Estrumate 250 micrograms/ml solution for injection

Not authorised

- Cloprostenol sodium

Product identification

Medicine name:
Estrumat vet. 0,25 mg/ml stungulyf, lausn, handa nautgripum, svínum og hestum
Estrumate 250 micrograms/ml solution for injection

Active substance:
Cloprostenol sodium

Target species:
Cattle
Equid
Goat

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 day
- **Equid**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Goat**
  - 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:
Surrendered

Authorised in:
Iceland

Package description:
1 x 10 ml bottle
1 x 20 ml bottle

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
1/01/1991

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
880024

Date of authorisation status change:
13/09/2011

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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Combined File of all Documents

English (PDF)
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