





TABLE OF CONTENTS

INTRODUCTION	1
OVERVIEW	2
Safe Food for Canadians Regulations (SFCR)	2
Types of Food Safety Hazards Controlled by a Preventive Control Plan	2
PURPOSE	5
How Do I Use This Guide?	5
Guide Layout	5
DETERMINING WHAT IS APPLICABLE TO YOUR OPERATION	6
GLOSSARY	8
PREVENTIVE CONTROLS, PART 4 OF THE SFCR	10
Division 4: Maintenance and Operation of Establishment	10
Subdivision A: Responsibility of Operator	10
Subdivision B: Sanitation, Pest Control and Non-Food Agents	11
Subdivision C: Conveyances and Equipment	16
Subdivision D: Conditions Respecting Establishment	18
Subdivision E: Unloading, Loading and Storing	26
Subdivision F: Competency	28
Subdivision G: Hygiene	29
Division 5: Investigation, Notification, Complaints and Recall	31
Division 6: Preventive Control Plan	34
TRACEABILITY, PART 5 OF THE SFCR	40
PACKAGING, PART 10 OF THE SFCR	43
LABELLING, PART 11 OF THE SFCR	44
GRADES AND GRADE NAMES, PART 12 OF THE SFCR	46
APPENDIX	48
How to Draft an SOP/Policy/Procedure	48
What to Expect During Your CFIA Inspection	49
How to Respond to a Non-Compliance	50
How to Include Cross-Contamination Points on the Hazard Analysis and CCP Determination Form	50
DOWNLOADABLE FORMS	52
RESOURCES	52





INTRODUCTION

Our Quality and Food Safety Team is pleased to present a guide that has been developed to support maple producers in Nova Scotia meet the requirements of the Safe Food for Canadians Regulations.

With this guide we hope to enhance your knowledge of the requirements and provide resources to assist you in developing and implementing a food safety management program that meets the Safe Food for Canadians requirements or to assist you in strengthening your current program.

The resources mentioned throughout this guide can be found on Perennia's website under Quality & Food Safety/Food Safety Resources.

As always, if you have questions, feel free to contact our Quality and Food Safety Specialists. We are here to help.

Perennia's Quality and Food Safety Team

Elaine Grant, Pam Laffin, Shelly MacDonald, Clarissa McIsaac, Cheryl Andrews

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OVERVIEW

SAFE FOOD FOR CANADIANS REGULATIONS (SFCR)

Consumers want to know their food is safe. The SFCR supports this with mandatory preventive controls, traceability, and better controls for imported foods, to name a few. In 2019, the SFCR consolidated 14 sets of food regulations into one. The intent is to improve consistency between all food businesses and types of food, reduce administrative burden and enable outcome-based provisions. However, it is important to understand that many commodities have food-specific requirements that must also be met. It is your responsibility to be familiar with your products' specific requirements. The Safe Food for Canadians Act (SFCA) and SFCR came into effect on January 15, 2019. The SFCR uses preventive controls and preventive control plans to control potential hazards associated with production. While preventive controls control hazards associated with the environment in which a product is processed, the preventive control plan is specific to controlling potential hazards associated with the process and production of food.

TYPES OF FOOD SAFETY HAZARDS CONTROLLED BY A PREVENTIVE CONTROL PLAN

Hazard concerns with food production and processing are biological, chemical, and physical hazards.

Biological hazards include bacteria, viruses, parasites, or fungi (yeast and molds) that can cause food-borne illness if they or their toxins are ingested. Biological hazards can be found anywhere when conditions favour their growth. Such conditions include temperature, humidity, pH, water activity and oxygen availability. Bacteria can be found in soil, mud, air (e.g., aerosols or dust suspended in air), water, decaying matter, fecal matter, sewage, the gut of warm-blooded animals, and even in our nose, mouth and on our skin. Fungi are typically found in warm and humid environments.

Chemical hazards can fall into four broad categories; naturally occurring toxins and allergens (e.g., shellfish toxins and mycotoxins); chemicals intentionally added to food (e.g., water, preservatives and additives); chemicals unintentionally added to food (e.g., chemicals from packaging material, chemicals used for cleaning or maintenance, and chemicals from equipment such as lead residues); and radiological hazards, although they are rarely encountered in food, when they do occur they can present a risk (e.g., from contaminated soil, water or air; packaging materials; ingredients with radionuclides). Chemicals that are intentionally added to food are not intended to be hazardous. However, higher than desired amounts may render them harmful to human health.

A physical hazard is any extraneous or foreign material or object that can cause injury or illness to a consumer such as (but not limited to) glass, plastic, metal, wood, and animal droppings or insects.





PURPOSE

HOW DO I USE THIS GUIDE?

This guide is meant to be a tool to assist maple producers in meeting the Safe Food for Canadians requirements. It can be used to assist you with interpreting the regulations and determining if you are currently meeting them or help identify areas for improvement in your current program. It is designed to include the current regulations (as of the edition date on the booklet cover) as they apply to maple production, packing, storing and transport within Nova Scotia and for those that ship between provinces.

This guide begins at Part 4 of the Safe Food for Canadians Regulations detailing the Preventive Controls (PCs) and Preventive Control Plan (PCP) that are required to be in place to control potential hazards at your operation. Depending on your activities, some aspects of the regulations and sections in this guide may not apply to you. Mark these sections as Not Applicable (N/A) and move on to the next section. This guide will complement those resources already posted in the Food Safety Resources section on the Perennia Food and Agriculture website. For those resources not on our website, an appendix has been created to capture this information. There are also sample forms/records you can use, or you can create your own and use the samples provided as a guide to help you draft your preventive control plan and supporting documents.

GUIDE LAYOUT

Each section is laid out to match the flow of the Safe Food for Canadians Regulations as they apply to Maple Production.

The **title** of each section matches that of the sections in the Safe Food for Canadians Regulations.

Requirements are the specific regulations as they are written in the Safe Food for Canadians Regulations.

Rationale is the reasoning (the why) these requirements must be met.

Interpretation is our interpretation of how you can meet these requirements.

The Checklist details what documents you need in your PCP to meet the requirements of each section. It can also be used as a "to-do list" while actively using this guide to check off what has been completed.

The Appendix has guidance and resources to frequently asked questions we have received from Nova Scotia maple producers and processors. It can be used like the Food Safety Resource section on our website.

The Forms are editable Excel spreadsheets that can be used as-is or as a guide when developing your own forms and customized records.

You will need to review the table below to determine what requirements you need to comply with in the Safe Food for Canadians Regulations as they apply to Maple Products.

DETERMINING WHAT IS APPLICABLE TO YOUR OPERATION

In order to determine which sections of the SFCR apply to your operation, we have streamlined the Timeline for Maple Products from the Canadian Food Inspection Agency (CFIA) website. This enables you to decide which applies to your situation, as the requirements are very specific to the activities conducted at your operation. For example, if your only activity is to sell maple products to consumers at retail, you only require traceability; you would then proceed directly to the Traceability section in this guide.



WHAT IS YOUR ACTIVITY?	DO I NEED A LICENSE?	DO I NEED A PREVENTIVE CONTROL PLAN?	DO I NEED PREVENTIVE CONTROLS?	DO I NEED TRACEABILITY?
I manufacture, process, treat, preserve, grade, package or label maple products for	Yes – deadline, January 15, 2019	Yes , if your gross annual food sales are greater than \$100,000 – deadline January 15, 2019 **	Yes – deadline, January 15, 2019	Yes – deadline, January 15, 2019
interprovincial trade		No , if your gross annual food sales are less than \$100,000 **		
I manufacture, process, treat, preserve, grade, package or label maple products for export	Yes – deadline, January 15, 2019	No , not required unless you would like an export certificate or other export permission from CFIA *	like ate January 15, 2019 Yes – deadline, January 15, 2019 Yes – deadline, January 15, 2019	
My only activity is to trade maple products interprovincially	No	No	No	Yes – deadline, January 15, 2019
I import maple products	Yes – deadline, January 15, 2019	Yes, if your gross annual food sales are greater than \$100,000 – deadline January 15, 2019 ** No, if your gross annual food sales are less than \$100,000 **	Yes – deadline, January 15, 2019	Yes – deadline, January 15, 2019
My only activity is to export maple products	No, not required unless you would like an export certificate or other export permission from CFIA	No, not required unless you would like an export certificate or other export permission from CFIA	No Yes – deadline, January 15, 2019	
I sell maple products to consumers at retail	No	No	No	Yes – deadline, January 15, 2019
My only activity is that I am a primary producer of maple sap for export or interprovincial trade	No	No	No	Yes – deadline, January 15, 2019
I manufacture, process, treat, preserve, grade, package, or label maple products for intraprovincial trade	No , not required in Nova Scotia	No , not required in Nova Scotia	No , not required in Nova Scotia	No , not required in Nova Scotia

^{*} excerpt from SFCR timelines – Honey and maple products. "You need a licence and PCP prior to obtaining the certificate or other permission from CFIA."

^{**}excerpt from SFCR timelines – Honey and maple products; visit the section "Calculating your gross annual food sales" to determine your gross annual food sales **SFCR timelines - Honey and maple products - Canadian Food Inspection Agency (canada.ca).**

GLOSSARY

The following is a list of terms with accompanying definitions. This glossary elaborates on specific terms you will see referenced throughout this guide.

Allergen: a protein that causes physiological reactions due to an immunological response. CFIA priority allergens are peanuts, tree nuts, sesame, soy, fish, crustaceans and molluscs, wheat and triticale, eggs, milk, mustard, and sulphites.

Calibration: measuring the accuracy of a device to a known standard and adjusting, as necessary, so that it conforms to the standard.

CCP - Critical Control Point: a step at which a control measure is essential to prevent, eliminate or reduce any biological, chemical, or physical hazard that presents a risk of contamination of a food to an acceptable level.

Consumer: the final user of the purchased goods.

Consumer Prepackaged: packaged in a container in the manner in which the food is ordinarily sold to or used or purchased by an individual - or in which the food may reasonably be expected to be obtained by an individual - without being repackaged, to be used for non-commercial purposes.

Corrective Action: any action or step taken to eliminate or resolve the cause of a deviation from the written food safety program.

Cross Contamination Points: points throughout the process where if control measures are not put in place, a hazard associated with one process, product, or item may contaminate another.

Deviation: a variation from a written procedure, specification, standard or regulation.

Establishment: any domicile where a food is manufactured, prepared, stored, packaged, or labelled.

Exporting: sending food to another country.

Food Contact Surface: any surface where opened (unpackaged) or packaged product will come into direct contact.

Hazard: any biological, chemical, or physical risk that has the potential to cause illness or injury to a consumer.

Hazard Analysis: the process of identifying and assessing potential hazards by determining the risk level associated with the likelihood and severity level.

Hazard Analysis Critical Control Point (HACCP): a systematic, scientific approach to the identification, evaluation, and control of food safety hazards during the production and manufacturing of food.

High-Risk: foods that are ready to eat, do not require further cooking and provide an environment for bacteria to live, grow and thrive.

Importing: bringing food into Canada from another country.

Interprovincial: the trade of food from one province or territory to another.

Intraprovincial: the trade of food within a home province or territory.

Labelling: the act of applying information on a product.

Licence: a permit from CFIA allowing an establishment to conduct registered activities.

Monitor: a designated individual who plans to observe or measure control parameters to assess whether a control measure is effective.

Non-Food Agent: chemicals that are not considered to be a food or food ingredient. These include cleaning chemicals, detergents, lubricants, agricultural chemicals, and pest control products.

Packing: physically placing product into packaging materials.

Personal Effects: refers to privately owned items (e.g., jewelry, earbuds, phone, lunches, clothing, medication, etc.).

Potable Water: water that is considered safe to drink and meets Health Canada's Drinking Water Quality Guidelines.

Preventive Controls (PCs): control measures (e.g., sanitation, maintenance, calibration, training, pest control, etc.) used to prevent risk of contamination to food and achieve compliance with regulatory requirements.

Preventive Control Plan (PCP): a written document that demonstrates how risks to food are identified and controlled through preventive controls/prerequisite programs/good manufacturing practices or critical control points (similar to a HACCP plan).

Recall: the removal from sale or use, or correction, of a marketed product that poses a health risk or is noncompliant with respect to legislation.

Retail: refers to the sale of food to consumers for consumption. Examples: supermarkets, farmers' markets, grocery stores, bakeries, and butcheries.

Risk Assessment: the documented process of identifying, measuring, and evaluating the level of risk associated with a product, process, procedure, or environmental factor to determine the appropriate control measures.

Safe Food for Canadians Regulations (SFCR): is a single set of rules for all food commodities to ensure that food prepared domestically, imported, or exported is safe for consumers, complies with grade requirements and is labelled and presented in a way that is not false or misleading. These regulations are enforced by CFIA.

Sanitation Standard Operating Procedure (SSOP): written procedures that address sanitation conditions and practices before, during and after processing.

Specification: a detailed description of a material, product or service that outlines important characteristics, attributes, or requirements.

Standard Operating Procedure (SOP): written procedures for preventive control measures, production and process controls designed to assure the production of safe food.

Storing: keeping product in a pre-determined location (ambient – dry storage, refrigerated storage, frozen storage) for a pre-determined amount of time.

Traceability: the ability to track the movement of product one step forward (person to whom you provided the food) and one step back (supplier) in the supply chain.

Verifier: a designated individual who ensures the monitor is effectively observing or measuring control parameters of a control measure. This individual is referred to as the double checker and should not be the same person as the monitor unless absolutely necessary.



PREVENTIVE CONTROLS — PART 4 OF THE SFCR

DIVISION 4: MAINTENANCE AND OPERATION OF ESTABLISHMENT

SUBDIVISION A: RESPONSIBILITY OF OPERATOR

Requirements: An operator must maintain and operate an establishment so that the requirements of sections 50 to 81 are met (for maintenance and operation of an establishment). 49

Rationale: Without senior management's commitment and adequate resources in place to support the development, implementation and ongoing maintenance of a food safety program, a successful Preventive Control Plan is not possible.

Interpretation: An operator's responsibilities of maintaining and operating an establishment to meet requirements are to:

- Ensure that the establishment complies with CFIA requirements: Safe Food for Canadians Regulations;
- Ensure that food safety is fully embedded in every level of their business (all members, including senior management, contractors, services providers, etc. are required to follow good manufacturing practices. There are no exceptions);
- Provide the necessary resources (e.g., staffing, training) and the time required for the development, implementation and effective maintenance of the Preventive Control Plan;
- Provide the necessary training of staff and clearly define their responsibilities;
- Provide financial resources to ensure that the facility's design (internal and external) and equipment meets regulatory and any customer requirements.

CHECKLIST
☐ Letter of commitment signed and dated by senior management
 Detail of how the company plans to provide safe, quality products, provide necessary resources, meet regulatory requirements, and appoint a trained and qualified food safety team leader
 Letter of commitment should be communicated to all staff members, signed, dated, and posted in a prominent location
☐ Regulatory Compliance:
 Applicable regulations are on file and up-to-date (e.g., Safe Food For Canadians Act and Regulations, Food and Drugs Act and Regulations)
 Ensure any updates or changes that affect the program as written are included
☐ Maintenance and Reassessment Procedure (SOP) (should have the following):
$\ \square$ Frequency of at least annually
Detail how the entire preventive control plan is reviewed and by whom:
 How the Food Safety/HACCP Plan, Preventive Controls, Regulatory and Administrative policies are updated
 Identification of the food safety hazards and details of how the control measures in place for all food safety hazards are reviewed
 A check to ensure that the plan conforms to current regulatory and program requirements as well as conforms to customer requirements
$\hfill \square$ Verification and Deviation details, as well as a list of records used
☐ Maintenance and Reassessment Record
☐ Change Log

SUBDIVISION B: SANITATION, PEST CONTROL AND NON-FOOD AGENTS

CLEAN AND SANITARY CONDITION/CLEANING AND SANITATION

Requirements: An establishment, and any conveyance or equipment in it that is used in connection with an activity that is regulated under the Act, must be clean and in a sanitary condition. 50(1)

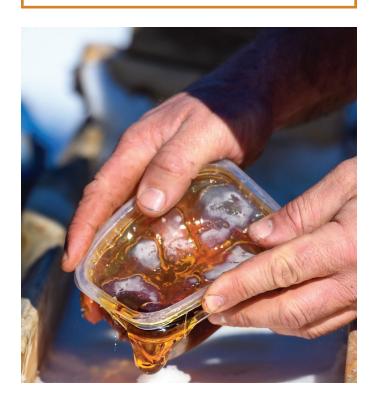
The cleaning and sanitation of the establishment and of any conveyance or equipment in it that is used in connection with an activity that is regulated under the Act must be conducted in a manner that does not present a risk of contamination of a food. 50(2)

Rationale: Improper or inadequate sanitation or use of improper chemical concentrations can lead to contamination of food, ingredients, packaging materials and food contact surfaces.

Interpretation: An establishment must be maintained in a clean manner. Cleaning and sanitation must be achieved through cleaning schedules and procedures, including chemical concentrations. To meet this requirement, you should:

- Ensure schedules and clear procedures are in place;
- Ensure chemicals are approved for use in a food establishment (e.g., processing equipment and sugar house);
- Clean equipment away from actively in use sap storages, processing, packaging and storage areas, remove/move packaging, and product away from area being cleaned prior to cleaning equipment and surrounding areas;
- Provide the necessary training to staff.

- ☐ Sanitation Program:
 - ☐ Sanitation schedule/frequency for all equipment and facility areas, utensils/tools, waste, work gear, carts, pans, pails, sap storage tanks, transfer hoses, tanks, barrels, reusable filters, RO, etc. (e.g., daily, weekly, monthly, pre and post-season)
 - ☐ Cleaning and sanitizing procedures (methods for dry cleaning, pressure washing, cleaning with steam/hot water, handling chemicals, chemical concentrations, PPE, proper handling, etc.)
 - ☐ Pre-operational/Post-operational inspection procedures: after cleaning and before product is handled, the processing, packing and storage areas, and equipment are clean and free from dust, dirt, food product, and excessive grease
- ☐ Sanitation Record(s)
- ☐ Employee training



PEST CONTROL - ANIMALS

Requirements: An establishment must be protected against the entry of any animal that presents a risk of contamination of a food, except if, in the case of any land that forms part of an establishment, there are no reasonably practicable measures that may be taken to prevent the entry of such animals onto the land. 51(1) An animal must not be in a facility or conveyance where a food is manufactured, prepared, stored, packaged, or labelled.

(a) a food that is intended to be manufactured, prepared, stored, packaged, or labelled in the facility or conveyance; 51(2)

Any measures that are taken for the purposes of complying with subsections (1) and (2) must not present a risk of contamination of a food. 51(3)

Rationale: Pests can contribute to biological and physical contamination of food, packaging materials and food contact surfaces. Inadequate biosecurity/biocontainment can lead to the contamination of products, facility, packaging materials or the spread of diseases.

Interpretation: Pest management is required for the entire site, including temporary or seasonal storage units.

- Where you have personal property that coexists with land that forms your establishment (e.g., house, camps and storage/out buildings are in the same vicinity), then measures need to be in place that are within reason/practical to prevent entry of domestic or wild animals into the sap, maple product, packaging and equipment storage areas and your establishment.
- Some practical options include but are not limited to keeping doors/entrances closed or screened, screening windows and outside air vents, fences, bird or animal scares to prevent nesting and den making, and limiting pest harbourages near the buildings (e.g., keep grass trimmed, unused equipment away from areas, no pooling of water, gravelling the perimeter of buildings to discourage rodent travel).
- Animals cannot be used as a form of pest control.



CHECKLIST:	
☐ In house/self-managed pest control:	☐ Contracted Pest Control, including name and contact information:
 □ Person responsible for pesticide application, pest control □ Copy of pesticide applicators certificate (if applicable see Non-Food Agents section for details) □ Schedule/frequency (monthly inspection at minimum, pre-season and during season or while product is stored) □ Device map signed and dated □ Training □ Corrective action reports □ Proper handwashing is completed after handling bait, devices, and pests 	 □ Contract, signed and dated, including scope (insects, rodents, birds, beetles, etc.) □ Schedule/frequency (monthly inspection at minimum, pre-season and during season or while product is stored) □ Device map signed and dated □ Copy of insurance □ Copy of pesticide applicators certificate □ Service reports, including pesticide usage, concentration, and trending □ Safety Data Sheets for pesticides
☐ Safety Data Sheets for pesticides	☐ Contractor training
 □ Pest Control Records must be kept □ Pest Control Products, labels/identification intact and legible if applicable (e.g., name of product, active ingredient(s), concentration, PCP#) □ Biosecurity and biocontainment program: □ Procedures in place to prevent the introduction or spread of pests and/or disease □ Proper and secure storage of pest control chemicals (keep away from raw materials, packaging and product) 	 □ Recommendations and corrective action reports □ Records must be kept □ Contractor reports findings to operator after inspection is completed. Operator must sign off and complete any corrective actions
For Both:	
DO NOT allow animals, either wild or domestic (including pets) or storages), or pests (e.g., birds, rodents) entrance into buildings
Traps are located against the wall on each side of entrances to unless inside a trap, if it's required then it is only used when no To prevent pests from spreading bait, avoid using bait in pellet	operation is taking place or when storage areas are empty.
Fly control (lights or strips) are not used directly over top/in clo	se proximity of product, packaging or food handling areas.
Old traps, bait, glue boards and pests that are disposed of are	done so in a sealed container and placed in the garbage.

Pest control devices are clearly numbered/labelled/identified on the trap and the wall above the trap location (this helps

Pest control products are registered for use in Canada and are used according to label directions

keep the trap in the proper location in case it gets moved).

NON-FOOD AGENTS — SANITIZERS AND NON-FOOD CHEMICAL AGENTS

Requirements: Any sanitizer or non-food chemical agent that is in an establishment must

- (a) be properly and clearly identified;
- (b) be suitable for its intended use and not present a risk of contamination of a food; and
- (c) be handled and used in a manner that does not present a risk of contamination of a food, and in accordance with any manufacturer's instructions. (52)

Rationale: Improper use of non-food agents, concentrations and/or improper application or rinsing procedures can lead to both chemical and biological contamination.

Interpretation: Non-food agents present a risk of chemical and biological contamination if they are not used, handled and/or stored properly. An establishment must demonstrate controls through procedure and employee training.

- Ensure procedures are in place;
- Ensure chemicals are approved for use in a food establishment. For example, oils and greases used to lubricate equipment where it comes in contact with food must be food grade; detergents and sanitizers used on food contact surfaces are approved for use on food contact surfaces.
- Provide the necessary training of staff.

□ Non-Food Agents Policy:
 Master list of approved chemicals, approval letters
\square Safety Data Sheets (SDS)
 Proper labelling (for example where applicable: clearly labelled, identified with product name, active ingredients, PCP#, concentration, manufacturer's instructions for use and identification of chemicals)
☐ Concentrations
 Designated storage area - separation of food- grade vs. non-food grade; stored separately, well labelled, secure from food products, packaging, and equipment
☐ Receipts are on file and signed
☐ If you apply commercial and restricted class pesticides, then Nova Scotia requires you to have a pesticide applicator certificate. If this is required then ensure you have a Pesticide Applicator License on file, and are formally trained (e.g., online, self-study, examinable course)
☐ Master List of Approved Chemicals (can be added to your Non-Food Agents Policy)
☐ Personnel Training Record



SUBDIVISION C: CONVEYANCES AND EQUIPMENT

CONVEYANCES AND EQUIPMENT — FOOD

Requirements: Any conveyance or equipment that is used in the manufacturing, preparing, storing, packaging, or labelling of a food must

- (a) be appropriate for the food, as the case may be, and for the activity being conducted;
- (b) be designed, constructed, and maintained to prevent contamination of the food;
- (c) be constructed of, and maintained using, materials that are suitable for their intended use and, if those materials present a risk of contamination of the food, that are
 - (i) corrosion-resistant,
 - (ii) durable,
 - (iii) capable of withstanding repeated cleaning and, if necessary to prevent contamination of the food, repeated sanitizing, unless the equipment is intended for single-use, and
 - (iv) free of any noxious constituent;
- (d) be equipped with instruments to control, indicate, and record any parameters that are necessary to prevent contamination of the food;
- (e) function as intended;
- (f) be accessible and, if necessary for its cleaning, sanitizing, maintenance, or inspection, able to be easily disassembled;
- (g) be used, maintained and, if necessary, calibrated in accordance with the manufacturer's instructions and in a manner that does not present a risk of contamination of the food; and
- (h) have surfaces that, if they come into contact with a food, are smooth, free from pitting, cracks, and flakes and non-absorbent, except when the surface does not present a risk of contamination of the food. (53)

OTHER CONVEYANCES AND EQUIPMENT

Any conveyance or equipment in an establishment that is used to handle any contaminated materials, any waste or any other thing that is inedible must, unless that conveyance or equipment does not come into contact with those materials, waste, or things,

- (a) be used only for that purpose;
- (b) be identified as being reserved for that purpose; and
- (c) meet the applicable requirements of section 53. (54)

Rationale: Well-constructed and maintained equipment minimizes the potential for biological, chemical, and physical hazards. Equipment must be calibrated to ensure accuracy of the device.

Interpretation: Conveyances and equipment may present a risk of biological, chemical and/or physical contamination if they are not maintained properly. Maintained and well-constructed equipment minimizes risks. Devices that require calibration must be accurate to ensure they are functioning as intended (e.g., reading the correct measurement, etc.)

- Ensure procedures are in place;
- Ensure proper chemicals (lubricants, oil, grease, etc.) are used on equipment and conveyances;
- Avoid using equipment or surfaces intended to be used for food contact that are made of wood, galvanized metal or lead, painted surfaces especially painted with lead-based paint, or lighting that is not protected or shatter proofed.
- Provide the necessary training of staff.

CHECKLIST ☐ Preventive Maintenance Program (details the following): ☐ Equipment design and construction is maintained and in good condition, allows for easy cleaning (e.g., no dead zones (hidden/hard to clean areas), open ends, away from walls, easy access) ☐ Food contact surfaces that are easy to clean, non-porous, smooth (free from cracks, crevices, pitting or flaking) (e.g., stainless steel, hard plastic, corrosion-free), gauges on equipment are shatter proof or protected from breakage ☐ Sap and syrup storage tanks, transfer hoses and containers are food grade, cleanable and in good condition ☐ No temporary repairs (e.g., string, cardboard, wire, tape), missing nuts, screws, bolts, cracks in surfaces ☐ Maintenance schedule ☐ Maintenance program and equipment list ☐ Equipment inspections ☐ Preventative Maintenance & Calibration **Record**, service reports from outside contractors, if applicable ☐ Calibration Program (can be combined with the Preventative Maintenance Program above): ☐ List of devices to be calibrated (may include but not limited to scales, thermometers, hydrometers (accuracy check), refractometers.) ☐ Calibration schedule ☐ Calibration procedures, calibration record (see above) ☐ Employee training



SUBDIVISION D: CONDITIONS RESPECTING ESTABLISHMENT

LAND

Requirements: If any land that forms part of an establishment presents a risk of contamination of a food, measures must be taken to eliminate the risk. 56(1)

LOCATION

Requirements: If an establishment is located near any place or thing that presents a risk of contamination of a food, measures must be taken to eliminate the risk. 56(2)

Rationale: Facility location can lead to contamination of food. For example, a facility located beside a farm or barn. Well-constructed and maintained facilities reduce the risk of contamination to food, ingredients, and packaging materials (e.g., free of debris, refuse, pooling water and pest harbourage).

Interpretation: The facility shall be an adequate size and located to reduce risk of contamination. The facility shall be constructed and maintained to reduce risk of contamination.

Fa	cility and Premises Inspection Procedure
Sit	e Assessment:
	Previous use (e.g., heavy metal contamination, sewage sludge, industrial activities)
	Location to roads or public pathways (you may need to install a buffer to prevent roadside debris and road salt or dust as well as traffic and domestic animals from entering)
	Flood zone and incompatible neighbouring properties
(pr	terior and Interior Inspection Checklist re-season and monthly during the season, est-season):
	Located away from or protected against potential sources of contamination (landfills, livestock barns, incompatible operations (such as fuel storage, generators, vacuum pumps, floodplains, etc.))
	Vegetation is controlled and not growing against building
	No debris, old pallets, equipment, etc. lying against buildings or stored close to buildings
	Adequate drainage, no pooling water around buildings
	No holes or cracks in the foundation
	Exterior entrances and windows are well-sealed, screened (e.g., no holes around door seals)
	Waste is contained with secure lids to ensure animals and pest entrance is deterred
	Waste is situated away from production areas and buildings to prevent attracting pests
	Exterior of building do not have evidence of nesting birds or areas for wildlife to make a den close to the buildings. Pest control devices are in place and secured

INTERIOR OF FACILITY OR CONVEYANCE

Requirements: The interior of any facility or conveyance where a food is manufactured, prepared, stored, packaged, or labelled must be

- (a) designed to prevent the accumulation of substances that present a risk of contamination of the food, including dust, dirt, micro-organisms, and food particles, and to permit effective maintenance, cleaning and sanitizing;
- (b) designed, constructed, and maintained in such a manner that
 - (i) the size and layout is adequate to accommodate the activity being conducted and the equipment used in the activity,
 - (ii) the entry of insects, rodents, and other vermin is prevented,
 - (iii) any floors, walls, ceilings, windows, and doors are smooth, non-absorbent and impervious to moisture, except if those floors, walls, ceilings, windows or doors do not present a risk of the contamination of the food, and
 - (iv) any floors provide or permit good drainage, except if there is no risk of liquid accumulation;
- (c) constructed of, and maintained using, materials that are $\,$
 - (i) suitable for their intended use,
 - (ii) appropriate for the food or the food animal, as the case may be, and for the activity being conducted,
 - (iii) durable,
 - (iv) capable of withstanding repeated cleaning and, if necessary, to prevent contamination of the food, repeated sanitizing, and
 - (v) free of any noxious constituent; and
- (d) of sound construction and in good repair. 57

Rationale: Well-constructed and maintained facilities reduce the risk of contamination to food, ingredients, and packaging materials.

Interpretation: The facility shall be constructed and maintained to reduce risk of contamination.

☐ Exterior and Interior Inspection Checklist (pre-season, monthly during the season and post-season):	
 Doors are self-closing and close-fitting (e.g., r gaps or visible light when closed) 	10
☐ Floors, walls, ceilings, windows, and doors are cleanable, constructed of appropriate materia junctions are smoothly sealed and designed t allow maintenance, cleaning, and sanitizing	ıls,
☐ Windows are screened and shatter proofed if located in food handling and storage areas	
☐ Lighting and windows near storage or maple production areas is protected from glass breakage/shatterproofed	
 Area is free of pests and animals (wild or domestic) 	
 Product, packaging, food contact items (such as filters, testing equipment, defoamer, diatomaceous earth, etc.) are kept off the floo covered to stay dry and clean where required 	or,
$\ \square$ Sound construction and good condition	
$\ \square$ Air intakes and vents are screened	
 Drains are accessible for cleaning and maintenance 	
 Floors are sloped to drain, no pooling of water no cracks, or crevices 	er,
\square No leaking of overhead pipes	

DESIGN, CONSTRUCTION, AND MAINTENANCE

Requirements: A facility or conveyance where a food is manufactured, prepared, stored, packaged, or labelled must be designed, constructed and maintained in such a manner that the movement of persons and things within, into and out of it is controlled. 59(1)

MOVEMENT — NO RISK OF CONTAMINATION

Requirements: The movement must not present a risk of contamination of the food. 59(2)

Rationale: Operational flows such as employee entry to the establishment and flow to work rooms, ingredient/product flows and/or adequate separation or control between incompatible operations will prevent biological, chemical, or physical contamination of the product.

Interpretation: Operational flows must be controlled to prevent the risk of contamination. Consider the following traffic flows:

- Employees
- Visitors
- Product (sap or market-ready)
- Waste/compost
- Ingredients
- Packaging
- Chemicals
- Allergens

- ☐ Operational flows do not pose a risk of cross-contamination. Identify points where they overlap and pose a potential source for cross-contamination (e.g., market-ready product and chemicals)
- ☐ Operational flows are identified on plant schematic
- ☐ Cross-contamination points are identified on facility/building schematic
- ☐ Handwashing sinks, employee welfare rooms (lunchroom, washrooms, change rooms, sleeping quarters), storages, chemical storages, equipment layout should be identified
- ☐ Cross-contamination points are identified on hazard analysis forms with control measures in place to minimize the risk of cross-contamination (Your PCP, see appendix)
- ☐ Seasonal storage use
- ☐ **Visitor's Record** and policy (can be combined with Hygiene Policy)
- ☐ Employee training



INCOMPATIBLE ACTIVITIES

Requirements: Physical or other effective means must be used to separate incompatible activities in order to prevent contamination of a food. 60

SEPARATION OF FOOD

Requirements: Physical or other effective means must be used to separate a food from

- (a) anything that presents a risk of contamination of the food:
- (b) any food that does not meet the requirements of the Act or these Regulations; and
- (c) anything that is manufactured, prepared, stored, packaged, or labelled in an establishment and not intended or sold for use as food. 61

ARRIVAL OF CERTAIN FOOD AT ESTABLISHMENT

Requirements: Any food that presents a risk of injury to human health, that is exempted under section 22 from the application of the import requirements that are set out in the Act and these Regulations or that does not meet the requirements that are set out in the Act or these Regulations must be identified as such and placed in a designated area when it arrives at an establishment. 62(1)

MEASURES TO PREVENT CONTAMINATION

Requirements: Any measures that are necessary to prevent the food described in subsection (1) from contaminating any other food that is in the establishment must be taken. 62(2)

Rationale: Incompatible activities may result in cross-contamination or cross-contact.

Interpretation: Incompatible activities must be controlled to prevent the risk of contamination. Consider the following:

- Allergens (Canada's Priority Allergens: eggs, milk, mustard, peanuts, crustaceans and molluscs, fish, sesame seeds, soy, sulphites, tree nuts, wheat, and triticale). Different countries have different priority allergens, make sure you check their government websites prior to exporting.
- Additives
- Preservatives
- Returned/suspect product

CHECKEIST
☐ Allergen Management Program:
 Proper production scheduling (e.g., process non-allergenic products first or on separate days, where applicable)
☐ Allergens in employee lunches (handwashing is key, leaving work effects at the worksite to avoid cross-contact, not eating or drinking in the sap collection and production areas, with the exception of water)
 Designate allergen storage areas (where applicable)
 Allergen-free clean up and inspections (where applicable)
☐ Employee training
 Defoamers and food grade lubricants used do not contain allergens
$\ \square$ Correct labelling and verifications
☐ Return Products/Non-conforming Items Policy
☐ Held items identified

LIGHTING/LIGHT FIXTURES

Requirements: An establishment must be equipped with natural or artificial lighting that is appropriate for the food and for the activity being conducted. 63(1)

Any light fixtures in the establishment must

- (a) be capable of withstanding repeated cleaning and, if necessary, to prevent contamination of a food, repeated sanitizing; and
- (b) not present a risk of contamination of the food in the event of breakage. 63(2)

VENTILATION SYSTEM

Requirements: A facility or conveyance where a food is manufactured, prepared, stored, packaged, or labelled must be equipped with a ventilation system that

- (a) provides natural or mechanical ventilation with sufficient air exchange to provide clean air and to remove unclean air and odours that might affect the food;
- (b) is accessible and, if necessary for its cleaning, maintenance, or inspection, can be disassembled;
- (c) is capable of withstanding repeated cleaning; and
- (d) functions as intended. 64

TEMPERATURE AND HUMIDITY

Requirements: The temperature and humidity level in a facility or conveyance where a food is manufactured, prepared, stored, packaged, or labelled must be maintained at levels appropriate for the food, as the case may be, and for the activity being conducted. 65(1)

HEATING, COOLING OR HUMIDITY-CONTROL SYSTEM

Requirements: If the facility or conveyance is equipped with a heating, cooling or humidity-control system, the system must

- (a) if necessary to prevent contamination of a food, be equipped with instruments to control, indicate, and record the temperature and humidity levels;
- (b) be accessible and, if necessary for its cleaning, maintenance, or inspection, is able to be disassembled;
- (c) be capable of withstanding repeated cleaning; and
- (d) functions as intended. 65(2)

Rationale: Inadequate lighting, ventilation, temperature, and humidity can lead to contamination of food, ingredients, packaging materials, and food contact surfaces.

Interpretation: Lighting, ventilation, temperature, and humidity must be in good condition and maintained to prevent the risk of contamination. When the colour of product is being assessed, the lighting must not alter or affect the natural colour of the food. Lighting located overhead or on equipment must be sufficient for the activity being conducted (e.g., inspection, grading). Lighting intensity guidelines are as follows: Storages 110 lux, General Production Areas 220 lux, Inspection Areas 540 lux.

☐ Exterior and Interior Inspection Checklist:
 Adequate lighting is provided (as per guidelines above), and lighting is shatterproof or protected from breakage
 Ventilation provides sufficient air exchange to prevent accumulation of steam, condensation, and dust, to remove contaminated air
$\hfill \Box$ Air intakes are screened and filtered, if applicable
☐ Maple products that are required to be kept at a particular temperature are maintained at appropriate levels for quality purposes or extending shelf life (refrigerated stored at 4°C or less, frozen -18°C or less.). Product stored in coolers or freezers needs to be spaced so that it does not restrict airflow to prevent product from reaching and staying at the appropriate temperature
 Ensure condensation is not an issue in production areas or storages
☐ Glass and Brittle Plastic Checklist (is combined on the Exterior and Interior Inspection Checklist)

REMOVAL AND DISPOSAL OF CONTAMINATED MATERIALS AND WASTE/FREQUENCY AND MANNER

Requirements: An establishment must have means for the removal and disposal of contaminated materials and waste and, if necessary, to prevent contamination of a food, be equipped with a drainage, sewage and plumbing system that functions as intended. 66(1)

Contaminated materials and waste must be removed and disposed of at a frequency that is sufficient to prevent contamination of a food and in a manner that does not present a risk of contamination of a food. 66(2)

Rationale: Effective procedures will prevent the accumulation of waste, inedible or food waste products and the potential contamination of food handling areas and minimize the attraction of pests.

Interpretation: Removal and disposal of contaminated materials and waste must ensure that the risk of contamination to products is minimized through the control of cross-contamination and control of pests.

- ☐ Waste management program:
 - ☐ Waste containers are identifiable (colour coded or labelled, covered)
 - ☐ Waste container cleaning and sanitizing
 - ☐ Emptied at appropriate frequencies (does not allow waste to overflow)
 - ☐ Drainage, sewage (plumbing blueprints/ drawings) adequate with no pooling of water, are trapped and vented to prevent backflow and have no cross-connections



CLEANING STATIONS, LAVATORIES, ETC.

Requirements: If necessary to prevent the contamination of a food, an establishment must be equipped with hand cleaning and sanitizing stations, lavatories, showers, drinking water stations, break rooms or change rooms that

- (a) are appropriately equipped and adequate in number and size for the number of persons using them;
- (b) are located so that they are readily accessible to the persons using them; and
- (c) are capable of withstanding repeated cleaning and, if necessary, to prevent contamination of a food, repeated sanitizing. 67(1)

HAND CLEANING AND SANITIZING STATIONS

Requirements: The hand cleaning and sanitizing stations must permit the effective cleaning of hands. 67(2)

LAVATORIES

Requirements: The lavatories must be located and maintained so that they do not present any risk of contamination of a food. 67(3)

Rationale: Cleaning stations and lavatory installations can become a source of contaminants if they are not properly maintained.

Interpretation: Cleaning stations and lavatories shall be maintained to prevent the risk of contamination.

- ☐ Washrooms are adequate in size, kept clean, and sufficient for the number of staff (i.e., meet applicable OH&S regulations). Washrooms are provided at the site or within a short walk or transportation is provided from the location. If located at the production site, they are situated so they do not create a food safety hazard to product or water sources. Work effects must not be worn into washrooms and a spot must be provided for these to be stored
 - ☐ Handwashing and sanitizing stations permit effective cleaning of hands in the facility, handwashing signage posted with clear steps for proper handwashing
 - ☐ Waste receptacle (covered)
 - ☐ Single use soap, single use hand towels and approved hand sanitizer (if applicable)
 - ☐ Potable water (warm)
- ☐ Cleaning stations and washrooms (lavatories) are on the sanitation schedule (Sanitation program and **Sanitation Record**)



WATER - CONTACT WITH FOOD

Requirements: Any water that might come into contact with a food must be potable, unless it does not present a risk of contamination of the food and must be protected against contamination. 70(1)

STEAM AND ICE

Requirements: Any steam or ice that might come into contact with a food must be made from water that meets the requirements of subsection (1) unless the steam or ice does not present a risk of contamination of the food. 70(2)

WATER

Requirements: Any system that supplies water that meets the requirements of subsection (1) must not be cross-connected with any other system unless measures are taken to eliminate any risk of contamination of a food as a result of the cross-connection. 70(3)

SUPPLY OF WATER, STEAM, AND ICE

Requirements: An establishment must be supplied, as appropriate for the food, and for the activity being conducted, with

- (a) water that is adequate in quantity, temperature, pH, and pressure to meet the needs of the establishment;
- (b) steam that is adequate in quantity and pressure to meet those needs; and
- (c) ice that is adequate in quantity to meet those needs. 71(1)

TREATMENT OF WATER, STEAM, OR ICE

Requirements: Any treatment of water, steam or ice must be applied in a manner that does not present a risk of contamination of a food. 71(2)

Rationale: Water, steam and ice can be a source of biological, chemical, or physical contaminants. Treated water can be a source of contaminants if the chemical treatment or treatment process is incorrectly performed and/or monitored.

Interpretation: Water, steam and ice must be of adequate supply and pose no risk of contamination. Things to consider:

- Water sources: municipal or well, permeate
- Intended use: handwashing, equipment washing and final rinse
- Delivery method
- Water storage tanks (need to be cleaned at the start of the season, after emptying (with approved chemicals or power washed) and added to the Sanitation Program)
- Tampering of water supply
- Adequate supply, temperature, and pressure
- Back-flow prevention, no cross-connections, vacuum breakers on hoses, if applicable

☐ Potable water, steam, and ice supply testing procedure (where applicable):
\square Source of the water, steam, and ice
☐ Treatments used and how they are maintained or monitored if applicable (e.g., UV, filters, chlorination)
$\hfill\Box$ Testing frequency and collection method
☐ Up-to-date schematic of water supply (where applicable)
☐ Potable water, steam, and ice analysis reports, where applicable
☐ Accredited lab used for testing, certificate of accreditation and scope of accreditation for lab
☐ Where permeate is produced, stored, and used, the tank must be kept clean and in good condition (included in Sanitation Program & Preventative Maintenance Program)

SUBDIVISION E: UNLOADING, LOADING AND STORING

CONVEYANCES

Requirements: Any conveyance that is used to convey a food to or from an establishment and that is unloaded or loaded at the establishment

- (a) must be designed, constructed, and maintained to prevent contamination of the food;
- (b) must be constructed of, and maintained using, materials that are suitable for their intended use and, if the materials present a risk of contamination of the food, that are
 - (i) durable,
 - (ii) capable of withstanding repeated cleaning and, if necessary, to prevent contamination of a food, repeated sanitizing, and
 - (iii) free of any noxious constituent;
- (c) must be capable of maintaining the temperature and humidity at levels that are appropriate for the food and, if necessary, to prevent contamination of the food, be equipped with instruments that control, indicate and record those levels;
- (d) must not contain any animal, other than an animal referred to in paragraph 51(2)(a), any pest control product as defined in subsection 2(1) of the Pest Control Products Act or any other material or substance that presents a risk of contamination of the food; and
- (e) must be clean and in a sanitary condition at the time of unloading or loading. 72

UNLOADING AND LOADING

Requirements: Any unloading and loading of a food, from or onto a conveyance at an establishment must be conducted in a manner that does not present a risk of contamination of a food. 73

STORING - FOOD

Requirements: Any storing of a food must be conducted in a manner that does not present a risk of contamination of the food. 74(1)

STORING — OTHER

Requirements: Any storing of conveyances, equipment, sanitizers, agronomic inputs, chemical agents, starter products, packaging material, labels or any other thing that is used in the manufacturing, preparing, storing, packaging, or labelling of a food must be conducted in a manner that does not present a risk of contamination of the food. 74(2)

Rationale: Improper loading, unloading, and storage practices can lead to biological, chemical, or physical contamination of food, ingredients, and packaging materials.

Interpretation: Loading (shipping), unloading, and storage practices must be controlled to ensure they do not pose a biological, chemical, or physical contamination risk to products, ingredients, and packaging materials. Things to consider:

Loading

- Product to be loaded should remain at proper storage temperatures until ready to load, should be neatly stacked, and securely wrapped.
- Product should not be placed directly on the transport vehicle floor.
- Temperature requirements must be maintained during shipping.
- Inspect the carrier for temperature, cleanliness, and physical conditions before loading.
- Ensure the carrier is not carrying any hazardous or incompatible materials - food products and packaging must be transported separately from incompatible food products and items (e.g., meat, fish, chemicals).
- Should be completed in a manner to avoid contamination of the product.

Unloading

- Inspect the carrier for temperature, cleanliness, and physical conditions before unloading.
- Ensure the carrier is not carrying any hazardous or incompatible materials - food products and packaging must be transported separately from incompatible food products and items (e.g., meat, fish, chemicals).
- Ensure materials received are from an approved supplier/source and that required information, such as, specifications, letter of guarantee and/ or certificates of analysis are on file to ensure any incoming materials do not pose a risk of contamination to your product.
- Inspect the product to ensure temperature requirements were met, and that there is no evidence of spoilage, damage, foreign material, off-odours, and best before date is adequate.
- Ensure unloading occurs in a timely manner and in an appropriate area which avoids unloading product, ingredients, and packaging material under inadequate conditions which could pose a risk to the product (e.g., rain).

Storage

- Product, ingredients, and packaging material should be protected from cross-contamination and damage during storage. Sap is to be stored separate from market-ready product.
- Food product and market-ready packaging is kept off the floor or ground such as on pallets or shelves.
 Shelving units and pallets must be kept away from the walls to facilitate inspection and cleaning.
- Practice First In, First Out (FIFO) stock rotation to ensure older product, product close to its shelf-life are used first.
- Production site equipment must be stored separate from packaging, product, oil, and non-food grade lubricants which can pose a risk of contamination.
- Chemicals must be stored separately from product, ingredients, and packaging material.
- Chemical storages must be clean, well ventilated and in a location where there is no cross-contamination risk to product, ingredients, and packaging. Access should be restricted to only personnel who have been trained on the proper handling and use of chemicals.

CHECKLIST
☐ Loading, Unloading and Storage Procedures:
☐ Approved supplier/source
 Non-conforming product SOP in place to address any product that does not meet the requirements
☐ Product inspections
☐ Food carrier inspections
Food carrier temperature inspections (where applicable)
 Loading/unloading is performed in a matter to prevent risk of contamination
$\ \square$ First In First Out stock rotation is followed
 Where and how product, packaging, chemicals, and equipment are stored
□ Receiving Record

☐ **Shipping Record**/Invoices/Shipping Logbook

SUBDIVISION F: COMPETENCY

COMPETENCIES AND QUALIFICATIONS

Requirements: Any person who is involved in the manufacturing, preparing, storing, packaging or labelling of a food must have the competencies and qualifications that are necessary to carry out their duties. 75

Rationale: Competencies and qualifications increase awareness of potential hazards and the responsibilities that personnel have to minimize contamination risks.

Interpretation: Personnel performing tasks shall be competent and qualified to carry out their roles and responsibilities. Personnel must be adequately trained on food hazards, food hygiene, clean/sanitary conditions, general hygienic practices, and complete technical training such as how to use equipment and monitoring devices to properly monitor critical control points (CCPs). Training is ongoing, must be effective and documented. Visitors and contractors will also require training on your hygienic practices if entering production and storage areas.

Training should occur at the start of the season for returning employees, and with each new hire prior to starting work. For non-seasonal operations, training or refresher training must occur at least annually and be conducted in a language understood by employees.

Training may include (but is not limited to):

- Grading
- Best Management Practices
- Preventative Maintenance and Calibration
- Pest Control/Pest Sightings
- Receiving, Transport, and Storage Activities
- Sanitation and chemical handling
- How to fill out records
- Good Manufacturing Practices
- Personnel Hygiene
- Allergens
- Visitor policy

- ☐ Personnel Training Program:
 - ☐ What kind of training is required and completed, who trains and at what frequency?
 - ☐ Describes how training effectiveness checks will be completed and recorded (written test, observing staff perform tasks correctly after training or another means of evaluation that is appropriate for the trained task)
- ☐ Employee Training Certificates, if applicable (e.g., pesticide applicators license)
- ☐ Personnel Training Records
 (date, who was trained, what they were trained on, by whom, effectiveness check)



SUBDIVISION G: HYGIENE

CLOTHING, FOOTWEAR, AND PROTECTIVE COVERINGS

Requirements: Any person who enters or is in an area where a food is manufactured, prepared, stored, packaged, or labelled must wear clothing, footwear and protective coverings, including gloves, a hairnet, a beard net and a smock that are in good condition, clean and in a sanitary condition and that are appropriate for the food and the activity being conducted. 76

PERSONAL CLEANLINESS

Requirements: Any person who enters or is in an area where a food is manufactured, prepared, stored, packaged, or labelled must maintain personal cleanliness to prevent contamination of the food, including by cleaning and, if necessary, by sanitizing their hands

- (a) immediately on entering the area;
- (b) immediately after using a lavatory;
- (c) immediately before beginning to conduct the activity;
- (d) at a frequency appropriate for the food and the activity being conducted. 77

SPITTING, CHEWING GUM AND OTHER ACTS

Requirements: Any person who enters or is in an area where a food is manufactured, prepared, stored, packaged, or labelled must refrain from spitting, chewing gum, using tobacco products, eating, having unnecessary contact with the food and doing any other act that presents a risk of contamination of the food. 78

OBJECTS AND SUBSTANCES - RISK OF CONTAMINATION

Requirements: Any person who enters or is in an area where a food is manufactured, prepared, stored, packaged, or labelled must refrain from wearing or using any object or substance that presents a risk of contamination of the food. 79

REPORTING OF DISEASE, ILLNESS, SYMPTOMS, AND LESIONS

Requirements: Any person who works in an area where a food is manufactured, prepared, stored, packaged, or labelled and who has a disease or illness, symptoms of a

disease or illness or an open or infected lesion must report them to the operator. 80

COMMUNICABLE DISEASE AND LESIONS – RISK OF CONTAMINATION

Requirements: The operator must prevent any person who is suffering from, or is a known carrier of, a communicable disease or who has an open or infected lesion from entering or being in an area of an establishment where a food is manufactured, prepared, stored, packaged, or labelled if the person's condition presents a risk of contamination of the food. 81

Rationale: Personnel play a key role in producing safe quality food; improper hygienic habits can lead to biological, chemical, or physical contamination of food, ingredients, packaging materials, and food contact surfaces. Visitors and contractors can also be a source of contamination and must be properly trained on food hazards, food hygiene, clean/sanitary conditions, and general hygienic practices to prevent the risk of contamination or spread of diseases.

Interpretation: Personnel hygienic standards minimize the risk of product contamination and shall be adopted by all personnel, temporary workers, contractors, and visitors. Good Manufacturing Practices (GMP) that should be in place at your operation for all of the approved personnel are:

- Showering/bathing regularly and wearing clean clothes/workwear and footwear
- Clean, short fingernails (e.g., no fingernail polish or artificial nails)
- No lash extensions or false eyelashes
- No jewelry in production areas (medical jewelry and plain wedding bands may be permitted)
- No smoking, chewing gum, eating, drinking, or spitting in the production or storage areas
- Wear hair and beard nets and suitable clothes/ gloves/footwear
- No personal items in production or storage areas
- No loose objects in the production area (e.g., pens, paper clips, pins, buttons, etc.)
- Cover cuts and wounds with a secure, waterproof covering that can be easily detected if it comes off (e.g., bright coloured, metal detectable)

- Employees of food-contact areas displaying any symptoms of an infectious disease or illness should refrain from coming to work and only return 48 hours after their last symptom has subsided as their disease or illness could be transmitted through the product
- Employees should be trained to sneeze or cough into their elbow or turn their head into their shoulder, stepping away from the processing and packing area to avoid contaminating the product and/or equipment
- Employees should be trained on and follow proper traffic flow and product flow throughout the facility in order to prevent risk of cross-contamination or cross-contact
- Handwashing sinks should not be used to clean food equipment and vice versa
- Doors must be kept closed at all times, including any applicable refrigerator and freezer doors
- Waste bins must be kept clean, in good condition and emptied whenever full
- Employees should be trained on how to manage blood and bodily fluid spills, glass breakage, spilled product, product that has fallen on the floor or has been exposed to dripping condensation
- Product packaging should only be used for the product, not for things such as equipment parts, temporary stands, tools, or garbage
- Handwashing, the single most important thing that people can do to prevent contamination of food.
 Hands need to be washed at the following times:
 - At the start of a shift
 - After breaks
 - After eating, smoking, or drinking
 - Each time an employee enters the production area
 - After handling anything dirty and going back to handle product, packaging, equipment, etc. (e.g., picking something up off the floor, handling tools, handling garbage, handling pallets, etc.)
 - After using the washroom
 - After coughing or sneezing
 - After touching their face, adjusting hair/beard nets, or using a tissue

Note: if your handwashing station is not hands-free, please make sure you use paper towel to turn off the taps when you finish.

Proper handwashing procedures can be found under the Quality & Food Safety resource tab of the Perennia website (www.perennia.ca)

CHECKLIST
☐ Personnel Training Program
☐ Personnel GMP and Hygiene Policy:
 Personnel Protective Equipment - clothing and footwear
 Personal Cleanliness – handwashing and sanitizing, where applicable
☐ Personal Conduct – no food, drinks, smoking, vaping, spitting in production or storage areas. If water is allowed or required for health and safety reasons, then it must be consumed and stored away from product areas, and proper handwashing following handling
 Personal objects – jewelry and personal effects. Personal effects must be stored away from the production and storage areas
☐ Reporting of health conditions
☐ Communicable disease and lesions
☐ Daily GMP Checks
$\ \square$ Proof of training effectiveness checks
☐ Personnel Training Records
□ Visitor's Record

DIVISION 5: INVESTIGATION, NOTIFICATION, COMPLAINTS AND RECALL

INVESTIGATION

Requirements: An operator who suspects on reasonable grounds that a food presents a risk of injury to human health or does not meet the requirements of the Act, or these Regulations must immediately investigate the matter. 82(1)

NOTIFICATION AND MITIGATION OF RISK

Requirements: If the investigation establishes that the food presents a risk of injury to human health, the operator must immediately notify the Minister and immediately take action to mitigate the risk. 82(2)

COMPLAINTS PROCEDURE

Requirements: An operator must prepare, keep and maintain a document that sets out a procedure for receiving, investigating and responding to complaints that are received in relation to a food. 83(1)

COMPLAINTS

Requirements: If a complaint is received, the operator must implement the procedure and prepare a document that sets out the details of the complaint, the results of the investigation and the actions taken based on those results and keep it for two years after the day on which the actions are completed. 83(2)

RECALL PROCEDURE

Requirements: An operator must prepare, keep and maintain a document that sets out a recall procedure that enables the effective recall of a food, the name of a contact person who is responsible for the procedure and the name of a contact person who is responsible for conducting recalls. 84(1)

RECALL SIMULATION

Requirements: The operator must, at least once every 12 months,

(a) conduct a recall simulation, based on the recall procedure, and

(b) prepare a document that sets out the details of how the recall simulation was conducted and the results of the simulation, and keep that document for two years after the day on which the recall simulation is completed. 84(2)

RECALL — NOTICE TO MINISTER

Requirements: If an operator determines that a food should be recalled because it presents a risk of injury to human health, the operator must immediately notify the Minister 84(3)

RECALL — IMPLEMENTATION

Requirements: If a food is the subject of a recall because it presents a risk of injury to human health, the operator must

- (a) immediately implement the recall procedure; and
- (b) prepare a document that sets out the details of the recall, including any information that substantiates its effectiveness and keep the document for two years after the day on which the recall is initiated. 84(4)

IMPORTED FOOD

Requirements: The holder of a licence to import must comply with sections 82 to 84 in respect of a food that is imported. 85

Rationale: Food recalls can be triggered by a number of hazards within or external to a facility. Quickly regaining control of implicated lots of products is crucial in preventing the risk of hazard to consumers.

Complaints from any source are important indicators of possible deficiencies in the system or the possible presence of a contaminant, pest, or disease. When the complaint handling system itself is deficient, it could result in failure to identify, control, and mitigate risks.

Interpretation: The company shall have a recall and complaints plan in place to manage complaints and recalls effectively should an issue arise.

Recall Plan/Mock Recall (Recall Simulation)

Testing your recall plan is just as important as having one. Since recalls can happen to anyone at anytime in any industry, you must be prepared at all times. A Mock Recall (Recall Simulation) is a means of testing the Recall Plan to ensure it continues to be effective in tracing all materials coming in and all products going out within a reasonable amount of time. It is also a good time to test the communication plan to ensure all contact lists are up to date.

Mock recalls must be completed at least once every 12 months and your finished product must be traceable to the customer (one step forward) and you must also show traceability through to the ingredient/primary input supplier (one step back). To do this, the date of receipt as well as use of incoming materials, food contact packaging or other processing inputs where reasonable will need to be recorded.

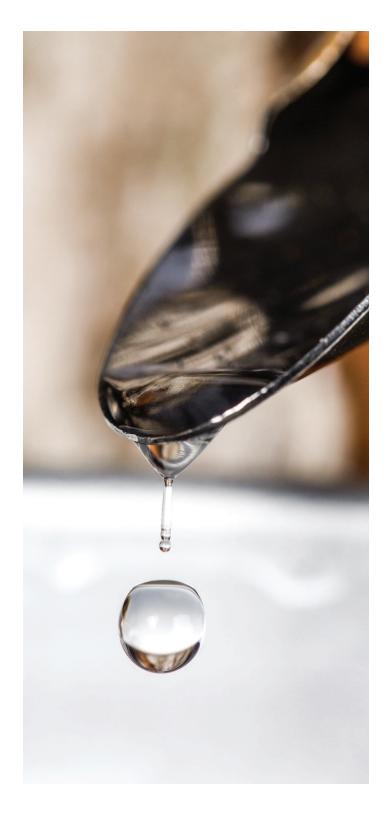
Use your mock recall/recall simulation record(s) to document the exercise:

- Determine a realistic scenario that would trigger a recall
- Select a product, ingredient/input that has been sent to a customer
- Trace the particular lot to the customers affected, or trace an ingredient/input back to the supplier
- Do not forget to include reworked products, product samples that were sent out, product samples sent for testing, or product taken for staff or personal use
- Keep copies of all documentation used for the exercise
- Do not let customers know you are performing a mock recall; this may unintentionally trigger an unwanted/unnecessary recall with that customer
- Determine if your recall plan is clear, if your customer/supplier contact list is up to date, if you were able to account for 100% of the product you produced and packaged, or ingredient/input used
- Determine if you need to improve your traceability/ recall plan and re-test the process
- Keep all documents for two years

Complaints

Complaints can vary from quality issues to food safety concerns or even legality issues (misrepresentation). A documented customer/consumer complaint plan enables a food business to quickly assess and investigate the complaint to determine severity. Complaints related to food safety, pests, and product misrepresentation must be investigated to determine the root cause and corrective actions. Complaint files allow a business to track the number and type of complaints and, therefore, can be used as a form of continuous improvement. Complaints must be assessed, investigated, and resolved in a timely manner. Should processes or procedures require changes based on the results of the assessment and investigation, ensure that changes are made and appropriate personnel are notified.

CHE	ECI	KLIST					
	Cu	stomer/Consumer Complaint Procedure:					
		Who records the complaint?					
		Who investigates and categorizes the complaint (must be a qualified individual)?					
		Brief procedure for receiving and conducting the investigation and follow-up with CFIA and or customer					
	Re	call and Mock/Simulation Recall Program:					
		Roles and responsibilities					
		Recall team and contact information					
		Recall steps/procedure. Steps should be brief and outline what needs to happen in a recall situation, list the records at each step that you will need to review to be able to find your product. Clear, concise steps will be easier to follow in a stressful recall situation. See CFIA's step by step recall procedure					
		CFIA contact information					
	☐ Customer/Consumer Complaint Records						
	Re	call Management Records					
		ock Recall/Recall Simulation Records st your recall program at least once per year)					
	Em	nployee training					



DIVISION 6: PREVENTIVE CONTROL PLAN

LICENCE HOLDERS

Requirements: A licence holder must prepare, keep and maintain a written preventive control plan that meets the requirements of section 89 for any activity identified in their licence that they conduct in respect of a food. 86(1)

EXCEPTION — FOOD TO BE EXPORTED

Requirements: Despite subsection (1), a preventive control plan is not required to be prepared, kept or maintained for any activity that the licence holder conducts in respect of a food, other than fish or a meat product, that is exported, unless a certificate or other document referred to in section 48 of the Act is sought in respect of the food. 86(2)

EXCEPTION — SALES OF \$100,000 OR LESS

Requirements: Despite subsection (1), if a licence holder's gross sales that are derived from food are \$100,000 or less for the 12 months before the day on which they most recently made an application for the issuance, renewal or amendment of a licence, a preventive control plan must be prepared, kept and maintained only for any activity that they conduct in respect of

- (a) a food animal, meat product, fish, dairy product, egg, processed egg product or processed fruit or vegetable product that is identified in their licence; and
- (b) a food in respect of which a certificate or other document referred to in section 48 of the Act is sought. 86(3)

IMPLEMENTATION

Requirements: Any person who is required to prepare, keep and maintain a preventive control plan must implement that plan. 88

CONTENT OF PREVENTIVE CONTROL PLAN

Requirements: The preventive control plan must include

(a) a description of the measures for ensuring that the applicable requirements of sections 201 and 205, subsection 206(1), sections 208, 218, 221, 296, 306, 307, 316, 317, 321, 322, 324 to 326 and 328 are met;

- (b) a description of the measures for ensuring that the food is packaged and labelled in a manner that does not contravene subsection 6(1) of the Act;
- (c) in relation to the applicable requirements of these Regulations,
 - (i) a description of the biological, chemical and physical hazards that are identified under subsection 47(1) as presenting a risk of contamination of a food, of the control measures for preventing or eliminating those hazards or reducing them to an acceptable level and of the evidence that the control measures are effective,
 - (ii) a description of the critical control points, of the related control measures and of the evidence that the control measures are effective,
 - (iii) a description of the critical limits for each critical control point,
 - (iv) the procedures for monitoring the critical control points in relation to their critical limits,
 - (v) the corrective action procedures for each critical control point,
 - (vi) the procedures for verifying that the implementation of the preventive control plan results in compliance with the provisions of the Act and these Regulations, and
 - (vii) documents that substantiate that the preventive control plan has been implemented with respect to subparagraphs (i) to (vi); and
- (d) in relation to the applicable requirements of sections 128 to 136, paragraphs 140(b) and (c) and sections 141 to 144,
 - (iii) a description of the performance criteria for evaluating the effectiveness of each of those measures,
 - (iv) the procedures for monitoring each of those measures,
 - (v) the corrective action procedures for each of those measures,
 - (vi) the procedures for verifying that the implementation of the preventive control plan results in compliance with the provisions of the Act and these Regulations,

(vii) the procedures for auditing, on a regular basis, the outcome of the implementation of the preventive control plan, and

(viii) documents that substantiate that the preventive control plan has been implemented with respect to subparagraphs (i) to (vii); and

(e) supporting documents that show evidence of the information recorded under paragraphs (a) and (b), subparagraphs (c)(i) to (vi) and (d)(i) to (vii). 89(1)

RETENTION PERIOD OF DOCUMENTS

Requirements: Each document referred to in subparagraphs (1)(c)(vii) and (d)(viii) must be kept for two years after the day on which it is prepared. 89(2)

ADDITIONAL CONTENT — IMPORT

Requirements: The preventive control plan of the holder of a licence to import must also include the information specified in subparagraphs (1)(c)(i) to (vii) in relation to the requirements of section 11. 89(4)

ADDITIONAL CONTENT — EXPORT

Requirements: The preventive control plan of the holder of a licence to export must also include the information specified in subparagraphs (1)(c)(i) to (vii) in relation to the requirements of subsection 15(1). 89(5)

Rationale: Preventive Control Plans identify potential hazards associated with the process and production of products. By identifying and assessing all potential hazards, control measures can be designed and implemented to reduce, prevent, or eliminate hazards.

Interpretation: A Preventive Control Plan shall be developed and implemented to reduce, prevent, or eliminate hazards associated with inputs, process, and production. A Preventive Control Plan is similar to a HACCP Plan but incorporates preventive controls/prerequisite programs including traceability and the HACCP plan.

The Preventive Control Plan will be documented through a hazard analysis process. For each different product/process type at your facility, there needs to be a Preventive Control Plan. Hazards will be identified for incoming materials and ingredients, cross-contamination points, and process

steps. Appropriate controls for each will be implemented to reduce the risk or eliminate the hazard. Those control measures need to be monitored to ensure they effectively control the hazard. The control measures can include your preventive controls or critical control point(s). If one of those control measures fails, then corrective actions need to be taken and a corrective action/deviation report documented. The root cause of the incident needs to be determined and addressed to prevent reoccurrence.

There are 12 steps when implementing a HACCP Plan.

These steps can be divided into the five preliminary steps and the seven basic principles of HACCP.

Five Preliminary Steps

- Assemble a Preventive Control Plan Team/HACCP Team
- 2. Describe the product
- 3. Identify its intended use
- Construct a process flow diagram and a plant schematic
- 5. On-site verification of the process flow diagram and plant schematic

Seven Basic Principles of HACCP

- List all potential hazards associated with each step, conduct a hazard analysis, and identify control measures
- 7. Determine critical control point(s) (CCPs)
- 8. Establish validated critical limits
- **9.** Establish a system to monitor control of the critical control points
- Establish corrective actions to be taken when monitoring indicates a deviation of the critical limit at a critical control point
- **11.** Validate the HACCP plan and then establish verification procedures to confirm the HACCP system is working as intended
- 12. Establish record keeping

When implementing a Preventive Control Plan/HACCP Plan, the first step is deciding who should be on the Preventive Control Plan/HACCP Team for each HACCP Plan. Secondly, determine how many HACCP Plans are needed. In other words, how many different products/ processes? For example, a maple producer that produces maple syrup and maple barbecue sauce will have two HACCP plans.

A Preventive Control Plan/HACCP Team can consist of senior management and production employees from various disciplines, such as:

- CEO, Owner, Operator
- Quality Assurance, HACCP Coordinator, PCP Manager
- Receiver/Shipper
- Production Supervisor/Manager
- Sanitation
- Maintenance
- Purchasing and/or Sales
- Product Development

A smaller company may only have two or three people that fill these roles. The person that deals with product safety and quality will have the proper food safety training and will be the one that leads the Preventive Control Plan/HACCP Team. A cross-functional team is important because food safety and quality is everyone's responsibility, and one person cannot successfully do it on their own. The rest of the HACCP steps will correspond with forms and are discussed in further detail in the following section.

The Forms

A HACCP Plan generally consists of 11 forms that are best completed in the following order:

- Form 1: Product Description
- Form 2: List of Product Ingredients and Incoming Materials
- Form 3: Process Flow Diagram
- Form 4: Facility Schematic
- Form 5: Biological Hazard Identification*
- Form 6: Chemical Hazard Identification*
- Form 7: Physical Hazard Identification*
- Form 8: CCP Determination and Other Control Measures (Decision Tree) *

- Form 9: Hazards Not Controlled by the Facility
- Form 10: Critical Control Point(s) (CCP)
- Form 11: Process Controls (PC)

*Forms 5 to 8 can be combined into a Hazard Identification and CCP Determination form.

Each form is in place to help guide the team through the hazard analysis process in order to eliminate or decrease the food safety risk in the products.

Form 1's basic purpose is to record the products produced, their specific requirements and special characteristics that need to be considered when completing a HACCP plan. Form 1 should include the product name, important product characteristics, how the product will be used (including who the product may be consumed by), packaging used, shelf-life of the product, where it will be sold, important labelling instructions, and special distribution controls. With regard to the intended use of the product, be very clear in the documentation and on the labels as to the intended final use of the product; for example, include a 'Ready to eat' statement.

The purpose of **Form 2** is to help identify all ingredients, processing aids, other inputs, and packaging materials used to produce the products that were recorded in **Form 1**.

Form 3 is documented to show the process steps for the products listed in Form 1. The process flow diagram gives you a snapshot of the process and helps determine what potential hazards could be present at each step in the process. It is important to include everything (inputs and outputs), including rework, store sales and waste where applicable. Each step in the process will be numbered, and a hazard category will be assigned to each (biological, chemical, and/or physical). This information will be used in the hazard analysis in Form 5-8.

The purpose of Form 4 is to document the facility and how ingredients, product, rework, allergens, chemicals, personnel, and waste move through it. This form is for all products that are made in the facility. It is important to include all processes in order to identify any crosscontamination and cross-contact points. Examples of crosscontamination or cross-contact are waste with finished product or raw ingredients, chemicals with product, people with product, and allergens with non-allergens. Draw a picture of the facility as if you were looking down onto it, make sure all equipment and rooms are included and labelled. To show the movement of ingredients, product, rework, allergens, chemicals, personnel, and waste, use different colours or different types of lines. Where the different lines cross are the potential cross-contamination points. The cross-contamination points identified here will be inputted into Form 5-8. Pest control trap numbers (bait stations, tin cats and fly traps) can also be included on this schematic, or you can create a separate map as part of the pest control program.

Forms 5, 6, 7 and 8 can be combined to make it easier to complete the hazard analysis. This is the form that pulls the information gathered in Forms 1-4 together and then looks at the process, hazards, and risk in detail. This form asks five questions to help guide you through the hazard analysis process and determining if the hazards identified are controlled under the documented preventive controls/ prerequisite programs or will need to be controlled as a process control (PC) or a critical control point (CCP) and what their risk is. Note that you cannot have a PC without a CCP, but you may have a CCP without a PC.

To assist with Hazard Analysis, CFIA has an online tool called the Reference Database for Hazard Identification (RDHI). It can be found on their website; the link is provided below (last modified 2020-09-16). CFIA periodically updates the online tool. However, if you produce an innovative item, it may not be listed in this reference tool.

Reference Database for Hazard Identification - Introduction (inspection.gc.ca)

Sometimes hazards are beyond the control of the facility. In those instances, **Form 9** comes into play. On this form, you record the hazard, what part of the process this affects (before or after receipt), and how the outside source will control it. For example, if using diatomaceous earth as a processing aid, the supplier is responsible for ensuring it is safe for use in food manufacturing, not the processor.

If it is determined that critical control points (CCPs) are required to control a hazard, then **Form 10** is used to describe the details of the CCP, who will monitor it, when and how it will be monitored, any critical limits that need to be adhered to and what records and standard operating procedures (SOPs) are used. Other details that need to be considered are how to verify that the hazard is being controlled properly and what happens if the critical limit(s) are not met.

The process controls (PC) that were determined through the hazard analysis are recorded on **Form 11**. Much like **Form 10**, you document the details and refer to the CCP that the process control is linked to.

Examples of HACCP Forms 1-11 are available in the CFIA Archive for "HACCP Generic Models and Commodity-Specific Food Safety Guidance Documents".

Verify the system is effective. Is the final product completely safe, or was something missed? Records should be simple and nonredundant. Feedback from employees on their usability will be beneficial as employees will be more likely to fill them out if they are easy to use. Records must be legible and filled out at the actual time the check was performed. They must be signed by the person who monitors the task and by the person who verifies that the record is complete, and the task was performed properly. Records must be kept for at least two years or for the duration of the product's shelf life plus one year unless otherwise requested.

Record Keeping/Document Control

Keeping track of the HACCP records and documentation of your quality and food safety system is critical. Records will demonstrate the application of the HACCP plan. Documents are an important tool for training existing and new employees as well as ensuring procedures are followed consistently by all. Documents demonstrate the effective implementation of your quality and food safety management program and allow for continuous improvement. They are most often a condition of registration, licensing, or certification and are evidence of due diligence on your company's behalf. Creating a basic template for documents such as procedures and work instructions will allow you to keep records organized, up to date, legible, accurate and readily accessible. Note that it is important to use appropriate languages in your documents.

CHECKLIST						
☐ Preventive Control Plan Team/HACCP Team☐ Product Description:	☐ Hazard Identification and Critical Control Point(s) determination:					
☐ Product Name(s)	☐ Incoming Materials					
☐ Source of Raw Material	☐ Cross Contamination Points					
☐ Important Final Characteristics	☐ Process Steps					
☐ Ingredients & Inputs	 Outputs (e.g., waste, compost, rework, returns, rejects, etc.) 					
☐ Product Packaging	☐ Preventive Controls Controlling Hazards					
 How the End Product is to be used (intended use and include who the product may be consumed by) 	☐ Critical Control Point(s), if applicable					
☐ Shelf Life	☐ Risk Assessment of the Hazards					
☐ Where the Product Will be Sold	☐ Hazards Not Controlled by the Operator/Company					
☐ Special Labelling Instructions.	☐ Critical Control Point Form (if applicable, see above):					
☐ Special Distribution Controls	☐ Process Step					
☐ Incoming Ingredients and Materials:	☐ Hazard Description and Control Measure					
☐ Raw Ingredients	☐ Critical Limit					
☐ Added Ingredients	☐ Monitoring Procedure					
 Packaging Materials (Primary Packaging, Secondary Packaging, Tertiary Packaging) 	☐ Deviation Procedure☐ Verification Procedure					
☐ Other Inputs (water, steam, ice)	☐ Record Keeping					
☐ Allergens	☐ Process Control Form, if applicable:					
☐ Process Flow:	 Process Control number and associated Critical Control Point 					
 Inputs, process steps, and outputs (e.g., waste, compost, rework, returns, rejects, etc.) 	☐ Hazard Description					
\square List of biological, chemical, and physical hazards.	\square Standards					
☐ Identification of Critical Control Point(s), if applicable	☐ Monitoring Procedure					
☐ Plant Schematics:	☐ Deviation Procedure					
☐ Traffic Flows	☐ Verification Procedure					
☐ Cross Contamination Points	☐ Record Keeping					
☐ Water Supply Lines	☐ Document Control Procedure:					
☐ Sewage and Drain Lines	☐ Retention of documents					
☐ Raw and Finished Product Flow	☐ Electronic or hard copies					
☐ Chemical Flow	☐ Maintenance and Reassessment Procedures					
☐ Waste Flow	☐ Corrective Action Plan					
☐ Pest Control Devices	☐ Corrective Action Report					
☐ Handwash Stations						
☐ Chemical Storage Areas						
☐ Equipment						



TRACEABILITY — PART 5 OF THE SFCR

DOCUMENTS

Requirements: Any person who sends or conveys a food from one province to another, or who imports or exports it, any holder of a licence, to manufacture, process, treat, preserve, grade, store, package or label a food in its imported condition and any person who grows or harvests fresh fruits or vegetables that are to be sent or conveyed from one province to another or exported must, if they provide the food to another person, prepare and keep documents that set out

- (a) the common name of the food, a lot code or other unique identifier that enables the food to be traced and the name and principal place of business of the person by or for whom the food was manufactured, prepared, produced, stored, packaged, or labelled;
- (b) except if they provide the food to another person as a sale at retail, the date on which it was provided and the name and address of the person to whom it was provided;
- (c) if they were provided the food by another person, the name and address of that person and the date on which it was provided; and
- (d) the name of any food commodity that they incorporated into the food or from which they derived the food and, if they were provided the food commodity by another person, the name and address of that person and the date on which it was provided. 90(1)

DOCUMENTS — RETAIL SALE DOCUMENTS

Requirements: Any person who sells a food at retail, other than a restaurant or other similar enterprise that sells the food as a meal or snack, must prepare, and keep documents that include the information specified in paragraphs (1)(a), (c) and (d). 90(2)

RETENTION PERIOD OF DOCUMENTS

Requirements: The documents referred to in subsections (1) and (2) must be kept for two years after the day on which the food was provided to another person or sold at retail and must be accessible in Canada. 90(3)

PRODUCTION OF DOCUMENTS

Requirements: Any person who has received a request from the Minister for a document referred to in section 90, or any part of such a document, must provide it to the Minister

- (a) within 24 hours after receipt of the request, or within
 - (i) any shorter period that is specified by the Minister, if the Minister believes that it is necessary in order to identify or respond to a risk of injury to human health associated with a food commodity, or
 - (ii) any longer period that is specified by the Minister, if the Minister believes that the document is not necessary for a recall that is or may be ordered under subsection 19(1) of the Canadian Food Inspection Agency Act; and
- (b) if provided electronically, in a single file and in plain text that is capable of being imported into and manipulated by standard commercial software. 91(1)

DEFINITION OF PLAIN TEXT

Requirements: In paragraph (1)(b), plain text means data that is not encrypted and whose semantic content is available. 91(2)

Rationale: Effective traceability programs allow companies to trace their product quickly and effectively to recall products that are suspect or identified as unsafe to protect the general public from preventable risks. Traceability documents and labelling/coding can make product recall situations less difficult when implemented correctly.

Interpretation: The operation shall have methods developed and implemented to trace product throughout the process until final packaging; one step back (to the immediate supplier) and one step forward (to the immediate customer). Traceability documents must contain the following information:

- Common name of the food
- A lot code or unique identifier (lot code for consumer prepackaged and either a lot code or unique identifier for prepackaged products other than consumer prepackaged such as a shipping container)
- Name and principal place of business of the person by or for whom the food was manufactured, prepared, produced, stored, packaged, or labelled
- The date which you provided the food to another person (does not include retail customers) and the name and address of the person (e.g., shipping record, invoices)
- The date on which the food was provided to you and the name and address of the person who provided it (e.g., receiving record)
- The name of any food commodity that you incorporated into the food or from which you derived the food and, if you were provided with the food commodity by another person, the name and address of that person, and the date on which it was provided

Take note that records must be kept, as hard copy or electronically, for at least two years after that lot was provided to you or you provided the food to a customer. They must be accessible in Canada.

- Processors must be able to trace ingredients back to the original supplier and trace all finished products to the first level of distribution.
- Importers must be able to trace imported products back to the foreign supplier and trace products to the first level of distribution.
- Exporters must be able to trace products back to the supplier and trace products to the buyer.
- Retailers must be able to trace product back to the supplier.



LABELLING

Requirements: Any person referred to in subsection 90(1) or (2) must ensure that a label that bears the information specified in paragraph 90(1)(a) is applied or attached to any food, or accompanies any food, that is provided to another person. 92(1)

CONSUMER PREPACKAGED FOOD

Requirements: In the case of consumer prepackaged food that is not packaged at retail, the unique identifier referred to in paragraph 90(1)(a) must be a lot code. 92(2)

EXCEPTION

Requirements: Subsections (1) and (2) do not apply in respect of

- (a) a food at the time of its export;
- (b) a food, other than a consumer prepackaged food, at the time of its sale at retail; or
- (c) a prepackaged food described in paragraphs 213(a) to (c) at the time of its sale at retail. 92(3)

EXCEPTION — FOODS DESCRIBED IN PARAGRAPHS 219(1)(A) AND (B)

Requirements: Despite subsection (1), any food described in paragraph 219(1)(a) or (b) is not required to be labelled with the common name of the food at the time of its sale at retail. 92(4)

EXCEPTION — FOODS DESCRIBED IN SECTION 220

Requirements: Despite subsection (1), any food described in section 220 is not required to be labelled with the name and principal place of business of the person by or for whom the food was manufactured, prepared, produced, stored, packaged or labelled. 92(5)

Rationale: Labelling/coding can make product recall situations less difficult when implemented correctly. Correct labelling and coding enable the next person to handle, display, store, and use the product safely.

Interpretation: The operation shall have methods developed and implemented to ensure that products will be correctly labelled and coded. Packaged products must be labelled with the common name, name and principal place of business, and a lot code or unique identifier for traceability purposes. Lot codes should be recorded on your production records as well as any shipping records/invoices.

Lot codes may be alphabetic, numeric, or alphanumeric. Examples of allowable lot coding options for maple products are as follows.

- Production/pack date
- Best before date (if applicable)
- Establishment number in combination with a unique identifier
- SFC license number in combination with a unique identifier
- Or any other unique code that can be used for traceability reasons

CHECKLIST
☐ Traceability Program, detailing the following:
☐ Product Coding
☐ Product Labelling
☐ Employee Training

PACKAGING — PART 10 OF THE SFCR

DIVISION 1: GENERAL

REQUIREMENTS FOR PACKAGES

Requirements: A prepackaged food that is sent or conveyed from one province to another or that is imported or exported must meet the following requirements:

- (a) its package
 - (i) must be suitable for its intended use and appropriate for the food,
 - (ii) must be capable of protecting the food against moisture, loss, damage, contamination and deterioration during normal handling, storing and conveying,
 - (iii) must be clean and in a sanitary condition,
 - (iv) must be of sound construction,
 - (v) must be free from odours that might affect the food,
 - (vi) must not impart any undesirable substance to the food,
 - (vii) must not have a design or mark, or be of a colour, that enhances the appearance of the food with respect to its quality or composition. 186

Rationale: Packaging must be appropriate and not contaminate or permit contamination of the product.

Interpretation: Ensure food packaging is suitable for its intended use, does not pose a risk of contamination to the product, and that your product is packaged correctly, the following checklist will help you confirm these requirements are met.

Lot codes for food-contact packaging should also be recorded on your receiving and production/packing records for traceability.

CHECKLIST

- ☐ Written specifications are on file for food-contact packaging
- ☐ Packaging material meets regulatory requirements
- ☐ Packing Procedure should detail the following:
 - ☐ Damaged or defective packaging material is not used during the packing process
 - ☐ Packaging material is protected from risks of contamination during receiving, storage and packing
- ☐ Employee Training
- □ Packing Records



LABELLING, PART 11 OF THE SFCR

DIVISION 3 – SPECIFIC REQUIREMENTS FOR CERTAIN FOODS

SUBDIVISION K: MAPLE PRODUCTS

NET QUANTITY

Requirements: The label of a prepackaged maple product must bear a declaration of net quantity in metric units. 280(1)

EXCEPTION

Requirements: Subsection (1) does not apply in respect of maple syrup unless it is graded in accordance with these Regulations. 280(2)

IMPORTED MAPLE PRODUCTS

Requirements: The label of the following maple products must bear the name of the foreign state of origin:

- (a) any imported prepackaged maple syrup whose net quantity is 5 L or less; and
- (b) any other imported prepackaged maple product whose net quantity is 5 kg or less. 281

Canadian Standards of Identity: Volume 6 - Maple Products

(from CFIA's documents incorporated by reference)

Maple Products

- Maple Syrup is syrup obtained exclusively by the concentration of maple sap or by the dilution or solution of a maple product in potable water.
- 2. Any maple product other than maple syrup must have a characteristic maple flavour.
- 3. Maple sugar must not contain more than 10% moisture.

Rationale: Labels provide buyers and consumers with basic product information as well as health, safety, and nutritional information. All information included on labels must not be false, misleading, or deceptive.

Interpretation: Below is a list of mandatory information for consumer prepackaged products (e.g., individual unit (jug) or case of product sold to a consumer). Keep in mind all mandatory information must be in both official languages on consumer prepackaged with the exception of the name and principal place of business. There are also requirements on the legibility, location, and type size fonts. It is recommended to have an expert review your label to ensure compliance.

- Common Name (must meet the standard outlined in Volume 6 of the Canadian Standards of Identity for Maple Syrup)
- Name and principal place of business of the person by or for whom the food was manufactured, prepared, produced, stored, packaged, or labelled
- Lot Code
- Declaration of Net Quantity in Metric Units
- Grade
- Colour Class (Canada Grade A or imported Grade A Maple Syrup Only)
- Country of Origin (Imported Only)
- Nutrition Labelling

Refer to CFIA's **Labelling Requirements for Maple Products** webpage for more information.

Keep in mind that retailers may require something specific regarding labelling to supply their stores (UPC codes, specific labelling format, etc.). Please check with those retailers prior to producing product and creating labels for them.

CHECKLIST: the following can be compiled into one Quality SOP
☐ Labelling Procedure should detail the following:
☐ Product coding
☐ Product labelling
☐ Grading Procedure, if applicable, should detail the following and how the requirements are met:
☐ Grading requirements
☐ Grading kit (that is not expired)
☐ Weights Procedures, if applicable, should detail the following and how they are met:
☐ Weight requirements
☐ Employee training
□ Labelling Checklist
□ Packing Records



GRADES AND GRADE NAMES — PART 12 OF THE SFCR

DIVISION 3 - GRADING

MANDATORY GRADING

Requirements: Any eggs, fish, fresh fruits or vegetables, processed fruit or vegetable products, honey, maple syrup or beef carcass in respect of which grades are prescribed by these Regulations that are sent or conveyed from one province to another or that are imported or exported must

- (a) be graded;
- (b) meet the requirements that are set out in the Compendium or the Grades Document in respect of the applicable grade of that food; and
- (c) be labelled, in accordance with the Compendium or the Grades Document, with the applicable grade name that is set out in the Compendium or the Grades Document. 306(1)

AUTHORIZED APPLICATION OR USE

Requirements: Subject to subsection (2), a licence holder is authorized to apply a grade name to, and use a grade name in connection with, a food that is identified in their licence if

- (a) the food meets the requirements of paragraphs 8(1) (a) to (d);
- (b) the food meets the requirements that are set out in the Compendium or the Grades Document in respect of the applicable grade of that food;
- (c) the food complies with any standards that are set out in the Standards of Identity Document;
- (d) in the case of a dairy product, an egg, fish, a processed fruit or vegetable product, honey or maple syrup, the food has been graded by a licence holder;
- (e) in the case of a livestock carcass or poultry carcass that is dressed or partially dressed, the food has been graded by a grader; and
- (f) the food is packaged and labelled in accordance with these Regulations. 308(1)

DIVISION 4 - PACKAGING AND LABELLING

SUBDIVISION G: MAPLE SYRUP

COLOUR CLASS

Requirements: Maple syrup that is graded Canada Grade A and is sent or conveyed from one province to another or exported, or that is graded Grade A and is imported, must be labelled with the applicable colour class that is set out in the Compendium. The colour class must be shown on the container in characters of at least the minimum character height that is set out in column 2 of Schedule 6 for the area of a principal display surface that is set out in column 1. 325

Canadian Grade Compendium: Volume 7 – Maple Syrup (from CFIA's documents incorporated by reference)

General Requirement

1. The grades and grade names for maple syrup that is not imported, and the applicable colour classes, may be determined under Maple Syrup Grade Requirements.

Grades and Grade Names

2. There are two grades of maple syrup that is not imported "Canada Grade A" and "Canada Processing Grade" (see **Volume 9, Import Grade Requirements** for grade names used for imported maple syrup).

Canada Grade A

- Maple syrup graded "Canada Grade A" must be maple syrup that
 - (a) has a minimum soluble solids content of 66% and a maximum soluble solids content of 68.9% as determined by a refractometer or hydrometer at 20°C;
 - (b) is free from fermentation;
 - (c) is uniform in colour and free from sediment and any cloudiness or turbidity;
 - (d) has a colour class, as determined in accordance with section 5; and
 - (e) has a maple flavour characteristic of its colour class and is free from any objectionable odour or taste.

Canada Processing Grade

- Maple syrup graded "Canada Processing Grade" must be maple syrup that
 - (a) has a minimum soluble solids content of 66% and a maximum soluble solids content of 68.9% as determined by a refractometer or hydrometer at 20°C; and
 - (b) does not meet the requirements of paragraphs 3(c) to (e).

Colour Classes of Canada Grade A Maple Syrup

- 5. (1) Canada Grade A maple syrup has the colour class set out in Column 1 of an item of the Table of Colour Classes of Canada Grade A Maple Syrup if it has the ability to transmit light, expressed as a percentage that is set out in Columns 2 and 3 of that item.
 - (2) The ability of Canada Grade A maple syrup to transmit light must be determined by means of
 - (a) a spectrophotometer using matched square optical cells having a 10 mm light path at a wavelength of 560 nm, the colour values being expressed in per cent of light transmission as compared to A.R. Glycerol fixed at 100% transmission; or
 - (b) a visual glass comparator, the optical specifications of which correspond to the method described in paragraph (a).

Rationale: Grades establish product quality requirements. Grading for maple is mandatory if the product is sent to another province or exported. Canada Grade A maple syrup sent to another province or exported or imported Grade A maple syrup must be labelled with the colour class. In order to use a grade name, the product must meet health and safety requirements, the grade requirements, the standard of identity, be packed and labelled in accordance with regulatory requirements, and graded by a licence holder.

Interpretation: Grades and grade names for maple syrup are set out in **Volume 7 of the Canadian Grade Compendium** which is incorporated by reference in the SFCR. In order to use a grade name, the product must meet the grade requirements above.

TABLE OF COLOUR CLASSES OF CANADA GRADE A MAPLE SYRUP							
ltem	Column 1 Colour Class	Column 2 Percentage of Light Transmission Less than	Column 3 Percentage of Light Transmission Not less than				
1	Golden, Delicate Taste (Doré, goût délicat)		75.0				
2	Amber, Rich Taste (Ambré, goût riche)	75.0	50.0				
3	Dark, Robust Taste (Foncé, goût robuste)	50.0	25.0				
4	Very Dark, Strong Taste (Très Foncé, goût prononcé)	25.0					

APPENDIX

HOW TO DRAFT AN SOP/POLICY/PROCEDURE

The purpose of documenting a standard operating procedure (SOP) or policy regardless of the size of the operation is for consistency, transparency, and to ensure everyone is clear on what is expected to occur in the run of a day.

Every document must have the company's name, the document title, issue/revision date, date of the previous version, who approved or documented the policy, and page numbers. Best practice for page number formatting is page 1 of X, so everyone knows how many pages are included.

The key is to include important information while keeping it short and simple to follow. It should be written so that a new employee can understand it.

It's important to include the reason why the procedure is in place (purpose), who is responsible, if there is a backup person (trained designate), and how often the task is to be completed. A simple step by step instruction is best to describe the procedure. You will need to include a procedure for when things do not go as planned (deviation), as well as who is going to check to make sure the task is done correctly (verification), and any corresponding paperwork to be filled out.

		Issue/Revision Date:
Company's Name or Logo	Title of SOP/Procedure	Supersedes Date:
		Approved By:

Purpose: What is to be accomplished?

Responsibility: Who is responsible – designated staff? Who is the alternate?

Frequency: How often is it to be done?

Procedure: What is to be done? How is it to be done? Detailed instructions

Deviation/Corrective Action: Action(s) to be taken if out of specification

Verification: Who is responsible for the double-check? How is it to be done?

Records: List associated records

WHAT TO EXPECT DURING YOUR CFIA INSPECTION

If you intend to ship outside of the province, you will need to apply for a license. When you apply for a license, CFIA will likely contact you to start the inspection process. The process involves three main steps; pre-inspection, during inspection, and post-inspection. The following steps are a general guideline of what to expect when you are inspected.

Pre-Inspection

- 1. You receive a CFIA Inspection Notice
- Inspector will meet with you and review the scope of the inspection. If you have any changes or updates to your company information or profile on My CFIA and/or your preventive control plan, this is the time to inform the inspector.
- The inspector will do an initial walkthrough of your operation and identify areas that do not meet requirements. It's in your best interest to always accompany an inspector and take notes during this process.
- **4.** The scope of the inspection is confirmed during the walkthrough.

During Inspection

- 5. The Preventive Control Plan is verified that it is complete and accurate. This is completed through onsite observations, employee interviews, reviewing the written program (policies, procedures, and records) and depending on your product grading and measuring.
- The inspector will make notes either manually or electronically.
- The inspector will determine if the operation meets regulatory requirements and completes the report.

Post-Inspection

- 8. The inspector will meet with you to review the results and give you the report.
- 9. Your operation completes the corrective actions for any non-conformances given for non-compliant aspects at your facility or written program. Have your corrective actions documented with supporting information ready for your next inspection or if your inspector requests them.
- 10. A follow-up inspection is scheduled by the inspector.



HOW TO RESPOND TO A NON-COMPLIANCE

For each non-compliance, you will need to draft a Corrective Action or Deviation Report. You will need to detail the area of non-compliance, any immediate corrective actions taken, document the root cause analysis, permanent (preventive) corrective action taken, and then follow up with an effectiveness check to ensure that the issue has been fully addressed and corrected. For tips on how to do a root cause analysis, please see the Perennia Quality & Food Safety Resource section of our website.

HOW TO INCLUDE CROSS-CONTAMINATION POINTS ON THE HAZARD ANALYSIS AND CCP DETERMINATION FORM

The purpose of documenting cross-contamination points on the hazard analysis and CCP determination form is to identify and assess all potential cross-contamination points. Start by listing all potential cross-contamination points, type of hazard, and how it will be controlled. Once these have been identified and assessed, you must determine how each one will be controlled to mitigate the risk of contamination. If the cross-contamination point can be fully controlled, the title of the preventive control(s) or standard operating procedure(s) should be listed.



CROSS CONTAMINATION POINT(S)	HAZARD	В	С	Р	DETERMINE IF FULLY CONTROLLED BY PREVENTIVE CONTROLS/ PREREQUISITE	Q1	Q2	Ø3	Q4	Q5	CCP OR PC
Employee Flow	1B – Risk of cross- contamination from employees not washing their hands when leaving washroom	х			Yes – Employee Training Program and Personnel GMP and Hygiene Policy						
Employee Flow	2B – Risk of cross- contamination from employees not washing their hands upon entering production/packing areas	x			Yes - Employee Training Program and Personnel GMP and Hygiene Policy						
Employee Flow	3B – Risk of cross- contamination of product and/or employees due to improper movement of waste & compost during processing/ packing	х			Yes - Employee Training Program and Waste Management Program						
Chemical Flow	1C – Risk of cross- contamination of packaging and people with cleaning chemicals		х		Yes - Employee Training and Non-Food Agents Policy						

- **Q1**. Could a control measure(s) be used by the establishment at any process step? If no, indicate how the hazard will be controlled before and after the process on form 9 (which outlines the hazards not controlled by the operator). Then proceed to the next identified hazard. If yes, describe the control measure and proceed to Q2.
- **Q2**. Is it likely that contamination with the identified hazard could occur in excess of the acceptable level or could increase to an unacceptable level? If no (not a CCP), identify reason(s) why it is not likely to occur and proceed to the next identified hazard. If yes, identify the acceptable level of the hazard in the finished product, wherever possible, then proceed to Q3.
- Q3. Is this process step specifically designed to prevent, eliminate, or reduce the likely occurrence of the identified hazard to an acceptable level? If yes, (CCP) enter the CCP number in the last column. If no, proceed to Q4.
- **Q4**. Will a subsequent step eliminate the identified hazard or reduce its likely occurrence to an acceptable level? If no, (CCP) enter the CCP number in the last column then proceed to the next identified hazard. If yes, (not a CCP) identify the subsequent controlling step and proceed to Q5.
- **Q5.** Does this step provide partial control of the identified hazard? If yes, (PC) enter the PC number in the last column and proceed to the next identified hazard. If no, proceed to the next identified hazard.

DOWNLOADABLE FORMS

Throughout this guide we have highlighted the names of each record. To download the records that are referred to in this guide please go to:

www.perennia.ca/foodsafetyresources/

PLEASE NOTE: Links will not download if using Internet Explorer. Please ensure your browser is up to date if you encounter issues downloading the files.

The following sample records can be used as is or used as a reference in creating your own documents. You may need to modify these records to fit your operation and some records may not apply. For example, the Shipping Record may not need to be used if you have an invoicing system that you can modify to include aspects that are required to meet the SFCR such as truck/trailer inspection, and/or lot codes shipped.

Letter of Commitment

Maintenance and Reassessment Record

Exterior and Interior Inspection Checklist (Annual Inspection)

Receiving Record

Sanitation Record

Preventative Maintenance and Calibration Record

Pest Control Record

Shipping Record

Packing Record

Farm Gate Sales

Visitor's Record

Daily GMP Checks

Personnel Training Record

Corrective Action Report

Customer/Consumer Complaint Record

Recall Management Record

Mock Recall Record

Labelling Checklist

Change Log

RESOURCES

Our Quality and Food Safety Team have created a variety of resources which can be found on our website at www.perennia.ca. We offer coaching, assessments and online or public training courses covering a variety of topics. We also have a monthly newsletter which you can sign up for **here**. Most recently, we have added a resources page, which includes Perennia's publications and fact sheets, recommended resources, and videos. If you have any questions, please do not hesitate to contact one of our Quality and Food Safety Team members. We are here to help.

Perennia's Quality and Food Safety Team

Elaine Grant, Pam Laffin, Shelly MacDonald, Clarissa McIsaac, Cheryl Andrews

Elain C Grant Par Laffin Som Clarissa M'Usaac Rhenyl Andrews



OTES	

NOTES	



OFFICE LOCATIONS

28 Aberdeen Street, Kentville, Nova Scotia B4N 2N1

Phone: 902.678.7722 **Fax**: 902.678.7266 Email: info@perennia.ca

90 Research Drive Bible Hill, Nova Scotia B6L 2H5

PERENNIA FOOD AND BEVERAGE **INNOVATION CENTRE**

173 Dr Bernie MacDonald Drive, Bible Hill, Nova Scotia B6L 2H5

Phone: **9**02.896.8782 Fax: 902.896.8781

Email: innovation@perennia.ca













