

# MYOMETRYL , 1000 U.I./100 ml, soluzione iniettabile per bovine, cavalle, scrofe, pecore, capre, cagne e gatte

Authorised

- Oxytocin

## Product identification

### Medicine name:

MYOMETRYL , 1000 U.I./100 ml, soluzione iniettabile per bovine, cavalle, scrofe, pecore, capre, cagne e gatte

### Active substance:

Oxytocin

### Target species:

Dog (bitch)

Goat

Sheep (ewe)

Horse (mare)

Cat (adult female)

Pig (female)

Cattle (cow)

### Route of administration:

Intramuscular use

Intravenous use  
Subcutaneous use

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## Product details

### **Active substance and strength:**

Oxytocin  
10000.00 international unit(s) / 100.00 millilitre(s)

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### **Pharmaceutical form:**

Solution for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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##### **Goat**

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

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##### **Sheep (ewe)**

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

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##### **Horse (mare)**

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

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##### **Pig (female)**

- Meat and offal. 0 day

#### **Intravenous use:**

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##### **Cattle (cow)**

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

**Subcutaneous use:**

- 

**Cattle (cow)**

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

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**Goat**

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

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**Sheep (ewe)**

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

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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:**

Italy

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**Package description:**

Available only in Italian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ternova S.r.l.

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**Marketing authorisation date:**

13/12/1954

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

1/01/2009

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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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