

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

Authorised

- Eprinomectin

## Product identification

**Medicine name:**

Unomec 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:**

Belgium

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**Package description:**

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

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 the following: 3L 'Flexi' packs. Pack sizes 3L.

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

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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:**

9/11/2017

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**Manufacturing sites for batch release:**

Chanelle Pharmaceuticals Manufacturing Limited

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V519395

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**Date of authorisation status change:**

9/11/2017

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0354/001

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**Concerned member states:**

Austria Belgium Bulgaria Czechia Denmark Finland France Germany  
Greece Hungary Italy Netherlands Norway Poland Portugal Romania Spain  
Sweden 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

English (PDF)

Published on: 12/04/2026

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### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Combined File of all Documents

### Labelling

This document does not exist in this language (English). You can find it in another language below.