

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

Authorised

- Paromomycin

Product identification

Medicine name:

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

Paroform crypto 140 000 IU/ml πόσιμο διάλυμα για βοοειδή πριν την έναρξη λειτουργίας της μεγάλης κοιλίας

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:

Cyprus

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:

14/04/2019

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00737V

Date of authorisation status change:

14/04/2019

Reference member state:

Ireland

Procedure number:

IE/V/0610/001

Concerned member states:

Austria Bulgaria Croatia Cyprus Czechia Denmark Estonia France Germany
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
www.adrreports.eu/vet

Documents

Combined File of all Documents

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