Prinovox 400 mg + 100 mg Spot-on Solution for Extra-Large Dogs

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:
Prinovox 400 mg + 100 mg spot-on solution for extra-large dogs
Prinovox 400 mg + 100 mg Spot-on Solution for Extra-Large Dogs

Active substance:
Moxidectin
Imidacloprid

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Moxidectin
100.00 milligram(s) / 4.00 millilitre(s)

Imidacloprid
400.00 milligram(s) / 4.00 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:
United Kingdom (Northern Ireland)

Package description:
(ID6) 84 millilitre(s): unspecified outer container with 21 Pipette (polypropylene) each with 4 millilitre(s)
(ID5) 24 millilitre(s): unspecified outer container with 6 Pipette (polypropylene) each with 4 millilitre(s)
(ID4) 16 millilitre(s): unspecified outer container with 4 Pipette (polypropylene) each with 4 millilitre(s)
(ID3) 12 millilitre(s): unspecified outer container with 3 Pipette (polypropylene) each with 4 millilitre(s)
(ID2) 8 millilitre(s): unspecified outer container with 2 Pipette (polypropylene) each with 4 millilitre(s)
(ID1) 4 millilitre(s): unspecified outer container with 1 Pipette (polypropylene) with 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 Europe Limited

Marketing authorisation date:
17/12/2014

Manufacturing sites for batch release:
KVP Pharma + Veterinär Produkte GmbH

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 00879/4157

Date of authorisation status change:
27/11/2024

Reference member state:
Germany

Procedure number:
DE/V/0196/006

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