NEOMICINA SULFAT FP 100 mg/g, pulbere pentru utilizare în apa de băut pentru cai, bovine, oi, capre, porci, găini, câini, pisici, nurci

Authorised

- Neomycin

Product identification

Medicine name:
NEOMICINA SULFAT FP 100 mg/g, pulbere pentru utilizare în apa de băut pentru cai, bovine, oi, capre, porci, găini, câini, pisici, nurci

Active substance:
Neomycin

Target species:
Cattle (calf)
Sheep (lamb)
Goat (kid)
Pig (piglet)
Chicken (layer hen)
Chicken (broiler)
Dog
Cat
Mink
Horse

Route of administration:
In drinking water use

Product details

Active substance and strength:
Neomycin
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
In drinking water use:

- **Cattle (calf)**
  - Meat and offal. 5 day
- **Sheep (lamb)**
  - Meat and offal. 5 day
- **Goat (kid)**
  - Meat and offal. 5 day
- **Pig (piglet)**
  - Meat and offal. 4 day
- **Chicken (layer hen)**
  - Eggs. 0 day
  - Meat and offal. 3 day
- **Chicken (broiler)**

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA01

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
Available only in Romanian
Available only in Romanian
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:
20/07/2005

Manufacturing sites for batch release:
Pasteur Filiala Filipesti S.A.

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

Authorisation number:
120151

Date of authorisation status change:
15/04/2025

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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