

# EURICAN L-MULTI SUSPENSION FOR INJECTION

Authorised

- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

## Product identification

### Medicine name:

EURICAN L-MULTI SUSPENSION FOR INJECTION  
EURICAN LMULTI SUSPENSION INYECTABLE

### Active substance:

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

### Target species:

Dog

### Route of administration:

Subcutaneous use

## Product details

### Active substance and strength:

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

1.00 Hamster protective Dose 80% / 1.00 Dose

Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated

1.00 Hamster protective Dose 80% / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

1.00 Hamster protective Dose 80% / 1.00 Dose

---

### Pharmaceutical form:

Suspension for injection

---

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

QI07AB01

---

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

### Authorisation status:

Valid

---

### Authorised in:

Spain

---

### Available in:

Spain

---

### Package description:

Plastic box of 25 vials (glass) of suspension (1 ml)

Plastic box of 50 vials (glass) of suspension (1 ml)

Plastic box of 10 vials (glass) of suspension (1 ml)

---

## Additional information

### Entitlement type:

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health Espana S.A.

---

**Marketing authorisation date:**

5/11/2015

---

**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

---

**Responsible authority:**

Spanish Agency For Medicines And Medical Devices

---

**Authorisation number:**

3324 ESP

---

**Date of authorisation status change:**

1/01/2017

---

**Reference member state:**

France

---

**Procedure number:**

FR/V/0288/001

---

**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta  
Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain  
Sweden 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.

### Package Leaflet

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.