Biocan DHPPi+LR Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Authorised

- Canine distemper virus, strain CDVU 39, Live
- Canine adenovirus 2, strain CAV-2-Bio 13, Live
- Canine parvovirus, strain OP-I/81, Live
- Canine parainfluenza virus, strain CPiV-2-Bio 15, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Grippotyphosa, Inactivated
- Rabies virus, strain SAD Vnukovo-32, Inactivated
- Algeldrate

Product identification

Medicine name:
Biocan DHPPi+LR Liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań

Active substance:
Canine distemper virus, strain CDVU 39, Live

Canine adenovirus 2, strain CAV-2-Bio 13, Live

Canine parvovirus, strain OP-I/81, Live

Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Grippotyphosa, Inactivated

Rabies virus, strain SAD Vnukovo-32, Inactivated

Algeldrate

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain CDVU 39, Live

Canine adenovirus 2, strain CAV-2-Bio 13, Live

Canine parvovirus, strain OP-I/81, Live

Canine parainfluenza virus, strain CPiV-2-Bio 15, Live

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Grippotyphosa, Inactivated

Rabies virus, strain SAD Vnukovo-32, Inactivated

Algeldrate
2.00 milligram(s) / 1.00 milligram(s)

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AJ02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Package description:
Available only in Polish
Available only in Polish
Available only in Polish

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:
Grabikowski-Grabikowska Przedsiebiorstwo Produkcyjno-Handlowo-Uslugowe Inex Sp. j.

Marketing authorisation date:
21/10/2010

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2011

Date of authorisation status change:
21/10/2010

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

Documents

Package Leaflet

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Summary of Product Characteristics

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