

# GABBROVET 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/MILK FOR PRE-RUMINANT CATTLE AND PIGS

Authorised

- Paromomycin sulfate

## Product identification

### **Medicine name:**

GABBROVET 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/MILK FOR PRE-RUMINANT CATTLE AND PIGS

### **Active substance:**

Paromomycin sulfate

### **Target species:**

Cattle (pre-ruminant)

Pig

### **Route of administration:**

In drinking water/milk use

In drinking water use

## Product details

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

Paromomycin sulfate

200.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Solution 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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#### **Cattle (pre-ruminant)**

- Meat and offal. 20 day

#### **In drinking water use:**

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#### **Pig**

- Meat and offal. 3 day

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

QA07AA06

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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:**

Czechia

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### **Package description:**

Box of 1 vial of 125 mL

Vial of 1000 mL

Vial of 500 mL

Vial of 250 mL

Vial of 125 mL  
Box of 1 vial of 1000 mL  
Box of 1 vial of 500 mL  
Box of 1 vial of 250 mL

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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:**

Ceva Sante Animale

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**Marketing authorisation date:**

16/03/2018

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**Manufacturing sites for batch release:**

Ceva Sante Animale

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/013/18-C

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**Date of authorisation status change:**

28/01/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0317/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Greece

Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands  
Poland Portugal Romania Slovakia Slovenia Spain  
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

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Labelling

This document does not exist in this language (English). You can find it in another language below.

### Combined File of all Documents

eu-puar-frv0317001-mr-rpe364-en.pdf