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Bodyglob injekčný roztok pre psy

Authorised

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

Product identification

Medicine name:
Bodyglob injekčný roztok pre psy

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

Target species:
Dog

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Immunoglobulins against Canine parvovirus, Canine
512.00 haemagglutination inhibiting unit(s) / 1.00 millilitre(s)

Immunoglobulins against Canine distemper virus, Canine
80.00 log2 virus neutralising unit(s) / 1.00 millilitre(s)

Immunoglobulins against Canine adenovirus 1 and Canine adenovirus 2, Canine
64.00 haemagglutination inhibiting unit(s) / 1.00 millilitre(s)

Immunoglobulins against Canine parainfluenza virus, Canine
32.00 haemagglutination inhibiting unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AM03

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak

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:
Pharmagal Bio spol. s r.o.

Marketing authorisation date:
26/04/2004

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

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

Authorisation number:
97/044/04-S

Date of authorisation status change:
29/11/2022

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

Documents

Combined File of all Documents

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