# OXYTOCIN 10 IU/ml injekčný roztok

Authorised

Oxytocin

## **Product identification**

#### **Medicine name:**

OXYTOCIN 10 IU/ml injekčný roztok

#### **Active substance:**

Oxytocin

## **Target species:**

Cattle (cow)

Sheep (ewe)

Pig (sow)

Dog (bitch)

#### **Route of administration:**

Subcutaneous use

Intramuscular use

Intravenous 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 (cow)

- All relevant tissues. no withdrawal period Zero days

## Sheep (ewe)

- All relevant tissues. no withdrawal period Zero days

## Pig (sow)

- All relevant tissues. no withdrawal period Zero days

### Dog (bitch)

#### Intramuscular use:

## Cattle (cow)

- All relevant tissues. no withdrawal period Zero days

## Sheep (ewe)

- All relevant tissues. no withdrawal period Zero days

## Pig (sow)

- All relevant tissues. no withdrawal period Zero days

## Dog (bitch)

#### Intravenous use:

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## Cattle (cow)

- All relevant tissues. 0 day

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## Sheep (ewe)

- All relevant tissues. 0 day

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#### Pig (sow)

- All relevant tissues. 0 day

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Dog (bitch)

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

OH01BB02

#### **Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Slovakia

## Package description:

Available only in Slovak

Available only in Slovak

## 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:

Biovet AD

## Marketing authorisation date:

13/05/2010

#### Manufacturing sites for batch release:

**Biovet AD** 

## **Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

#### **Authorisation number:**

96/008/10-S

#### Date of authorisation status change:

13/05/2010

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

## **Documents**

Combined File of all Documents

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