Eprecis 5 mg/ml pour-on solution for cattle, sheep and goats

Authorised

- Eprinomectin

Product identification

Medicine name:
Eprecis 5 mg/ml pour-on solution for cattle, sheep and goats
EPRECIS 5 MG/ML SOLUTION POUR POUR-ON POUR BOVINS, OVINS ET CAPRINS

Active substance:
Eprinomectin

Target species:
Cattle
Goat
Sheep

Route of administration:
Pour-on use

Product details

Active substance and strength:
Eprinomectin
5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Pour-on solution

Withdrawal period by route of administration:

Pour-on use:

- **Cattle**
  - Meat and offal. 15 day
  - Milk. 0 hour
- **Goat**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 2 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:
France

Available in:
France

Package description:
HISTORICAL Squeeze-measure pour-on system:250 ml translucent high density polyethylene (HDPE) bottle including 10 ml dispenser graduated each 5 ml, with

removablealuminium/PE seals and PE screw cap.
Back pack:1 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap or 1 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap included in acardboard box.
Back pack:2.5 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap or 2.5 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap included in acardboard box.
Back pack:5 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap or 5 L white HDPE bottles, with a removable aluminium/PE seal and a polypropylene (PP) screw cap included in acardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
13/05/2015

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1482427 6/2015

Date of authorisation status change:
28/07/2020

Reference member state:
Ireland

Procedure number:
IE/V/0343/001

Concerned member states:
Belgium Denmark France Germany Hungary Italy Netherlands Poland Portugal Spain 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: 30/03/2025
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Package Leaflet and Labelling

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Combined File of all Documents