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Colfive 5,000,000 IU/ml solution for use in drinking water/milk for calves, pigs, lambs, chickens and turkeys

Authorised

- COLISTIN SULFATE

Product identification

Medicine name:
Colfive 5,000,000 IU/ml solution for use in drinking water/milk for calves, pigs, lambs, chickens and turkeys
Colfive 5 000 000 IU/ml koncentrát na perorálny roztok pre teľatá, ošípané, jahňatá, kurčatá a morky

Active substance:
COLISTIN SULFATE

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

Route of administration:
In drinking water/milk use

Oral use

Product details

Active substance and strength:
COLISTIN SULFATE
5000000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for oral solution

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

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

Oral use:

-

Pig

- Meat and offal. 1 day

•

Turkey

- Meat and offal. 1 day
- Eggs. 0 day

•

Cattle (calf)

- Meat and offal. 1 day

•

Sheep (lamb)

- 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

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Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

bottle of 5 l

bottle of 1 l

box containing 1 bottle of 100 ml

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
26/06/2015

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
aniMedica GmbH

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/039/DC/15-S

Date of authorisation status change:
26/06/2015

Reference member state:
Spain

Procedure number:
ES/V/0221/001

Concerned member states:
Austria Belgium Czechia France Germany Greece Hungary Ireland Italy Netherlands Poland Portugal Romania Slovakia Slovenia

United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
Published on: 24/07/2025
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Summary of Product Characteristics

Package Leaflet

Labelling