Ecomectin 10 mg/ml Solution for Injection

Authorised

- Ivermectin

Product identification

Medicine name:
Ecomectin 10 mg/ml Solution for Injection
Ecomectin 10 mg/ml Oplossing voor injectie
Ecomectin 10 mg/ml Solution injectable
Ecomectin 10 mg/ml Injektionslösung

Active substance:
Ivermectin

Target species:
Cattle
Sheep
Pig

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 42 day
- **Sheep**
  - Meat and offal. 42 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 50 ml
HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 200 ml.
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 500 ml
HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 500 ml.
HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 50 ml.
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 250 ml

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:
Eco Animal Health Europe Limited

Marketing authorisation date:
29/03/2004

Manufacturing sites for batch release:
Produlab Pharma B.V.
Divasa Farmavic S.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V261091

Date of authorisation status change:
29/03/2004

Reference member state:
Ireland

Procedure number:
IE/V/0144/001

Concerned member states:
Austria Belgium Estonia France Germany Greece Italy Latvia Lithuania Luxembourg Netherlands 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: 11/02/2022
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Package Leaflet

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Labelling

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