# 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 инжекционен разтвор за говеда, овце и прасета

#### **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

<b>Authorisation status:</b> Valid	
<b>Authorised in:</b> Bulgaria	
<b>Available in:</b> Bulgaria	
Package description: box containing 1 vial of 250 ml box containing 1 vial of 100 ml	
Additional information	
<b>Entitlement type:</b> Marketing Authorisation	
<b>Legal basis of product author</b> Generic application (Article 13(1)	
Marketing authorisation hold Livisto Int'l S.L.	er:
Marketing authorisation date 23/04/2017	:

# Responsible authority:

Animedica GmbH

Bulgarian Agency For Food Safety

# **Authorisation number:**

0022-2727

#### Date of authorisation status change:

29/05/2022

#### **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)

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

#### **Documents**

Summary of Product Characteristics

English (PDF)

Published on: 22/12/2023

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Package Leaflet and Labelling

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Package Leaflet
Labelling
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