

Nobivac Pi, Lyophilisate and Solvent for Suspension for Injection for Dogs

Authorised

- Canine parainfluenza virus, strain Cornell, Live

Product identification

Medicine name:

Nobivac Pi, Lyophilisate and Solvent for Suspension for Injection for Dogs
Nobivac Pi, Lyofilizát a ředidlo k naředění k injekční aplikaci

Active substance:

Canine parainfluenza virus, strain Cornell, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine parainfluenza virus, strain Cornell, Live
316228.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD08

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Package description:

(ID4) 50 Dose; 50 millilitre(s): Box (Cardboard) with 50 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 50 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID3) 25 Dose; 25 millilitre(s): Box (Cardboard) with 25 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 25 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID2) 10 Dose; 10 millilitre(s): Box (Cardboard) with 10 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 10 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID1) 5 Dose; 5 millilitre(s): Box (Cardboard) with 5 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 5 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID16) 50 Dose; 50 millilitre(s): Box (Plastic) with 50 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Plastic) with 50 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID15) 25 Dose; 25 millilitre(s): Box (Plastic) with 25 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Plastic)

with 25 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID14) 10 Dose; 10 millilitre(s): Box (Plastic) with 10 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Plastic) with 10 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID13) 5 Dose; 5 millilitre(s): Box (Plastic) with 5 Vial (Glass type I) each with 1 Dose, closed with Lid and Stopper (Aluminium, Halobutyl Rubber) and Box (Plastic) with 5 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID12) 50 Dose; 50 millilitre(s): Box (Cardboard) with 50 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Cardboard) with 50 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID11) 25 Dose; 25 millilitre(s): Box (Cardboard) with 25 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Cardboard) with 25 Vial (Glass type I) each with 1 Dose, closed with Lid and Stopper (Aluminium, Halobutyl Rubber)

(ID10) 10 Dose; 10 millilitre(s): Box (Cardboard) with 10 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and Box (Cardboard) with 10 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID9) 5 Dose; 5 millilitre(s): Box (Cardboard) with 5 Vial (Glass type I) each with 1 Dose, closed with Lid and Stopper (Aluminium, Halobutyl Rubber) and Box (Cardboard) with 5 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID8) 50 Dose; 50 millilitre(s): Box (Plastic) with 50 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 50 Vial (Glass type I) each with 1 millilitre(s), closed with Lid and Stopper (Aluminium, Halobutyl Rubber)

(ID7) 25 Dose; 25 millilitre(s): Box (Plastic) with 25 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 25 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium)

(ID6) 10 Dose; 10 millilitre(s): Box (Plastic) with 10 Vial (Glass type I) each with 1 millilitre(s), closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 10 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber,

Aluminium)

(ID5) 5 Dose; 5 millilitre(s): Box (Plastic) with 5 Vial (Glass type I) each with 1 Dose, closed with Stopper and Lid (Halobutyl Rubber, Aluminium) and 5 Vial (Glass type I) each with 1 millilitre(s), closed with Lid and Stopper (Aluminium, Halobutyl Rubber)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

6/08/2004

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/128/04-C

Date of authorisation status change:

6/08/2004

Reference member state:

Germany

Procedure number:

DE/V/0286/001

Concerned member states:

Austria Belgium Cyprus Czechia Denmark Estonia Finland Greece Hungary
Ireland Latvia Lithuania Luxembourg Netherlands Norway Portugal Slovenia
Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 28/01/2022

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Package Leaflet

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Labelling

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