

# 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

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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:**

Luxembourg

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**Available in:**

Luxembourg

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**Package description:**

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

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

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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:**

23/04/2001

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 817/01/04/2119

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**Date of authorisation status change:**

23/04/2001

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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