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
Enzaprost Bovis 12,5 mg/ml Injektionslösung für Rinder

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:
Medicinal product on medical prescription for non-renewable delivery

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Authorisation status:
Revoked

Authorised in:
Austria

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

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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:
5/02/2021

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
840506

Date of authorisation status change:
5/02/2021

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

Documents

Labelling

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

Package Leaflet

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

Summary of Product Characteristics

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