

# O.T.C. 10 + P.V.P. pro inj., oplossing voor injectie voor toediening aan koe, kalf, schaap, lam

Not  
authorised

- Oxytetracycline hydrochloride

## Product identification

### **Medicine name:**

O.T.C. 10 + P.V.P. pro inj., oplossing voor injectie voor toediening aan koe, kalf, schaap, lam

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### **Active substance:**

Oxytetracycline hydrochloride

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### **Target species:**

Cattle (cow)  
Sheep  
Cattle (calf)  
Sheep (lamb)  
Pig (piglet)

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### **Route of administration:**

Intramuscular use

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## Product details

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

Oxytetracycline hydrochloride  
100.00 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 (cow)**

- Milk. 7 day
- Meat and offal. 23 day

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#### **Sheep**

- Milk. 7 day
- Meat and offal. 23 day

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#### **Cattle (calf)**

- Meat and offal. 23 day

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#### **Sheep (lamb)**

- Meat and offal. 23 day

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#### **Pig (piglet)**

- Meat and offal. 35 day

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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:**

Surrendered

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**Authorised in:**

Netherlands

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**Package description:**

Available only in Dutch

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Eurovet Animal Health B.V.

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**Marketing authorisation date:**

8/10/1991

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**Manufacturing sites for batch release:**

Eurovet Animal Health B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 1418

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**Date of authorisation status change:**

20/03/2023

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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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