

# CADOREX 300 mg/ml Solution for injection for cattle, pigs and sheep

Authorised

- Florfenicol

## Product identification

**Medicine name:**

CADOREX 300 mg/ml Solution for injection for cattle, pigs and sheep

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**Active substance:**

Florfenicol

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**Target species:**

Cattle  
Sheep  
Pig

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**Route of administration:**

Intramuscular use  
Subcutaneous use

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## Product details

**Active substance and strength:**

Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

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

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**Sheep**

- Meat and offal. no withdrawal period      Meat and offal: im 39 Days

- Milk. no withdrawal period

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

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**Pig**

- Meat and offal. no withdrawal period

Meat and offal: Intramuscular use: 18 days

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

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

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**Sheep**

- Meat and offal. no withdrawal period Meat and offal: im 39 Days
- Milk. no withdrawal period

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

**Subcutaneous use:**

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

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

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**Cattle**

- Meat and offal. no withdrawal period

Meat and offal: IM: 30 dAYS / SC: 44 Days

- Milk. no withdrawal period

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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01BA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Romania

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**Available in:**

Romania

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**Package description:**

box containing 1 vial of 250 ml  
box containing 1 vial of 100 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Industrial Veterinaria S.A.

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**Marketing authorisation date:**

11/09/2017

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.  
aniMedica GmbH

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

220063

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**Date of authorisation status change:**

29/01/2026

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0246/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland

France Germany Greece Hungary Ireland Italy Latvia Lithuania  
Netherlands Poland Portugal Romania Slovakia Slovenia  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Package Leaflet

Labelling

Combined File of all Documents

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