COLIPRO 2 000 000 IU/ML CONCENTRATE FOR ORAL SOLUTION FOR PIGS AND POULTRY

Not authorised

- COLISTIN SULFATE

Product identification

Medicine name:
COLIPRO 2 000 000 IU/ML CONCENTRATE FOR ORAL SOLUTION FOR PIGS AND POULTRY

Active substance:
COLISTIN SULFATE

Target species:
Poultry
Pig

Route of administration:
Oral use

Product details

Active substance and strength:

COLISTIN SULFATE
2.00 million international units / 1.00 millilitre(s)

Pharmaceutical form:
Concentrate for oral solution

Withdrawal period by route of administration:
Oral use:

- **Poultry**
  - Meat and offal. 1 day
  - Eggs. 0 day
- **Pig**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA10

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

Authorisation status:
Surrendered

Authorised in:
Austria

Package description:
250 ml high density polyethylene bottle with polypropylene screw cap fitted with a polypropylene dosing device and a polyethylene seal.
5 litres high density polyethylene bottle with polypropylene screw cap fitted with a polypropylene dosing device and a polyethylene seal.
2 litres high density polyethylene bottle with polypropylene screw cap fitted with a polypropylene dosing device and a polyethylene seal.
1 litre high density polyethylene bottle with polypropylene screw cap fitted with a polypropylene dosing device and a polyethylene seal.

500 ml high density polyethylene bottle with polypropylene screw cap fitted with a polypropylene dosing device and a polyethylene seal.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
28/08/2007

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-00710

Date of authorisation status change:
22/05/2025

Reference member state:
France

Procedure number:
FR/V/0184/001

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

Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

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.

Combined File of all Documents