RECEPTAL

Authorised

- Buserelin acetate

Product identification

Medicine name:
RECEPTAL

Active substance:
Buserelin acetate

Target species:
Pig (sow for reproduction)
Horse (mare)
Rabbit (female for reproduction)
Cattle (cow)

Route of administration:
Intramuscular use
Intravenous 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 (sow for reproduction)**

  - Meat and offal. 0 day

- **Horse (mare)**

  - Meat and offal. 0 day

- **Rabbit (female for reproduction)**

  - Meat and offal. 0 day

- **Cattle (cow)**

  - Meat and offal. 0 day

Intravenous use:

- **Pig (sow for reproduction)**

  - Meat and offal. 0 day

- **Horse (mare)**

  - Meat and offal. 0 day

- **Rabbit (female for reproduction)**

  - Meat and offal. 0 day

- **Cattle (cow)**

  - Meat and offal. 0 day

Subcutaneous use:

- **Pig (sow for reproduction)**

  - Meat and offal. 0 day

- **Horse (mare)**

  - Meat and offal. 0 day

- **Rabbit (female for reproduction)**

  - Meat and offal. 0 day

- **Cattle (cow)**

  - 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:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
24/06/1996

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
The Spanish Agency Of Medicines And Medical Devices

Authorisation number:
1106 ESP

Date of authorisation status change:

24/06/1996

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.

Package Leaflet

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

Labelling

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