

Oestrophan 0,25 mg/ml injekčný roztok

Authorised

- Cloprostenol sodium

Product identification

Medicine name:

Oestrophan 0,25 mg/ml injekčný roztok

Active substance:

Cloprostenol sodium

Target species:

Cattle (cow)
Cattle (heifer)
Horse (mare)
Pig (sow)

Route of administration:

Submucosal use
Intramuscular use

Product details

Active substance and strength:

Cloprostenol sodium
263.16 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

-

Cattle (cow)

- Meat. 1 day Milk - Zero hours

-

Cattle (heifer)

- Meat. 1 day

Intramuscular use:

-

Cattle (cow)

- Meat. 1 day Milk - Zero hours

-

Cattle (heifer)

- Meat. 1 day

-

Horse (mare)

- Meat. 1 day Milk - Zero hours

-

Pig (sow)

- Meat. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Available only in Slovak

Available only in Slovak

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

23/12/1994

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

99/238/80-C/S

Date of authorisation status change:

23/12/1994

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

Documents

Combined File of all Documents

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