OXY-100, 100 mg/ml, injekcinis tirpalas galvijams, veršeliams, avims, ožkoms ir kiaulėms

Authorised

- Oxytetracycline

Product identification

Medicine name:
OXY-100, 100 mg/ml, injekcinis tirpalas galvijams, veršeliams, avims, ožkoms ir kiaulėms

Active substance:
Oxytetracycline

Target species:
Cattle
Goat
Sheep
Pig

Route of administration:
Intramuscular use
Subcutaneous 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:
Intramuscular use:

- **Cattle**
  - Meat. 18 day
  - Milk. 5 day
- **Goat**
  - Meat. 15 day
  - Milk. no withdrawal period

Not authorized for use in goats, whose milk is meant for human consumption.

- **Sheep**
  - Meat. 15 day
  - Milk. no withdrawal period

Not authorized for use in sheep, whose milk is meant for human consumption.

- **Pig**
  - Meat. 15 day

Subcutaneous use:

-

Cattle

- Meat. 18 day
- Milk. 5 day

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Goat

- Meat. 15 day
- Milk. no withdrawal period

Not authorized for use in goats, whose milk is meant for human consumption.

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Sheep

- Meat. 15 day
- Milk. no withdrawal period

Not authorized for use in sheep, whose milk is meant for human consumption.

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Pig

- Meat. 15 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

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Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Interchemie werken De Adelaar LT UAB

Marketing authorisation date:
26/03/2015

Manufacturing sites for batch release:
Interchemie Werken De Adelaar Eesti AS

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/15/2281/001-004

Date of authorisation status change:
13/11/2025

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

Documents

Combined File of all Documents

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