



DEPARTMENT OF HEALTH AND HUMAN SERVICES

HFE-35

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Food and Drug Administration
Cincinnati District Office
Central Region
6751 Steger Drive
Cincinnati, OH 45237-3097
Telephone: (513) 679-2700

**CERTIFIED MAIL
RETURN RECEIPT REQUESTED**

WARNING LETTER CIN-WL-6934-01

March 23, 2001

Randall A. Hegenderfer
President
The Centerburg Mill and General Store, Inc
108 North Hartford Avenue, Box 207
Centerburg, OH 43011

Dear Mr. Hegenderfer:

An inspection of your feed mill was conducted by a Food and Drug Administration (FDA) investigator on March 1, 2001. This inspection found significant deviations from the requirements set forth in Title 21, Code of Federal Regulations, Part 589.2000 – Animal Proteins Prohibited in Ruminant Feed. This regulation is intended to prevent the establishment and amplification of Bovine Spongiform Encephalopathy (BSE). Such deviations cause products being manufactured and distributed by your facility to be adulterated within the meaning of Section 402(a)(4) of the Federal Food, Drug, and Cosmetic Act (the Act).

Our investigator was verbally told that you have separate processing equipment for products containing prohibited materials, but the equipment is not identified, there are no written procedures to address separation of the products, and there are no processing records for products which contain, or may contain, prohibited material.

Your records must allow your firm to track products that contain, or may contain, prohibited material throughout their receipt, processing, and distribution. This includes records of cash sales.

This letter is not intended to be an all-inclusive list of deficiencies at your facility. As a manufacturer of materials intended for animal feed use, you are responsible for assuring that your overall operation and the products you manufacture and distribute are in compliance with the law. We have enclosed a copy of the FDA's Small Entity Compliance Guide to assist you with complying with the regulation.

You should take prompt action to correct these violations, and you should establish a system whereby such violations do not recur. Failure to promptly correct these violations may result in regulatory action without further notice, such as seizure and/or injunction.

You should notify this office in writing within 15 working days of receipt of this letter, of the steps you have taken to bring your firm into compliance with the law. Your response should include an explanation of each step being taken to correct the violations, and prevent their recurrence. If corrective action cannot be completed in 15 days, state the reason for the delay and the date by which the

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corrections will be completed. Include copies of any available documentation demonstrating that corrections have been made.

Your reply should be directed to Deborah Grelle, Director of Compliance, U.S. Food and Drug Administration, 6751 Steger Drive, Cincinnati, Oh 45237, telephone (513) 679-2700 extension 160.

Sincerely yours,



Henry L. Fielden
District Director

Enclosure: Small Entity Compliance Guide

Cc: David Schleich, Chief Plant Industry Division
Ohio Department of Agriculture
8995 East Main Street
Reynoldsburg, OH 43068-3399