Nobivac Respira Bb vet. suspension for injection for dogs

Authorised

• Bordetella bronchiseptica, strain Bb7 92932, fimbriae

Product identification

Medicine name:

Nobivac Respira Bb vet. suspension for injection for dogs

Nobivac Respira Bb Suspensie voor injectie

Nobivac Respira Bb Suspension injectable

Nobivac Respira Bb Injektionssuspension

Active substance:

Bordetella bronchiseptica, strain Bb7 92932, fimbriae

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Bordetella bronchiseptica, strain Bb7 92932, fimbriae 88.00 enzyme-linked immunosorbent assay unit / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration: **Subcutaneous use:** • Dog Anatomical therapeutic chemical veterinary (ATCvet) codes: **QI06AE02** Legal status of supply: Veterinary medicinal product subject to veterinary prescription **Authorisation status:** Valid Authorised in: Belgium Package description: Available only in Danish 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: 23/10/2020 Manufacturing sites for batch release: INTERVET INTERNATIONAL B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number: BE-V573013
Date of authorisation status change: 23/10/2020
Reference member state: Denmark
Procedure number: DK/V/0123/002
Concerned member states: Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics
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Package Leaflet

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