

Fertigest 0.004 mg/ml solution for injection

Authorised

- Buserelin

Product identification

Medicine name:

Fertigest 0.004 mg/ml solution for injection

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. no withdrawal period zero days
- Meat and offal. no withdrawal period zero days

-

Horse

- Milk. no withdrawal period zero days
- Meat and offal. no withdrawal period zero days

-

Rabbit

- Meat and offal. no withdrawal period zero days

-

Pig

- Meat and offal. no withdrawal period zero days

Subcutaneous use:

-

Cattle (cow)

- Milk. no withdrawal period zero days
- Meat and offal. no withdrawal period zero days

-

Horse

- Milk. no withdrawal period zero days

- Meat and offal. no withdrawal period zero days

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Rabbit

- Meat and offal. no withdrawal period zero days

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Pig

- Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

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.

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.

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.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402369.00.00

Date of authorisation status change:

3/02/2023

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

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.