

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

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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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