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# 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

**Active substance:**
Florfenicol

**Target species:**
Cattle
Sheep
Pig

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
Florfenicol

300.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**
**Intramuscular use:**

- **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.

- **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.

- **Pig**
  - Meat and offal. no withdrawal period

Meat and offal: Intramuscular use: 18 days

- **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.

- **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:**

- **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.

- **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.

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01BA90

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

**Authorisation status:**
Valid

**Authorised in:**
Czechia

**Package description:**
box containing 1 vial of 100 ml
box containing 1 vial of 250 ml

## 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:**
Industrial Veterinaria S.A.

**Marketing authorisation date:**
26/01/2018

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

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

**Authorisation number:**
96/006/18-C

**Date of authorisation status change:**
25/04/2022

**Reference member state:**
Spain

**Procedure number:**
ES/V/0246/001

**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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## Documents

Summary of Product Characteristics

English (PDF)
Published on: 20/10/2025
Updated on: 13/03/2026
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Package Leaflet

English (PDF)
Published on: 20/10/2025
Updated on: 13/03/2026
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Labelling

English (PDF)
Published on: 20/10/2025
Updated on: 13/03/2026
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