NEOMYCINE 50 COOPHAVET POUDRE POUR SOLUTION BUVABLE

Authorised

- NEOMYCIN SULFATE

Product identification

Medicine name:
NEOMYCINE 50 COOPHAVET POUDRE POUR SOLUTION BUVABLE

Active substance:
NEOMYCIN SULFATE

Target species:
Cattle
Pig (piglet)
Rabbit
Sheep (lamb)
Sheep
Goat (kid)
Goat
Cattle (calf)
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
NEOMYCIN SULFATE
500000.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for oral solution

Withdrawal period by route of administration:
Oral use:

- **Cattle**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine

- **Pig (piglet)**
  - Meat and offal. 14 day

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

- **Sheep (lamb)**
  - Meat and offal. 14 day

- **Sheep**
  - Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine

-

Goat (kid)

- Meat and offal. 14 day

•

Goat

- Milk. no withdrawal period

Ne pas utiliser chez les animaux producteurs de lait destiné à la consommation humaine

•

Cattle (calf)

- Meat and offal. 14 day

•

Poultry

- Meat and offal. 14 day
- Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA01

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

---

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
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:
Dopharma France S.A.S.

Marketing authorisation date:
18/06/1992

Manufacturing sites for batch release:
Dopharma France

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

Authorisation number:
FR/V/0359766 8/1992

Date of authorisation status change:
18/06/2012

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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