

# CALIMICINA 200 C.L., 200 mg/ml soluzione iniettabile per bovini, suini, ovini e caprini

Authorised

- Oxytetracycline

## Product identification

### Medicine name:

CALIMICINA 200 C.L., 200 mg/ml soluzione iniettabile per bovini, suini, ovini e caprini

### Active substance:

Oxytetracycline

### Target species:

Cattle

Goat

Sheep

Pig

### Route of administration:

Intramuscular use

Intravenous use

Subcutaneous use

## Product details

### Active substance and strength:

Oxytetracycline

200.00 milligram(s) / 1.00 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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##### Cattle

- Meat and offal. 42 day
- Milk. 288 hour

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##### Goat

- Meat and offal. 42 day
- Milk. 288 hour

- 

##### Sheep

- Meat and offal. 42 day
- Milk. 288 hour

- 

##### Pig

- Meat and offal. 42 day

#### Intravenous use:

- 

##### Cattle

- Meat and offal. 42 day
- Milk. 288 hour

- 

### **Goat**

- Meat and offal. 42 day
- Milk. 288 hour

- 

### **Sheep**

- Meat and offal. 42 day
- Milk. 288 hour

## **Subcutaneous use:**

- 

### **Cattle**

- Meat and offal. 42 day
- Milk. 288 hour

- 

### **Goat**

- Meat and offal. 42 day
- Milk. 288 hour

- 

### **Sheep**

- Meat and offal. 42 day
- Milk. 288 hour

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

QJ01AA06

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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](#)

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:**

Laboratorios Calier S.A.

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**Marketing authorisation date:**

14/10/1993

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**Manufacturing sites for batch release:**

Laboratorios Calier S.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:**

14/10/1993

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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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