Enzaprost Bovis 12.5 mg/ml solution for injection for cattle

Not authorised

- Dinoprost trometamol

Product identification

Medicine name:
Enzaprost Bovis 12.5 mg/ml solution for injection for cattle

Active substance:
Dinoprost trometamol

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Dinoprost trometamol
12.50 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Surrendered

Authorised in:

Finland

Package description:

Cardboard box containing 1 translucent multi layer (polypropylene _ethylene vinyl alcohol _polypropylene) vial of 100 closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule
Cardboard box containing 1 colourless type I glass vial of 20 ml closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule
Cardboard box containing 1 colourless type I glass vial of 2 ml closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule
Cardboard box containing 1 colourless type I glass vial of 10 ml closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule
Cardboard box containing 1 translucent multi layer (polypropylene _ethylene vinyl alcohol _polypropylene) vial of 50 closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule
Cardboard box containing 10 colourless type I glass vial of 2 ml closed with a bromobutyl rubber stopper and aluminium and plastic flip capsule

---

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:
19/09/2019

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Finnish Medicines Agency

Authorisation number:
36317

Date of authorisation status change:
11/12/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0256/001

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