File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000031087

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

Authorised

- Paromomycin sulfate

Product identification

Medicine name:
GABBROVET 140 MG/ML SOLUTION FOR USE IN DRINKING WATER/MILK FOR PRE-RUMINANT CATTLE AND PIGS

Active substance:
Paromomycin sulfate

Target species:
Cattle (pre-ruminant)
Pig

Route of administration:
In drinking water/milk use
In drinking water 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:
In drinking water/milk use:

- **Cattle (pre-ruminant)**
  - Meat and offal. 20 day

In drinking water use:

- **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:
France

Available in:
France

Package description:
Available only in French

Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
2/03/2018

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5151167 8/2018

Date of authorisation status change:
10/01/2023

Reference member state:
France

Procedure number:
FR/V/0317/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia 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

eu-puar-frv0317001-mr-rpe364-en.pdf