

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

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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:

Bulgaria

Available in:

Bulgaria

Package description:

cardboard box containing 1 glass vial of 20 ml

cardboard box containing 5 glass vials 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:

5/06/2016

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2653

Date of authorisation status change:

5/06/2016

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

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Labelling

English (PDF)

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Package Leaflet

English (PDF)

Published on: 30/10/2025

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Combined File of all Documents

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