

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

Authorisation status:

Valid

Authorised in:

Croatia

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. Subsidiary In The Republic Of Croatia

Marketing authorisation date:

17/08/2021

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/17-01/162

Date of authorisation status change:

18/06/2024

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

Documents

Summary of Product Characteristics

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