KANAMICINA FP 250 mg/ml, soluție injectabilă pentru cai, bovine, oi, capre, porci, găini, câini și pisici

Authorised

- Kanamycin

Product identification

Medicine name:
KANAMICINA FP 250 mg/ml, soluție injectabilă pentru cai, bovine, oi, capre, porci, găini, câini și pisici

Active substance:
Kanamycin

Target species:
Cattle
Pig
Sheep
Goat
Chicken (hen)
Dog
Cat
Horse

Route of administration:

Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Kanamycin
250.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 36 hour
  - Meat and offal. 10 day
- **Pig**
  - Meat and offal. 10 day
- **Sheep**
  - Milk. 36 hour
  - Meat and offal. 10 day
- **Goat**
  - Meat and offal. 10 day
  - Milk. 36 hour
- **Chicken (hen)**

- Meat and offal. 10 day

Nu este permisă utilizarea la păsările ouătoare care produc ouă pentru consum uman.

- **Horse**
  - Meat and offal. 10 day

Subcutaneous use:

- **Cattle**
  - Milk. 36 hour
  - Meat and offal. 10 day
- **Pig**
  - Meat and offal. 10 day
- **Sheep**
  - Milk. 36 hour
  - Meat and offal. 10 day
- **Goat**
  - Meat and offal. 10 day
  - Milk. 36 hour
- **Chicken (hen)**
  - Meat and offal. 10 day

NU este permisă utilizarea la păsările ouătoare care produc ouă pentru consum uman.

- **Horse**

- Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01GB04

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
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:
Pasteur Filiala Filipesti S.A.

Marketing authorisation date:
11/06/2007

Manufacturing sites for batch release:

Pasteur Filiala Filipesti S.A.

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

Authorisation number:
150291

Date of authorisation status change:
18/11/2025

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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