# 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
VETEGLAN 0,075 MG/ML SOLUTION INJECTABLE POUR VACHES TRUIES ET JUMENTS

**Active substance:**
Cloprostenol sodium

**Target species:**
Cattle (cow)
Pig
Horse (mare)

**Route of administration:**
Solution for injection

## Product details

**Active substance and strength:**
Cloprostenol sodium
0.08 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Solution for injection:**

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 1 day
- **Horse (mare)**
  - Meat and offal. 2 day
  - Milk. 0 hour

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QG02AD90

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

**Authorisation status:**
Valid

**Authorised in:**
France

**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.

## 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 Calier S.A.

**Marketing authorisation date:**
18/09/2020

**Manufacturing sites for batch release:**
Laboratorios Calier S.A.

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**
FR/V/6536868 4/2020

**Date of authorisation status change:**
16/12/2021

**Reference member state:**
Portugal

**Procedure number:**
PT/V/100/001

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

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

## Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

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