DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION

DISTRICT ADDRESS AND PHONE NUMBER	DATE(S) OF INSPECTION
22215 26th Ave SE Suite 210 Bothell, WA 98021 (425)302-0340 Fax:(425)302-0404	4/20/2026-4/21/2026
	FEI NUMBER 1000187936

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Larry J. Boss, Owner

FIRM NAME	STREET ADDRESS
Van Beek Dairy	213 W 400 S
CITY, STATE, ZIP CODE, COUNTRY Jerome, ID 83338-5952	TYPE ESTABLISHMENT INSPECTED 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 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 07/11/2025, a dairy cow (ear tag (b) (4) sale tag (b) (4) from your dairy was processed for the purpose of human food at a slaughterhouse in (b) (4) At the time of slaughter on (b) (4) this dairy cow was sampled by USDA FSIS and found to contain levels of the drug Desfuroylceftiofur (Ceftiofur metabolite) in the kidney at a level of 1.28 ppm that exceeds the established tolerance of 0.4 ppm in the kidney.

OBSERVATION 2

You did not keep treatment records.

Specifically,

You stated that this dairy cow identified with ear tag (b) (4) was administered (b) (4) (b) (4) Ceftiofur Sodium) sometime early in the month of July 2025 however you did not keep the treatment records that documented the administrations of veterinary drugs to this animal for the minimum required time.

SEE REVERSE OF THIS PAGE

EMPLOYEE(S) SIGNATURE
Antonetta M Colacchio, Investigator

X Antonetta M Colacchio
Investigator
Signed By: ANTONETTA COLACCHIO -S
Date Signed: 04-21-2026 08:14:32

DATE ISSUED
4/21/2026

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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."