Nobivac KC

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

Active substance:
Canine parainfluenza virus, strain Cornell, Live

Bordetella bronchiseptica, strain B-C2, Live

Target species:
Dog

Route of administration:
Nasal use

Product details

Active substance and strength:
Canine parainfluenza virus, strain Cornell, Live
5.80 log10 50% tissue culture infectious dose / 0.40 millilitre(s)

Bordetella bronchiseptica, strain B-C2, Live
9.70 log10 colony forming unit(s) / 0.40 millilitre(s)

Pharmaceutical form:
Nasal spray, lyophilisate and solvent for suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AF

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

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:
15/11/1999

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 06376/3014

Date of authorisation status change:
14/11/2024

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 Spain Sweden United Kingdom (Northern Ireland)

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