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	2/17/2026-2/25/2026*
	FEI NUMBER 3041047522

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Matthew A. Leo, Partner

FIRM NAME	STREET ADDRESS
Matthew A. Leo	606 Figmond Ave
CITY, STATE, ZIP CODE, COUNTRY Snelling, CA 95369-9750	TYPE ESTABLISHMENT INSPECTED Beef Ranch

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, you delivered a steer to be slaughtered into food for human consumption identified with your ear tag number (b) (4) This animal, identified on USDA Form (b) (4) , was found by USDA tests to contain meloxicam in its kidney. There is no tolerance for the presence of meloxicam in the edible tissues of cattle.

OBSERVATION 2

You did not maintain complete treatment records.

Specifically, your treatment records for the steer identified on USDA Form (b) (4) does not show a treatment for meloxicam. However, USDA tests revealed the presence of meloxicam in the kidney of this steer. Additionally, your records do not include the route of administration and the withdrawal times for slaughter, and the dates those withdrawal times will be observed.

*DATES OF INSPECTION

2/17/2026(Tue), 2/25/2026(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: 02-25-2026 10:14:32	2/25/2026

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."