

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, 250 microgram/ml oplossing voor injectie

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:

Netherlands

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 Sante Animale B.V.

Marketing authorisation date:

10/05/2022

Manufacturing sites for batch release:

Vetem S.p.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 127799

Date of authorisation status change:

10/05/2022

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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www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

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