Nobivac Ducat Lyophilisate and solvent for suspension for injection

Not authorised

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

Product identification

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

Active substance:
Feline calicivirus, strain F9, Live
Felid herpesvirus 1, strain G2620A, Live

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Feline calicivirus, strain F9, Live
0.66 plaque forming unit / 1.00 Dose

Felid herpesvirus 1, strain G2620A, Live
0.68 plaque forming unit / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI06AD03

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

Authorisation status:
Surrendered

Authorised in:
Estonia

Package description:
(ID44): 1 Box with 50 Box with (1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)) (50 Dose, 50 millilitre(s))
(ID34): 1 Box with 25 Box with (1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)) (25 Dose, 25 millilitre(s))
(ID24): 1 Box with 10 Box with (1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)) (10 Dose, 10 millilitre(s))
(ID14): 1 Box with 5 Box with (1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)) (5 Dose, 5 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 International B.V.

Marketing authorisation date:
31/10/2008

Manufacturing sites for batch release:
INTERVET INTERNATIONAL B.V.

Responsible authority:
State Agency Of Medicines

Authorisation number:
1531

Date of authorisation status change:
31/08/2023

Reference member state:
Germany

Procedure number:
DE/V/0207/001

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 13/03/2023

Download

Package Leaflet
