L.S. INJECTION

Authorised

- Spectinomycin dihydrochloride pentahydrate
- Lincomycin hydrochloride monohydrate

Product identification

Medicine name:
L.S. INJECTION

Active substance:
Spectinomycin dihydrochloride pentahydrate
Lincomycin hydrochloride monohydrate

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

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Spectinomycin dihydrochloride pentahydrate

100.00 milligram(s) / 1.00 millilitre(s)

Lincomycin hydrochloride monohydrate
50.00 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
- **Dog**

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:
3/06/2010

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2528

Date of authorisation status change:
3/06/2010

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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