

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

•

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Available in:

Spain

Package description:

Available only in Spanish

Available only in Spanish

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:

11/07/2017

Manufacturing sites for batch release:

Mevet S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

3572 ESP

Date of authorisation status change:

11/07/2017

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

English (PDF)

Published on: 27/06/2024

[Download](#)

Package Leaflet

English (PDF)

Published on: 8/05/2026

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Labelling

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