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# Vetimec 5 mg/ml Pour-on Solution for Cattle

Not authorised

Ivermectin

# **Product identification**

#### **Medicine name:**

Imec 5 mg/ml pour-on solution for cattle Vetimec 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:**

Expired

#### **Authorised in:**

United Kingdom (Northern 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 Limited
Marketing authorisation date: 29/07/2008
Manufacturing sites for batch release: Safapac Limited Acme Drugs S.r.l.
Responsible authority: The Veterinary Medicines Directorate
Authorisation number: Vm 13277/4023
<b>Date of authorisation status change:</b> 16/10/2025
Reference member state: Ireland
Procedure number: IE/V/0178/001
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