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

- Eprinomectin

Authorised

Product identification

Medicine name:
EPRECIS 20 mg/ml ??????? ???????
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:

- Greece

Available in:

- Greece

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 Hellas LLC

Marketing authorisation date:

- 15/03/2016

Manufacturing sites for batch release:

- CEVA Santé Animale

Responsible authority:

- National Organization For Medicines

Authorisation number:

- 37817/06-05-2021/K-0207201

Date of authorisation status change:

- 5/05/2021

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

Product information

Summary of Product Characteristics

English (PDF)
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