Gonavet Veyx 50 micrograms/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 micrograms/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)

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:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

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:
31/03/2015

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 27569/4005

Date of authorisation status change:

1/10/2020

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

Combined File of all Documents