Parofor 70000 IU/g Powder for use in drinking water/milk

Authorised

- Paromomycin

Product identification

Medicine name:
Parofor 70000 IU/g Powder for use in drinking water/milk
Parofor 70000 IU/g Poeder voor toediening in het drinkwater/in de melk
Parofor 70000 IU/g Poudre pour administration dans le lait ou l'eau de boisson
Parofor 70000 IU/g Pulver zum Eingeben über das Trinkwasser/die Milch

Active substance:
Paromomycin

Target species:
Cattle
Pig

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
Paromomycin
70000.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:
In drinking water/milk use:

- **Cattle**
  - Meat and offal. 20 day — 20 days for pre-ruminant cattle
- **Pig**
  - Meat and offal. 3 day — 3 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
1000g: sachet (PE/ALU/PET) with 1000g powder
500g: sachet (PE/ALU/PET) with 500g powder
250g: sachet (PE/ALU/PET) with 250g powder
25g: box (cardboard) with 40 sachets (PE/ALU/PP) each with 25g powder

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
28/08/2014

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
28/08/2014

Reference member state:
Belgium

Procedure number:
BE/V/0027/001

Concerned member states:
Austria Bulgaria Cyprus Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

Generic of:
600000085987

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Labelling

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