EURICAN L-MULTI SUSPENSION FOR INJECTION

Not 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 Injektionsvätska, suspension

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:

Surrendered

Authorised in:

Sweden

Package description:

Available only in Swedish
Available only in Swedish
Available only in Swedish

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 Denmark A/S

Marketing authorisation date:
3/09/2015

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
52109

Date of authorisation status change:
22/12/2025

Reference member state:
France

Procedure number:
FR/V/0288/001

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.