Novosol 500 000 IU/g powder for use in drinking water/milk for cattle, chickens, pigs, ducks, turkeys, geese, quail and partridges

Authorised

- NEOMYCIN SULFATE

Product identification

Medicine name:
Novosol 500 000 IU/g powder for use in drinking water/milk for cattle, chickens, pigs, ducks, turkeys, geese, quail and partridges

Active substance:
NEOMYCIN SULFATE

Target species:
Quail
Pig
Cattle (pre-ruminant)
Partridge
Goose
Duck
Chicken
Turkey

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 use in drinking water/milk

Withdrawal period by route of administration:
Oral use:

- **Quail**
  - Meat and offal. 14 day
  - Eggs. 0 day
- **Pig**
  - Meat and offal. 3 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 14 day
- **Partridge**
  - Meat and offal. 14 day
  - Eggs. 0 day
- **Goose**
  - Meat and offal. 14 day
  - Eggs. 0 day

- **Duck**
  - Meat and offal. 14 day
  - Eggs. 0 day
- **Chicken**
  - Meat and offal. 14 day
  - Eggs. 0 day
- **Turkey**
  - 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:
Ireland

Package description:
100 g sachet
1 kg bag

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
18/07/2025

Manufacturing sites for batch release:
Huvepharma S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10782/048/001

Date of authorisation status change:
18/07/2025

Reference member state:
France

Procedure number:
FR/V/0501/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

Generic of:
600000004401

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

Documents

Combined File of all Documents

eu-puar-frv0501001-mr-rpe907-en.pdf