

# Oxytobel 10 IU/ml solution for injection for Horses, Cattle, Pigs, Sheep, Goats, Dogs and Cats

Authorised

- Oxytocin

## Product identification

### **Medicine name:**

Oxytobel 10 IU/ml solution for injection for Horses, Cattle, Pigs, Sheep, Goats, Dogs and Cats

Oxytobel 10 NE/ml oldatos injekció lovak, szarvasmarhák, sertések, juhok, kecskék, kutyák és macskák részére

---

### **Active substance:**

Oxytocin

---

### **Target species:**

Cattle

Dog

Goat

Sheep

Horse

Cat

Pig

---

### **Route of administration:**

Intramuscular use

Intravenous 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**

- Meat and offal. 0 day
- Milk. 0 hour

•

**Goat**

- Meat and offal. 0 day
- Milk. 0 hour

•

**Sheep**

- Meat and offal. 0 day
- Milk. 0 hour

•

**Horse**

- Meat and offal. 0 day
- Milk. 0 hour

•

**Pig**

- Meat and offal. 0 day

### **Intravenous use:**

- 

#### **Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

- 

#### **Goat**

- Meat and offal. 0 day
- Milk. 0 hour

- 

#### **Sheep**

- Meat and offal. 0 day
- Milk. 0 hour

- 

#### **Horse**

- Meat and offal. 0 day
- Milk. 0 hour

- 

#### **Pig**

- Meat and offal. 0 day

### **Subcutaneous use:**

- 

#### **Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

- 

#### **Goat**

- Meat and offal. 0 day

- Milk. 0 hour

- 

### **Sheep**

- Meat and offal. 0 day
- Milk. 0 hour

- 

### **Horse**

- Meat and offal. 0 day
- Milk. 0 hour

- 

### **Pig**

- Meat and offal. 0 day

---

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

QH01BB02

---

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

Veterinary medicinal product subject to veterinary prescription

---

### **Authorisation status:**

Valid

---

### **Authorised in:**

Hungary

---

### **Available in:**

Hungary

---

### **Package description:**

Type I (10 ml) brown glass vial closed with bromobutyl rubber stopper and sealed with aluminium cap.1 x 10 ml in a cardboard box

Type I (10 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.5 x 10 ml in a cardboard box

Type I (10 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.12 x 10 ml in a cardboard box

Type I (25 ml) brown glass vial closed with bromobutyl rubber stopper and sealed with aluminium cap.1 x 25 ml in a cardboard box

Type I (25 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.10 x 25 ml in a cardboard box

Type II (50 ml) brown glass vial closed with bromobutyl rubber stopper and sealed with aluminium cap.1 x 50 ml in a cardboard box

Type II (50 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.12 x 50 ml in a cardboard box

Type II (50 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.6 x (1 x 50 ml) wrapped with clear foil (multipack)

Type II (100 ml) brown glass vial closed with bromobutyl rubber stopper and sealed with aluminium cap.1 x 100 ml in a cardboard box

Type II (100 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.12 x 100 ml in a cardboard box

Type II (100 ml) brown glass vials closed with bromobutyl rubber stopper and sealed with aluminium caps.6 x (1 x 100 ml) wrapped with clear foil (multipack)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Bela-Pharm GmbH & Co. KG

---

**Marketing authorisation date:**

10/12/2014

---

**Manufacturing sites for batch release:**

Bela-Pharm GmbH & Co. KG

---

**Responsible authority:**

Directorate Of Veterinary Medicinal Products

---

**Authorisation number:**

**Date of authorisation status change:**

10/12/2014

---

**Reference member state:**

Ireland

---

**Procedure number:**

IE/V/0313/001

---

**Concerned member states:**

Croatia Czechia Denmark Estonia France Hungary Iceland Norway Poland  
Slovakia United Kingdom (Northern Ireland)

---

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

## Documents

Summary of Product Characteristics