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Ecomectin 10 mg/ml Solution for Injection

Authorised

- Ivermectin

Product identification

Medicine name:
Ecomectin 10 mg/ml Solution for Injection

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. 49 day
  - Milk. no withdrawal period

Do not use in lactating cows producing milk for human consumption. Do not use in non-lactating dairy cows, including pregnant dairy heifers, within 60 days of calving.

- **Sheep**
  - Meat and offal. 42 day
  - Milk. no withdrawal period

Do not use in lactating ewes producing milk for human consumption. Do not use in sheep which are intended to produce milk for human consumption within 60 days of lambing.

- **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:

Ireland

Package description:

Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 250 ml
HDPE 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: 500 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: 200 ml.
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 50 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:
1/10/1999

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22693/001/001

Date of authorisation status change:

1/10/1999

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