# 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
Apravet, 552 RÜ/mg pulber joogivees/piimas manustamiseks sigadele, vasikatele, kanadele ja küülikutele

**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)

**Pharmaceutical form:**
Powder for use in drinking water/milk

**Withdrawal period by route of administration:**
**In drinking water/milk use:**

- **Pig (weaned piglet)**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 28 day
- **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.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QA07AA92

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Estonia

**Available in:**
Estonia

**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

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
HuVepharma

**Marketing authorisation date:**
8/10/2018

**Manufacturing sites for batch release:**
Biovet AD

**Responsible authority:**
State Agency Of Medicines

**Authorisation number:**
2114

**Date of authorisation status change:**
8/10/2018

**Reference member state:**
Spain

**Procedure number:**
ES/V/0252/001

**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)

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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 21/12/2023
Download

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