

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 roztwór do podania w wodzie do picia/mleku dla bydła z nierozwiniętą funkcją przedżołądków i świń

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:

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Cattle (calf)

- Meat and offal. 20 day

Cattle (pre-ruminant cattle and newborn calves): - Colibacillosis: Dosage: 25-50 mg/kg/day for 3 to 5 days. Meat and offal: 20 days

- Meat and offal. 110 day

Cattle (pre-ruminant cattle and newborn calves): - Cryptosporidiosis: Dosage: 150 mg/kg/day for 5 days. Meat and offal: 110 days

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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:

Poland

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 Animal Health Polska Sp. z o.o.

Marketing authorisation date:

8/08/2022

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3203

Date of authorisation status change:

8/08/2022

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000096461>