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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:
Norway

Available in:
Norway

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 10 x 20 ml vials. Type I colourless glass vials sealed with bromo-butyl rubber stoppers closed by aluminium flip-off caps.
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.

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:
15/07/2022

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

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:
20-13808

Date of authorisation status change:
15/07/2022

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

Combined File of all Documents

Summary of Product Characteristics

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Package Leaflet

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