

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

Active substance:

Eprinomectin

Target species:

Cattle

Goat

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Eprinomectin

20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Cattle

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

-

Goat

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

-

Sheep

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

18/06/2015

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Danish Medicines Agency

Authorisation number:

53767

Date of authorisation status change:

18/06/2015

Reference member state:

Ireland

Procedure number:

IE/V/0340/001

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)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2026

[Download](#)