

Nobivac DHP

Authorised

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

Product identification

Medicine name:

Nobivac DHP

Active substance:

Canine distemper virus, strain Onderstepoort, Live

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine parvovirus, strain 154, 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)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

Package description:

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 10 single dose vials of diluent.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 50 single dose vials of diluent.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 50 single dose vials of vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 10 single dose vials of vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 50 single dose vials. The diluent is packed together with the vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Plastic box containing 10 single dose vials. The diluent is packed together with the vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 10 single dose vials of diluent.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 50 single dose vials of diluent.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 50 single dose vials of vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 10 single dose vials of vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 50 single dose vials. The diluent is packed together with the vaccine.

Clear, Glass Type I (Ph. Eur.) single vials with halogenobutyl rubber stopper, closed with a colour coded aluminium cap. Cardboard box containing 10 single dose vials. The diluent is packed together with the vaccine.

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:

28/07/2005

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 06376/3036

Date of authorisation status change:

26/02/2024

Reference member state:

Ireland

Procedure number:

IE/V/0161/001

Concerned member states:

Netherlands Norway United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics