Buserelin aniMedica 0,004 mg/ml solution for injection for cattle, horses, rabbits

Authorised

- Buserelin acetate

Product identification

Medicine name:
Buserelin aniMedica 0,004 mg/ml solution for injection for cattle, horses, rabbits
Buserelin animedica 0,004mg/ml-Injektionslösung für Rinder, Pferde und Kaninchen

Active substance:
Buserelin acetate

Target species:
Cattle
Horse
Rabbit

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous 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**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day

- **Rabbit**
  - Meat and offal. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 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:
Germany

Available in:
Germany

Package description:
(ID14) 50 millilitre(s): unspecified outer container with 5 Vial (Glass) each with 10 millilitre(s)

(ID4) 5000 millilitre(s): unspecified outer container with 100 unspecified outer container each with 5 Vial (Glass) each with 10 millilitre(s)
(ID3) 2500 millilitre(s): unspecified outer container with 50 unspecified outer container each with 5 Vial (Glass) each with 10 millilitre(s)
(ID2) 1000 millilitre(s): unspecified outer container with 20 unspecified outer container each with 5 Vial (Glass) each with 10 millilitre(s)
(ID1) 500 millilitre(s): unspecified outer container with 10 unspecified outer container each with 5 Vial (Glass) each with 10 millilitre(s)

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:
18/01/2005

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
400702.00.00

Date of authorisation status change:
31/03/2010

Reference member state:
Germany

Procedure number:

DE/V/0110/001

Concerned member states:
Austria France Poland United Kingdom (Northern Ireland)

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: 28/01/2025
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