

# Epromec 5 mg/ml Pour-on Solution for beef and dairy cattle

Authorised

- Eprinomectin

## Product identification

**Medicine name:**

Epromec 5 mg/ml Pour-on Solution for beef and dairy cattle

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**Active substance:**

Eprinomectin

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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:**

Eprinomectin

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. 15 day
- Milk. 0 hour

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

QP54AA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Portugal

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**Package description:**

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:5L 'Flexi' packs.Pack sizes 5 L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:3L 'Flexi' packs.Pack sizes 3L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of the following:2.5 L 'Flexi' packs.Pack sizes 2.5L.

High density polyethylene container with a polypropylene tamper evident screw cap which consists of:1L 'Squeeze pour' packs.Pack sizes 1L.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Marketing authorisation date:**

4/04/2016

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Directorate General For Food And Veterinary

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**Authorisation number:**

996/01/16RFVPT

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**Date of authorisation status change:**

18/08/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0355/001

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**Concerned member states:**

France Italy Netherlands Poland Portugal Spain

United Kingdom (Northern Ireland)

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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

## Combined File of all Documents

English (PDF)

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