

# 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

CADOREX 300 mg/ml ενέσιμο διάλυμα για βοοειδή, χοίρους και πρόβατα

### 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

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**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
- IM 18 days

**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.

**• 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**

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

QJ01BA90

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**Legal status of supply:**

Medicinal product subject to medical prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Available in:**

Greece

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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:**

Livisto Int'l S.L.

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**Marketing authorisation date:**

22/08/2017

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**Manufacturing sites for batch release:**

Industrial Veterinaria S.A.

Animedica GmbH

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

59996/15/23-08-2017/K-0218401

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**Date of authorisation status change:**

6/06/2021

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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

eu-PUAR-esv0246001-dcp-cadorex-300-mg-ml-solution-for-injection-for-cattle--pigs-and-sheep-en.pdf

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