BIOCAN LR - solution for injection

Authorised

- Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated
- Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated
- Leptospira interrogans, serogroup Canicola, serovar Canicola, strain MSLB 1010, Inactivated
- Rabies virus, strain SAD Vnukovo-32, Inactivated

Product identification

Medicine name:
БИОКАН LR - инжекционен разтвор
BIOCAN LR - solution for injection

Active substance:
Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated

Leptospira interrogans, serogroup Canicola, serovar Canicola, strain MSLB 1010, Inactivated

Rabies virus, strain SAD Vnukovo-32, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serogroup Grippotyphosa, serovar Grippotyphosa, Inactivated
10.00 million organisms / 1.00 Dose

Leptospira interrogans, serogroup Icterohaemorrhagiae, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated
10.00 million organisms / 1.00 Dose

Leptospira interrogans, serogroup Canicola, serovar Canicola, strain MSLB 1010, Inactivated
10.00 million organisms / 1.00 Dose

Rabies virus, strain SAD Vnukovo-32, Inactivated
2.00 international unit(s) / 1.00 Dose

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**

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:

Bulgaria

Package description:

Available only in Bulgarian

Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
9/07/2008

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1964

Date of authorisation status change:
9/07/2008

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.
