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

Authorised



## Product identification

## Medicine name:

CADOREX 300 mg/ml Solution for injection for cattle, pigs and sheep Cadorex 300 mg/ml injekčný roztok pre hovädzí dobytok, ovce a ošípané

## **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 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

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

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

## Authorisation status:

Valid

Authorised in: Slovakia

## **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:

Livisto Int'l S.L.

## Marketing authorisation date:

4/08/2017

## Manufacturing sites for batch release:

Industrial Veterinaria S.A. Animedica GmbH

## **Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

# Authorisation number: 96/035/MR/17-S

## Date of authorisation status change:

6/04/2023

#### **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 <u>www.adrreports.eu/vet</u>

## Documents

Combined File of all Documents

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

Package Leaflet

Summary of Product Characteristics

Labelling

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

**Source URL:** https://medicines.health.europa.eu/veterinary/60000039727