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FACILPART 10 U.I./ ml SOLUCION INYECTABLE

Authorised

- OXYTOCIN SYNTHETIC

Product identification

Medicine name:
FACILPART 10 U.I./ ml SOLUCION INYECTABLE

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

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

Subcutaneous use:

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

- Meat and offal. 0 day

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Sheep

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

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Goat (adult female)

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

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
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:
Laboratorios Syva S.A.

Marketing authorisation date:
5/10/1992

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
528 ESP

Date of authorisation status change:
5/10/1992

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

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