

Prinovox 250 mg + 62.5 mg spot-on solution for large dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Prinovox 250 mg + 62.5 mg spot-on solution for large dogs

Prinovox 250 mg + 62,5 mg solução para unção punctiforme para cães grandes

Active substance:

Moxidectin

Imidacloprid

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Moxidectin

62.50 milligram(s) / 2.50 millilitre(s)

Imidacloprid
250.00 milligram(s) / 2.50 millilitre(s)

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:

Portugal

Package description:

(ID6) 52.5 millilitre(s): unspecified outer container with 21 Pipette (polypropylene) each with 2.5 millilitre(s)

(ID5) 15 millilitre(s): unspecified outer container with 6 Pipette (polypropylene) each with 2.5 millilitre(s)

(ID4) 10 millilitre(s): unspecified outer container with 4 Pipette (polypropylene) each with 2.5 millilitre(s)

(ID3) 7.5 millilitre(s): unspecified outer container with 3 Pipette (polypropylene) each with 2.5 millilitre(s)

(ID2) 5 millilitre(s): unspecified outer container with 2 Pipette (polypropylene) each with 2.5 millilitre(s)

(ID1) 2.5 millilitre(s): unspecified outer container with 1 Pipette (polypropylene) with 2.5 millilitre(s)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Elanco Animal Health GmbH

Marketing authorisation date:

27/02/2017

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1091/05/17RFVPT

Date of authorisation status change:

22/07/2025

Reference member state:

Germany

Procedure number:

DE/V/0196/005

Concerned member states:

Ireland Italy Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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