

# Busol 0.004 mg/ml solution for injection for cattle, horses, rabbits

Authorised

- Buserelin acetate

## Product identification

**Medicine name:**

BUSOL 0,0040mg/ml ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

Busol 0.004 mg/ml solution for injection for cattle, horses, rabbits

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**Active substance:**

Buserelin acetate

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**Target species:**

Cattle

Horse

Rabbit

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**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

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## Product details

**Active substance and strength:**

Buserelin acetate

0.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Meat and offal. 0 day

- Milk. 0 day

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**Horse**

- Meat and offal. 0 day

- Milk. 0 day

•

**Rabbit**

- Meat and offal. 0 day

**Intravenous use:**

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**Horse**

- Meat and offal. 0 day

- Milk. 0 day

**Subcutaneous use:**

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**Horse**

- Meat and offal. 0 day

- Milk. 0 day

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**Rabbit**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH01CA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Available in:**

Greece

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**Package description:**

Pack of 5 injection vials (glass type I) each containing 10 ml in a cardboard carton. Injection vial closed with a bromobutyl rubber stopper and sealed with an aluminium crimp cap.

Pack of 50 (10x5) injection vials (multipack). Injection vial closed with a bromobutyl rubber stopper and sealed with an aluminium crimp cap.

Pack of 100 (20x5) injection vials (multipack). Injection vial closed with a bromobutyl rubber stopper and sealed with an aluminium crimp cap.

Pack of 250 (50x5) injection vials (multipack). Injection vial closed with a bromobutyl rubber stopper and sealed with an aluminium crimp cap.

Pack of 500 (100x5) injection vials (multipack). Injection vial closed with a bromobutyl rubber stopper and sealed with an aluminium crimp cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Well-established use application (Article 13a of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

T. P. Whelehan Son & Co. Limited

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**Marketing authorisation date:**

16/10/2008

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**Manufacturing sites for batch release:**

aniMedica GmbH

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

99126/03-10-2023/K-0174301

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**Date of authorisation status change:**

2/10/2023

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0213/001

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**Concerned member states:**

Belgium Bulgaria Croatia Cyprus Czechia Estonia Finland Greece Hungary  
Iceland Italy Latvia Lithuania Portugal Romania Slovakia Slovenia Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 11/01/2026

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Package Leaflet

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