

# Gonavet Veyx 50 µg/ml solution for injection for cattle, pigs and horses

Authorised

- Gonadorelin (6-D-phenylalanine) acetate

## Product identification

### **Medicine name:**

Gonavet Veyx 50 µg/ml solution for injection for cattle, pigs and horses

### **Active substance:**

Gonadorelin (6-D-phenylalanine) acetate

### **Target species:**

Cattle

Pig

Horse

### **Route of administration:**

Subcutaneous use

Intramuscular use

## Product details

### **Active substance and strength:**

Gonadorelin (6-D-phenylalanine) acetate

52.40 microgram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

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

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**Pig**

- Meat and offal. 0 day

**Intramuscular use:**

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**Cattle**

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

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**Horse**

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

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**Pig**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH01CA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Luxembourg

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**Available in:**

Luxembourg

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**Package description:**

(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)

(ID2) 20 millilitre(s): unspecified outer container with 1 Vial (glass) with 20 millilitre(s)

(ID1) 10 millilitre(s): unspecified outer container with 1 Vial (glass) with 10 millilitre(s)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Veyx Pharma GmbH

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**Marketing authorisation date:**

17/06/2015

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**Manufacturing sites for batch release:**

Veyx Pharma GmbH

Veyx-Pharma B.V.

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 840/16/07/1513

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**Date of authorisation status change:**

17/06/2015

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0158/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents