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 Powder for use in drinking water/milk
PAROFOR 70000 UI/G POUDRE POUR ADMINISTRATION DANS L'EAU DE BOISSON / LE LAIT POUR BOVINS (PRERUMINANTS) ET PORCS

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:
France

Available in:
France

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:
10/09/2014

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/8668669 1/2014

Date of authorisation status change:
24/07/2019

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

Summary of Product Characteristics

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

Package Leaflet and Labelling

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