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A.A. Oxytocine, 10 I.E./ml oplossing voor injectie

Authorised

- Oxytocin

Product identification

Medicine name:
A.A. Oxytocine, 10 I.E./ml oplossing voor injectie

Active substance:
Oxytocin

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

Route of administration:
Subcutaneous use
Intramuscular 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:
Subcutaneous use:

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

- Meat and offal. no withdrawal period Nul dagen

Intramuscular use:

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

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:
Netherlands

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
A.A.-Vet Diergeneesmiddelen N.V.

Marketing authorisation date:
10/06/2013

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 113626

Date of authorisation status change:
2/09/2019

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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