

# Prinovox 40 mg + 4 mg spot-on solution for small cats and ferrets

Authorised

- Moxidectin
- Imidacloprid

## Product identification

**Medicine name:**

Prinovox 40 mg + 4 mg spot-on solution for small cats and ferrets

**Active substance:**

Moxidectin

Imidacloprid

**Target species:**

Cat

Ferret

**Route of administration:**

Cutaneous use

## Product details

**Active substance and strength:**

Moxidectin

4.00 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:**

Portugal

---

**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 Animal Health GmbH

---

**Marketing authorisation date:**

27/02/2017

---

**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

---

**Responsible authority:**

Directorate General For Food And Veterinary

---

**Authorisation number:**

1091/01/17RFVPT

---

**Date of authorisation status change:**

22/07/2025

---

**Reference member state:**

Germany

---

**Procedure number:**

DE/V/0196/001

---

**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](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

Published on: 13/03/2026

Download

2402349-paren-20210426.pdf