

Bioestrovet 0.250 mg/ml solution for injection for cattle

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

Bioestrovet 0.250 mg/ml solution for injection for cattle

Active substance:

Cloprostenol sodium

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Cloprostenol sodium

0.26 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Intramuscular use:**

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Cattle

- Meat and offal. 1 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

Package description:

Type 1 (colourless) glass vial closed with bromobutyl rubber stopper coated with a FluroTec film (ETFE) and a polypropyleneflip-off cap.Pack size:Box with 1 vial of 20 ml

Type 1 (colourless) glass vial closed with bromobutyl rubber stopper coated with a FluroTec film (ETFE) and a polypropyleneflip-off cap.Pack size:Box with 1 vial of 50 ml

Type 1 (colourless) glass vial closed with bromobutyl rubber stopper coated with a FluroTec film (ETFE) and a polypropyleneflip-off cap.Pack size:Box with 1 vial of 100 ml

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:

Vetoquinol S.A.

Marketing authorisation date:

3/05/2017

Manufacturing sites for batch release:

Vetoquinol S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

36015/07-04-2022/K-0220201

Date of authorisation status change:

6/04/2022

Reference member state:

Ireland

Procedure number:

IE/V/0359/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France
Germany Greece Hungary Italy Latvia Lithuania Luxembourg Malta
Netherlands Poland Portugal Romania Slovakia Slovenia Spain
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents