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
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:
Ireland

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 Vetmedica GmbH

Marketing authorisation date:
9/12/2016

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10454/034/001

Date of authorisation status change:
9/12/2016

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)

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

Documents

Summary of Product Characteristics

Combined File of all Documents