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Canigen DHA2PPi/LR liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań dla psów

Authorised

- Canine parvovirus, strain Cornell 780916, Live
- Canine adenovirus 2, strain Manhattan, Live
- Canine distemper virus, strain Lederle VR128, Live
- Leptospira interrogans, serovar Canicola, Inactivated
- Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated
- Canine parainfluenza virus, strain Manhattan, Live
- Rabies virus, strain VP12, Inactivated

Product identification

Medicine name:
Canigen DHA2PPi/LR liofilizat i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań dla psów

Active substance:
Canine parvovirus, strain Cornell 780916, Live

Canine adenovirus 2, strain Manhattan, Live

Canine distemper virus, strain Lederle VR128, Live

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine parainfluenza virus, strain Manhattan, Live

Rabies virus, strain VP12, Inactivated

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine parvovirus, strain Cornell 780916, Live

Canine adenovirus 2, strain Manhattan, Live

Canine distemper virus, strain Lederle VR128, Live

Leptospira interrogans, serovar Canicola, Inactivated

Leptospira interrogans, serovar Icterohaemorrhagiae, Inactivated

Canine parainfluenza virus, strain Manhattan, Live

Rabies virus, strain VP12, Inactivated

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI

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

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

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

Marketing authorisation date:
30/04/2004

Manufacturing sites for batch release:
Virbac

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

Authorisation number:
1484

Date of authorisation status change:
30/04/2004

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.