Oxytocine NCP 10 NE/ml oldatos injekció lovak, szarvasmarhák, juhok, sertések, kutyák és macskák számára

Authorised

- Oxytocin

Product identification

Medicine name:
Oxytocine NCP 10 NE/ml oldatos injekció lovak, szarvasmarhák, juhok, sertések, kutyák és macskák számára

Active substance:
Oxytocin

Target species:
Cattle
Sheep
Pig
Horse
Dog
Cat

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:

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

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

- **Horse**
  - Meat and offal. 0 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day

- **Sheep**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

- **Horse**
  - 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:
Hungary

Package description:
Available only in Hungarian

Available only in Hungarian
Available only in Hungarian
Available only in Hungarian

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:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
17/08/1998

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
17/08/1998

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

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