Gabbrostim 2 mg/ml Oplossing voor injectie

Authorised

- Alfaprostol

Product identification

Medicine name:
Gabbrostim 2 mg/ml Oplossing voor injectie

Active substance:
Alfaprostol

Target species:
Horse
Pig
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Alfaprostol
2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Horse**
  - Meat and offal. 24 hour

Animals aimed for human consumption can only be slaughtered 24h after last treatment.

- Milk. 24 hour

Milk from treated animals can only be used for human consumption 24h after treatment.

- **Pig**
  - Meat and offal. 24 hour

Animals aimed for human consumption can only be slaughtered 24h after last treatment.

- **Cattle**
  - Meat and offal. 24 hour

Animals aimed for human consumption can only be slaughtered 24h after last treatment.

- Milk. 24 hour

Milk from treated animals can only be used for human consumption 24h after treatment.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD94

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:
Belgium

Package description:
Gabbrostim 12 Vials with 50 ml of Solution for injection
Gabbrostim 12 Vials with 20 ml of Solution for injection
Gabbrostim 12 Vials with 4 ml of Solution for injection
Gabbrostim 1 Vial with 50 ml of Solution for injection
Gabbrostim 1 Vial with 20 ml of Solution for injection
Gabbrostim 1 Vial with 4 ml of Solution for injection

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
27/06/1994

Manufacturing sites for batch release:
Vetem SPA
CEVA Santé Animale

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V166101

Date of authorisation status change:
3/07/2019

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

Documents

Summary of Product Characteristics

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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.