

# Imec 5 mg/ml pour-on solution for cattle

Not  
authorised

- Ivermectin

## Product identification

**Medicine name:**

Imec 5 mg/ml pour-on solution for cattle

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**Active substance:**

Ivermectin

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**Target species:**

Cattle

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**Route of administration:**

Pour-on use

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## Product details

**Active substance and strength:**

Ivermectin

5.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Pour-on solution

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

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**Cattle**

- Meat and offal. 31 day

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

QP54AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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**Authorisation status:**

Surrendered

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**Authorised in:**

France

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Eco Animal Health Europe Limited

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**Marketing authorisation date:**

9/03/2006

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**Manufacturing sites for batch release:**

Safapac Limited  
Acme Drugs S.r.l.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5294397 1/2006

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**Date of authorisation status change:**

20/11/2025

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0178/001

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Summary of Product Characteristics

English (PDF)

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