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

Authorised

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

Product identification

Medicine name:
Nobivac KC nasal drops, lyophilisate and solvent for suspension for dogs
Nobivac KC

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

Target species:
Dog

Route of administration:
Nasal use

Product details

Active substance and strength:
Bordetella bronchiseptica, strain B-C2, Live
9.70 log10 colony forming unit(s) / 0.40 millilitre(s)

Canine parainfluenza virus, strain Cornell, Live
5.80 log10 tissue culture infective dose 50 / 0.40 millilitre(s)

Pharmaceutical form:
Nasal spray, lyophilisate and solvent for suspension

Withdrawal period by route of administration:

Nasal use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AF

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Cardboard or plastic boxes with 50 glass (tiype 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 (tiype I) vial closed with a halogenobutyl rubber stopper and aluminium cap, 5 x 1 dose of vaccine and solvent

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:
This information is not available for this product.

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
Ministry Of Health

Authorisation number:
103962

Date of authorisation status change:
25/07/2012

Reference member state:
Italy

Procedure number:
IT/V/0134/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Greece Ireland Luxembourg Netherlands Norway Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)

Published on: 13/02/2024

Download

Source URL: *https://medicines.health.europa.eu/veterinary/600000085510*