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L.S. Injection solution for injection for calves (pre-ruminant), sheep, goats, pigs, birds and dogs

Authorised

- Lincomycin hydrochloride monohydrate
- Spectinomycin dihydrochloride pentahydrate

Product identification

Medicine name:
L.S. Injection solution for injection for calves (pre-ruminant), sheep, goats, pigs, birds and dogs

Active substance:
Lincomycin hydrochloride monohydrate
Spectinomycin dihydrochloride pentahydrate

Target species:
Cattle (pre-ruminant)
Sheep
Goat
Pig
Dog
Chicken

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Lincomycin hydrochloride monohydrate
50.00 milligram(s) / 1.00 millilitre(s)

Spectinomycin dihydrochloride pentahydrate
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (pre-ruminant)**
  - Meat. 14 day
  - Liver. 21 day
  - Kidney. 21 day
- **Sheep**
  - Meat. 14 day

Не се разрешава за употреба при животни, чието мляко е предназначено за човешка консумация.

  - Liver. 21 day
  - Kidney. 21 day

- **Goat**
  - Meat. 14 day

Не се разрешава за употреба при животни, чието мляко е предназначено за човешка консумация.

  - Liver. 21 day

- Kidney. 21 day

- **Pig**
  - Meat. 14 day
  - Liver. 21 day
  - Kidney. 14 day

Subcutaneous use:

- **Chicken**
  - Meat. 14 day

Не се разрешава употребата при птици, чиито яйца са предназначени за човешка консумация

  - Liver. 21 day
  - Kidney. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF52

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Bibliographical application (stand-alone)

Marketing authorisation holder:
Kepro B.V.

Marketing authorisation date:
2/06/2010

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2528

Date of authorisation status change:
2/06/2010

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

Documents

Summary of Product Characteristics

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Labelling

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.