Oxytocine Kela 10 IU/ml Oplossing voor injectie

Authorised

- Oxytocin

Product identification

Medicine name:
Oxytocine Kela 10 IU/ml Oplossing voor injectie
Oxytocine Kela 10 IU/ml Solution injectable
Oxytocine Kela 10 IU/ml Injektionslösung

Active substance:
Oxytocin

Target species:
Sheep
Goat
Dog
Cat
Pig
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Oxytocin
10.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Milk. 0 day
  - Meat and offal. 0 day

Intravenous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH01BB02

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Oxytocine Kela 12 x 100 ml Vial Solution for injection
Oxytocine Kela 12 x 50 ml Vial Solution for injection
Oxytocine Kela 12 x 30 ml Vial Solution for injection
Oxytocine Kela 12 x 10 ml Vial Solution for injection
Oxytocine Kela 1 x 100 ml Vial Solution for injection
Oxytocine Kela 1 x 50 ml Vial Solution for injection
Oxytocine Kela 1 x 30 ml Vial Solution for injection
Oxytocine Kela 1 x 10 ml Vial Solution for injection

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Kela Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
16/08/2006

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V286072

Date of authorisation status change:
28/05/2024

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.