Fertigest 0.004 mg/ml solution for injection

Authorised

- Buserelin

Product identification

Medicine name:
Fertigest 0.004 mg/ml solution for injection
FERTIGEST 0.004 mg/ml Injektionslösung

Active substance:
Buserelin

Target species:
Cattle (cow)
Horse
Rabbit
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Buserelin
0.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle (cow)**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle (cow)**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Germany

Available in:
Germany

Package description:
1 vial of 20 ml product in type I colourless glass vials closed with bromobutyl rubber stopper (type I) and sealed with aluminium cap in a cardboard box.
5 vials of 20 ml product in type I colourless glass vials closed with bromobutyl rubber stopper (type I) and sealed with aluminium cap in a cardboard box.

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:
Vetpharma Animal Health S.L.

Marketing authorisation date:
14/08/2017

Manufacturing sites for batch release:
Mevet S.A.U.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402369.00.00

Date of authorisation status change:
14/08/2017

Reference member state:
Netherlands

Procedure number:
NL/V/0212/001

Concerned member states:
Austria Belgium Croatia Czechia Germany Hungary Ireland Poland Portugal Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

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