

Engemycin, 100 mg/ml, injekcinis tirpalas galvijams, kiaulėms, arkliams, avims, šunims ir katėms

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:

Engemycin, 100 mg/ml, injekcinis tirpalas galvijams, kiaulėms, arkliams, avims, šunims ir katėms

Active substance:

Oxytetracycline hydrochloride

Target species:

Cattle
Horse
Sheep
Pig
Dog
Cat

Route of administration:

Intravenous use
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Oxytetracycline hydrochloride
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

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Cattle

- Meat and offal. 27 day At low doses.
- Milk. 4 day At low doses. 4 days or 8 milkings.
- Meat and offal. 18 day At high doses.

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Sheep

- Meat and offal. 18 day
- Milk. 4 day 4 days or 8 milkings.

Intramuscular use:

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Cattle

- Meat and offal. 27 day At low doses.
- Milk. 4 day 4 days or 8 milkings.
- Meat and offal. 18 day At high doses.

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Pig

- Meat and offal. 8 day At low doses.

- Meat and offal. 7 day At high doses.

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Sheep

- Meat and offal. 18 day

- Milk. 4 day 4 days or 8 milkings.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Available in:

Lithuania

Package description:

Available only in Lithuanian

Available only in Lithuanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

21/12/1995

Manufacturing sites for batch release:

Intervet International GmbH

Intervet Productions S.r.l.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/95/0263/001-002

Date of authorisation status change:

25/04/2024

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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