Engemycin 100 mg/ml raztopina za injiciranje za govedo, prašiče, konje, ovce, pse in mačke

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
Engemycin 100 mg/ml raztopina za injiciranje za govedo, prašiče, konje, ovce, pse in mačke

Active substance:
Oxytetracycline hydrochloride

Target species:
Cattle
Horse
Pig
Sheep
Dog
Cat

Route of administration:
Intramuscular use
Intravenous 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:
Intramuscular use:

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

Meso in organi: zdravljenje z majhnimi odmerki: 24 dni

  - Meat and offal. 16 day zdravljenje z velikimi odmerki: 16 dni
  - Milk. 3 day Mleko: 3 dni (6 molž)

- **Horse**
  - Meat and offal. no withdrawal period

Mi dovoljena uporaba pri konjih, ki so namenjeni za prehrano ljudi. Not for use in food producing horses.

- **Pig**
  - Meat and offal. 7 day Meso in organi: 7 dni

- **Sheep**
  - Meat and offal. 12 day Meso in organi: 12 dni

- Milk. 3 day Mleko: 3 dni (6 molž)

Intravenous use:

- **Cattle**
  - Meat and offal. 24 day zdravljenje z majhnimi odmerki: 24 dni
  - Meat and offal. 16 day zdravljenje z velikimi odmerki: 16 dni
  - Milk. 3 day 3 dni (6 molž)
- **Sheep**
  - Meat and offal. 12 day Meso in organi: 12 dni
  - Milk. 3 day Mleko: 3 dni (6 molž)

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:
Slovenia

Available in:
Slovenia

Package description:
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
12/12/2002

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0109/001

Date of authorisation status change:
12/12/2002

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

Documents

Summary of Product Characteristics

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Package Leaflet

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.