

# Bravecto 150 mg/ml - Powder and solvent for suspension for injection

Authorised

- Fluralaner

## Product identification

**Medicine name:**

Bravecto 150 mg/ml - Powder and solvent for suspension for injection

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**Active substance:**

Fluralaner

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Fluralaner

150.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Powder and solvent for suspension for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP53BE02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Available in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Vial (glass), Package\_size:1 vial of powder + 1 vial of solvent, 1 vent needle

Packaging:Vial (glass), Package\_size:2 vials of powder + 2 vials of solvent, 2 vent needles

Packaging:Vial (glass), Package\_size:10 vials of powder + 10 vials of solvent, 10 vent needles

Packaging:Vial (glass), Package\_size:5 vials of powder + 5 vials of solvent, 5 vent needles

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

11/02/2014

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**Manufacturing sites for batch release:**

Intervet Ges.m.b.H.

Intervet Productions S.A.

Vet Pharma Friesoythe GmbH

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

25/01/2024

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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