Nobivac Ducat Lyophilisate and solvent for suspension for injection

Authorised

- Felid herpesvirus 1, strain G2620A, Live
- Feline calicivirus, strain F9, Live

Product identification

Medicine name:
Nobivac Ducat Lyophilisate and solvent for suspension for injection

Active substance:
Felid herpesvirus 1, strain G2620A, Live

Feline calicivirus, strain F9, Live

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Felid herpesvirus 1, strain G2620A, Live
4.80 log10 50% tissue culture infectious dose / 1.00 Dose

Feline calicivirus, strain F9, Live
4.60 log10 plaque forming unit(s) / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI06AD03

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
(ID44): 1 Cardbox with 50 Bottle (Glass) with 1 Dose and 50 Bottle (Glass) with 1 millilitre(s)) (50 Dose, 50 millilitre(s))
(ID34): 1 Cardbox with 25 Bottle (Glass) with 1 Dose and 25 Bottle (Glass) with 1 millilitre(s)) (25 Dose, 25 millilitre(s))
(ID24): 1 Cardbox with 10 Bottle (Glass) with 1 Dose and 10 Bottle (Glass) with 1 millilitre(s)) (10 Dose, 10 millilitre(s))
(ID14): 1 Cardbox 5 Bottle (Glass) with 1 Dose and 5 Bottle (Glass) with 1 millilitre(s)) (5 Dose, 5 millilitre(s))
(ID45): 1 Plasticbox 5 Bottle (Glass) with 1 Dose and 5 Bottle (Glass) with 1 millilitre(s)) (5 Dose, 5 millilitre(s))
(ID46): 1 Plasticbox with 10 Bottle (Glass) with 1 Dose and 10 Bottle (Glass) with 1 millilitre(s)) (10 Dose, 10 millilitre(s))
(ID47): 1 Plasticbox with 25 Bottle (Glass) with 1 Dose and 25 Bottle (Glass) with 1 millilitre(s)) (10 Dose, 10 millilitre(s))
(ID48): 1 Plasticbox with 50 Bottle (Glass) with 1 Dose and 50 Bottle (Glass) with 1 millilitre(s)) (10 Dose, 10 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Nederland B.V.

Marketing authorisation date:
4/12/2009

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10235

Date of authorisation status change:
8/02/2022

Reference member state:
Germany

Procedure number:
DE/V/0207/001

Concerned member states:
Austria Belgium Czechia Denmark Estonia Finland France Greece Ireland Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal 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

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

Summary of Product Characteristics

Package Leaflet