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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

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)

Pharmaceutical form:
Solution for use in drinking water/milk

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA06

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

Authorisation status:
Valid

Authorised in:
Netherlands

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

Marketing authorisation date:
19/07/2022

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 128521

Date of authorisation status change:
28/03/2024

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)

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Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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