

NEOSOL 500 000 IU/G POWDER
FOR USE IN DRINKING
WATER/MILK FOR CATTLE,
CHICKENS, PIGS, DUCKS,
TURKEYS, GEESE, QUAIL AND
PARTRIDGES

Authorised

- Neomycin

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

Target species:

Quail

Pig

Cattle (calf)

Partridge

Goose

Duck

Chicken

Turkey

Route of administration:

Oral use

Product details

Active substance and strength:

Neomycin

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 (calf)

- 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:

Lithuania

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:

27/10/2025

Manufacturing sites for batch release:

Huvepharma S.A.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/25/2906/001-002

Date of authorisation status change:

27/10/2025

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

English (PDF)

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