LINCO-SPECTIN SOLUTION INJECTABLE POUR BOVINS (VEAUX) OVINS CAPRINS PORCINS VOLAILLES CHATS ET CHIENS

Authorised

- Lincomycin hydrochloride monohydrate
- Spectinomycin sulfate tetrahydrate

Product identification

Medicine name:
LINCO-SPECTIN SOLUTION INJECTABLE POUR BOVINS (VEAUX) OVINS CAPRINS PORCINS VOLAILLES CHATS ET CHIENS

Active substance:
Lincomycin hydrochloride monohydrate
Spectinomycin sulfate tetrahydrate

Target species:
Cattle (calf)
Pig
Cat
Sheep
Goat
Dog
Poultry

Route of administration:
Intramuscular use
Subcutaneous use

Product details

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

Spectinomycin sulfate tetrahydrate
151.17 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 14 day
- **Pig**
  - Meat and offal. 14 day
- **Sheep**
  - Meat and offal. 14 day
  - Milk. no withdrawal period

En l'absence de détermination d'un temps d'attente pour le lait, ne pas utiliser chez les femelles productrices de lait de consommation, en lactation ou en période de tarissement ni chez les futures productrices de lait de consommation dans les 2 mois précédant la mise-bas.

- **Goat**

- Meat and offal. 14 day

Subcutaneous use:

- **Poultry**
  - Meat and offal. 14 day
  - Eggs. no withdrawal period

En l'absence de LMR pour les œufs, ne pas utiliser chez les espèces pondeuses productrices d'œufs de consommation

- **Goat**
  - Milk. no withdrawal period

En l'absence de détermination d'un temps d'attente pour le lait, ne pas utiliser chez les femelles productrices de lait de consommation, en lactation ou en période de tarissement ni chez les futures productrices de lait de consommation dans les 2 mois précédant la mise-bas.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF52

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

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Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis France

Marketing authorisation date:
30/06/1992

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG
Zoetis Belgium

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7501878 8/1992

Date of authorisation status change:
30/06/2012

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

This document does not exist in this language (English). You can find it in another language below.