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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

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Authorisation status:

Valid

Authorised in:

Norway

Available in:

Norway

Package description:

10 x 1 dose vials in a carton or plastic box
5 x 1 dose vials in a carton or plastic box
50 x 1 dose vials in carton or plastic box
25 x 1 dose vials in a carton or plastic box

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:
27/09/2001

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
2000-03491

Date of authorisation status change:
6/12/2010

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

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Package Leaflet

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

PI.pdf