Ecomectin 10 mg/ml Solution for Injection

Authorised

- Ivermectin

Product identification

Medicine name:
Ecomectin 10 mg/ml Solution for Injection
VETOMECTIN 10 MG/ML SOLUTION INJECTABLE

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

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Authorisation status:

Valid

Authorised in:

France

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

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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:
5/02/2004

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/4934184 5/2004

Date of authorisation status change:
5/02/2009

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)
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Package Leaflet and Labelling

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