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# MASTICEN SECADO, intramaminė suspensija

Authorised

• Cloxacillin hemibenzathine

### **Product identification**

### **Medicine name:**

MASTICEN SECADO, intramaminė suspensija

### **Active substance:**

Cloxacillin hemibenzathine

### **Target species:**

Cattle (cow)

Sheep

Goat

### **Route of administration:**

Intramammary use

### **Product details**

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

Cloxacillin hemibenzathine 1000.00 milligram(s) / 1.00 Syringe

### **Pharmaceutical form:**

## Withdrawal period by route of administration: Intramammary use:

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### Cattle (cow)

- Meat and offal. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

- Milk. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

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### Sheep

- Meat and offal. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

- Milk. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

Goat

- Meat and offal. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

- Milk. 0 day

If the drug was administered 28 days before giving birth. If the drying period is less than 4 weeks, milk has to be tested for antibiotics residue.

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

QJ51CF02

Authorization status	
<b>Authorisation status:</b> Valid	
<b>Authorised in:</b> Lithuania	
Package description:	
Available only in <u>Lithuanian</u>	
Available only in <u>Lithuanian</u>	
Additional information	
Entitlement type:	
Marketing Authorisation	
Legal basis of product authorisation:	
Legal basis not covered by Directive 2001/82/EC	
Marketing authorisation holder: Vetmarket UAB	
Marketing authorisation date: 10/06/2007	
Manufacturing sites for batch release: Cenavisa S.L.	

State Food And Veterinary Service

### **Authorisation number:**

LT/2/02/1410/001-002

### Date of authorisation status change:

10/06/2007

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

### **Documents**

Combined File of all Documents

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