

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:

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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.

•

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.

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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.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01BA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

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

22/08/2017

Manufacturing sites for batch release:

Industrial Veterinaria S.A.
aniMedica GmbH

Responsible authority:

National Organization For Medicines

Authorisation number:

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

Date of authorisation status change:

6/06/2021

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)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 20/10/2025

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Package Leaflet

English (PDF)

Published on: 20/10/2025

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Labelling

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