



Food and Drug Administration  
Baltimore District Office  
Central Region  
900 Madison Avenue  
Baltimore, MD 21201-2199  
Telephone: (410) 962-3396  
FAX: (410) 962-2219

Refer to: CFN 1124873

December 4, 1998

**WARNING LETTER**

**CERTIFIED MAIL**  
**RETURN RECEIPT REQUESTED**

Mr. William Won Huang, President  
Won Trading, Inc.  
350 Morse Street NE  
Washington, DC 20002

Dear Mr. Huang:

During a Food and Drug Administration inspection of your food warehouse conducted November 24 through December 2, 1998, our investigator observed significant insanitary conditions. At the conclusion of the inspection you were issued a Form FDA-483, Inspectional Observations. The FDA-483 listed the following insanitary conditions observed in your warehouse at the time of the inspection that cause food products stored in your to be adulterated within the meaning of Section 402(a) (4) of the Food, Drug, and Cosmetic Act (the Act):

1. At least 1 (one) live mouse on a glue board in your warehouse.
2. At least 8 (eight) dead mice on glue boards or in traps at various locations throughout your warehouse.
3. One lot of Tempura Batter Mix was found to have fluorescent stains indicative of rodent urine and apparent rodent gnawed holes.
4. Two lots of Mung Beans were found to have fluorescent stains indicative of rodent urine and apparent rodent gnawed holes.
5. A bag of Barley was found to have fluorescent stains indicative of rodent urine.
6. Over 450 rodent excreta pellets were observed in various locations throughout the warehouse.
7. Various food products were improperly stored directly on the floor against the walls in a manner which prohibit adequate cleaning and control of rodents or insects.

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This letter is not intended to be an all-inclusive list of deficiencies at your facility. It is your responsibility to ensure adherence to each requirement of the Act and regulations.

Your voluntary corrections of some of the above deviations by voluntarily destroying the lots of tempura mix, mung beans, and barley is appreciated and has been documented in our records.

You must notify this office in writing, within 15 working days of receipt of this letter, of the specific steps you have taken to correct the noted violations. If corrective action cannot be completed within 15 days, state the reason for the delay and the time within which the corrections will be completed. Further violations may result in regulatory actions such as injunction and/or prosecution.

Your reply should be sent to the Food and Drug Administration, Gerald W. Miller, Compliance Officer, U.S. Food and Drug Administration, 101 West Broad Street (Suite 400), Falls Church, Virginia 22046-4200, phone number 703-235-8440, extension 504.

Sincerely,



Elaine Knowles Cole  
Director, Baltimore District