

# 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)

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### Pharmaceutical form:

Powder for use in drinking water/milk

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### Withdrawal period by route of administration:

#### Oral use:

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#### **Turkey**

- Meat and offal. 1 day

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#### **Pig**

- Meat and offal. 1 day

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#### **Sheep (lamb)**

- Meat and offal. 1 day

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#### **Cattle (calf)**

- Meat and offal. 1 day

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#### **Chicken**

- Meat and offal. 1 day

- Eggs. 0 day

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### Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA07AA10

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### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Belgium

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**Available in:**

Belgium

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Dopharma Research B.V.

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**Marketing authorisation date:**

13/01/2017

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**Manufacturing sites for batch release:**

Dopharma B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

5/01/2022

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**Reference member state:**

France

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**Procedure number:**

FR/V/0299/001

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**Concerned member states:**

Belgium Denmark Germany Hungary Italy Lithuania Netherlands Poland  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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.