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	6/9/2022-7/6/2022* FEI NUMBER 3022591166

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Paulo Jorge De Sousa Silva, Owner

FIRM NAME	STREET ADDRESS
S S 2 Dairy	5324 Clayton Rd
CITY, STATE, ZIP CODE, COUNTRY	TYPE ESTABLISHMENT INSPECTED
Turlock, CA 95380-9421	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

Your extralabel use of an approved human or animal drug caused a residue above an established safe level, safe concentration, or tolerance.

Specifically, on or about, (b) (4), your dairy delivered a bob veal calf to be slaughtered into food for human consumption. This calf was identified with a Back Tag (b) (6), (b) (7)(C) and by a (b) (4) (b) (4). (b) (4) tests revealed the presence of Neomycin in the kidney of this calf at (b) (4). A tolerance for neomycin in the kidneys of cattle has been established at 7.20 ppm.

OBSERVATION 2

You did not use the VFD feed in accordance with the approved, conditionally approved, or indexed conditions of use and/or labeling.

Specifically, your veterinarian has provided to you a Veterinary Feed Directive Order (VFD Number: (b) (4)) for a feed containing Neomycin and Oxytetracycline at the rate of (b) (4) of product. The section of the Veterinary Feed Directive titled "Caution" states in part "Do not use in calves to be processed for veal."

You stated an employee fed milk replacer containing Neomycin and Oxytetracycline to a bull calf, and then after feeding this calf this medicated product, sent the calf to be slaughtered into food for human consumption.

OBSERVATION 3

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: 07-06-2022 07:39:33	7/6/2022

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You did not maintain complete treatment records.

Specifically, you are failing to maintain complete medication treatment records for your lactating dairy cattle, and failing to maintain any medication treatment records for your replacement heifer calves. The records you do maintain for your lactating dairy cattle consist of the date of treatment, the identity of the animal treated, and the drug administered. You are failing to record the dosage of the drug administered, the route of administration, the individual administering the drug, the milk and slaughter withdrawal times, and the dates those withdrawal times will have been observed.

You stated you are not maintaining medication treatment records for your lactating dairy cattle for more than a few months. Currently, you are only able to provide treatment records back to early March, 2022. You stated you discarded the treatment records prior to March 2022. You were also unable to provide any treatment records for your heifer calves.

OBSERVATION 4

You did not keep records regarding the identity of the animal(s) that you and delivered for sale at a slaughter plant.

Specifically, you delivered a bob veal calf to be slaughtered into food for human consumption on or about (b) (4) . You have stated that this calf was sold to a calf hauler/buyer. When asked for the record of sale containing the identity of this calf, you did not provide one. You stated that you discard all of these sales records received from your calf buyer.

***DATES OF INSPECTION**

6/09/2022(Thu), 7/06/2022(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: 07-06-2022 07:39:33	7/6/2022

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."