

Prinovox 40 mg + 10 mg spot-on solution for small dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Prinovox 40 mg + 10 mg spot-on solution for small dogs

Active substance:

Moxidectin

Imidacloprid

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Moxidectin

10.00 milligram(s) / 0.40 millilitre(s)

Imidacloprid

40.00 milligram(s) / 0.40 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:

Italy

Package description:

(ID1) 0.4 millilitre(s): unspecified outer container with 1 Pipette (polypropylene) with 0.4 millilitre(s)

(ID2) 0.8 millilitre(s): unspecified outer container with 2 Pipette (polypropylene) each with 0.4 millilitre(s)

(ID3) 1.2 millilitre(s): unspecified outer container with 3 Pipette (polypropylene) each with 0.4 millilitre(s)

(ID4) 1.6 millilitre(s): unspecified outer container with 4 Pipette (polypropylene) each with 0.4 millilitre(s)

(ID5) 2.4 millilitre(s): unspecified outer container with 6 Pipette (polypropylene) each with 0.4 millilitre(s)

(ID6) 8.4 millilitre(s): unspecified outer container with 21 Pipette (polypropylene) each with 0.4 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 Italia S.p.A.

Marketing authorisation date:

22/08/2017

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Ministry Of Health

Authorisation number:

105081

Date of authorisation status change:

22/08/2017

Reference member state:

Germany

Procedure number:

DE/V/0196/003

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

English (PDF)

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