**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000064626

# Oxytetracycline Injection 10%

Authorised

Oxytetracycline hydrochloride

## Product identification

### **Medicine name:**

Oxytetracycline Injection 10%

#### **Active substance:**

Oxytetracycline hydrochloride

## **Target species:**

Cattle

#### Route of administration:

Intramuscular use

# **Product details**

## **Active substance and strength:**

Oxytetracycline hydrochloride 100.00 milligram(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

Solution for injection

# Withdrawal period by route of administration:

#### Intramuscular use:

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#### **Cattle**

- Meat and offal. 21 day
- Milk. 3 day

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

**QJ01AA06** 

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

## **Authorised in:**

Ireland

## Package description:

Amber type II glass 100 ml vial, sealed with a butyl rubber stopper and non reusable aluminium closure.

# Additional information

# **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

# Marketing authorisation holder:

Alfasan Nederland B.V.

# Marketing authorisation date:

1/10/1988

# Manufacturing sites for batch release:

Alfasan Nederland B.V.

Responsible authority: Health Products Regulatory Authority
Authorisation number: VPA10980/001/001
Date of authorisation status change: 1/10/1988
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Summary of Product Characteristics