File downloaded on 2026-11-03
Source URL: https://medicines.health.europa.eu/veterinary/en/600000062670

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:
Germany

Available in:
Germany

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 GmbH

Marketing authorisation date:
30/03/2017

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402351.00.00

Date of authorisation status change:
29/06/2021

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)
Published on: 14/03/2026
Download

2402351-paren-20211015.pdf