LINCO-SPECTIN 50 mg/ml + 100 mg/ml, soluție injectabilă pentru porci, bovine, găini, curcani, oi, capre, câini și pisici

Authorised

- Spectinomycin
- Lincomycin

Product identification

Medicine name:
LINCO-SPECTIN 50 mg/ml + 100 mg/ml, soluție injectabilă pentru porci, bovine, găini, curcani, oi, capre, câini și pisici

Active substance:
Spectinomycin
Lincomycin

Target species:
Pig
Cattle (calf)
Sheep
Goat
Dog
Cat
Chicken
Turkey (cockerel)

Route of administration:
Intramuscular use
Subcutaneous use
Oral use

Product details

Active substance and strength:
Spectinomycin
100.00 milligram(s) / 1.00 millilitre(s)
Lincomycin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 14 day

- **Cattle (calf)**
  - Meat and offal. 21 day

- **Sheep**
  - Meat and offal. 15 day

Nu se va administra la animale care produc lapte pentru consumul uman.

- **Goat**
  - Meat and offal. 15 day

Nu se va administra la animale care produc lapte pentru consumul uman.

- **Dog**
- **Cat**

Subcutaneous use:

- **Chicken**
  - Meat and offal. 15 day

Nu se va administra la găinile ce produc ouă pentru consumul uman.

- **Turkey (cockerel)**
  - Meat and offal. 15 day

Oral use:

- **Chicken**
  - Meat and offal. 15 day

Nu se va administra la găinile ce produc ouă pentru consumul uman.

- **Turkey (cockerel)**
  - Meat and offal. 15 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF52

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Available in:

Romania

Package description:
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
25/01/2015

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
150304

Date of authorisation status change:
5/12/2024

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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