

**Packaging and Ingredients
Supplier Handbook
(Self-Manufacturing Plants)
Supplemental Guide**



LucerneFOODSSM

AVAILABLE AT:

[ALBERTSONS FOR SUPPLIERS](#)

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WELCOME

Thank you for your interest in exploring business opportunities with Albertsons Companies. We are pleased that you are showing interest in doing business with us. Albertsons Companies has continually upheld its commitment to provide added value on world-class quality products, combined with award-winning service to all our customers. We are very excited to share this dynamic vision with our business partners as well.

A. REQUIRED BUSINESS DOCUMENTS

1. A completed MNDA
2. A completed [W-9](#) form with Supplier's tax identification number and physical address.
3. A [Broad Form Vendor's Endorsement \(ISO Form CG 2015\)](#)
4. Certificate of Insurance (COI)
5. Sample Supplier invoice showing net payment terms.
6. Award Letter
7. Supply Agreement (SSA)
8. Product Undertaking Agreement (PUA) – specific to awarded business

B. ITEM SET-UP & MAINTENANCE

Albertsons Item Codes

Albertsons Companies uses M3 manufacturing systems. All items will have a unique 6-digit numerical code, which will correlate to the ingredient or packaging material ("materials") being ordered. Suppliers must reference Albertsons' M3 number in the pricing, invoicing, and shipping processes. Suppliers will be required to provide their item code and description (if any) to Albertsons. Albertsons will then assign item codes and share back to Supplier. *Example below:*

Albertsons M3 Data			Supplier Item Data	
Supplier Name	Albertsons M3 Item #	Albertsons M3 Item Description	Supplier Item #	Supplier Item Description
Supplier XYZ	522550	ingredient item abc	xxxxx-q	ing item ab54
Supplier XYZ	522550	packaging item abc	xxxxx-q	pack item ab54

Required Item Information

Suppliers will be required to complete the Supplier Item Information file. *Example below:*

All Suppliers								Packaging Suppliers				Printed Packaging Suppliers		
Supplier	Item Group	Item Number	Item Description	UOM	Order Lead Time (must match RFP)	Minimum Order QTY (must match RFP)	Minimum Production QTY (if any)	Common or Unique Size	Common or Unique Material	Material Type	Minimum Run QTY	Printing Type	# Colors	Minimum Print QTY
Supplier A			packaging abc	M (000)	14 days	1,000.0	stock	Common	Common	HDPE resin	stock			
Supplier B			packaging efg	Each	14 days	50,000.0	100,000.0	Common	Unique	x% PCR HDPE resin	100,000.0			
Supplier C			packaging xyz	Each	14 days	25,000.0	stock	Common	Common					
Supplier D			ingredient abc	LB	21 days	100.0	N/A			PP resin	25,000.0	dry-offset	6	25,000.0

Artwork Design

Specific to printed packaging and labels, Suppliers will be required to provide dielines and complete PWO (project work order) files at the start of any business, and as requested from time-to-time as the business may require. Example below:

PLEASE NOTE THE BELOW:						Final Materials Notes:	
• Please submit editable dielines in Adobe Illustrator file format at 100% size. Make sure they are vector files. • After completing the below, please fill out Printer Spec Form for EACH printer listed. (i.e., if you have 5 total printers, you need to submit 5 printer specs). • Final files from Albertsons are NOT to be altered without consulting with the Albertsons team. • Artwork reviewer is usually a press or prepress persons familiar with artwork and print process						• Shipping direct to printer is HIGHLY recommended to prevent any delay. • If you decide to have press ready files sent to Supplier instead of Printer, you are responsible for relaying files to your printer to ensure there are no delays. • For physical proofs, a choice of "Send to Both Supplier and Printer contacts" will result in additional charges	
SUPPLIER TO COMPLETE ALL SECTIONS BELOW WHICH ARE SPECIFIC TO ARTWORK REVIEWS AND RECEIVING FINAL PRESS READY FILES AND PRINTED TARGET PROOFS						SUPPLIER TO COMPLETE ALL SECTIONS BELOW	
ARTWORK ROUTING CONTACTS						FINAL MATERIALS SHIPPING	
SUPPLIER ARTWORK CONTACT INFORMATION - Required		PRINTER #1 ARTWORK CONTACT INFORMATION - Required (DO NOT PLACE SUPPLIER INFO HERE)		PRINTER #2 ARTWORK CONTACT INFORMATION		PRINTER #1	
SUPPLIER COMPANY NAME:		PRINTER COMPANY NAME:		PRINTER COMPANY NAME:		PRINTER #2	
SUPPLIER CONTACT: (Artwork Reviewer)		PRINTER CONTACT NAME: (Artwork Reviewer)		PRINTER CONTACT NAME: (Artwork Reviewer)		MUST CHOOSE ONE	
SUPPLIER ARTWORK REVIEWER PHONE #:		PRINTER ARTWORK REVIEWER PHONE #:		PRINTER ARTWORK REVIEWER PHONE #:		MUST CHOOSE ONE	
SUPPLIER ARTWORK ADDRESS, CITY, STATE, ZIP:		PRINTER ARTWORK ADDRESS, CITY, STATE, ZIP:		PRINTER ARTWORK ADDRESS, CITY, STATE, ZIP:		IF AN ALTERNATE SHIP-TO IS NEEDED PLEASE PROVIDE CONTACT ADDRESS	
SUPPLIER ARTWORK CONTACT EMAIL:		PRINTER CONTACT EMAIL:		PRINTER CONTACT EMAIL:		EXTRA CHARGES ARE INCURRED IF SENDING MORE THAN ONE PHYSICAL PROOF	
WILL BOTH SUPPLIER AND PRINTER REVIEW ARTWORK? (If supplier only they will be reviewing on behalf of both printer and supplier)		OKAY TO CONTACT PRINTER?		OKAY TO CONTACT PRINTER?		MUST CHOOSE ONE	
MUST CHOOSE ONE		FTP SITE INFORMATION: (If Applicable)		FTP SITE INFORMATION: (If Applicable)		MUST CHOOSE ONE	

SUPPLIER TO COMPLETE ALL SECTIONS BELOW									
SUBSTRATE TYPE	DIE LINE FILE NAME (Specify dieline file name PEP packaging artwork component) MUST MATCH EXACT FILE NAME OF PROVIDED DIE LINES	DIE LINE DIMENSIONS IN INCHES	PRINTER NAME AS LISTED ABOVE	SIN # - This is REQUIRED (SIN is the Plant Approval # provided by Albertsons QA) Format: "SXXXX"	SIN Placement (SIN MUST be inkjetted or embedded)	COO - Country of Origin (Must list each product as a separate line for each COO) Only fill this out if COO	COO Placement (If needed, must inkjet or embed on art) Embedded info will appear on the artwork	Supplier/Manufactures Codes (supplier's item codes to put on the artwork - Optional)	Supplier / Printer item notes (any additional special notations per SKU - Optional) (put USDA EST here, if required)
Foil	LID.ai	2.25"X4"	Printer Name	S3452	Embed on Art	Product of USA	Inkjet	12465974	Do not type in yellow marked box on dieline
Plastic	CUP.ai	12"X4"	Printer Name	S3452	Inkjet	Product of Chile	N/A	R36974	Add supplier code on left dust flap
MUST CHOOSE ONE					MUST CHOOSE ONE		MUST CHOOSE ONE		


An important aspect of the design process is the utilization of BLUE, the electronic project, proofing and data management system. All projects are run through this system allowing for total control over the flow and tracking of information and access of up to the minute scheduling tools by anyone dedicated to the project including our Suppliers and printers. The system is quick and easy to use and very intuitive. Example of landing page below:

Paste example here

C. QUALITY ASSURANCE REQUIREMENTS AND PROCEDURES

All suppliers must complete Albertsons Companies required 8020_0570 Supplier Questionnaire and QA paperwork, which will be received during the RFP process / prior to any business being conducted. All ingredient suppliers must adhere to SafetyChain requirements.

8020 Sample

									
Albertsons Companies, as part of their Supplier Approval process, would like to know more about your company's personnel, manufacturing processes, programs and procedures. Please answer the following questions and provide materials									
General Information									
Date									
Company Name:									
Address:									
City:									
Type of Company*:					Manufacturer <input type="checkbox"/> Packager <input type="checkbox"/> Broker <input type="checkbox"/> Distributor <input type="checkbox"/>				
Regulatory Registration # (ie. IMS, USDA estab #, ICSSL)									
Site Location(s) Address if different from above:									
<small>*Note if Manufacturer, Packager, Broker or Distributor. If Broker or Distributor, a separate form must be submitted for each manufacturing or co-packing location.</small>									
Personnel Information									
SALES					Name:				
					Title:				
					Email:				
					Phone:				
QA/TECHNICAL					Name:				
					Title:				
					Located at Manufacturing Site or Corporate Location?				
					Email:				
CUSTOMER SERVICE					Name:				
					Title:				
					Email:				
					Phone:				
RECALL CONTACT					Name:				
					Title:				
					Email:				
					Phone:				
Item Information: Please identify all items supplied to Albertsons. Attach a separate list if necessary.									
DESCRIPTION/SIZE									
1									
2									
3									
4									
5									
6									
7									
8									
9									
10									
Certifications: Please list certifications (i.e., SQF, BRC, Organic) that you currently have or are in the process of attaining.									
Certification					Date	Score	Comments		

Manufacturing Location:				Manufacturing Location:	Question	Answer	Additional		
Sourcing	Question	Answer	Additional Comments	Foreign Material Control	Does the facility have a documented foreign material program? List the Foreign Material Control Devices and procedures.				
	Is the supplied material sold directly from the manufacturing company? If no, please clarify if a broker/3 rd party agent is used.				Are all materials (raw, WIP-work in progress, or finished product) prior to packaging protected from contamination with tight seals?				
	Is the supplied material sourced only from the US? If NO, list the countries				Does the facility have a map or list, which is audited periodically of all glass, ceramics, brittle or hard plastics locations?				
	Is your facility and outside warehouse if used registered under the FDA Food Facility Registration program?				Does the facility have a written procedure & documentation for routine calibration of all measuring devices including thermometers and other instruments used?				
	Are you following the Foreign Supplier Verification Program for the product(s) you are supplying?				Does the facility have a written physical, microbiological and or chemistry testing protocols for the handling of ingredients or products that are not within specifications?				
Quality Systems	Indicate which regulatory jurisdiction and code the products fall under. For non-US please indicate regulatory body and country.			Testing	Does the facility produce any items for the customer (Albertsons) that have a history of EMA (Economically Motivated Adulteration)? If yes, list your control measures.				
	Does the facility have a Food Safety and Quality Manual?				Food Defense	Does the facility have a written Food Security Program based on site specific hazards with risk assessments?			
	Does the facility have a documented training program?					Does the facility restrict access to all areas of the plant to only authorized personnel?			
	Does the facility have an internal systems audit program in place? List audits and frequency.					Sanitation	Does the facility have a Master Sanitation Schedule?		
	Does the facility have a supplier and raw material approval program?						Does the facility have written Standard Sanitation		
Does the facility have an effective document control system to ensure correct versions of documents and records are available and in use?			Is there a system in place for the prevention of cross contamination and contact with cleaning equipment & tools, such as labeling or color coding?						
HACCP/Food Safety System	Does the facility have a written and documented customer complaint program?			Equipment	Are all refrigeration units maintaining proper temperatures according to product type and equipped with thermometers being monitored and documented on a schedule?				
	Does the facility have an operational Food Safety/Haccp plan?				Maintenance	Is all food contact equipment clean and in good repair with no temporary repairs that may affect food safety and quality?			
	Does the facility have a Preventive Controls Qualified Individual (PCQI) responsible for the Food Safety Plan? Describe the credentials of the PCQI. If no PCQI, provide the credentials of the person responsible for the HACCP or Food Safety Plan.					Does the facility have a written Preventative Maintenance Program which covers all equipment and facilities to keep them in a positive and operative condition?			
	Have all critical control points (CCP), preventive controls, critical limits and monitoring procedures been identified, documented and implemented? List the CCP/Preventive Controls					Following maintenance or repairs, is the equipment cleaned and sanitized before food			
	Do you have a documented GMP Program?					Environmental Monitoring Program	Is there an Environmental Monitoring Program and is reviewed annually? List test organisms (e.g. Salmonella, Listeria, etc.)		
Does the facility maintain a documented Pest Control Program covering all areas of the facility and grounds? Please note in Additional information section if the facility is serviced by an outside provider or by in-house personnel.			Are results tracked and trended?						
Are there corrective actions taken to eliminate the cause of an environmental			Worker Hygiene	Does the facility have a written personal hygiene policy that is adopted by all personnel, including visitors?					
Does the facility have a written recall plan for all products and contact packaging?				Are employees with symptoms of illness or open wounds excluded from working in processing areas or around exposed product?					
Do you conduct mock recalls? If so, how often?				Are there adequate accessible hand sinks at the entrances of and in food handling and processing areas?					
Recall and Traceability	Can you trace a shipment of your product back to specific batch or lot#?			Storage and Receiving	Are all packaging materials and raw ingredients received and stored properly in a manner to protect from cross contamination from product to product?				
	Does the facility have a documented rework policy and does it ensure all rework and carryover use is recorded and traceable?				Is all in-process product and finished product properly protected to prevent contamination and preserve quality?				
	Does the facility have a formal written HOLD program for nonconforming product?				Is there a system in place to ensure that materials are only purchased from approved sources?				
	Are other allergens manufactured in the same facility as the supplied ingredient? If yes, list those allergens (Wheat, Milk, Egg, Soybean, Peanut, Tree nuts, Fish, Crustacean Shellfish, Sesame).				Does the facility have a written and documented program with training for receiving inbound and outbound food, packaging materials and finished good to ensure trailers are adequate, suitable, clean				
	Are other allergens manufactured on the same production line as the supplied ingredient? If yes, list those allergens (Wheat, Milk, Egg, Soybean, Peanut, Tree nuts, Fish, Crustacean Shellfish).				Transportation	Is the supplied material shipped "Direct" from the supplying facility to the manufacturing facility? If No, describe the process and justification of third party.			
Are sensitive or high risk products processed in the same facility as the supplied ingredients? List the products			Is the transport vehicle used to transport food items only? If NO, how is the supplied raw material adequately protected from						
Are sensitive or high risk products processed on the same production line as the supplied ingredient? List the products			Is the supplied material fully packaged, completely enclosed in a container?						
Does any packaging or packaging component (including but not limited to closures, coatings, inks, dyes, labels, pigments, adhesives, stabilizers or other additives) that you supply contain intentionally added perfluoroalkyl substances ("PFAS") or PFAS above 100 ppm? PFAS is defined as a class of fluorinated organic chemicals that contain at least one fully fluorinated carbon atom.			Will trailers be secured with seals or a padlock during transit?						
Is product produced using or has been exposed to radiation?			Does your supplied material fall under the FDA FSMA Sanitary Transportation for Human and Animal Food rule?						
Manufacturing Exposure				Food Safety and Regulatory History	Does the supplied material require time and temperature control for food safety (TCS)? If yes, explain how time/temperature is controlled, monitored and documented.				
					Have you had any recalls or withdrawals within the last 2 years? If yes, please explain.				
					Have you received any FDA Form 483/violation letters or other Food Safety Regulatory Violations, import bans or warning letters within the last 24 months?				
					Has the facility had any food safety complaints within the last 24 months? If yes, explain.				
					Have critical nonconformance/hold issues been identified in the last two years? If YES provide relevant details.				
				Did your last GFSI or Food Safety audit result in any significant deficiencies?					
				Did you complete the corrective actions in response to significant deficiencies identified above (Regulatory/Food Safety Audits)? Attach the corrective actions					



To All Ingredient Suppliers,

Below is a list of documents that are required by our regulatory and certifying agencies to be kept on hand for all ingredients that are used in this manufacturing facility.



1 Specification Sheet – including location/site of manufacturing



2 Ingredient Statement – including all sub-ingredients



3 Example label on packaging or picture of individual package with legible label



4 Country of Origin – for all components of the raw material



5 Allergen Statement – list of all allergens in the item and potential exposure to other allergens



6 Allergen Control Policy



7 Regulatory Compliance Statement - guaranteeing compliance with FD&C Act, Title 21 CFR Parts 170-189 for food-contact packaging materials, and regulated food additives comply with FDA, Food Chemical Codex, or equivalent international regulation



8 PFAS disclosure statement for packaging – guaranteeing that there are no intentionally added PFAS or PFAS greater than 100 ppm



9 Foreign Supplier Verification Program Compliance and Program Description including verification activities with records if applicable



10 Confirmation statement of compliance to FDA Facility Registration Requirements for foreign and domestic facilities including warehouses and distributors that hold food. Owners, operators, or agents in charge of either a domestic or foreign facility that is engaged in manufacturing/processing, packing or holding of food must register with the FDA .



11 Lot Coding Explanation



12 COA Example, Testing Method, Laboratory



13 Third Party Audit Report – including most recent full audit report documentation, corrective actions and certificate, GFSI audits



14 Flow Diagram and CCPs/Preventive Controls Summary Plan



15 Foreign Material Policy – including final screen size, magnet pull strengths and critical limits for metal detection



16 Food Safety Programs – including brief description ; Food Safety Controls for Food Packaging Materials. Include rework/repackaging programs.



17 Transportation and Storage Practices



18 Environmental Testing Policy and Corrective Action Procedure



19 Pesticide Testing and Monitoring Program – for agricultural, fish, fishery products ,where applicable



20 Heavy Metal Testing and Monitoring Program – for agricultural, fish, fishery products , where applicable



21 Verification Practices to Control Mycotoxins (e.g., aflatoxin, patulin) – for agricultural products with history/potential for mycotoxins



22 Evidence of Good Agricultural Practices (if applicable)



23 Annual Letter of Authenticity and any relevant testing /frequency – for any ingredient with history of EMA (Economically Motivated Adulteration)



24 Current Kosher Certificate (if applicable)



25 Current Halal Certificate (if applicable)



26 Organic Certification/Natural Status/Non GMO Certification (if applicable)



(1) GMO status statement



(2) Organic Certificate, System Plan, Fraud Prevention Plan and NOP Import Certificate



(3) Non irradiation compliance



(4) Non usage of sewage sludge compliance



(5) Disclosure of usage of any synthetic compounds

General Quality Requirements

The material shall conform to the quality parameters specified herein and shall be produced, packaged and stored in accordance with current good manufacturing practices (cGMPs) under sanitary conditions. Material failing to perform satisfactorily under plant conditions is subject to rejection.

The seller shall warrant that the material shall comply with all applicable provisions of the Federal Food, Drug and Cosmetic Acts, all other applicable federal laws and regulations and State and Local codes as amended and that such material is neither adulterated nor misbranded. A Continuing Product Guaranty shall be furnished per the preceding each year.

Albertsons Companies reserves the right to inspect supplier's facilities. Qualification as an approved supplier and/or retention of status may be contingent upon inspection of supplier's facilities.

No change in composition of the material or grade substitution may be permitted without our prior evaluation and authorization. Quality Assurance and Strategic Sourcing must be notified two (2) months in advance of any changes in product makeup. Supplier must submit a sample of changed product for approval by Albertsons Companies along with such notification.

Acceptance of the shipment shall be contingent upon compliance with specifications. Material failing to meet all specifications is subject to rejection and return at the supplier's expense. Each shipment must be uniform in character, exhibit no signs of deterioration, and perform in a satisfactory manner. Reasonable uniformity of chemical and physical characteristics from shipment to shipment is expected.

The supplier will be held responsible for informing the Strategic Sourcing Manager and Quality Assurance Department of any real or expected deviations from these specifications. Performance and analytical testing are required by Quality Assurance prior to approval of any changes.

A bill of lading shall accompany each shipment. Shelf Life should be in accordance with "Minimum Acceptable Shelf Life" as detailed in the Packaging and Ingredients Supplier Handbook (PISH-SM V.1).

Delivery to Albertsons Companies locations shall be made via clean trucks, trailers or rail cars, suitable for transportation and protection of its contents with respect to integrity and quality, and in keeping with good manufacturing practices.

Packaged material shall be packed in suitable containers to preserve and protect product integrity and quality during customary conditions of handling, shipping and storage. Containers must be labeled in compliance with all applicable regulations including clear identification of the material, manufacturer's name, lot number, applicable special designations, and net f, and the Albertsons Companies Item Code number. Damaged containers which fail to provide protection to product integrity and/or quality will be rejected at the time of receiving.

Albertsons Companies requires that a Certificate of Analysis (COA) attesting to the conformance to specific specification requirements of each lot be submitted by the supplier. The COA must be communicated to the plant receiving the shipment prior to or accompanying the delivery. Results may be communicated by phone and email to ensure timely reporting, however, they must be confirmed in writing. The scope and content of the COA is part of this specification.

This material must conform to Kosher dietary laws and regulations where applicable. Specific materials require the certifying agency's symbol appear on every unit or container received at our plants. Kosher certificates must be updated annually.

In addition to all of the above, if the material is imported, each lot(s) must have clearance from U.S. customs and pass all F.D.A./U.S.D.A. inspections prior to shipment to Albertsons Companies locations. Imported raw materials may require, at the discretion of Albertsons Companies, independent laboratory analysis prior to receipt, with the expense borne by the supplier.