## DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration

# Food Process Filing for Low-Acid Aseptic Systems (Form FDA 2541g)

**Note:** There are separate process filing forms for each of the following: Food Process Filing for Low-Acid Retorted Method (Form FDA 2541d); Food Process Filing for Acidified Method (Form FDA 2541e); Food Process Filing for Water Activity/Formulation Control Method (Form FDA 2541f); and Food Process Filing for Low-Acid Aseptic Systems (Form FDA 2541g).

USE FDA INSTRUCTIONS ENTITLED "Instructions for Paper Submission of Form FDA 2541a (Food Process Filing for Low-Acid Aseptic Systems)" FDA USE ONLY Date Received by FDA: / (MM/DD/YYYY) Food Canning Establishment (FCE) Number (Enter number assigned by FDA) Submission Identifier (SID) (YYYY-MM-DD/SSS) 20 - - / A. Product Information A.1 (Food Product Group) (Continued) Note: Section A.1 (Food Product Group) requests optional information. **Mixed Vegetables** Mixed Vegetables (e.g., carrots and peas, etc.) 1. (Optional) Select one Food Product Group. If there is no single best Food Product Mixed Vegetables as a Juice or Drink (e.g., carrot and green bean juice, etc.) Group that applies, select Other. Baby Food (infant/junior foods including infant formula) □ Nut Spread and Nut Topping □ Other Vegetables ☐ Bakery Products (canned brown bread, bakery glazes) Rice, Wheat, Oat or Grain (liquid form – ready-to-eat such as grits) Berry/Citrus/Core Fruit as a Jam, Jelly, Preserve, Drink, Syrup, Topping **Root and Tuber Vegetables** Beverage Base Breakfast Foods (liquid form – ready-to-eat, such as porridge, gruel) Root/Tuber Vegetables (e.g., carrots, leeks, potatoes, etc.) Cheese (does not include soy cheese or imitation dairy) Root/Tuber Vegetables as a Juice or Drink (e.g., carrot juice, etc.) ☐ Cocoa ☐ Coffee/Teas (excluding herbal and botanical teas) ☐ Dairy (milk-based) ☐ Dietary Supplement and/or herbal and botanical teas Sweet Goods/Dessert (liquid form – ready-to-eat, such as pudding) ☐ Soup Dressings/Condiments (e.g., salad dressing, chutney, salsa, pepper sauce, etc.) ☐ Vine/Other Fruit as a Jam, Jelly, Preserve, Drink, Syrup, Topping ☐ Wine Cooler Fruit as a Vegetable ☐ Other (Specify below) Fruit as a Vegetable (e.g., eggplant, pumpkin, etc.) Fruit as a Vegetable Juice or Drink (e.g., eggplant juice, pumpkin juice, etc.) 2. Enter Product Name (e.g., Cheese Sauce (with Jalapeno Pieces), Pudding (Vanilla or Gelatin, Pudding Filling for Pies, Pie Filling (liquid form ready-to-eat such as apple pie Strawberry), etc.). filling, etc.) Gravies/Sauces (spaghetti sauce, mushroom gravy) ☐ Imitation Dairy (includes soy-based products) Imitation/Pit/Mixed/Subtropical Fruit as a Jam, Jelly, Preserve, Drink, Syrup, Topping 3. What is the form of the product? (Select all that are applicable) Liquid (i.e., all liquid no solids) Liquid with Solids (e.g., diced, chunks, pieces, etc.) Leafy/Stem Vegetables ☐ Paste/Puree ☐ Leafy/Stem Vegetable Other (Enter product form) Leafy/Stem Vegetable as a Juice or Drink (e.g., spinach juice, etc.) Meal Replacement/Medical Foods (e.g., supplemental liquid nutrition, etc.)

Food Process Filing for Low-Acid Aseptic Systems (Form FDA 2541g)						
A. Product Information (Continued)	C. Container Type (Continued)					
4. What is the packing medium? (Select all that are applicable)	2.  Flexible Pouch					
☐ Brine ☐ Cream/Sauce/Gravy ☐ Oil ☐ Syrup ☐ Water ☐ None	a) What is the shape of the container? (Select one)					
Other (Enter packing medium)	☐ Flat pouch ☐ Gable top ☐ Gable top/side gusseted ☐ Gusseted					
Continue to Section B.	☐ Irregular (Attach a picture or schematic. Provide name or a brief description of attachment below.)					
B. Governing Regulation (Refer to the precursor questions in the instructions)	Other (Attach a picture or schematic. Provide name or a brief description of attachment below.)					
X Low-acid (21 CFR 108.35 and 21 CFR Part 113)						
Continue to Section C.	3. Semi-Rigid					
	a) What is the shape of the container? (Select one)					
C. Container Type (Select one)	☐ Bowl ☐ Cylindrical ☐ Oval ☐ Rectangular ☐ Tray					
Note: If the product is not packaged in one of the container types identified below, select Other.	☐ Irregular (Attach a picture or schematic. Provide name or a brief description of attachment below.)					
1. Aluminum/Tinplate/Steel Can						
a) What is the shape of the container? (Select one)	Other (Attach a picture or schematic. Provide name or a brief description of					
☐ Cylindrical ☐ Oval ☐ Rectangular	attachment below.)					
☐ Irregular (Attach a picture or schematic. Provide name or a brief description of attachment below.)	b) Is this a single piece container?					
	Yes (Continue to d) No (Continue to c)					
<ul> <li>Other (Attach a picture or schematic. Provide name or a brief description of attachment below.)</li> </ul>	c) Is this a compartmentalized container?					
uttuomment selemy	Yes How many compartments? No					
b) How many pieces are used to construct the container? (Select one or more choices, as applicable)  i.	d) What is the predominant material used to make the body of the container? <i>(Select one)</i> HDPE (high-density polyethylene) HDPP (high-density polypropylene)  Paperboard PET (polyethylene teraphthalate)  Other <i>(Enter material)</i>					
☐ Cemented ☐ Welded	Note: If "Yes" is selected as a single piece container in question 3.b, continue to Section D.					

#### C. Container Type: 3. Semi-Rigid (Continued) D. Container Size e) What is the predominant material used to make the lid of the container? (**Select one**) Note: Section D.1 (dimensions) is required information; however, volume is acceptable for container size in lieu of container dimensions if package sterilizer □ Aluminum/Steel HDPE (high-density polyethylene) does not depend on the container dimensions. Section D.3 (net weight) is optional information. HDPP (high-density polypropylene) ☐ Nylon PET (polyethylene teraphthalate) ■ Not Applicable 1. Dimensions: a) \_\_\_\_ Diameter \_\_\_ Height (Use for cylindrical shapes) (see accompanying Other (Enter material) instructions for proper coding) b) \_\_\_\_ Length \_\_\_ Width \_\_\_ Height/Thickness (Use retangular shapes, f) How is the lid sealed to the body of the container? (Select one) pouches, or irregular shapes) (see accompanying instructions for proper coding) ☐ Double Seam ☐ Induction Weld 2. Volume: \_ \_ \_. (Select one) ☐ Snap On □ Threaded Closure ☐ Press Twist Fluid Ounces Gallons Liters Milliliters ☐ Ultrasonic Seal Not Applicable 3. Net Weight (Optional): \_\_\_\_\_ (enter in ounces) Other (Enter seal type) Continue to Section E. 4. Other (Enter container type) E. Product Processing Method: Thermally Processed using Aseptic Systems **Product Sterilization** a) Attach schematic or picture of container. (Provide name or a brief description of a) What is the finished equilibrium pH of the product after processing? \_ \_ . \_ \_ attachment below.) b) Heating Method Direct Heating i. (Select one) Indirect Heating b) Specify the material that, based on weight, is the predominant material used to make the ii. What is the Thermal Expansion Coefficient? \_\_\_\_ container stock. This is the material that constitutes the highest weight value of the iii. Where is the product flow rate controlled? (Select one) container stock. ☐ Before the heater (Continue to b.iii.1) ☐ After the heater (Continue to c) (1) Volume Expansion Factor: . (Direct Heating Only) c) Specify the predominant material used to make the lid. This is the material that constitutes the highest weight value of the lid stock. If the container does not have a lid, c) What is the Manufacturer's name and the model number of the Product Sterilization specify Not Applicable. System? d) Specify the method used to seal the lid to the body of the container. If the container does d) What is the Process Source of the Product Sterilization System? not have a lid, specify Not Applicable. (Attach Process Source Document. Provide name or a brief description of attachment below.) Continue to Section D. e) What is the date of the Process Source Document of the Product Sterilization System (mm/dd/yyyy)? \_\_ / \_\_ / \_\_\_\_ Continue to Section F.

Food Process Filing for Low-Acid Aseptic Systems (Form FDA 2541g)

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## F. Product Critical Factors: (Complete all product critical factor questions as delineated by process authority to assure commercial sterility.)

1. Does the product contain particulates?
☐ Yes
(Attach supporting documentation and validation reports. Provide name or a brief description of attachment below.)
(Continue to a)
☐ No (Continue to F.2)
a) Is controlling particulate size a critical factor?
☐ Yes (Continue to b) ☐ No (Continue to F.2)
b) What is the shape and dimension of the particulate size to be controlled? If more than one, list all that apply.
2. Does the product contain any dry ingredients that are hydrated before processing the product?
☐ Yes (Continue to a) ☐ No (Continue to F.3)
a) What is the minimum % moisture of the hydrated dry ingredients before processing?
3. Does the % total solids affect the heating of the product during processing?
☐ Yes (Continue to a) ☐ No (Continue to F.4)
a) What is the % total solids?
4. Is the finished equilibrium pH of the product after processing (identified in Section E) critical to the process?
☐ Yes ☐ No
5. What is the flow correction factor used during the scheduled process? (Select one)
a) 0.5 (Laminar) (Continue to Section G)
b) 0.83 (Turbulent) (Continue to F.6)

### F. Product Critical Factors (Continued)

6. Answer the following questions if the flow correction factor you identified in question F.5 is 0.83 (Turbulent)									
a) What is the instrument used to measure the consistency/viscosity?									
b) What is the temperature when you measure the consistency/viscosity?(enter in Fahrenheit)									
c) What is the consistency/viscosity?									
What is the unit of measure? (Select one) Centipoise									
Other (Enter the units of measure)									
d) What is the specific gravity?									
7. Is starch added to maintain consistency/viscosity of the product?									
☐ Yes (Continue to a-b) ☐ No (Continue to F.8)									
a) What is the maximum % starch added?									
b) What type of starch is added?									
8. Are other binders added?									
☐ Yes (Continue to a-b) ☐ No (Continue to F.9)									
a) What is the maximum % binder?									
b) What is the type of binder added?									
9. Is syrup strength a critical factor that needs to be controlled during processing?									
☐ Yes (Continue to a) ☐ No (Continue to Section G)									
a) What is the brix measurement?									
Continue to Section G.									

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## G. Package Sterilization System and Supplemental Information

1.	Sterilization System									
a) What is the Manufacturer name and the model number of the sterilization system use to sterilize the packaging of the product?										
	b) What is the Process Source of the Package Sterilization System?									
	c) What is the date of the Process Source of the Package Sterilization System (mm/dd/yyyy) / / /									
d) Supplemental Submission Identifier (SUP SID)										
	(Attach Supplemental Information. Provide name or a brief description of attachment below.) (See accompanying instructions.)									
2.	Sterilization System									
	a) What is the Manufacturer name and the model number of the sterilization system used to sterilize the packaging of the product?									
	b) What is the Process Source of the Package Sterilization System?									
	c) What is the date of the Process Source of the Package Sterilization System (mm/dd/yyyy) / / /									
	d) Supplemental Submission Identifier (SUP SID)									
	(Attach Supplemental Information. Provide name or a brief description of attachment below.) (See accompanying instructions.)									
_										
3.	Sterilization System									
	a) What is the Manufacturer name and the model number of the sterilization system used to sterilize the packaging of the product?									
	b) What is the Process Source of the Package Sterilization System?									

<b>G. Package Sterilization System and Supplemental Information:</b> 3. Sterilization System (Continued)
c) What is the date of the Process Source of the Package Sterilization System (mm/dd/yyyy) / / /
d) Supplemental Submission Identifier (SUP SID)
(Attach Supplemental Information. Provide name or a brief description of attachment below.) (See accompanying instructions.)
4. Sterilization System
a) What is the Manufacturer name and the model number of the sterilization system used to sterilize the packaging of the product?
b) What is the Process Source of the Package Sterilization System?
c) What is the date of the Process Source of the Package Sterilization System (mm/dd/yyyy) / / /
d) Supplemental Submission Identifier (SUP SID)
(Attach Supplemental Information) (see accompanying instructions)

Continue to Section H.

### H. Scheduled Process

In the section below, please do NOT enter decimal points. They are already on the form. No blank spaces are allowed, therefore, enter leading zeros, where necessary.

Col. 1	Col. 2	Col. 3	Col. 4	Col. 5	Col. 6 Col. 7		Col. 8	Col. 9	
Process No	Hold Tube Section	Inside Diameter of Hold Tube Section	Hold Tube Section Length	Initial Temperature  (*only for heating with control of flow rate before the heater)	Process Time	Temperature (at exit of final hold tube section)	Fo (F18/250)	Maximum Product Flow Rate	
Number	Number	Inches	Inches	°Fahrenheit	Seconds	°Fahrenheit	Minutes	Gal/min	
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I Note: Under the terms and provisions of Title 18, Section 1001, United States Code, in any matter within the jurisdiction of the executive branch of the Government of the United States it is a criminal offense to falsify, conceal, or cover up a material fact, make any materially false, fictious, or fraudulent statement or representation; or make or use any false writing or document knowing the same to contain any materially false, fictious, or fraudulent statement or entry.  If your process filing appears to be fabricated, the product on this form will not be in compliance with 21 CFR 108.35(c)(2). A process filing appears fabricated  Signature  Signature  Country (other than U.S.)  Date  Telephone No.	Food Process Filing for Low-Acid Aseptic Systems (Form FD	OA 2541g)							
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Establishment Name State or Province Country (other than U.S.) Date Telephone No.	ruii Name (Please Type or Print)		Signature						
	Establishment Name	State or Province	1	Country (other than U.S.)	Date	Telephone No.			

#### **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) Human Foods Program 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 45 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 PRAStaff@fda.hhs.gov

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