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Nobivac Tricat Trio, Lyophilisate and Solvent for Suspension for Injection for Cats

Authorised

- Feline panleucopenia virus, strain MW-1, Live
- Felid herpesvirus 1, strain G2620A, Live
- Feline calicivirus, strain F9, Live

Product identification

Medicine name:
Nobivac Tricat Trio, Lyophilisate and Solvent for Suspension for Injection for Cats

Active substance:
Feline panleucopenia virus, strain MW-1, Live

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:

Feline panleucopenia virus, strain MW-1, Live
4.30 log10 50% tissue culture infectious dose / 1.00 Dose

Felid herpesvirus 1, strain G2620A, Live
5.20 log10 plaque forming unit(s) / 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:
QI06AD04

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

Authorisation status:
Valid

Authorised in:
Finland

Available in:
Finland

Package description:
(ID4) 50 Dose; 50 millilitre(s): Box (Cardboard) with 50 Box (Cardboard) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)
(ID2) 10 Dose; 10 millilitre(s): Box (Cardboard) with 10 Box (Cardboard) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)
(ID1) 5 Dose; 5 millilitre(s): Box (Cardboard) with 5 Box (Cardboard) each with 1 Bottle (Glass) with 1 millilitre(s) and 1 Bottle (Glass) with 1 Dose
(ID3) 25 Dose; 25 millilitre(s): Box (Cardboard) with 25 Box (Cardboard) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)
(ID8) 50 Dose; 50 millilitre(s): Box (Kunststoff) with 50 Box (Kunststoff) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)
(ID7) 25 Dose; 25 millilitre(s): Box (Kunststoff) with 25 Box (Kunststoff) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)

(ID6) 10 Dose; 10 millilitre(s): Box (Kunststoff) with 10 Box (Kunststoff) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 millilitre(s)
(ID5) 5 Dose; 5 millilitre(s): Box (Kunststoff) with 5 Box (Kunststoff) each with 1 Bottle (Glass) with 1 Dose and 1 Bottle (Glass) with 1 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:
5/03/2009

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

Responsible authority:
Finnish Medicines Agency

Authorisation number:
22805

Date of authorisation status change:
5/03/2009

Reference member state:
Germany

Procedure number:
DE/V/0240/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France Greece Hungary 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

Package Leaflet

Summary of Product Characteristics

2603424-paren-20251101.pdf