

# Eprecis 20 mg/ml solution for injection for cattle, sheep and goats

Authorised

- Eprinomectin

## Product identification

**Medicine name:**

Eprecis 20 mg/ml solution for injection for cattle, sheep and goats

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**Active substance:**

Eprinomectin

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**Target species:**

Cattle

Goat

Sheep

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Eprinomectin

20.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Subcutaneous use:**

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**Cattle**

- Meat and offal. 63 day
- Milk. 0 hour

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**Goat**

- Meat and offal. 42 day
- Milk. 0 hour

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**Sheep**

- Meat and offal. 42 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:**

Austria

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**Available in:**

Austria

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**Package description:**

Nature of immediate packaging: Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper and aluminium

cap and plastic flip-off disc in a cardboard box.Pack size: 50 ml vial

Nature of immediate packaging:Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper and aluminium cap and plastic flip-off disc in a cardboard box.Pack size: 100 ml vial

Nature of immediate packaging:Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper and aluminium cap and plastic flip-off disc in a cardboard box.Pack size: 250 ml vial

Nature of immediate packaging:Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper and aluminium cap and plastic flip-off disc in a cardboard box.Pack size: 500 ml vial

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

8/07/2015

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

836330

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**Date of authorisation status change:**

8/07/2015

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0340/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France  
Germany Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands  
Poland Portugal Romania Slovakia Slovenia 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: 30/06/2024

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### Package Leaflet

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

### Labelling

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