

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 injekčný roztok

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
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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

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:

29/01/2018

Manufacturing sites for batch release:

Mevet S.A.U.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/062/DC/17-S

Date of authorisation status change:

15/02/2023

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

Documents

Combined File of all Documents

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