BIOCAN LR Λυοφιλοποιημένο υλικό και διαλύτης για ενέσιμο εναιώρημα

Authorised

- Leptospira interrogans, serovar Grippotyphosa, Inactivated
- Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
- Rabies virus, Inactivated

Product identification

Medicine name:
BIOCAN LR Λυοφιλοποιημένο υλικό και διαλύτης για ενέσιμο εναιώρημα

Active substance:
Leptospira interrogans, serovar Grippotyphosa, Inactivated
Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
Rabies virus, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Grippotyphosa, Inactivated
32.00 log2 geometric mean titre / 1.00 Dose

Leptospira interrogans, serovar Canicola, strain 16070, Inactivated
32.00 log2 geometric mean titre / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 16069, Inactivated
32.00 log2 geometric mean titre / 1.00 Dose

Rabies virus, Inactivated
2.00 international unit(s) / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - Not applicable. no withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Greece

Package description:

Available only in Greek

Available only in Greek

Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
18/10/2021

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
National Organization For Medicines

Authorisation number:
69253/19-06-2025/K-0267501

Date of authorisation status change:
18/06/2025

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.