VETERELIN 0.004 mg/ml solution for injection for cattle, horses, pigs, and rabbits

Authorised

- Buserelin

Product identification

Medicine name:
VETERELIN 0.004 mg/ml solution for injection for cattle, horses, pigs, and rabbits

Active substance:
Buserelin

Target species:
Cattle
Horse
Rabbit
Pig

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Buserelin
0.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

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

Subcutaneous use:

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

Horse

- Milk. 0 day
- Meat. 0 day

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Rabbit

- Meat. 0 day

Intravenous use:

•

Cattle

- Meat. 0 day
- Milk. 0 day

•

Horse

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

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Rabbit

- Meat and offal. 0 day

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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:
Denmark

Available in:
Denmark

Package description:
Colourless glass vials of 10 ml, with bromobutyl rubber closure and aluminium capsules with Flip-off opening ring in PP of blue colour. 5 Bottles of 10 ml.
Colourless glass vials of 20 ml, with bromobutyl rubber closure and aluminium capsules with Flip-off opening ring in PP of blue colour. 1 Bottle of 20 ml.
Colourless glass vials of 10 ml, with bromobutyl rubber closure and aluminium capsules with Flip-off opening ring in PP of blue colour. 1 Bottle of 10 ml.
Colourless glass vials of 10 ml, with bromobutyl rubber closure and aluminium capsules with Flip-off opening ring in PP of blue colour. 1 Bottles of 100 ml.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Calier S.A.

Marketing authorisation date:
12/08/2022

Manufacturing sites for batch release:
Laboratorios Calier S.A.

Responsible authority:
Danish Medicines Agency

Authorisation number:
67165

Date of authorisation status change:
12/08/2022

Reference member state:
Portugal

Procedure number:
PT/V/0104/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Denmark France Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Poland Romania Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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Combined File of all Documents