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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:
Slovenia

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:
24/08/2022

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
DC/V/0614/002

Date of authorisation status change:
24/08/2022

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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

Combined File of all Documents

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