

# Terramycin LA 200 mg/ml Solution for Injection

Not  
authorised

- Oxytetracycline dihydrate

## Product identification

**Medicine name:**

Terramycin LA 200 mg/ml Solution for Injection

---

**Active substance:**

Oxytetracycline dihydrate

---

**Target species:**

Cattle

Sheep

Deer

Pig

---

**Route of administration:**

Intramuscular use

Intravenous use

Subcutaneous use

---

## Product details

**Active substance and strength:**

Oxytetracycline dihydrate

215.60 milligram(s) / 1.00 millilitre(s)

---

**Pharmaceutical form:**

Solution for injection

---

**Withdrawal period by route of administration:**

**Intramuscular use:**

•

**Cattle**

- Meat and offal. 21 day

- Milk. 7 day

•

**Sheep**

- Meat and offal. 21 day

•

**Deer**

- Meat and offal. 30 day

•

**Pig**

- Meat and offal. 21 day

**Intravenous use:**

•

**Cattle**

- Meat and offal. 21 day

- Milk. 72 hour

**Subcutaneous use:**

•

**Cattle**

- Meat and offal. 21 day

•

**Sheep**

- Meat and offal. 21 day

•

**Deer**

- Meat and offal. 30 day

•

**Pig**

- Meat and offal. 21 day

---

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

QJ01AA06

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Surrendered

---

**Authorised in:**

Ireland

---

**Package description:**

Multi dose Type II amber coloured glass vial of 100 ml capacity stoppered with a red butyl rubber bung and capped with a centre tear off aluminium crimp seal and grey flick-off cap - containing a sterile, clear yellow to amber solution.

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

Zoetis Belgium S.A.

---

**Marketing authorisation date:**

13/08/2015

---

**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain S.L.

---

**Responsible authority:**

Health Products Regulatory Authority

---

**Authorisation number:**

VPA10387/077/001

---

**Date of authorisation status change:**

26/07/2022

---

To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics