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# Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

Authorised

- Cloprostenol

## Product identification

**Medicine name:**
Syncroprost, 0.250 mg/ml solution for injection for cattle, horses, pigs and goats

**Active substance:**
Cloprostenol

**Target species:**
Cattle (cow)
Cattle (heifer)
Horse (mare)
Pig (female)
Goat (adult female)

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**

Cloprostenol
0.25 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 day
- **Cattle (heifer)**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Horse (mare)**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Pig (female)**
  - Meat and offal. 1 day
- **Goat (adult female)**
  - Meat and offal. 1 day
  - Milk. 0 day

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

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

**Authorisation status:**
Valid

**Authorised in:**
Luxembourg

**Available in:**
Luxembourg

**Package description:**
Box with one 50 ml vial. Type I colourless glass vial sealed with bromo-butyl rubber stopper closed by aluminium flip-off cap.
Box with one 100 ml vial. Type I colourless glass vial sealed with bromo-butyl rubber stopper closed by aluminium flip-off cap.
Box with one 10 ml vial. Type I colourless glass vial sealed with bromo-butyl rubber stopper closed by aluminium flip-off cap.
Box with one 20 ml vial. Type I colourless glass vial sealed with bromo-butyl rubber stopper closed by aluminium flip-off cap.
Box with 10 x 20 ml vials. Type I colourless glass vials sealed with bromo-butyl rubber stoppers closed by aluminium flip-off caps.

## 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:**
Ceva Sante Animale

**Marketing authorisation date:**
10/02/2023

**Manufacturing sites for batch release:**
Vetem S.p.A.

**Responsible authority:**

Ministry Of Health And Social Security

**Authorisation number:**
V 915/09/02/2023

**Date of authorisation status change:**
10/02/2023

**Reference member state:**
Italy

**Procedure number:**
IT/V/0147/001

**Concerned member states:**
Austria Belgium Bulgaria Czechia Denmark Estonia Finland France Germany Hungary Ireland Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Slovakia Spain Sweden

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

## Documents

Package Leaflet

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Combined File of all Documents