Porceptal 4µg/ml solution for injection for pigs

Not authorised

- Buserelin acetate

Product identification

Medicine name:
Porceptal 4µg/ml solution for injection for pigs

Active substance:
Buserelin acetate

Target species:
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Buserelin acetate
4.20 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. no withdrawal period: zero days

Subcutaneous use:

- **Pig**
  - Meat and offal. no withdrawal period: zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01CA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Cyprus

Package description:

Cardboard box containing colourless glass (type II) vials of 50 ml closed with a bromobutyl rubber stopper and an aluminium crimp cap.
Cardboard box containing colourless glass (type I) vials of 5 ml closed with an ETFE laminated bromobutyl rubber stopper (type I) and an aluminium crimp cap.
Cardboard box containing colourless glass (type I) vials of 10 ml closed with a bromobutyl rubber stopper and an aluminium crimp cap.
5x 10 ml - Cardboard box containing colourless glass (type I) vials of 10 ml closed with a bromobutyl rubber stopper and an aluminium crimp cap.
10x 5 ml - Cardboard box containing colourless glass (type I) vials of 5 ml closed with an ETFE laminated bromobutyl rubber stopper (type I) and an aluminium crimp cap.
10x 2,5 ml - Cardboard box containing colourless glass (type I) vials of 2.5 ml closed with an ETFE laminated bromobutyl rubber stopper (type I) and an aluminium crimp

cap.

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:
Intervet International B.V.

Marketing authorisation date:
12/08/2014

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00476V

Date of authorisation status change:
2/11/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0176/001

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

Documents

Summary of Product Characteristics

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