**DEPARTMENT OF HEALTH AND HUMAN SERVICES**
**FOOD AND DRUG ADMINISTRATION**

| DISTRICT ADDRESS AND PHONE NUMBER | DATE(S) OF INSPECTION |
|---|---|
| 250 Marquette Ave, Ste. 600<br>Minneapolis, MN 55401<br>(612)334-4100 Fax:(612)334-4134 | 5/4/2026-5/7/2026 |
| | FEI NUMBER<br>3011307479 |

NAME AND TITLE OF INDIVIDUAL TO WHOM REPORT ISSUED
Siebrand Johannes Miedema, Owner

| FIRM NAME | STREET ADDRESS |
|---|---|
| Miedema Dairy Farm Llc | 4431 County Road C |
| CITY, STATE, ZIP CODE, COUNTRY<br>Pulaski, WI 54162-9739 | TYPE ESTABLISHMENT INSPECTED<br>Grower |

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,

1. The cow identified by USDA/FSIS Sample Form No (b) (4) ear tag (b) (4) tag (b) (4) and back tag (b) (4) was collected at slaughter on (b)(4) and tested positive for ampicillin at 0.0561 ppm in kidney tissue and 0.0168 ppm in muscle tissue. The established tolerance for this drug in edible tissues is 0.010 ppm.

   Your auction sale records indicate that this animal weighed (b) (4) This cow was initially treated for vaginal tears with 30 cc o(b) (4) ampicillir(b) (4) administered via (b) (4) on 08/27/2025. The animal was subsequently sold for slaughter on(b) (4)

2. The cow identified by USDA/FSIS Sample Form No (b) (4), back tag (b) (4), and subsequently identified with ear ta (b) (4) was collected at slaughter or (b) (4) and tested positive for ampicillin at 0.0176 ppm. The tolerance for this drug in edible tissues is 0.010 ppm.

| SEE REVERSE OF THIS PAGE | EMPLOYEE(S) SIGNATURE | | DATE ISSUED |
|---|---|---|---|
| | Hesham A Elgaali, Investigator | X Hesham A Elgaali Investigator Signed By: 2004279840 Date Signed: 05-07-2026 10:19:21 | 5/7/2026 |

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Your auction sale records indicate that this animal weighed (b) (4) The cow was first treated for mastitis on 09/01/2025 with 30 cc o (b) (4) (ampicillin) (b) (4) administered in (b)(4) This animal was sold for slaughter on (b) (4)

The veterinary instructions written on the (b) (4) label state to administer (b) (4) intramuscular (IM) injection once daily for a minimum of 3–5 days, with a maximum of 7 days. The label indicates use for metritis, mastitis, pneumonia, and post-operative care in cattle. Milk withdrawal: (b) (4)

**OBSERVATION 2**

You did not maintain complete treatment records.

Specifically, your treatment records do not include the following information: the route of administration, identity of the person who administered the drug to the animal.

**SEE REVERSE OF THIS PAGE**

EMPLOYEE(S) SIGNATURE
Hesham A Elgaali, Investigator

X Hesham A Elgaali
Investigator
Signed By: 2004279840
Date Signed: 05-07-2026
10:19:21

DATE ISSUED
5/7/2026

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."