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 vet. Frystorkat pulver och vätska till näsdroppar, suspension

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 50% tissue culture infectious dose / 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:
Sweden

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:
9/05/2008

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
25263

Date of authorisation status change:
9/05/2008

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

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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