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Receptal 4 microgram/ml solution for injection

Authorised

- Buserelin

Product identification

Medicine name:
Receptal 4 microgram/ml solution for injection

Active substance:
Buserelin

Target species:
Cattle
Horse
Rabbit
Pig
Trout

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:

Buserelin
4.00 microgram(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
- **Rabbit**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Trout**
  - Meat and offal. 0 day

Intravenous use:

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

- Meat and offal. 0 day

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

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

Subcutaneous use:

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

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

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

- **Pig**
  - 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:
Norway

Package description:
Colourless type I glass vial of 2.5 ml, closed with a laminated, halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 1 vial of 2.5 ml.
Colourless type I glass vial of 5 ml, closed with an laminated, halogenated butyl rubber stopper and an aluminium cap.Pack size:Cardboard box with 1 vial of 5 ml.
Colourless type I glass vial of 10 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack size:Cardboard box with 1 vial of 10 ml.
Colourless type II glass vial of 50 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack size:Cardboard box with 1 vial of 50 ml.
Colourless type I glass vial of 2.5 ml, closed with an laminated, halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 5 vials of 2.5 ml.
Colourless type I glass vial of 5 ml, closed with a laminated, halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 5 vials of 5 ml.
Colourless type I glass vial of 10 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 5 vials of 10 ml.
Colourless type II glass vial of 50 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 5 vials of 50 ml.
Colourless type I glass vial of 2.5 ml, closed with an laminated, halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 10 vials of 2.5 ml.
Colourless type I glass vial of 5 ml, closed with an laminated, halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 10 vials of 5 ml.
Colourless type I glass vial of 10 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 10 vials of 10 ml.
Colourless type II glass vial of 50 ml, closed with a halogenated butyl rubber stopper and an aluminium cap.Pack sizes:Cardboard box with 10 vials of 50 ml.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
18/06/1982

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
0000-06737

Date of authorisation status change:
18/06/2007

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Package Leaflet

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