

# Nobivac KC nasal drops, lyophilisate and solvent for suspension for dogs

Authorised

- Canine parainfluenza virus, strain Cornell, Live
- Bordetella bronchiseptica, strain B-C2, Live

## Product identification

### **Medicine name:**

Nobivac KC nasal drops, lyophilisate and solvent for suspension for dogs  
NOBIVAC KC GOUTTES NASALES, LYOPHILISAT ET SOLVANT POUR SUSPENSION POUR  
CHIENS

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### **Active substance:**

Canine parainfluenza virus, strain Cornell, Live  
Bordetella bronchiseptica, strain B-C2, Live

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### **Target species:**

Dog

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### **Route of administration:**

Nasal use

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## Product details

### **Active substance and strength:**

Canine parainfluenza virus, strain Cornell, Live  
5.80 log<sub>10</sub> 50% tissue culture infectious dose / 0.40 millilitre(s)  
Bordetella bronchiseptica, strain B-C2, Live  
9.70 log<sub>10</sub> colony forming unit(s) / 0.40 millilitre(s)

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**Pharmaceutical form:**

Nasal spray, lyophilisate and solvent for suspension

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

QI07AF

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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:**

France

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**Available in:**

France

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**Package description:**

Cardboard or plastic boxes with 50 glass (type I) vial closed with a halogenobutyl rubber stopper and aluminium cap, 25 x 1 dose of vaccine and solvent  
Cardboard or plastic boxes with 10 glass (type I) vial closed with a halogenobutyl rubber stopper and aluminium cap, 5 x 1 dose of vaccine and solvent

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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

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**Marketing authorisation date:**

2/02/2001

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/9909423 4/2001

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**Date of authorisation status change:**

2/02/2006

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**Reference member state:**

Italy

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**Procedure number:**

IT/V/0134/001

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**Concerned member states:**

Austria Belgium Denmark Finland France Germany Greece Ireland

Luxembourg Netherlands Norway Poland Portugal Spain Sweden

United Kingdom (Northern Ireland)

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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

## Summary of Product Characteristics

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

## Package Leaflet and Labelling

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