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:
Belgium

Available in:
Belgium

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:
T. P. Whelehan Son & Co. Limited

Marketing authorisation date:

31/01/2018

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V525342

Date of authorisation status change:
31/01/2018

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

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

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