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
Syncroprost, 0,250 mg/ml, soluzione iniettabile per bovini, cavalli, suini e capre

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:
Italy

Package description:
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 20 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 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.

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 Salute Animale S.p.A.

Marketing authorisation date:
25/05/2022

Manufacturing sites for batch release:

Vetem S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/05/2022

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

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Documents

Combined File of all Documents

English (PDF)
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