Nobivac DHPPi

Authorised

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

Product identification

Medicine name:
Nobivac DHPPi

Active substance:
Canine distemper virus, strain Onderstepoort, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine parvovirus, strain 154, Live

Canine parainfluenza virus, strain Cornell, Live

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Canine distemper virus, strain Onderstepoort, Live

10000.00 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 parvovirus, strain 154, Live
10000000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parainfluenza virus, strain Cornell, Live
316228.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:
Lyophilisate for solution 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:
Ireland

Available in:
Ireland

Package description:
Clear, glass (Type I Ph.Eur) vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap.Cartons or plastic box with 50 single dose vials of vaccine lyophilisate
Clear, glass (Type I Ph.Eur) vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap.Cartons or plastic box with 10 single dose vials of vaccine lyophilisate

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 (Ireland) Limited

Marketing authorisation date:
3/06/2005

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/166/001

Date of authorisation status change:
3/06/2005

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

Documents

Summary of Product Characteristics