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Baytril vet. 100 mg/ml Injektionsvätska, lösning

Authorised

- Enrofloxacin

Product identification

Medicine name:
Baytril vet. 100 mg/ml Injektionsvätska, lösning

Active substance:
Enrofloxacin

Target species:
Sheep
Goat
Cattle
Pig

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Sheep**
  - Milk. 3 day
  - Meat and offal. 4 day
- **Goat**
  - Milk. 4 day
  - Meat and offal. 6 day
- **Cattle**
  - Milk. 3 day Efter intravenös injektion
  - Milk. 4 day Efter subkutan injektion
  - Meat and offal. 12 day Efter subkutan injektion
  - Meat and offal. 5 day Efter intravenös injektion
- **Pig**
  - Meat and offal. 13 day

Intravenous use:

- **Sheep**
  - Milk. 3 day
  - Meat and offal. 4 day

- **Goat**
  - Milk. 4 day
  - Meat and offal. 6 day
- **Cattle**
  - Milk. 3 day Efter intravenös injektion
  - Milk. 4 day Efter subkutan injektion
  - Meat and offal. 5 day Efter intravenös injektion
  - Meat and offal. 12 day Efter subkutan injektion
- **Pig**
  - Meat and offal. 13 day

Subcutaneous use:

- **Sheep**
  - Milk. 3 day
  - Meat and offal. 4 day
- **Goat**
  - Milk. 4 day
  - Meat and offal. 6 day
- **Cattle**
  - Milk. 4 day Efter subkutan injektion
  - Milk. 3 day Efter intravenös injektion

- Meat and offal. 12 day Efter subkutan injektion
- Meat and offal. 5 day Efter intravenös injektion

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Pig

- Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

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Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Elanco Animal Health GmbH

Marketing authorisation date:
27/10/1989

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
11019

Date of authorisation status change:
27/10/1989

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

Documents

Package Leaflet

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.