COLDOSTIN 4 800 000 UI/G POWDER FOR ADMINISTRATION IN DRINKING WATER/MILK

Authorised

- COLISTIN SULFATE

Product identification

Medicine name:
COLDOSTIN 4 800 000 UI/G POWDER FOR ADMINISTRATION IN DRINKING WATER/MILK
Coldostin 4800000 IU/g Poeder voor toediening in het drinkwater/in de melk
Coldostin 4800000 IU/g Poudre pour administration dans le lait ou l'eau de boisson
Coldostin 4800000 IU/g Pulver zum Eingeben über das Trinkwasser/die Milch

Active substance:
COLISTIN SULFATE

Target species:
Turkey
Pig
Sheep (lamb)
Cattle (calf)
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
COLISTIN SULFATE
4800000.00 international unit(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:
Oral use:

- **Turkey**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day
- **Sheep (lamb)**
  - Meat and offal. 1 day
- **Cattle (calf)**
  - Meat and offal. 1 day
- **Chicken**
  - Meat and offal. 1 day
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Composite can of 1 kg containing a measuring spoon (3 g)
Securitainer of 1 kg
Securitainer of 100 g
Bucket of 1 kg

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
13/01/2017

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

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

Date of authorisation status change:
5/01/2022

Reference member state:
France

Procedure number:
FR/V/0299/001

Concerned member states:
Belgium Denmark Germany Hungary Italy Lithuania Netherlands Poland United Kingdom (Northern Ireland)

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

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.
