

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

Authorised

- Paromomycin sulfate

## Product identification

### Medicine name:

Gabbrovet Multi 140 mg/ml Oplossing voor gebruik in het drinkwater/in de melk  
Gabbrovet Multi 140 mg/ml Solution pour administration dans l'eau de boisson/le lait  
Gabbrovet Multi 140 mg/ml Lösung zum Eingeben über das Trinkwasser/die Milch  
GABBROVET MULTI 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/ MILK FOR  
PRE-RUMINANT CATTLE AND PIGS

### Active substance:

Paromomycin sulfate

### Target species:

Cattle (calf)

Pig

### Route of administration:

Oral 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:

#### Oral use:

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#### Cattle (calf)

- Meat and offal. 110 day

Cryptosporidiosis: Dosage: 150 mg/kg/day for 5 days.

- Meat and offal. 20 day

Colibacillosis: Dosage: 25-50 mg/kg/day for 3 to 5 days.

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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:

Belgium

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### Package description:

Cardboard box containing 1 bottle of 250 ml (HDPE/EVOH/HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 500 ml (HDPE/EVOH/HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 1 L (HDPE/EVOH/HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 125 ml (HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 250 ml (HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 500 ml (HDPE) containing a dosing device of 30 ml

Cardboard box containing 1 bottle of 1 L (HDPE) containing a dosing device of 30 ml

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## Additional information

### **Entitlement type:**

Marketing Authorisation

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### **Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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### **Marketing authorisation holder:**

Ceva Sante Animale

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### **Marketing authorisation date:**

30/08/2022

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### **Manufacturing sites for batch release:**

Ceva Sante Animale

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### **Responsible authority:**

Federal Agency For Medicines And Health Products

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### **Authorisation number:**

This information is not available for this product.

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### **Date of authorisation status change:**

30/08/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0429/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark 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.

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