Source URL: https://medicines.health.europa.eu/veterinary/en/600000064178

Receptal 0.004 mg/ml Solution for injection

Authorised

• Buserelin

Product identification

Medicine name:

Receptal 0.004 mg/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:

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

•

Rabbit

- 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

Subcutaneous use:

•

Cattle

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

•

Horse

- 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:

Ireland

Package description:

Colourless type I glass vial of 2.5 ml closed with an ETFE laminated type I bromobutyl rubber stopper and an aluminium crimp cap.Pack sizes:Cardboard box with 1 vial of 2.5 ml.

Colourless type I glass vial of 5 ml, closed with an ETFE laminated type I bromobutyl rubber stopper and an aluminium crimp cap.Pack size:Cardboard box with 1 vial of 5 ml.

Colourless type I glass vial of 10 ml closed with a type I bromobutyl rubber stopper and an aluminium crimp cap.Pack size:Cardboard box with 5 ml

Colourless type II glass vial of 50 ml, closed with a type I bromobutyl rubber stopper and an aluminium crimp cap.Pack size:Cardboard box with 1 vial of 50 ml. Colourless type I glass vials of 2.5 ml closed with an ETFE laminated type I bromobutyl rubber stopper and an aluminium crimp cap.Pack sizes:Cardboard box with 5 vials of 2.5 ml

Colourless type I glass vials of 5 ml, closed with an ETFE laminated type I bromobutyl rubber stopper or a type I bromobutyl rubber stopper and an aluminium crimp cap.Pack sizes:Cardboard box with 5 vials of 5 ml

Colourless type I glass vials of 10 ml, closed with a type I bromobutyl rubber stopperand an aluminium crimp cap.Pack sizes:Cardboard box with 5 vials of 10 ml Colourless type II glass vials of 50 ml, closed with a type I bromobutyl rubber stopper and an aluminium crimp cap.Pack sizes:Cardboard box with 5 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 (Ireland) Limited

Marketing authorisation date:

1/10/1999

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10996/123/001

Date of authorisation status change:

1/10/1999

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

Documents	
Summary of Product Characteristics	
Combined File of all Documents	