

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

Authorised

- Buserelin acetate

Product identification

Medicine name:

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

Active substance:

Buserelin acetate

Target species:

Cattle

Horse

Rabbit

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

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

-

Horse

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

-

Rabbit

- Meat and offal. 0 day

Intravenous use:

-

Horse

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

Subcutaneous use:

-

Horse

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

-

Rabbit

- 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:

Czechia

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.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

aniMedica GmbH

Marketing authorisation date:

14/11/2017

Manufacturing sites for batch release:

aniMedica GmbH

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

96/051/17-C

Date of authorisation status change:

31/03/2022

Reference member state:

Ireland

Procedure number:

IE/V/0213/001

Concerned member states:

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

To consult adverse reactions on veterinary medicinal products please go to
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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Labelling

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