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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