File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000085988

Gabbrovet 140 140 mg/g Poeder voor toediening in het drinkwater/in de melk

Authorised

- Paromomycin

Product identification

Medicine name:
Gabbrovet 140 140 mg/g Poeder voor toediening in het drinkwater/in de melk

Active substance:
Paromomycin

Target species:
Cattle (pre-ruminant)
Pig

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Paromomycin
140.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder 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
- **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:
Belgium

Package description:
Gabbrovet 140 1 kg Jar Powder for use in drinking water/milk

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
14/05/1985

Manufacturing sites for batch release:
Unione Commerciale Lombarda S.p.A.
Ceva Vetem S.p.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V131616

Date of authorisation status change:
25/02/2019

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

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

Labelling

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