DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
1201 Harbor Bay Parkway Alameda, CA 94502-7070 (510)337-6700 Fax:(510)337-6702	5/3/2022-5/11/2022* FEI NUMBER 3003541463

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
William Carr, Owner

FIRM NAME	STREET ADDRESS
Costa View Farms #2	16800 Road 15
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Madera, CA 93637-9075	Dairy

This document lists observations made by the FDA representative(s) during the inspection of your facility. They are inspectional observations, and do not represent a final Agency determination regarding your compliance. If you have an objection regarding an observation, or have implemented, or plan to implement, corrective action in response to an observation, you may discuss the objection or action with the FDA representative(s) during the inspection or submit this information to FDA at the address above. If you have any questions, please contact FDA at the phone number and address above.

DURING AN INSPECTION OF YOUR FIRM I OBSERVED:

OBSERVATION 1

You offered a food-producing animal(s) for interstate commerce that was found to bear or contain a drug residue in an edible tissue above an established safe level, safe concentration, or tolerance.

Specifically, on (b) (4), you delivered a cull dairy cow, identified with your ear tag (b) (6), (b) (7)(C), to be slaughtered into food for human consumption. This animal was identified by a USDA retain tag (b) (6), (b) (7)(C). The tissues of this animal (kidney) was found by USDA tests to contain (b) (4) (b) (4) at 2.51 parts per million (ppm). The tolerance for (b) (4) in the edible tissues of cattle has been established at (b) (4)ppm.

OBSERVATION 2

You did not maintain complete treatment records.

Specifically, your medication treatment records consist of electronic entries into your (b) (4) Cow File for each treated animal. The records consist of the ear tag number of the treated animal, the initial date of treatment, and the medication administered. The record does not contain the dosage administered, the route of administration, and the individual administering the medication.

*DATES OF INSPECTION

5/03/2022(Tue), 5/11/2022(Wed)

SEE REVERSE OF THIS PAGE	EMPLOYEE(S) SIGNATURE		DATE ISSUED
	Thomas W Gordon, Investigator	X Thomas W Gordon Investigator Signed By: Thomas W. Gordon -S Date Signed: 05-11-2022 07:42:21	5/11/2022

The observations of objectionable conditions and practices listed on the front of this form are reported:

1. Pursuant to Section 704(b) of the Federal Food, Drug and Cosmetic Act, or
2. To assist firms inspected in complying with the Acts and regulations enforced by the Food and Drug Administration.

Section 704(b) of the Federal Food, Drug, and Cosmetic Act (21 USC 374(b)) provides:

"Upon completion of any such inspection of a factory, warehouse, consulting laboratory, or other establishment, and prior to leaving the premises, the officer or employee making the inspection shall give to the owner, operator, or agent in charge a report in writing setting forth any conditions or practices observed by him which, in his judgment, indicate that any food, drug, device, or cosmetic in such establishment (1) consists in whole or in part of any filthy, putrid, or decomposed substance, or (2) has been prepared, packed, or held under insanitary conditions whereby it may have become contaminated with filth, or whereby it may have been rendered injurious to health. A copy of such report shall be sent promptly to the Secretary."