

Biocan LR, Injekční suspenze

Authorised

- *Leptospira interrogans*, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated
- *Leptospira interrogans*, serovar Canicola, strain MSLB 1010, Inactivated
- *Leptospira kirschneri*, serovar Grippotyphosa, strain MSLB 1009, Inactivated
- Rabies virus, strain SAD Vnukovo-32, Inactivated

Product identification

Medicine name:

Biocan LR, Injekční suspenze

Active substance:

Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated

Leptospira interrogans, serovar Canicola, strain MSLB 1010, Inactivated

Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1009, Inactivated

Rabies virus, strain SAD Vnukovo-32, Inactivated

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Leptospira interrogans, serovar Icterohaemorrhagiae, strain MSLB 1008, Inactivated
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Leptospira interrogans, serovar Canicola, strain MSLB 1010, Inactivated
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

Leptospira kirschneri, serovar Grippotyphosa, strain MSLB 1009, Inactivated
32.00 Antibody microagglutination-lytic reaction / 1.00 Dose

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

Pharmaceutical form:

Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

22/02/2004

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/008/04-C

Date of authorisation status change:

13/03/2009

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.

Labelling

This document does not exist in this language (English). You can find it in another language below.