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CANIGEN CHP/L LYOPHILISAT ET SUSPENSION POUR SUSPENSION INJECTABLE

Authorised

- Canine adenovirus 2, strain Manhattan, Live
- Canine parvovirus, strain Cornell 780916, Live
- Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated
- Leptospira interrogans, serovar Canicola, strain 601903, Inactivated
- Canine distemper virus, strain Lederle, Live

Product identification

Medicine name:
CANIGEN CHP/L LYOPHILISAT ET SUSPENSION POUR SUSPENSION INJECTABLE

Active substance:
Canine adenovirus 2, strain Manhattan, Live

Canine parvovirus, strain Cornell 780916, Live

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated

Canine distemper virus, strain Lederle, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine adenovirus 2, strain Manhattan, Live
10000.00 50% cell culture infectious dose / 1.00 Dose

Canine parvovirus, strain Cornell 780916, Live
100000.00 50% cell culture infectious dose / 1.00 Dose

Leptospira interrogans, serovar Icterohaemorrhagiae, strain 601895, Inactivated
4250.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Leptospira interrogans, serovar Canicola, strain 601903, Inactivated
4350.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Canine distemper virus, strain Lederle, Live
1000.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AI03

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
30/05/1985

Manufacturing sites for batch release:
Virbac

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1998212 2/1985

Date of authorisation status change:
5/01/2023

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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