

# 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

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**Active substance:**

Oxytocin

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**Target species:**

Cattle

Goat (adult female)

Sheep

Horse

Pig

Dog

Cat

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**Route of administration:**

Subcutaneous use

Intramuscular use

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## Product details

### Active substance and strength:

Oxytocin

10.00 international unit(s) / 1.00 millilitre(s)

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### Pharmaceutical form:

Solution for injection

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### Withdrawal period by route of administration:

#### Subcutaneous use:

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#### Cattle

- Milk. no withdrawal period Nul dagen
- Meat and offal. no withdrawal period Nul dagen

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#### Goat (adult female)

- Milk. no withdrawal period Nul dagen
- Meat and offal. no withdrawal period Nul dagen

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#### Sheep

- Milk. no withdrawal period Nul dagen
- Meat and offal. no withdrawal period Nul dagen

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#### Horse

- Meat and offal. no withdrawal period Nul dagen

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#### Pig

- Meat and offal. no withdrawal period  
Nul dagen

**Intramuscular use:**

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**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

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**Horse**

- Meat and offal. no withdrawal period  
Nul dagen

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**Pig**

- Meat and offal. no withdrawal period  
Nul dagen

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QH01BB02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Netherlands

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**Package description:**

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Informed consent application (Article 13c of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

A.A.-Vet Diergeneesmiddelen N.V.

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**Marketing authorisation date:**

10/06/2013

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 113626

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**Date of authorisation status change:**

2/09/2019

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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