APSALIQ COLISTIN 3 000 000 IU/ml solution for use in drinking water/milk for pig, cattle, sheep, chickens and turkeys

Authorised

- COLISTIN SULFATE

Product identification

Medicine name:
APSALIQ COLISTIN 3 000 000 IU/ml solution for use in drinking water/milk for pig, cattle, sheep, chickens and turkeys

Active substance:
COLISTIN SULFATE

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

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:
COLISTIN SULFATE
3000000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water/milk

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
- **Chicken**
  - Meat and offal. 1 day
  - Eggs. 0 day
- **Cattle (calf)**
  - Meat and offal. 1 day
  - Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

- **Sheep (lamb)**
  - Meat and offal. 1 day

- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA10

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Package description:
bottle of 1 L
bottle of 5 L

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:
Andres Pintaluba S.A.

Marketing authorisation date:
31/01/2018

Manufacturing sites for batch release:
Andres Pintaluba S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2738

Date of authorisation status change:
31/01/2018

Reference member state:
Spain

Procedure number:
ES/V/0270/001

Concerned member states:
Bulgaria Croatia Cyprus Czechia Greece Hungary Italy Poland Portugal Romania Slovakia

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 22/12/2023
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Package Leaflet

English (PDF)
Published on: 22/12/2023

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Labelling

Combined File of all Documents

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