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
Eprecis 20 mg/ml, injekcinis tirpalas galvijams, avims ir ožkoms

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:
Lithuania

Available in:
Lithuania

Package description:

Nature of immediate packaging:Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper andaluminium cap and plastic flip-off disc in a cardboard box.Pack size: 500 ml vial
Nature of immediate packaging:Amber multilayer plastic vial (polypropylene / ethylene vinyl alcohol / polypropylene) with bromobutyl rubber stopper andaluminium 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 andaluminium 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 andaluminium cap and plastic flip-off disc in a cardboard box.Pack size: 50 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:
17/07/2015

Manufacturing sites for batch release:
CEVA Santé Animale

Responsible authority:
State Food And Veterinary Service

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
29/12/2020

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

RV2291.pdf