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
Gonavet Veyx 50 µg/ml Injektionslösung für Rinder, Schweine und Pferde

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)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01CA01

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

Authorisation status:
Valid

Authorised in:
Austria

Available in:
Austria

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)

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:
Veyx Pharma GmbH

Marketing authorisation date:
10/03/2015

Manufacturing sites for batch release:
Veyx Pharma GmbH
Veyx-Pharma B.V.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
836076

Date of authorisation status change:

10/03/2015

Reference member state:
Germany

Procedure number:
DE/V/0158/001

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)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 14/02/2022
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Package Leaflet

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Labelling

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