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

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:**

Valid

#### Authorised in:

Ireland

## Package description:

5.0 L white non-fluorinated or fluorinated high-density polyethylene back-pack with polypropylene strap and vented capClosure: 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:

.5/01/2001
Manufacturing sites for batch release: Acme Drugs - S.r.l. Safapac (Cambridge) Limited
Responsible authority: Health Products Regulatory Authority
Authorisation number: /PA22693/006/001
Date of authorisation status change: 23/04/2004
Reference member state: reland
Procedure number: E/V/0177/001
Concerned member states: France Italy United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Summary of Product Characteristics

**Source URL:** https://medicines.health.europa.eu/veterinary/600000048699