

# NEOMYCINE 40 LAPIN-VOLAILLE- PORC ET AGNEAU-CHEVREAU SEVRES

Authorised

- NEOMYCIN SULFATE

## Product identification

**Medicine name:**

NEOMYCINE 40 LAPIN-VOLAILLE-PORC ET AGNEAU-CHEVREAU SEVRES

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**Active substance:**

NEOMYCIN SULFATE

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**Target species:**

Quail  
Pig  
Rabbit  
Sheep (lamb)  
Sheep  
Goat (kid)  
Goat  
Pheasant  
Partridge  
Guinea fowl  
Goose  
Duck  
Chicken

Turkey

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

NEOMYCIN SULFATE

40000.00 international unit(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Premix for medicated feeding stuff

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**Withdrawal period by route of administration:**

**Oral use:**

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**Quail**

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

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**Pig**

- Meat and offal. 14 day

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**Rabbit**

- Meat and offal. 14 day

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**Sheep (lamb)**

- Meat and offal. 14 day

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**Sheep**

- Milk. no withdrawal period

La spécialité n'est pas destinée aux femelles laitières productrices de lait de consommation.

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**Goat (kid)**

- Meat and offal. 14 day

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**Goat**

- Milk. no withdrawal period

La spécialité n'est pas destinée aux femelles laitières productrices de lait de consommation.

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**Pheasant**

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

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**Partridge**

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

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**Guinea fowl**

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

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**Goose**

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

- 

**Duck**

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

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#### **Chicken**

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

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#### **Turkey**

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

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#### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA07AA01

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#### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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#### **Authorisation status:**

Valid

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#### **Authorised in:**

France

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#### **Available in:**

France

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#### **Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## **Additional information**

#### **Entitlement type:**

## Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Huvepharma S.A.

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**Marketing authorisation date:**

1/10/1990

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**Manufacturing sites for batch release:**

Huvepharma S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/6391074 8/1990

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**Date of authorisation status change:**

1/10/2010

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.