Luteosyl 0.075 mg/ml Solution for injection for cows and sows

Authorised

- D-cloprostenol

Product identification

Medicine name:
Luteosyl 0.075 mg/ml Solution for injection for cows and sows

Active substance:
D-cloprostenol

Target species:
Cattle (cow)
Pig (sow)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
D-cloprostenol
0.08 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle (cow)**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Pig (sow)**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
cardboard box containing 5 glass vials of 20 ml
cardboard box containing 1 glass vial of 20 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:
Laboratorios Syva S.A.

Marketing authorisation date:
11/07/2018

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
86392/22-09-2021/K-0233101

Date of authorisation status change:
14/06/2023

Reference member state:
Spain

Procedure number:
ES/V/0143/001

Concerned member states:
Belgium Bulgaria Czechia France Germany Greece Hungary Italy Netherlands Poland Portugal Romania Slovakia

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: 30/10/2025
Download

Package Leaflet

English (PDF)
Published on: 30/10/2025
Download

Labelling

Combined File of all Documents

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