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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

Active substance:
Fluralaner

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Fluralaner
150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53BE02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

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)

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)

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
11/02/2014

Manufacturing sites for batch release:
Intervet Ges.m.b.H.
Intervet Productions S.A.
Vet Pharma Friesoythe GmbH

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/01/2024

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 6/07/2026
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