

Nobivac DHPPi Vet. lyophilisate and solvent for suspension for injection for dogs

Authorised

- Canine parainfluenza virus, strain Cornell, Live
- Canine adenovirus 2, strain Manhattan LPV3, Live
- Canine distemper virus, strain Onderstepoort, Live
- Canine parvovirus, strain 154, Live

Product identification

Medicine name:

Nobivac DHPPi Vet. lyophilisate and solvent for suspension for injection for dogs

Active substance:

Canine parainfluenza virus, strain Cornell, Live
Canine adenovirus 2, strain Manhattan LPV3, Live
Canine distemper virus, strain Onderstepoort, Live
Canine parvovirus, strain 154, 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 millilitre(s)

Canine adenovirus 2, strain Manhattan LPV3, Live

10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine distemper virus, strain Onderstepoort, Live

10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, strain 154, Live

10000000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Available in:

Slovenia

Package description:

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

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:

Intervet International B.V.

Marketing authorisation date:

2/10/1999

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

MR/V/0750/001

Date of authorisation status change:

2/10/1999

Reference member state:

Denmark

Procedure number:

DK/V/0103/001

Concerned member states:

Norway Slovenia

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.

PI.pdf