

Date of Approval: April 8, 2026

FREEDOM OF INFORMATION (FOI) SUMMARY
ORIGINAL ABBREVIATED NEW ANIMAL DRUG APPLICATION (ANADA)

ANADA 200-848

Florfinasone™ Otic

(florfenicol, terbinafine, mometasone furoate)

Solution

Dogs

Florfinasone™ Otic is indicated for the treatment of otitis externa in dogs associated with susceptible strains of yeast (*Malassezia pachydermatis*) and bacteria (*Staphylococcus pseudintermedius*).

Sponsored by:

Cronus Pharma Specialities India Private Ltd.

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I. GENERAL INFORMATION

A. File Number

ANADA 200-848

B. Sponsor

Cronus Pharma Specialities India Private Ltd.
Plot No.9(B), Survey No. 99/1, GMR Hyderabad Aviation SEZ Ltd.
Mamidipalle Village, Balapur Mandal, Shamshabad
Rangareddy, Hyderabad, Telangana, 500108, India

Drug Labeler Code: 069043

U.S. Agent Name and Address:

Jodi Beaudry, MSc
J² Consulting LLC
45 Bugling Elk Ln
Columbus, MT 59019

C. Proprietary Name

Florfinasone™ Otic

D. Drug Product Established Name

florfenicol, terbinafine, mometasone furoate

E. Pharmacological Category

Antibacterial, antifungal, and anti-inflammatory

F. Dosage Form

Solution

G. Amount of Active Ingredient

16.6 mg/mL florfenicol, 14.8 mg/mL terbinafine (equivalent to 16.6 mg/mL terbinafine hydrochloride) and 2.2 mg/mL mometasone furoate

H. How Supplied

Florfinasone™ Otic is supplied in a single-use dropperette. Each dropperette contains one 1 mL dose. Available in cartons of two or ten dropperettes.

I. Dispensing Status

Prescription (Rx)

J. Dosage Regimen

Administer one dose (1 dropperette) per affected ear.

K. Route of Administration

Otic

L. Species/Class

Dogs

M. Indication

Florfinasone™ Otic is indicated for the treatment of otitis externa in dogs associated with susceptible strains of yeast (*Malassezia pachydermatis*) and bacteria (*Staphylococcus pseudintermedius*).

N. Reference Listed New Animal Drug (RLNAD)

CLARO™; florfenicol, terbinafine, mometasone furoate; NADA 141-440; Elanco US Inc.

II. BIOEQUIVALENCE

The Federal Food, Drug, and Cosmetic Act (FD&C Act), as amended by the Generic Animal Drug and Patent Term Restoration Act (GADPTRA) of 1988, allows for an abbreviated new animal drug application (ANADA) to be submitted for a generic version of an approved new animal drug (RLNAD). The ANADA sponsor is required to show that the generic product is bioequivalent to the RLNAD, which has been shown to be safe and effective. Effectiveness, target animal safety and human food safety data (other than tissue residue data) are not required for approval of an ANADA. If bioequivalence is demonstrated through a clinical endpoint study in a food-producing animal, then a tissue residue study to establish the withdrawal period for the generic product is also required. For certain dosage forms, the agency will grant a waiver from the requirement to perform *in vivo* bioequivalence studies (biowaiver) (55 FR 24645, June 18, 1990; Fifth GADPTRA Policy Letter; Bioequivalence Guideline, October 9, 2002).

Based on the formulation characteristics of the generic product, Cronus Pharma Specialities India Pvt Ltd., was granted a biowaiver for the generic product Florfinasone™ Otic (florfenicol, terbinafine, mometasone furoate) otic solution. The generic drug product is a solution, contains the same active ingredient in the same concentration and dosage form as the RLNAD, and contains no inactive ingredients that may significantly affect the bioavailability of the active ingredient. The RLNAD is CLARO™ (florfenicol, terbinafine, mometasone furoate) otic solution, sponsored by Elanco US Inc., under NADA 141-440, and was approved for use in dogs on September 20, 2015.

III. HUMAN FOOD SAFETY

This drug is intended for use in dogs. Because this new animal drug is not intended for use in food-producing animals, FDA did not require data pertaining to drug residues in food (i.e., human food safety) for approval of this ANADA.

IV. USER SAFETY

The product labeling contains the following information regarding safety to humans handling, administering, or exposed to Florfinasone™ Otic:

Florfinasone™ Otic should be administered by veterinary personnel.

Wear eye protection when administering Florfinasone™ Otic.

Splatter may occur if the dog shakes its head following administration. Persons near the dog during administration should also take steps to avoid ocular exposure.

Florfinasone™ Otic may cause eye injury and irritation.

If contact with eyes occurs, flush copiously with water for at least 15 minutes. If irritation persists, contact a physician.

Humans with known hypersensitivity to any of the active ingredients in Florfinasone™ Otic should not handle this product.

Not for use in humans. Keep this and all drugs out of the reach of children. Avoid skin contact. In case of accidental ingestion by humans, contact a physician immediately.

Wear eye protection when administering Florfinasone™ Otic and restrain the dog to minimize post application head shaking. Reducing the potential for splatter of product will help prevent accidental eye exposure in people and dogs and help to prevent ocular injury.

V. AGENCY CONCLUSIONS

The data submitted in support of this ANADA satisfy the requirements of section 512(c)(2) of the FD&C Act. The data demonstrate that Florfinasone™ Otic, when used according to the label, is safe and effective for the conditions of use in the General Information Section above.