

EXAMPLE

United States Food and Drug Administration
Los Angeles District Office

Notice of FDA Action¹

Entry Number: 112-9861457-6

Notice Number: 2
November 6, 1996

Filer:
FBN Freight Services Attention: George
500 Canal St.
New Orleans LA 70130

>

Port of Entry: 2704, Los Angeles,
Carrier: NOL RUBY
Entry Date: November 2, 1996
Arrival Date: November 4, 1996

Importer of Record: Shipley's Donut Shop Inc., Lafayette, LA
Consignee: a: Shipley's Donut Shop Inc., Lafayette, LA
 b: Specialty Commodities Inc., Fargo, ND

HOLD DESIGNATED

Notify FDA of Availability

Summary of Current Status of Individual Lines

Document: 1

Invoice: PRAC004

@ LINE	Product Description	Quantity	Current Status
ACS/FDA * a 001/001	PINEAPPLE, DEHYDRATED	500 CT	RELEASED 11-6-96
* a 002/001	DEHYDRATED GINGER SLICES	10 KG	Product Collected by FDA 11-06-96
* b 003/001	PAPAYA, DEHYDRATED	10 KG	Detained 11-06-96

* = Status change since the previous notice. Read carefully the sections which follow for important information regarding these lines.

@ = Consignee id

¹This example of a Notice of FDA Action is a model and should not be considered all inclusive. The format and wording in the actual Notice of FDA Action issued by districts from the Operational and Administrative System for Import Support (OASIS) may appear different.

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FDA will not request redelivery for examination or sampling, if the products not released by FDA are moved, following USCS conditional release to a location within the local metropolitan area or to a location approved by the FDA office at the number below.

All products in this entry not listed above may proceed without FDA examination. This notice does not constitute assurance the products involved comply with provisions of the Food, Drug, and Cosmetic Act or other related acts, and does not preclude action should the products later be found violative.

Please provide documentation concerning all products in this entry to the FDA office below. Include the USCS document (e.g., CF-3461 or CF-7501) and commercial invoice for these products, annotated to show the ACS/FDA line numbers sent electronically.

Also, advise FDA upon actual availability, and include date, location, and warehouse control number, where applicable, for all lines in this entry.

Jennifer A Thomas, Inspector
 U.S. Food & Drug Administration (213) 555-1212
 2nd and Chestnut Streets (HFR-MA100)
 Philadelphia, PA 19106

DETENTION WITHOUT EXAMINATION

The following products are subject to refusal pursuant to the Federal Food Drug and Cosmetic Act (FD&CA), Public Health Service Act (PHSA), or other related acts in that they appear to be adulterated, misbranded or otherwise in violation as indicated below:

LINE	Product Description	Respond	By
003/001 DEHYDRATED	Product: November 26, 1996		PAPAYA,

FD&CA Section 402(a)(1), 801(a)(3); ADULTERATION

The article appears to be held in a container containing a poisonous or deleterious substance which may render it injurious to health.

FD&CA Section 402(a)(2)(B), 801(a)(3); ADULTERATION

The article appears to be a raw agricultural commodity that bears or contains a pesticide chemical which is unsafe within the meaning of Section 408(a). The article appears to contain quinalphos.

Jennifer A Thomas, Inspector
 U.S. Food & Drug Administration (213) 555-1212
 2nd and Chestnut Streets (HFR-MA100)
 Philadelphia, PA 19106

You have the right to provide oral or written testimony, to the Food & Drug Administration, regarding the admissibility of the article(s) or the manner in which the article(s) can be brought into compliance. This testimony must be provided to FDA on or before the dates shown above.

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SAMPLES COLLECTED

LINE ACS/FDA	Product Description	Est. Cost
001/001 \$ 15.00	PINEAPPLE,	DEHYDRATED

Sample: 10 KG Collected 1 KG from each of 10 cartons

LINE ACS/FDA	Product Description	Est. Cost
002/001 \$.23	DEHYDRATED GINGER SLICES	

Sample: .1 KG Collected approximately 4 ounces from one carton.

LINES RELEASED

LINE ACS/FDA	Product Description
001/001	PINEAPPLE, DEHYDRATED

These products are released. This notice does not constitute assurance that the product released complies with all provisions of the Food, Drug, and Cosmetic Act, or other related Acts, and does not preclude action should the product later be found violative.

Notice Prepared by: Thomas J. DiNunzio (QA5)
 U. S. Food and Drug Administration