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

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

Package description:

Plastic box of 10 vials (glass) of suspension (1 ml)
Plastic box of 50 vials (glass) of suspension (1 ml)
Plastic box of 25 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 Of 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

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.