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 IE/ml Lösung zum Eingeben für Saugkälber

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:
Medicinal product on medical prescription for non-renewable delivery

Authorisation status:
Valid

Authorised in:
Austria

Available in:
Austria

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:
30/04/2019

Manufacturing sites for batch release:
Biovet J.S.C.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
838890

Date of authorisation status change:
30/04/2019

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

Labelling

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

Package Leaflet

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

Summary of Product Characteristics

English (PDF)
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