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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

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:
Norway

Available in:
Norway

Package description:
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
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

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:
25/05/2009

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
07-4878

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 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

Summary of Product Characteristics

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Package Leaflet

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