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

Authorised

- Eprinomectin

Product identification

Medicine name:
Eprinex Multi 5 mg/ml pour-on solution for cattle, sheep and goats

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:
Sweden

Available in:
Sweden

Package description:
5 L HDPE back packSealed foil and tamper evident HDPE screw cap with polypropylene liner5L back-pack with a dispensing capOne back-pack per cardboard

box.The 5 litre back-packs are designed for use with a suitable automatic dispensing gun.
2.5 HDPE back packSealed foil and tamper evident HDPE screw cap with polypropylene liner2.5L back-pack with a dispensing capOne bottle or one back-pack per cardboard box.The 2.5 litre back-packs are designed for use with a suitable automatic dispensing gun.
1 L HDPE bottleSealed foil and tamper evident HDPE screw cap with polypropylene liner1L bottle with 2 measuring devices (1 of 60 ml for cattle, 1 of 25 ml for sheep/goat)One bottle per cardboard box.
250 ml HDPE bottleSealed foil and tamper evident HDPE screw cap with polypropylene liner250 ml bottle with 2 measuring devices of 25 ml (1 for cattle, 1 for sheep/goat)One bottle per cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health Denmark A/S

Marketing authorisation date:
15/10/2018

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
57807

Date of authorisation status change:
15/10/2018

Reference member state:
Ireland

Procedure number:
IE/V/0347/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Greece Hungary Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Package Leaflet

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Summary of Product Characteristics

English (PDF)
Published on: 11/05/2025
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Labelling

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Combined File of all Documents