

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

•

Rabbit

- Meat and offal. no withdrawal period zero days

•

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:

Ireland

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:

1/09/2017

Manufacturing sites for batch release:

Mevet S.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10516/016/001

Date of authorisation status change:

1/09/2017

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