

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

FOOD CANNING ESTABLISHMENT REGISTRATION

FOR FDA USE ONLY	
FCE No.	Date Received by FDA

1. TYPE OF SUBMISSION

- Initial Registration
- Relocation (*new registration required*) Enter Current FCE: (*If applicable*) _____
- Change of Registration Information Enter Current FCE: (*If applicable*) _____

Specify Type of Change: _____

2. FOOD PROCESSING PLANT LOCATION

Establishment Name _____

Number and Street _____

City and State or Province (or other Subdivision) _____

Zip (or other Postal Code) _____ Country (if other than U.S.) _____

Telephone No. _____ Telefax No. _____

3. PREFERRED MAILING ADDRESS

Same as Plant Location

Establishment Name _____

Number and Street _____

City and State or Province (or other Subdivision) _____

Zip (or other Postal Code) _____ Country (if other than U.S.) _____

Telephone No. _____ Telefax No. _____

4. LOW ACID AND/OR ACIDIFIED FOODS PROCESSED AT THIS LOCATION

Food Product Name, Form or Style, and Packing Medium
Listing products produced at this location is not a process filing.

(Do not list meat and poultry foods under the jurisdiction of the Food Safety and Inspection Service of the U.S. Department of Agriculture.)

(Check One)

	Low-Acid	Acidified
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>
_____	<input type="checkbox"/>	<input type="checkbox"/>

PLEASE NOTE THE FOLLOWING:

- The requester hereby presents and acknowledges that the company is aware that in making this request the company is subject to the terms and provisions of Title 18, Section 1001, United States Code which makes it a criminal offense to falsify, conceal, or cover up a material fact; make any false, fictitious, or fraudulent statement or representation; or make or use any false writing or document knowing the same to contain any materially false, fictitious, or fraudulent statement or entry.
- Subject to the terms of 21 CFR 108.25 (c)(1) and (2) and 108.35 (c)(1) and (2), no commercial processor shall engage in the processing of low-acid or acidified foods until the completed forms FDA 2541 and 2541d, 2541e, 2541f or 2541g have been filed with the FDA within the applicable time frames specified in these regulations.
- Forms, Instructions, regulations, and information can be secured online at <http://www.fda.gov/Food/GuidanceRegulation/GuidanceDocumentsRegulatoryInformation/default.htm>
- For more information, contact the LACF Registration Coordinator by e-mail at lacf@fda.hhs.gov

**Food and Drug Administration
LACF Registration Coordinator (HFS-303)
Center for Food Safety & Applied Nutrition
5001 Campus Drive
College Park, Maryland 20740-3835**

5. ESTABLISHMENT CONTACT PERSON

Name of Contact and Business Address: _____

Position: Owner Technologist Manager Director President/Vice President Other Employee Authorized Third Party Phone Number: _____

FAX Number: _____ E-mail Address: _____ Signature: _____ Date: _____

LACF Contact Information

For more information, contact the LACF Registration Coordinator by e-mail at LACF@FDA.HHS.GOV or phone: 240-402-2411.

For paper submissions, send completed forms to:

Food and Drug Administration
LACF Registration Coordinator (HFS-303)
Center for Food Safety and Applied Nutrition
5001 Campus Drive
College Park, MD 20740-3835

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF ADDRESS BELOW.

PAPERWORK REDUCTION ACT NOTICE

The time required to complete this collection of information is estimated to average 10 minutes per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

FDA PRA Staff
Office of Operations
Food and Drug Administration
email to PRASStaff@fda.hhs.gov

“An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number.”