VIRGOCILLINE, soluție injectabilă

Not authorised

- COLISTIN SULFATE

Product identification

Medicine name:
VIRGOCILLINE, soluție injectabilă

Active substance:
COLISTIN SULFATE

Target species:
Horse
Sheep
Pig
Rabbit
Chicken
Cattle

Route of administration:
Intramuscular use
Subcutaneous use
Intraperitoneal use

Product details

Active substance and strength:
COLISTIN SULFATE
500000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Horse**
  - Meat and offal. 21 day
- **Sheep**
  - Milk. 2 day
  - Meat and offal. 21 day
- **Pig**
  - Meat and offal. 21 day
- **Rabbit**
  - Meat and offal. 21 day
- **Chicken**
  - Meat and offal. 21 day

nu se utilizeaza in timpul perioadei de ouat pentru oua consum uman

- **Cattle**
  - Milk. 2 day

- Meat and offal. 21 day

Subcutaneous use:

- **Sheep**
  - Meat and offal. 21 day
  - Milk. 2 day
- **Pig**
  - Meat and offal. 21 day
- **Horse**
  - Meat and offal. 21 day
- **Rabbit**
  - Meat and offal. 21 day
- **Chicken**
  - Meat and offal. 21 day

nu este permisa utiizarea in perioada de ouat pentru oua consum uman

- **Cattle**
  - Milk. 2 day
  - Meat and offal. 21 day

Intraperitoneal use:

- **Sheep**
  - Meat and offal. 21 day
  - Milk. 2 day
-

Pig

- Meat and offal. 21 day

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Horse

- Meat and offal. 21 day

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Rabbit

- Meat and offal. 21 day

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Chicken

- Meat and offal. 2 day

nu este permisa utilizarea in perioada de ouat pentru oua consum uman

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Cattle

- Milk. 2 day
- Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01XB01

Legal status of supply:
This information is not available for this product.

Authorisation status:
Surrendered

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian

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:
Dopharma B.V.

Marketing authorisation date:
8/06/2006

Manufacturing sites for batch release:
Dopharma France

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
190237

Date of authorisation status change:
4/06/2024

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.