

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

Parofor crypto 140 000 TV/ml, geriamasis tirpalas avims ir ožkoms

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

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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:

Lithuania

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:

6/08/2019

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/19/2545/001-004

Date of authorisation status change:

6/08/2019

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

RV2545.pdf