

Prinovox 100 mg + 25 mg spot-on solution for medium dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Prinovox 100 mg + 25 mg spot-on solution for medium dogs

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Active substance:

Moxidectin

Imidacloprid

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Moxidectin

25.00 milligram(s) / 1.00 millilitre(s)

Imidacloprid
100.00 milligram(s) / 1.00 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:

Ireland

Available in:

Ireland

Package description:

(ID6) 21 millilitre(s): unspecified outer container with 21 Pipette (polypropylene) each with 1 millilitre(s)

(ID5) 6 millilitre(s): unspecified outer container with 6 Pipette (polypropylene) each with 1 millilitre(s)

(ID4) 4 millilitre(s): unspecified outer container with 4 Pipette (polypropylene) each with 1 millilitre(s)

(ID3) 3 millilitre(s): unspecified outer container with 3 Pipette (polypropylene) each with 1 millilitre(s)

(ID2) 2 millilitre(s): unspecified outer container with 2 Pipette (polypropylene) each with 1 millilitre(s)

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

Marketing authorisation date:

5/05/2017

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22020/057/002

Date of authorisation status change:

5/05/2017

Reference member state:

Germany

Procedure number:

DE/V/0196/004

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