

# Prinovox 250 mg + 62.5 mg Spot-on Solution for Large Dogs

Not  
authorised

- Moxidectin
- Imidacloprid

## Product identification

**Medicine name:**

Prinovox 250 mg + 62.5 mg Spot-on Solution for Large Dogs

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**Active substance:**

Moxidectin

Imidacloprid

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**Target species:**

Dog

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Moxidectin

62.50 milligram(s) / 2.50 millilitre(s)

Imidacloprid

250.00 milligram(s) / 2.50 millilitre(s)

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**Pharmaceutical form:**

Spot-on solution

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP54AB52

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

United Kingdom (Northern Ireland)

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Elanco Europe Limited

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**Marketing authorisation date:**

17/12/2014

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**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

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**Responsible authority:**

The Veterinary Medicines Directorate

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**Authorisation number:**

Vm 00879/4154

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**Date of authorisation status change:**

27/11/2024

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0196/005

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)