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
Flodoex 300 mg/ml Oplossing voor injectie
Flodoex 300 mg/ml Solution injectable
Flodoex 300 mg/ml Injektionslösung

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

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

Meat and offal: im 39 Days

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

Meat and offal: Intramuscular use: 18 days

- **Cattle**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

- **Sheep**
  - 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

- **Cattle**
  - 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:
Belgium

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:
13/11/2017

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V519546

Date of authorisation status change:
13/11/2017

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

English (PDF)
Published on: 22/12/2023
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Labelling

English (PDF)
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