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POLYGLOB, Injekční roztok

Not authorised

- Immunoglobulins against Canine distemper virus, Canine
- Immunoglobulins against Canine parvovirus, Canine
- Immunoglobulins against Canine parainfluenza virus, Canine
- Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine

Product identification

Medicine name:
POLYGLOB, Injekční roztok

Active substance:
Immunoglobulins against Canine distemper virus, Canine
Immunoglobulins against Canine parvovirus, Canine
Immunoglobulins against Canine parainfluenza virus, Canine
Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine

Target species:
Dog

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:

Immunoglobulins against Canine distemper virus, Canine
160.00 virus neutralising unit(s) / 1.00 Dose

Immunoglobulins against Canine parvovirus, Canine
1024.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Immunoglobulins against Canine parainfluenza virus, Canine
64.00 haemagglutination inhibiting unit(s) / 1.00 Dose

Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine
160.00 virus neutralising unit(s) / 1.00 Dose

Pharmaceutical form:

Solution for injection

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
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:
29/03/2005

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/014/05-C

Date of authorisation status change:
23/05/2022

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