

Nobivac Parvo-C

Authorised

- Canine parvovirus, strain 154, Live

Product identification

Medicine name:

Nobivac Parvo-C

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Active substance:

Canine parvovirus, strain 154, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

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:

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

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

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

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

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

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

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 the 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 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 of 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. Plastic box containing 50 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 (Ireland) Limited

Marketing authorisation date:

5/03/2004

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10996/167/001

Date of authorisation status change:

5/03/2004

Reference member state:

Ireland

Procedure number:

IE/V/0160/001

Concerned member states:

Norway

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

Documents

Summary of Product Characteristics