Pituisan "s" pro inj. (oxycotin), 10 I.E./ml, oplossing voor injectie

Authorised

- Oxytocin

Product identification

Medicine name:
Pituisan "s" pro inj. (oxycotin), 10 I.E./ml, oplossing voor injectie

Active substance:
Oxytocin

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

Route of administration:
Intramuscular 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**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Goat (adult female)**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Sheep**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Horse**
  - Meat and offal. no withdrawal period zero days
- **Pig**

- Meat and offal. no withdrawal period zero days

Subcutaneous use:

- **Cattle**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Goat (adult female)**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Sheep**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days
- **Horse**
  - Meat and offal. no withdrawal period zero days
- **Pig**
  - Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH01BB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch
Available only in Dutch
Available only in Dutch
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
15/04/1996

Manufacturing sites for batch release:
Alfasan Nederland B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 2113

Date of authorisation status change:
16/12/2021

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

Documents

Combined File of all Documents

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