

# VETEGLAN 0.075 mg/ml Solution for injection for cows, sows and mares

Authorised

- Cloprostenol sodium

## Product identification

**Medicine name:**

VETEGLAN 0.075 mg/ml Solution for injection for cows, sows and mares

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**Active substance:**

Cloprostenol sodium

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**Target species:**

Cattle (cow)

Pig

Horse (mare)

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Cloprostenol sodium

0.08 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle (cow)**

- Meat and offal. 0 day
- Milk. 0 hour

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**Pig**

- Meat and offal. 1 day

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**Horse (mare)**

- Meat and offal. 2 day
  - Milk. 0 hour
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG02AD90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Lithuania

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**Package description:**

20 ml amber coloured Type I glass vials, with Teflon-coated chlorobutyl rubber closures and aluminium seals with blue coloured plastic flip-offs, packaged singly in a cardboard box.

10 ml amber coloured Type I glass vials, with Teflon-coated chlorobutyl rubber closures and aluminium seals with blue coloured plastic flip-offs, packaged singly in a cardboard box.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Calier S.A.

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**Marketing authorisation date:**

1/02/2024

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**Manufacturing sites for batch release:**

Laboratorios Calier S.A.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/24/2796/001-002

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**Date of authorisation status change:**

1/02/2024

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**Reference member state:**

Portugal

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**Procedure number:**

PT/V/0100/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Denmark France Germany Greece  
Hungary Ireland Latvia Lithuania Netherlands Romania Spain  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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Combined File of all Documents