File downloaded on 2026-11-10
Source URL: https://medicines.health.europa.eu/veterinary/en/600000053044

Parofor crypto 140 000 IU/ml oral solution for sheep and goats

Authorised

- Paromomycin sulfate

Product identification

Medicine name:
Parofor crypto 140 000 IU/ml oral solution for sheep and goats

Active substance:
Paromomycin sulfate

Target species:
Goat
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Paromomycin sulfate
162310.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Goat**
  - Meat and offal. 24 day
- **Sheep**
  - Meat and offal. 24 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Greece

Package description:

White high density polyethylene bottle with tamper-evident screw polypropylene closure. Bottle size:125 ml
White high density polyethylene bottle with tamper-evident screw polypropylene closure. Bottle size:250 ml
White high density polyethylene bottle with tamper-evident screw polypropylene closure. Bottle size:500 ml
White high density polyethylene bottle with tamper-evident screw polypropylene closure. Bottle size:1 L.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
15/07/2019

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
National Organization For Medicines

Authorisation number:
129450/21-10-2026/K-0241201

Date of authorisation status change:
20/10/2026

Reference member state:
Ireland

Procedure number:
IE/V/0412/001

Concerned member states:
Austria Bulgaria Croatia Cyprus Czechia Denmark Estonia Greece Hungary Italy Latvia Lithuania Luxembourg Malta 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

English (PDF)
Published on: 3/05/2024
Updated on: 14/03/2026
Download