

NEOSOL 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:

NEOSOL 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:

Denmark

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:

4/11/2025

Manufacturing sites for batch release:

Huvepharma S.A.

Responsible authority:

Danish Medicines Agency

Authorisation number:

72189

Date of authorisation status change:

4/11/2025

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