

# Fatromectin 5 mg/ml pour-on solution for cattle.

Not  
authorised

- Ivermectin

## Product identification

**Medicine name:**

Fatromectin 5 mg/ml pour-on solution for cattle.

Fatromectin 5 mg/ml pour-on solution for cattle.

**Active substance:**

Ivermectin

**Target species:**

Cattle

**Route of administration:**

Pour-on use

## Product details

**Active substance and strength:**

Ivermectin

5.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Pour-on solution

**Withdrawal period by route of administration:****Pour-on use:**

- 

**Cattle**

- Meat and offal. 31 day

---

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

QP54AA01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Surrendered

---

**Authorised in:**

Ireland

---

**Package description:**

5.0 L white non-fluorinated or fluorinated high-density polyethylene back-pack with polypropylene strap and vented cap Closure: White polypropylene screw-cap

2.5 L white non-fluorinated or fluorinated high-density polyethylene back-pack with polypropylene strap and vented cap Closure: White polypropylene screw-cap

1.0 L white non-fluorinated or fluorinated high-density polyethylene bottle with a drawing tube and measuring device. Closure: White polypropylene screw-cap

250 ml white non-fluorinated or fluorinated high-density polyethylene bottle with a drawing tube and measuring device. Closure: White polypropylene screw-cap

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Eco Animal Health Europe Limited

---

**Marketing authorisation date:**

23/04/2004

---

**Manufacturing sites for batch release:**

Safapac Limited

Acme Drugs S.r.l.

---

**Responsible authority:**

Health Products Regulatory Authority

---

**Authorisation number:**

VPA22693/006/001

---

**Date of authorisation status change:**

1/11/2025

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0177/001

---

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

## Documents

Summary of Product Characteristics