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GESLIN 0.0040 mg/ml solution for injection for cattle, horse, pig and rabbit

Authorised

- Buserelin acetate

Product identification

Medicine name:
GESLIN 0.0040 mg/ml solution for injection for cattle, horse, pig and rabbit

Active substance:
Buserelin acetate

Target species:
Cattle (cow)
Horse (mare)
Cattle (heifer)
Pig (sow)
Pig (sow, nullipar)
Rabbit (adult female)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Buserelin acetate
4.20 microgram(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 hour
- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cattle (heifer)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig (sow)**
  - Meat and offal. 0 day
- **Pig (sow, nullipar)**
  - Meat and offal. 0 day
- **Rabbit (adult female)**

- Meat and offal. 0 day

Intravenous use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Cattle (heifer)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig (sow)**
  - Meat and offal. 0 day
- **Pig (sow, nullipar)**
  - Meat and offal. 0 day
- **Rabbit (adult female)**
  - Meat and offal. 0 day

Subcutaneous use:

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

Horse (mare)

- Meat and offal. 0 day
- Milk. 0 hour

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Cattle (heifer)

- Meat and offal. 0 day
- Milk. 0 hour

•

Pig (sow)

- Meat and offal. 0 day

•

Pig (sow, nullipar)

- Meat and offal. 0 day

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Rabbit (adult female)

- 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:

Cyprus

Available in:

Cyprus

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:
7/01/2024

Manufacturing sites for batch release:
Mevet S.A.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00962V

Date of authorisation status change:
7/01/2024

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

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
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Summary of Product Characteristics

Package Leaflet

eu-PUAR-esv0434001-mrp-geslin-en.pdf