

zoetis



MODIFIED
LIVE
VACCINE



FELINE FIP

Intranasal

25 Vials (1 dose)

STERILE DILUENT

25 Vials (0.5 mL)

Feline Infectious Peritonitis Vaccine

MODIFIED LIVE VIRUS

To Reconstitute:
Rehydrate the vaccine
with the sterile diluent
for vaccination.

**VANGUARD®**



FELINE FIP Intranasal

This product has been shown to be effective for the vaccination of healthy cats 16 weeks of age or older against feline infectious peritonitis virus. Duration of immunity has not been established. For more information regarding efficacy and safety data, go to productdata.aphis.usda.gov.

Directions: Aseptically rehydrate the freeze-dried vaccine with the sterile diluent provided. Mix well. Use dropper to inoculate entire volume into nasal passages (1/2 volume into each nasal passage). Cats may sneeze or shake their heads at the time of administration. Healthy cats 16 weeks of age or older should receive 2 IN doses administered 3–4 weeks apart. Historically, annual revaccination with this product was recommended. The need for annual booster vaccinations has not been established; consultation with a veterinarian or the manufacturer is recommended.

Precautions: Store at 2°–8°C. Do not freeze. Inactivate unused contents before disposal. Contains gentamicin as preservative. Droppers should be used to administer this vaccine. As with many vaccines, anaphylaxis may occur after use. Initial antidote of epinephrine is recommended and should be followed with appropriate supportive therapy. This product has not been tested in pregnant animals. Do not mix with other products, except as specified on this label. In case of human exposure, contact a physician.

Technical inquiries should be directed to Zoetis Inc. Veterinary Services, (888) 963-8471.

This product has been shown to be efficacious in healthy animals. A protective immune response may not be elicited if animals are incubating an infectious disease, are malnourished or parasitized, are stressed due to shipment or environmental conditions, are otherwise immunocompromised, or the vaccine is not administered in accordance with label directions.