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Eurican L4 (--) - Suspension for injection

Authorised

- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
- Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated
- Leptospira interrogans, serovar Bratislava, strain 16785, Inactivated

Product identification

Medicine name:
Eurican L4 (--) - Suspension for injection

Active substance:
Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
Leptospira interrogans, serovar Grippotyphosa, strain Grippo Mal 1540, Inactivated
Leptospira interrogans, serovar Bratislava, strain 16785, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated

Presentation_strength:≥ 80% Reference:Ph.Eur. 0447 Index:0

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated

Presentation_strength:≥ 80% Reference:Ph.Eur. 0447 Index:1

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

Presentation_strength:≥ 80% Reference:Ph.Eur. 0447 Index:2

Leptospira interrogans, serovar Bratislava, strain 16785, Inactivated

Presentation_strength:≥ 80% Reference:Ph.Eur. 0447 Index:3

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Czechia , France , Luxembourg , Poland , Slovakia , Spain

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Package description:

Packaging:Vial (glass), Package_size:50 vials, Content:1 ml

Packaging:Vial (glass), Package_size:10 vials, Content: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 Vetmedica GmbH

Marketing authorisation date:
31/03/2023

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
31/03/2023

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)

Published on: 14/03/2026

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