Application for Food Establishment Permit

Alaska Department of Environmental Conservation
Division of Environmental Health
Food Safety and Sanitation Program

Permit ID: _____

Section 1: GENERAL INFORMATION *(All applicants complete entire section – please print)*

Purpose (check one) ☒ **New** ☐ **Information Change** ☐ **Extensive Remodel** ☐ **Change of owner/operator** ☐ **Reactivate**

Owner/Business Information

Name of Entity or Owner Responsible for Food Service: ISG, Inc.		AK Business License #: 1050003	
Business/Corporate Mailing Address: 2915 Jackson Rd.	City: Juneau	State: AK	Zip: 99801
Business/Corporate Phone: (907)723-1178	Email: paulyburrowsak@gmail.com		
Owner(s) or Corporate Officer(s) & Title(s) or Responsible Party: Paul Burrows		Fax:	

Type of Entity ☐ **Individual** ☐ **Partnership** ☒ **Corporation** ☐ **Other:**

Establishment Information

Establishment Name: Juneau's Green Market	Physical Location: 263 Marine Way	Nearest Community: Downtown Juneau	
Establishment Mailing Address: 2915 Jackson Rd.	City: Juneau	State: AK	Zip: 99801
Establishment Phone: (907)723-1178	Fax:	Contact Person: Paul Burrows	
Establishment Physical Address: 263 Marine Way	City: Juneau	State: AK	Zip: 99801

SEATING: (Food Service Only) ☐ **N/A** ☐ **25 or less** ☐ **26-100** ☐ **> 101**

TYPE OF OPERATION *Please describe the type of facility you plan to open below (i.e. restaurant, bar, grocery store, etc.)*

Retail Dispensary of edibles and concentrates

SECTION 2 – NEW OR EXTENSIVELY REMODELED FACILITIES

a. A plan review will be required if your facility has never been permitted by the Alaska's Food Safety and Sanitation Program; has not had an active permit in the last five years; will be extensively remodeled; or is a new construction. If any of these apply, a Plan Review Application is required to process your application. Have you attached the Plan Review Application? ☒ **Yes** ☐ **No**

SECTION 3 – COMPLETE FOR ALL FOOD ESTABLISHMENTS *(Check all that apply)*

FOOD SERVICE ESTABLISHMENTS

a. A copy of your menu will be required. Have you attached a copy of the proposed menu? ☐ **Yes** ☒ **No**

b. Attach appropriate label, placard, or menu notation for the consumer advisories if you serve:
☐ Wild Mushrooms ☐ Unpasteurized juices ☐ Farmed halibut, salmon, or sablefish
☐ Raw/undercooked animal foods such as beef, shell eggs, lamb, pork, poultry, seafood, and shellfish.

c. Methods of food preparation (check the one that most closely describes the establishment:
☒ Assembly of Ready to Eat Foods ☐ Cook and Serve
☐ Hot or cold Service for 2 hours or more is done
☐ Complex (Preparation 1 day or more in advance, cooling and reheating is done).

d. Style of Service: ☒ Counter Service ☐ Self Service (i.e. buffet line, salad bar) ☐ Table Service
☒ Other: Pre-packaged goods to be sold from behind counter

e. Do you plan to operate as a caterer? ☐ **Yes** ☒ **No**
If **yes**, list all the equipment used to protect food from contamination and maintain product temperature during:
Transportation: Hot or Cold Holding:

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Responsibilities

Individual recall activities should be assigned prior to a recall event to avoid confusion during a recall. Assignment of the recall responsibilities are found in Appendix D (assign an individual to each activity).

Evaluation of the Complaint or Condition

Complaint receipt, processing, and evaluation are the first steps in the recall process. The steps involved in the evaluation process are:

- Receive the complaint – A file should be maintained containing any product complaints the company receives. Information that should be maintained in the product complaint file is:
    - i. Complainant contact information
    - ii. Reported problem with the product
    - iii. Product Identification
    - iv. Product Storage
    - v. Product purchase date and location
    - vi. Illness and Injury details
- Provide the complaint to knowledgeable staff for initial evaluation. If an initial assessment indicates a recall may be necessary, the Recall Coordinator assembles the Recall Committee for a full evaluation.
- Determine the hazard and evaluate the safety concerns with the product.
- Determine the product removal strategy appropriate to the threat and location in commerce.
- Contact the appropriate regulatory authorities.
- Alert legal counsel, insurance, etc. as appropriate.
- Maintain a log of the events of the recall including information such as dates, actions, communications, and decisions.

Identification of Implicated Products

It is Juneau's Green Market responsibility to ensure the identification of all products and quantities of products implicated in the recall. In addition, determination should be made if any other codes, brands or sizes of product handled by the company are affected.

A distribution list should be prepared as part of the Identification process. The distribution list should at minimum identify:

- Account name (consignees) that received the recalled product(s)
- Account addresses
- Contact names
- Contact telephone numbers
- Type of account (e.g., manufacturer, distributor, retailer)

Additional information relating to product information may include:

- Amount of product received/shipped
- Product ship date(s)
- Amount of product returned
- Amount of product consumed

A link to the FDA Distribution Template can be found in Appendix B.

Notification of Affected Parties

Notifications during a recall must be done in a timely manner and should include the appropriate regulatory agencies, the product distribution chain, and consumers when necessary. Recall notices are typically used to notify regulatory agencies and those businesses in the distribution chain. Press releases are generally oriented to consumers, but may be used to notify any affected party.

- Regulatory Agencies should be notified at the earliest opportunity after the decision has been made to conduct a recall. Regulatory guidance may be found at: http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129259.htm.
    - Subsequent to the initial notification, the regulatory authority should be updated throughout the recall process.
- Distribution Chain contacts will be notified by appropriate means (telephone, fax, email, letter, etc.). It is recommended that a written recall notice be provided to all consignees. The Recall Notice **must** include all relevant recall information (see Appendix B, section (a)(ii)).
    - Confirm receipt of the Notice of Recall with all accounts. A record of all account communications should be maintained.
- Consumers should be notified by the most effective method available. If appropriate, a press release can be used to notify consumers. Considerations for preparing a press release include:
    - Issuance of a press release should be the highest priority and should be issued promptly.
    - The local FDA District Recall Coordinator should be consulted before issuance of a press release whenever possible.
    - All relevant information should be included in the press release (seesample templates in Appendix B).

Removal of Affected Product

The procedure for product removal can be divided into five components including: removal, control, and disposition of affected product, recall effectiveness, and recall termination.

Removal

All reasonable efforts must be made to remove affected products from commerce.

- Products in commerce should be detained, segregated, and handled in a manner determined by the recalling firm.
- Products that are still in the recalling firm's control (e.g. inventory located onsite, in transit, in off-site storage, and in offsite distribution) should be detained, and segregated.
- All quantities and identification codes shall be documented to assist in the reconciliation of product amounts.

Control of Recalled Product

When Juneau's Green Market chooses to retain recalled product, control must be regained to prevent reentry of the product into commerce.

- All affected product returned will be clearly marked, not for sale or distribution, and stored in an area that is separated from any other foodproducts.

All quantities and identification codes shall be documented to assist in the reconciliation of product amounts.

Product Disposition

The final disposition of the recovered product must be determined. The final disposition must be reviewed and approved by the regulatory agency. Options include:

- Redirection – Products may be redirected for uses other than human consumption.
- Destruction - Products determined to be unsafe for human consumption may be destroyed or denatured, and disposed by appropriate means.
- Recondition – Products may be reworked to remove the safety risk. For example, would be relabeling a product to declare an allergen originally omitted from the label.

All quantities, identification codes, and disposition shall be documented.

Recall Effectiveness

Juneau's Green Market is responsible for determining whether the recall is effective. Recall Effectiveness Checks verify that all consignees have been notified and have taken the appropriate action. Steps include:

- Verifying that all consignees have received the notification.
- Verifying that consignees have taken appropriate action.
- If the response from our consignees is less than 100%, then the recall should be deemed ineffective and the recall strategy should be reassessed. Certain circumstances (e.g. amount of product actually returned vs. expected, potential for consumption, shelf-life, etc.) may also require a reassessment of the recall strategy.

All verifications shall be documented.

Termination of a recall

Termination of the recall may be considered after all reasonable efforts have been made to remove the affected products from commerce, including reconciliation, recall effectiveness, and disposition.

A termination of the recall may be requested by submitting a written request to the regulatory authorities. Guidance for the termination of a recall is provided in Appendix C.

Mock Recall

In addition to an annual verification of the recall plan, Juneau's Green Market will conduct a mock recall annually or whenever there are significant changes to the plan or personnel. The mock recall will include the following elements:

- Selecting a product which has reached the consumer market.
- Tracing the product from the raw ingredient (e.g. source) level to the finished product in the marketplace.
- Verifying communications systems (e.g. contact information, test emails and faxes, etc.) to outside contacts.
- Modifying the recall plan to correct any problems encountered during the test.

Records of these mock recalls will be documented and filed appropriately.

Appendix A – Contact Information

Recall Committee and Key Personnel Contact Information

The contact information including phone number, fax number, email address, and alternate 24/7 information of all committee members, their alternates, and "outside" key personnel should be confirmed and updated as often as necessary to assure accuracy.

Contact Information

- Recall Committee (24/7)
    i. Recall Committee Members (907)343-4200

Permit ID(s)_____ Establishment Name(s)_____

f.	Will your food establishment be a kiosk or mobile unit?	☐ Yes	☒ No
	Are employee toilets available within 200 feet? *If you have an agreement with another business to use their restrooms, please attach written verification.*	☒ Yes	☐ No
	Portable water tanks, plumbing, and hoses are NSF or FDA approved components?	☐ Yes	☐ No
	If you have a kiosk, is it located outside of a building?	☐ Yes	☐ No
	Will you have a service provide water or remove wastewater? *If yes, provide a letter of agreement from water hauler or wastewater hauler outlining services provided and frequency.*	☐ Yes	☐ No
g.	Will another permitted food establishment (commissary) provide support to your facility? If yes, attach a copy of the Commissary Agreement.	☐ Yes	☒ No

FOOD PROCESSORS

a.	A copy of a label for each type of product you will produce is required. Have you attached food labels of each product to be produced?	☐ Yes	☒ No
b.	Describe who you will be distributing your product to (i.e. grocery stores, etc):		
c.	Will you be doing any of the following processes? Check all that apply. ☐ Reduced Oxygen Packaging ☐ Low Acid Canned Foods ☐ Shelf Stable Acidified Foods ☐ Smoking ☐ Curing ☐ Dehydrating ☐ Other: *Be sure to check with your local Environmental Health Officer for any applicable forms and FDA requirements.*		
d.	Do you have a HACCP Plan? *Required for high hazard food processors such as smoking, curing, acidifying, dehydrating, thermally processing low acid foods, reduced oxygen packaging, etc.*	☐ Yes	☒ No ☐ N/A
e.	You are required to have a product coding system and a recall plan. Have you attached a copy of the coding system and recall procedures?	☒ Yes	☐ No

MOBILE RETAIL VENDOR SELLING SEAFOOD

a.	A list of products that you will be selling is required. Have you attached a copy of the list of products?	☐ Yes	☒ No
b.	Provide names of suppliers where you will be purchasing your product: ISG Inc. DBA Evergreen Extracts		
c.	Will *all* of your product be prepackaged?	☒ Yes	☐ No
d.	Will another permitted food establishment (commissary) provide support to your facility? If yes, attach a copy of the Commissary Agreement.	☐ Yes	☒ No

MACHINES VENDING POTENTIALLY HAZARDOUS FOODS

a.	Have you attached the label that will be affixed to the front of each machine with name, physical address, and phone number of the permitted food establishment servicing the machine?	☐ Yes	☒ No

SECTION 4 – Food Managers Certification/Alaska Safe Food Worker Card

a.	Have you attached a copy of a Food Manager's Certification? *The operator of a food establishment that serves and prepares unwrapped or unpackaged food, except for a bar, tavern, or limited food service, must have at least one Certified Food Protection Manager who is involved in the daily operations of the establishment.*	☐ Yes	☒ No	☐ N/A
b.	Does everyone who works or will work at the food establishment have a Food Worker Card? *An operator of a food establishment shall keep on file a copy of the Food Worker Card issued by the department for each employed food worker and make the copy available to the Department upon request.*	☒ Yes	☐ No	☐ N/A

I declare, under penalty of unsworn falsification, that this application (including any accompanying statements) has been examined by me and to the best of my knowledge and belief is true, correct, and complete. I agree to pay all fees before operating.

Applicant's Signature [signature] **Date** 6/16/17

Applicant's Printed Name Paul Burrows **Title**

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Appendix B – Templates

1. Communication Templates
    i. Model Press Releases(FDA)
        a. Allergens (Allergy Alert)
           http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129262.htm
        b. Listeria monocytogenes
           http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129267.htm
        c. Clostridium botulinum
           http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129273.htm
        d. Salmonella (all serotypes)
           http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129275.htm
        e. E. coli 0157:H7
           http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129287.htm
    ii. FDA Guidance for Written Recall Notification Letters
        http://www.fda.gov/Safety/Recalls/IndustryGuidance/default.htm
    iii. FDB Distribution List Template
        http://www.cdph.ca.gov/pubsforms/Guidelines/Documents/fdb%20eru%20Gde%20DistListTemp.xls
2. Recall Events Log (should include the following information):
    i. Name of the person creating the action
    ii. Dates
    iii. Actions
    iv. Communications
    v. Decisions
    vi. Product disposition
3. Recalled Product Information Data Sheet (should include the following information):
    i. Product description: brand, product name, size, etc.
    ii. Lot codes
    iii. Quantity of recalled product
    iv. Date of the action
    v. Action taken for each product

Appendix C – Additional Resources

1. Industry Guidance: Information on Recalls of FDA Regulated Products
   http://www.fda.gov/Safety/Recalls/IndustryGuidance/default.htm

2. Recall policy – 21 CFR Recall Regulations Sec. 7.40
   http://edocket.access.gpo.gov/cfr_2004/aprqtr/21cfr7.40.htm

3. US Food and Drug Administration
   http://www.fda.gov/

4. USDA (FSIS)
   http://www.fsis.usda.gov/wps/portal/fsis/home

5. Center for Disease Control
   http://www.cdc.gov/

6. FDA District Recall Coordinators
   http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129334.htm

7. FDA Guidance: Action levels for Poisonous or Deleterious Substances in Human and Animal Feed
   http://www.fda.gov/food/guidanceregulation/guidancedocumentsregulatoryinformation/chemicalcontaminantsmetalsnaturaltoxinspesticides/ucm077969.htm

8. FDA Defect Levels Handbook
   http://www.fda.gov/food/guidanceregulation/guidancedocumentsregulatoryinformation/sanitationtransportation/ucm056174.htm

APPENDIX D – Assigned Responsibilities

Sample Assignments (may include, but not limited to the following)

Assignment

1. Management of the Recall – (TBD) is responsible for the coordination of all recall activities.
2. Assemble the Recall Committee – (TBD) is responsible for communicating the decision to recall to the members of the Recall Committee and that each member knows their responsibilities.

Evaluation

1. Management Approval of the Recall – (TBD) is responsible to decide if the recall should go forward.

Identification

1. Create a Product Recall Log – (TBD) is responsible to create and maintain a product recall log to document all events, when they occur and the company's response to each.
2. Identify all Products to be Recalled – (TBD) is responsible for identifying all products which need to be recalled.

Notification

1. Notify the Appropriate Regulatory Authority – (TBD) is responsible for notifying the appropriate regulatory authority (use the contact information in the Recall Plan).
   Contacts shall only be made through the designated committee member. Recommended information to be submitted can be found in the FDA guidance document at: http://www.fda.gov/Safety/Recalls/IndustryGuidance/ucm129259.htm
2. Prepare the Press Release (if required) – (TBD) is responsible for the recall press release if the decision to prepare a press release is made. Considerations for preparing a press release include:
   a. Issuance of a press release should be the highest priority and it should be issued promptly.
   b. Consult with your local District Recall Coordinator before issuance of a press release whenever possible.
   c. If the company decides to prepare the press release, include all relevant information (see sample templates in AppendixB)
3. Prepare the Distribution List – (TBD) is responsible for preparing the recalled product distribution list. The FDB distribution list template requests account type, name, address, phone number, and contactname.
4. Prepare the Notice of Recall – (TBD) is responsible for preparing the written notice includes all recall relevant information (see FDA Guidance for Written Recall Notification Letters inAppendix).
5. Distribute the Notice of Recall – (TBD) is responsible for distribution of the Notice of Recall to all accounts that received the recalled product. Responsibilities include:
   d. Confirm receipt of the Notice of Recall with allaccounts.
   e. Contact accounts that have not responded to the request for conformation.
   f. Maintain records of the accountcommunications.

Removal

1. Detain and Segregate all Products to be Recalled which are in your Firm's Control – (**TBD**) – is responsible to ensure that all products to be recalled in the firm's control are not distributed (identify, detain, and segregate products on-site, in transit, off-site storage, and off–site distribution).
2. Control the recalled product(s) – (**TBD**) is responsible to ensure that recalled products do not re-enter commerce. Responsibilities include:
    a. Quarantine and clearly identify recalled products.
    b. Reconcile quantities, identification codes, and monitor recalled products.
    c. Document the returned products.
3. Decide what to do with the recalled product(s) – (**TBD**) is responsible for determining the action to be taken on the recalled product (destruction, reworking, and redirection). Other related responsibilities include:
    a. Determine if the regulatory authority requires actions such as witnessing destruction of the recalled product.
    b. Verify that the action taken has been effective.
    c. Document the action(s) taken.
4. Verify Recall Effectiveness – (**TBD**) – is responsible for verifying the effectiveness of the recall. Responsibilities include:
    a. Verify that distribution of recalled products has ceased.
    b. Verify that all consignees at the recall depth specified by the recall strategy have received notification about the recall.
    c. Verify that consignees have taken appropriate action.
    d. Document all verifications.