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# 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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA07AA06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Bulgaria

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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

**Marketing authorisation holder:**
HuVepharma

**Marketing authorisation date:**
7/07/2019

**Manufacturing sites for batch release:**
Biovet AD

**Responsible authority:**
Bulgarian Food Safety Authority

**Authorisation number:**
0022-2904

**Date of authorisation status change:**
7/07/2019

**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
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Package Leaflet

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Labelling

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