

# Nobivac DHPPi

Authorised

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

## Product identification

**Medicine name:**

Nobivac DHPPi

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**Active substance:**

Canine adenovirus 2, strain Manhattan LPV3, Live

Canine parvovirus, strain 154, Live

Canine distemper virus, strain Onderstepoort, Live

Canine parainfluenza virus, strain Cornell, Live

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**Target species:**

Dog

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Canine adenovirus 2, strain Manhattan LPV3, Live

4.00 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

Canine parvovirus, strain 154, Live

7.00 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

Canine distemper virus, strain Onderstepoort, Live

4.00 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

Canine parainfluenza virus, strain Cornell, Live

5.50 log<sub>10</sub> 50% tissue culture infectious dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate and solvent for solution for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI07AD04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Bulgaria

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**Available in:**

Bulgaria

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**Package description:**

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

27/02/2013

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-1960

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**Date of authorisation status change:**

27/02/2013

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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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.