Engemycin 10%, 100 mg/ ml roztwór do wstrzykiwań dla bydła, świń, koni, owiec

Authorised

- Oxytetracycline

Product identification

Medicine name:
Engemycin 10%, 100 mg/ ml roztwór do wstrzykiwań dla bydła, świń, koni, owiec

Active substance:
Oxytetracycline

Target species:
Sheep
Horse
Pig
Cattle

Route of administration:
Subcutaneous use
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Oxytetracycline
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Sheep**
  - Meat and offal. 12 day
  - Milk. 3 day
- **Horse**
  - All relevant tissues. no withdrawal period

Do not use in horses whose tissues are intended for human consumption.

- **Pig**
  - Meat and offal. 7 day
- **Cattle**
  - Meat and offal. 16 day
  - Meat and offal. 24 day
  - Milk. 3 day

Intravenous use:

- **Sheep**

- Meat and offal. 12 day
- Milk. 3 day

- **Horse**
  - All relevant tissues. no withdrawal period

Do not use in horses whose tissues are intended for human consumption.

- **Pig**
  - Meat and offal. 7 day

- **Cattle**
  - Meat and offal. 16 day
  - Meat and offal. 24 day
  - Milk. 3 day

Intramuscular use:

- **Sheep**
  - Milk. 3 day
  - Meat and offal. 12 day

- **Horse**
  - All relevant tissues. no withdrawal period

Do not use in horses whose tissues are intended for human consumption.

- **Pig**
  - Meat and offal. 7 day

- **Cattle**
  - Meat and offal. 24 day

- Meat and offal. 16 day
- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
11/03/1999

Manufacturing sites for batch release:
Intervet International GmbH
Intervet Productions S.r.l.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0662

Date of authorisation status change:
11/03/1999

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

Documents

Labelling

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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.

Package Leaflet

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