LABIPITUIN

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:
LABIPITUIN

Active substance:
OXYTOCIN SYNTHETIC

Target species:
Horse (mare)
Cattle (cow)
Pig (female)
Sheep
Goat (adult female)
Cat (adult female)
Dog (bitch)

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
OXYTOCIN SYNTHETIC

10.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig (female)**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat (adult female)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cat (adult female)**
- **Dog (bitch)**

Intravenous use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig (female)**
  - Meat and offal. 0 day
- **Sheep**

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

- **Goat (adult female)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cat (adult female)**
- **Dog (bitch)**

Subcutaneous use:

- **Horse (mare)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig (female)**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat (adult female)**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Cat (adult female)**
- **Dog (bitch)**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
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:
Labiana Life Sciences S.A.

Marketing authorisation date:
16/11/1992

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
585 ESP

Date of authorisation status change:
16/11/1992

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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