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Bravecto Plus 250 mg + 12.5 mg - Spot-on solution

Authorised

- Fluralaner
- Moxidectin

Product identification

Medicine name:
Bravecto Plus 250 mg + 12.5 mg - Spot-on solution

Active substance:
Fluralaner

Moxidectin

Target species:
Cat

Route of administration:
Spot-on use

Product details

Active substance and strength:
Fluralaner

Presentation_strength:250 mg Reference:Hse Comments:micronised or non-micronised Index:0

Moxidectin

Presentation_strength:12.5 mg Reference:Ph. Eur. Index:1

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB52

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 , Croatia , Cyprus , Czechia , Estonia , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Pipette (PP/Alu/PP), Package_size:1 pipette, Content:0.89 ml

Packaging:Pipette (PP/Alu/PP), Package_size:2 pipettes, Content:0.89 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:
8/05/2018

Manufacturing sites for batch release:
Intervet Productions S.A.

Responsible authority:
European Commission

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

Date of authorisation status change:
8/05/2018

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: 28/02/2024
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