal Products (CVMP) adopted a positive opinion[2], recommending the granting of a variation to the terms of the marketing authorisation for the veterinary medicinal product Bravecto TriUNO. The marketing authorisation holder for this veterinary medicinal product is Intervet International B.V.

Bravecto TriUNO is currently authorised as chewable tablets for use in dogs. The variation concerns change(s) to therapeutic indication(s) - addition of a new therapeutic indication or modification of an approved one: treatment of infections with *Angiostrongylus vasorum*. Additionally, the product information has been aligned with version 9.1 of the QRD template.

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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