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Canigen L4 (--) - Suspension for injection

Authorised

- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
- Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated
- Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated

Product identification

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

Active substance:
Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated
Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

Presentation_strength:3550-7100 U Reference:Hse Index:0

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Presentation_strength:290-1000 U Reference:Hse Index:1

Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated

Presentation_strength:500-1700 U Reference:Hse Index:2

Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated

Presentation_strength:650-1300 U Reference:Hse 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:

France , Norway , Poland , Spain

Package description:

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

Packaging:Vial (glass), Package_size:10 vials, Content:1 ml (1 dose)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
3/07/2015

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
3/07/2015

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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