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Nobivac Respira Bb vet. suspension for injection for dogs

Not authorised

- Bordetella bronchiseptica, strain Bb7 92932, fimbriae

Product identification

Medicine name:
Nobivac Respira Bb vet. suspension for injection for dogs

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AB03

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

Authorisation status:
Surrendered

Authorised in:
Cyprus

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:
3/02/2021

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

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00825V

Date of authorisation status change:
2/11/2025

Reference member state:
Denmark

Procedure number:
DK/V/0123/002

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

Documents

Combined File of all Documents

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