Bimectin vet 10 mg/ml, solution for injection

Authorised

- Ivermectin

Product identification

Medicine name:
Bimectin vet 10 mg/ml, solution for injection

Active substance:
Ivermectin

Target species:
Cattle
Reindeer
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

Lactating cows producing milk for human consumption should not be treated. Dry cows and heifers should not be treated within 60 days prior to calving.

- **Reindeer**
  - Meat and offal. 28 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:

Finland

Package description:

Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
25/11/2001

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Finnish Medicines Agency

Authorisation number:
16727

Date of authorisation status change:
25/11/2001

Reference member state:
Sweden

Procedure number:
SE/V/0111/001

Concerned member states:
Finland

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

Documents

Package Leaflet

English (PDF)
Published on: 29/04/2024
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Summary of Product Characteristics

English (PDF)
Published on: 29/04/2024
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Labelling