

# Stronghold 240 mg - Spot-on solution (12% - dogs)

Authorised

- Selamectin

## Product identification

**Medicine name:**

Stronghold 240 mg - Spot-on solution (12% - dogs)

---

**Active substance:**

Selamectin

---

**Target species:**

Dog

---

**Route of administration:**

Spot-on use

---

## Product details

**Active substance and strength:**

Selamectin

240.00 milligram(s) / 1.00 Pipette

---

**Pharmaceutical form:**

Spot-on solution

---

**Withdrawal period by route of administration:**

**Spot-on use:**

- 

**Dog**

---

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

QP54AA05

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

---

**Available in:**

Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

---

**Package description:**

Packaging:Single-dose pipette (polypropylene), Package\_size:6 pipettes, Content:2 ml (120 mg/ml)

Packaging:Single-dose pipette (polypropylene), Package\_size:3 pipettes, Content:2 ml (120 mg/ml)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Complete application (stand-alone) - Council Directive 81/851/EEC

---

**Marketing authorisation holder:**

Zoetis Belgium SA

---

**Marketing authorisation date:**

25/11/1999

---

**Manufacturing sites for batch release:**

Zoetis Belgium

---

**Responsible authority:**

European Commission

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

25/11/1999

---

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: 5/03/2025

[Download](#)

ema-puar-stronghold-v-050-var-x-0051-g-en.pdf

ema-puar-stronghold-v-050-par-en.pdf

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000000563>