**Source URL:** <a href="https://medicines.health.europa.eu/veterinary/en/600000051336">https://medicines.health.europa.eu/veterinary/en/600000051336</a>

# Parofor crypto 140 000 IU/ml oral solution for pre-ruminant cattle

Authorised

• Paromomycin

# Product identification

#### **Medicine name:**

Parofor crypto 140 000 IU/ml oral solution for pre-ruminant cattle Parofor crypto 140 000 IU/ml oral solution for pre-ruminant cattle

#### **Active substance:**

Paromomycin

## **Target species:**

Cattle

#### Route of administration:

Oral use

# **Product details**

# **Active substance and strength:**

Paromomycin

140000.00 international unit(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

Oral solution

# Withdrawal period by route of administration:

#### Oral use:

**Cattle** 

- Meat and offal. 62 day

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

**QA07AA06** 

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### Authorised in:

Malta

## Package description:

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

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:250 ml

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

# 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:

14/09/2023

# Manufacturing sites for batch release:

**Biovet AD** 

## **Responsible authority:**

Veterinary And Phytosanitary Regulation Department

#### **Authorisation number:**

**VMA 120** 

# Date of authorisation status change:

14/09/2023

## **Reference member state:**

Ireland

## **Procedure number:**

IE/V/0610/001

#### **Concerned member states:**

Austria Bulgaria Croatia Cyprus Czechia Estonia Greece Hungary Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>