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

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

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Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

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:
27/06/2006

Manufacturing sites for batch release:
aniMedica GmbH

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1665

Date of authorisation status change:
27/06/2006

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

Labelling

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

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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