

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

•

Rabbit

- Meat. 0 day

Intravenous use:

•

Cattle

- Meat. 0 day
- Milk. 0 day

•

Horse

- Meat and offal. 0 day
- Milk. 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:

France

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. 1 Bottle 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. 5 Bottles 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:

9/06/2011

Manufacturing sites for batch release:

Laboratorios Calier S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/2023370 3/2011

Date of authorisation status change:

29/11/2020

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

English (PDF)

Published on: 17/11/2021

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Package Leaflet and Labelling

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

Combined File of all Documents