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
VETERELIN 0,004 MG/ML SOLUTION INJECTABLE POUR BOVINS EQUINS PORCINS ET LAPINS

Active substance:
Buserelin

Target species:
Cattle
Horse
Pig
Other Leporids

Route of administration:
Solution for injection

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:

Solution for injection:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Horse**
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Other Leporids**
  - 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. 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.

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:
National Veterinary Medicines Agency

Authorisation number:
FR/V/2023370 3/2011

Date of authorisation status change:
29/11/2020

Reference member state:
Portugal

Procedure number:
PT/V/104/001

Concerned member states:
Austria Belgium Bulgaria Croatia Denmark France Germany Hungary Ireland Italy 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

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