

# CYDECTIN LA 20 MG/ML SOLUTION FOR INJECTION FOR SHEEP

Authorised

- Moxidectin

## Product identification

**Medicine name:**

CYDECTIN LA 20 MG/ML SOLUTION FOR INJECTION FOR SHEEP  
CYDECTIN LARGA ACCION 20 mg/ml PARA OVINO

**Active substance:**

Moxidectin

**Target species:**

Sheep

**Route of administration:**

Subcutaneous use

## Product details

**Active substance and strength:**

Moxidectin  
20.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Solution for injection

**Withdrawal period by route of administration:****Subcutaneous use:**

- 

**Sheep**

- Meat and offal. 104 day
- Milk. no withdrawal period

Not permitted for use in dairy sheep, at any stage of life.

---

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

QP54AB02

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Spain

---

**Available in:**

Spain

---

**Package description:**

Carton of a natural high density polyethylene vial with Flurotec coated chlorinated butyl rubber stopper and aluminium flip off seal (50 ml)

Carton of a natural high density polyethylene vial with Flurotec coated chlorinated butyl rubber stopper and aluminium seal (500 ml).

Carton of a natural high density polyethylene vial with Flurotec coated chlorinated butyl rubber stopper and aluminium seal (200 ml).

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Zoetis Spain S.L.

---

**Marketing authorisation date:**

18/12/2008

---

**Manufacturing sites for batch release:**

Zoetis Manufacturing & Research Spain, S.L.

---

**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

---

**Authorisation number:**

1959 ESP

---

**Date of authorisation status change:**

18/12/2008

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0188/001

---

**Concerned member states:**

Ireland Italy Spain United Kingdom (Northern Ireland)

---

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

This document does not exist in this language (English). You can find it in another language below.

### Labelling

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

---

**Source URL:** <https://medicines.health.europa.eu/veterinary/600000044720>