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
- **Dog**

- **Cat**

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
- **Dog**
- **Cat**

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

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