

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:

Topical 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:

Topical 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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

5 L HDPE back pack Sealed foil and tamper evident HDPE screw cap with polypropylene liner 5L back-pack with a dispensing cap One back-pack per cardboard box. The 5 litre back-packs are designed for use with a suitable automatic dispensing gun.

2.5 HDPE back pack Sealed foil and tamper evident HDPE screw cap with polypropylene liner 2.5L back-pack with a dispensing cap One 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 bottle Sealed foil and tamper evident HDPE screw cap with polypropylene liner 1L 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 UK Limited

Marketing authorisation date:

15/12/2016

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

VM 08327/4282

Date of authorisation status change:

15/12/2016

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000047609>