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:

- **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:
Poland

Available in:
Poland

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 Biowet Sp. z o.o.

Marketing authorisation date:
2/08/2017

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2683

Date of authorisation status change:
2/08/2017

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

English (PDF)
Published on: 28/09/2025
Download

Package Leaflet

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

Labelling

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

Combined File of all Documents