CADOREX 100 mg/ml SOLUTION FOR USE IN DRINKING WATER FOR PIGS AND CHICKENS

Authorised

- Florfenicol

Product identification

Medicine name:
CADOREX 100 mg/ml SOLUTION FOR USE IN DRINKING WATER FOR PIGS AND CHICKENS

Active substance:
Florfenicol

Target species:
Pig
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Florfenicol
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

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

- **Pig**
  - Meat and offal. 20 day

- **Chicken**
  - Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption.

- Meat and offal. 8 day
- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption.

- Meat and offal. 8 day

- **Chicken**
  - Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption.

- Meat and offal. 8 day
- Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption.

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01BA90

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

Authorisation status:
Valid

Authorised in:
Poland

Package description:
High density polyethylene (HDPE) containers with a sealing foil made of paper/Alu/PET/PE and HDPE) screw cap.
High density polyethylene (HDPE) containers with a sealing foil made of paper/Alu/PET/PE and HDPE) screw cap.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application - change in active substance(s) (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Industrial Veterinaria S.A.

Marketing authorisation date:
24/11/2025

Manufacturing sites for batch release:
Industrial Veterinaria S.A.
aniMedica GmbH
aniMedica Herstellungs GmbH
Industria Italiana Integratori Trei S.p.A.

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

Authorisation number:
3457

Date of authorisation status change:
24/11/2025

Reference member state:
Spain

Procedure number:
ES/V/0445/001

Concerned member states:
Czechia Greece Hungary Italy Poland Portugal Romania

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 19/02/2026
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Package Leaflet and Labelling

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

Package Leaflet

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