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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, otopina za injekciju

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:
Croatia

Available in:
Croatia

Package description:
Available only in Croatian
Available only in Croatian

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:
4/08/2017

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/22-01/472

Date of authorisation status change:
8/05/2025

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

Summary of Product Characteristics

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