Form Approved: OMB No. 0910-0498; Expiration Date: 3/31/2015

	Department of Health and Human Services Food and Drug Administration Center for Food Safety and Applied Nutrition		FOO	FOOD EXPORT CERTIFICATE APPLICATION			te	
1.	Food Manufacturer Information							
	Manufacturer name			Doing business as name (If other than "Manufacturer left, and you wish this name to appear on the export c				
	State License/Registration number		Address					
	Contact person name							
	Contact phone/fax		City		State or Pi	rovince	ZIP/postal code	
	Contact email				Country			
2.	2. Exporting Company Information (if applicable)							
	Export company name							
	State License/Registration number	Address						
	Contact person name	ontact person name		City		rovince	ZIP/postal code	
	Contact phone/fax/or email	Contact phone/fax/or email					<u>I</u>	
3.	Shipment Description			l	I			
	Product	Common Nam	е	Manufacturer		Descrip	tion/Comments	
С	ontinue on additional page(s) as nee	ded.						
4.	Intended Destination of Shipme	nt (Country)						
	Name of country							
5.	5. Send Certificate To Manufacturer			Distributor Other (provide the following information)				
	Firm name	Address						
			Citv	City		ate	ZIP/postal code	
	Contact person name		Country				, , , , , , , , , , , , , , , , , , ,	
6.	. Send Certificate Via							
	Carrier name (U.S. Mail, FedEx, etc.)			Account number (If application	able)			
7.	Fees			1				
	Fees are \$10 per certificate, a billed upon receipt of this app		Copies of ce	ertificate: x	Fee/copy	= Total \$		

o. Labello)							
Attach an original or an electronic copy of any applicable product label(s). A fax copy is acceptable only if it is readable.							
9. Verification							
The undersigned verifies that all ingredients are approved for use by FDA or appear on the GRAS list, and each product is intended for human consumption and is available for sale in the U.S. without restriction.							
Signature	Name and Title	Date					

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FOOD EXPORT CERTIFICATE APPLICATION Instructions

For Manufacturers/Distributors

- The Manufacturer/Distributor fills out the application information describing the consignment, manufacturer (note that different processing facilities of the manufacturer may be listed on the table describing the foods), where and how to send the certificate, optional information as needed, and applicant signature, name, and date.
- 2. The Manufacturer/Distributor submits the application (by mail, fax, email), along with labels as applicable. For contacts, refer to http://www.fda.gov/Food/InternationalActivities/Exports/ExportCertificates/UCM151486.htm

For FDA Officials

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- 3. FDA Official reviews the application to be sure all the blanks are filled in properly, verifies manufacturer's license or registration, and investigates inspection data on the listed products.
- 4. The Official may require an inspection prior to issuance of the export certificate.

- 5. The Official prints the Certificate on watermarked Department letterhead, assigns a unique registration number and expiration date, signs, dates, seals, and issues the Certificate as indicated.
- 6. The Official maintains in his records an identical copy of the signed Certificate, marked "Copy" for a period of at least two years.
- 7. In the event that the Manufacturer fails to comply with the law as stated on the Certificate, the Official will reject the application and promptly notify the Manufacturer that the Certificate cannot be issued.

After the Certificate Has Been Issued

- 8. The Manufacturer/Distributor forwards the Certificate to the foreign Importer and verifies that it is acceptable.
- 9. If the Certificate is not acceptable, the Exporter notifies the FDA Official that the certificate has not been accepted by the Importer, and the Official will promptly attempt to reconcile the issue with the Importer.

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.

The burden time for this collection of information is estimated to average 1 hour 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:

Department of Health and Human Services Food and Drug Administration Office of Chief Information Officer Paperwork Reduction Act (PRA) Staff 1350 Piccard Drive, Room 400 Rockville, MD 20850

"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 control number."