

# Apravet 552 000 IU/g powder for use in drinking water/milk for pigs, calves, chickens and rabbits

Authorised

- Apramycin sulfate

## Product identification

### **Medicine name:**

Apravet 552 000 IU/g powder for use in drinking water/milk for pigs, calves, chickens and rabbits

### **Active substance:**

Apramycin sulfate

### **Target species:**

Pig (weaned piglet)

Rabbit

Cattle (pre-ruminant)

Chicken (broiler)

### **Route of administration:**

In drinking water/milk use

## Product details

### **Active substance and strength:**

Apramycin sulfate

1.00 milligram(s) / 1.00 milligram(s)

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**Pharmaceutical form:**

Powder for use in drinking water/milk

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**Withdrawal period by route of administration:**

**In drinking water/milk use:**

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**Pig (weaned piglet)**

- Meat and offal. 0 day

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**Rabbit**

- Meat and offal. 0 day

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**Cattle (pre-ruminant)**

- Meat and offal. 28 day

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**Chicken (broiler)**

- Meat and offal. 0 day

- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

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

QA07AA92

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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:**

Lithuania

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**Package description:**

box containing 50 sachets of 1,812 g

box containing 25 sachets of 1,812 g

bottle containing 90,58 g

bag containing 1811,6 g

1 Sachet with 1,812g

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

HuVepharma

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**Marketing authorisation date:**

17/10/2018

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**Manufacturing sites for batch release:**

Biovet AD

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/18/2489/001-003

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**Date of authorisation status change:**

24/07/2025

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0252/001

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**Concerned member states:**

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta  
Netherlands Poland Portugal Romania Slovakia Slovenia  
United Kingdom (Northern Ireland)

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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

Combined File of all Documents

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eu-PUAR-esv0252001-dcp-apravet-soluble-powder-en.pdf