GESLIN

Authorised

- Buserelin acetate

Product identification

Medicine name:
GESLIN
Geslin 0,004 mg/ml Roztwór do wstrzykiwań

Active substance:
Buserelin acetate

Target species:
Cattle (cow)
Pig (sow for reproduction)
Horse (mare)
Rabbit (female for reproduction)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig (sow for reproduction)**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Rabbit (female for reproduction)**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig (sow for reproduction)**
  - Meat and offal. 0 day
- **Horse (mare)**
  - Meat and offal. 0 day

- Milk. 0 day

- **Rabbit (female for reproduction)**
  - Meat and offal. 0 day

Subcutaneous use:

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

- **Pig (sow for reproduction)**
  - Meat and offal. 0 day

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day

- **Rabbit (female for reproduction)**
  - 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:
Poland

Available in:

Poland

Package description:
Box with 5 vials of 20 ml
Box with 1 vial of 20 ml

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:
Mevet S.A.

Marketing authorisation date:
14/02/2024

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3302

Date of authorisation status change:
14/02/2024

Reference member state:
Spain

Procedure number:
ES/V/0434/001

Concerned member states:

Cyprus Estonia Poland Romania

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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

Combined File of all Documents