# 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 SOLUTION FOR USE IN DRINKING WATER/ MILK FOR PRE-RUMINANT CATTLE AND PIGS
GABBROVET MULTI 140 mg/ml soluție pentru utilizare în apa de băut sau lapte pentru bovinele pre-rumegătoare și porci

**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:**

- **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.

- **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:**
Romania

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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

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Ceva Sante Animale Romania S.R.L.

**Marketing authorisation date:**
19/06/2022

**Manufacturing sites for batch release:**
Ceva Sante Animale

**Responsible authority:**
Institute For Control Of Biological Products And Veterinary Medicines

**Authorisation number:**
220092

**Date of authorisation status change:**

19/06/2022

**Reference member state:**
France

**Procedure number:**
FR/V/0429/001

**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)

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

## Documents

Summary of Product Characteristics

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