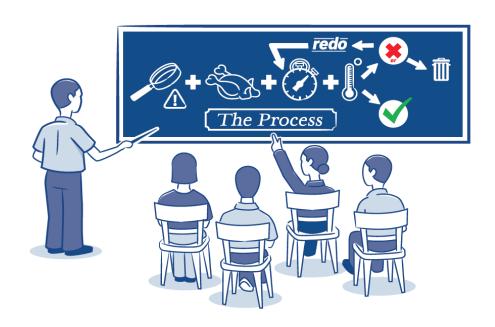
Sample Food Safety Plan MEETS BC REGULATORY REQUIREMENTS

DUCK POT PIE





Product Description

Pr	oduct Description	
1.	What is your product name and weight/volume?	Duck pot pie (500 g)
2.	What type of product is it (e.g., raw, ready-to-eat, ready-to-cook, or ready for further processing, etc.)?	Baked Ready to eat
3.	What are your product's important food safety characteristics (e.g., acidity, A _w , salinity, etc.)?	None
4.	What allergens does your product contain?	Egg, milk, sulphite and wheat.
5.	What restricted ingredients (preservatives, additives, etc.) does your product contain, and in what amounts (e.g., grams)?	None
6.	What are your food processing steps (e.g., cooking, cooling, pasteurization, etc.)?	Receiving incoming materials, ambient storage, cool refrigerator storage, freezer storage, packaging materials in a separate location, weighing, sauce cooking, sauce cooling, mixing, hopper, sheeting, pressing, filling, sealing, docking, spraying, baking, cooling, depanning, transfer on packaging board, metal detecting, aluminium tray packaging and labeling, case packaging and labeling, palletizing, freezer storage, shipping.
7.	How do you package your product (e.g., vacuum, modified atmosphere, etc.) and what packaging materials do you use?	Individual pies are packaged in aluminium trays. Packaged pie trays are packed in cardboard boxes.
8.	How do you store your product (e.g., keep refrigerated, keep frozen, keep dry) in your establishment and when you ship your product?	Keep frozen. Frozen pies are shipped in a clean, temperature-controlled truck (less than or equal to -18°C)
9.	What is the shelf-life of your product under proper storage conditions?	Frozen pie shelf life is 3 months at freezer temperatures (less than or equal to -18°C). 4 days shelf life after thawing at refrigerated temperatures (less than or equal to 4°C)
10.	How is the best before date to be noted on your product? (When product shelf life is more than 3 month, lot code or manufacturing date is to be printed on product label.)	The best before date is printed on the cardboard box as YY MM DD. Example: 15 JA 04 (January 04, 2015)

Product Description					
11. Who will consume your product (e.g., the	Ready to eat for the general population.				
general public, the elderly, the immunocompromised, infants)?	Note: Duck pot pie is not suitable for people with egg, milk, sulphite or wheat allergies or gluten intolerance.				
	Frozen product must be thawed before eating.				
	Preparation instructions, such as for thawing, are provided on the label.				
12. How might the consumer mishandle your product, and what safety measures will prevent this?	1. Products not stored at correct temperatures can cause illness and can have quality defects – storage and handling instructions are on the label.				
	2. Products that have passed the best before date can cause illness and can have quality defects – the best before date is printed on the cardboard box.				
	3. Refreezing can cause quality defects – storage and handling instructions are on the label.				
13. Where will the product be sold?	Food service, retail, wholesale and distributor.				
14. What information is on your product label?	Individual cardboard box label contains information such as product name, weight, ingredients listing including allergens, nutritional table, storage and handling instructions, best before date, preparation instructions, manufacturing company name, address and contact information.				
	Corrugated box label contains information such as product name, best before date, quantity, storage and handling instructions, preparation instructions, manufacturing company name, address and contact information.				

Incoming Materials

Ingredients	
All purpose flour	Pasteurized cream
Cake flour	Salt
Pastry flour	Corn starch
Cinnamon powder	Vegetable oil
Nutmeg powder	Diced raw duck pieces
Parsley flakes	Diced vegetables (carrots, celery, mushrooms)
Black pepper	Frozen peas
Liquid pasteurized eggs	Minced garlic
Butter	Water
Food contact processing aid materials	
Baking spray	
Food contact packaging materials	
Cardboard boards	Aluminum trays and lids
Non-food contact packaging materials	
Pre-printed cardboard boxes	Plain labels
Corrugated boxes	Shrink wrap
Ink	Wooden pallets
Tape	
Chemicals (hand washing, sanitation and mainten	ance)
Hand soap	Sanitizer
Hand sanitizer	Lubricant
Degreaser	

Food Safety Plan Table: Meets BC Regulatory Requirements

1. Identifying Hazards	2. Identifying	3 Establishing Critical Limits	4	Establishing Monitoring Procedures	5	Establishing Corrective Actions		6 Establishing Verification	7 Keeping
(Regulatory Requirement*)	Critical Control	(Regulatory Requirement*)		(Regulatory Requirement*)		(Regulatory Requirement*)		Procedures	Records
	Points (Regulatory							(Pending Regulatory Requirement)	(Pending
	Requirement*)								Regulatory
									Requirement)
Biological hazard:	CCP # 1	The internal temperature of the	1.	Measure the product's internal	W	hen critical limits are not being	1.	At the end of each production	Daily Baking
Pathogen survival due to improper	Baking	product must be at least 74°C		temperature from different areas of the		et for one or more product		day, review the "Daily Baking	Record
temperature distribution and time /		(165°F) for at least 15 seconds.		oven rack (top, middle, and bottom)		mples		Record" to ensure that it has	
temperature applications (e.g.				during each baking session.	1.	The product must be baked for a		been properly completed.	
Salmonella spp., Clostridium			2.	Insert the thermometer into the centre		longer period of time until the	2.	Once per week, ensure that the	
botulinum, Clostridium perfringens,				of the product and wait until the		product's internal temperature		temperature check follows the	
Campylobacter jejuni, Escherichia				thermometer reading is steady.		reaches at least 74°C for at least		written monitoring procedure.	
coli, Escherichia coli 0157:H7,			3.	Record the each result on the "Daily		15 seconds, or the product must	3.	If non-conformance is found	
Yersinia spp., Listeria				Baking Record" including the date, the		be destroyed.		during the verification	
monocytogenes, Vibrio vulnificus,				time, and the operator's initials.	2.	Immediately investigate the		procedure, immediately	
Staphylococcus aureus and						cause of the non-conformance		investigate the cause of the	
enterotoxin)						and take necessary corrective		non-conformance and take	
						actions to prevent reoccurrence.		necessary corrective actions to	
					3.	Record all non-conformances and		prevent reoccurrence.	
						corrective actions taken on the	4.	Record all observations on the	
						"Daily Baking Record," including		"Daily Baking Record," including	
						the date, the time, and their		the date, the time, and the	
						initials.		technician's initials. (e.g.,	
								temperature readings, non-	
								conformances, and corrective	
								actions taken).	

DUCK POT PIE FOOD SAFETY PLAN

1. Identifying Hazards	2. Identifying	3 Establishing Critical Limits	4	Establishing Monitoring Procedures	5	Establishing Corrective Actions		6 Establishing Verification	7 Keeping
(Regulatory Requirement*)	Critical Control Points (Regulatory Requirement*)	(Regulatory Requirement*)		(Regulatory Requirement*)		(Regulatory Requirement*)		Procedures (Pending Regulatory Requirement)	Records (Pending Regulatory Requirement)
Biological hazard:	CCP # 2	During cooling, the product's	1.	Measure the product's internal	W	hen critical limits are not being	1.	Review the "Daily Cooling	Daily Cooling
Pathogen contamination due to	Cooling	internal temperature must not		temperature every hour during cooling.	m	et for one or two or all samples		Record" to ensure that it has	Record
inadequate cooling (e.g., Clostridium		remain between 60°C (140°F)	2.	Calibrate the thermometer to ensure it	1.	Immediately place all products		been properly completed.	
perfringens, Listeria monocytogenes)		and 20°C (70°F) for more than 2		is working correctly before measuring		that do not meet the critical limit	2.	Once per week, ensure that the	
		hours. The product's internal		the internal temperature of the		on hold.		temperature check follows the	
		temperature must not remain		product.	2.	Products put on hold must be re-		written monitoring procedure.	
		between 60°C (140°F) and 4°C	3.	Measure the product's internal		baked and re -cooled to meet the	3.	If non-conformance is found	
		(40°F) for more than 4 hours.		temperature from different trays of the		critical limit or if the critical limit		during the verification	
				trolley (top, middle, and bottom) at		cannot be met, product must be		procedure, investigate the	
				each check.		destroyed.		cause of the non-conformance	
			4.	Insert the thermometer into the centre	3.	Investigate the cause of the non-		and take necessary corrective	
				of the product and wait until the		conformance and take necessary		actions to prevent	
				thermometer reading is steady.		corrective actions to prevent		reoccurrence.	
			5.	Record the results from the three		reoccurrence.	4.	Record all observations (e.g.,	
				readings from different trays on the	4.	Record all non-conformances and		temperature readings, non-	
				"Daily Cooling Record," including the		corrective actions taken on the		conformances, and corrective	
				date, the time, and initials.		"Daily Cooling Record," including		actions) on the "Daily Cooling	
						the date, the time, and initials.		Record," including the date, the	
								time, and initials.	

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(Regulatory Requirement*)	Critical Control	(Regulatory Requirement*)		(Regulatory Requirement*)		(Regulatory Requirement*)		Procedures	Records
	Points (Regulatory							(Pending Regulatory Requirement)	(Pending
	Requirement*)								Regulatory
									Requirement)
Physical hazard:	CCP #3	Metal detector must detect 3.0	1.	Test the metal detector at the start,	A	. When the metal detector fails to	1.	At the end of each production	Daily Metal
Presence of hazardous extraneous	Metal detecting	mm ferrous, 3.0 mm non-		every hour during packaging, and at the	de	etect a metal test sample		day, review the "Daily Metal	Detector Check
metallic material in the finished		ferrous, and 3.5 mm stainless		end of each packaging run.	1.	Immediately stop the line and		Detector Check Record" to	Record
product due to the failure of the		steel test samples when the	2.	Test the metal detector by passing a		place all products processed since		ensure that it has been properly	
metal detector to detect metal and		test samples are passed		sample piece of metal through the		the last successful check on hold.		completed.	
reject the product when metal is		through the detector with the		detector to ensure that it is operating	2.	All products processed while the	2.	Once per week, ensure that the	
detected.		product. The metal detector		effectively and able to detect metal		metal detector was not functional		monitoring of the metal	
		must reject the product.		present in the product.		must be held until they can be		detector follows the written	
			3.	Check metal samples of 3.0 mm ferrous,		passed through a functional		monitoring procedure.	
				3.0 mm non-ferrous, and 3.5 mm		metal detector.	3.	If non-conformance is found	
				stainless steel, one at a time. Each	В.	When a product is rejected by the		during the verification	
				check must include all three sample	m	etal detector		procedure, investigate the	
				tests.	1.	Inspect the product for the metal		cause of the non-conformance	
			4.	Insert the metal sample into the middle		piece.		and take necessary corrective	
				of the product and then pass the				actions to prevent	
				product package through the metal	Fo	or above listed non-conformances (A		reoccurrence.	
				detector. A properly operating metal	&	B) investigate the cause of the non-	4.	Record all observations (e.g.,	
				detector must detect the metal sample	cc	onformance and take necessary		whether or not the detector is	
				in the product.	cc	prrective actions to prevent		operating effectively, non-	
			5.	Each time a metal contaminant is	re	eoccurrence.		conformances, and corrective	
				detected, the metal detector belt must				actions taken) on the "Daily	
				retract and the rejected product must	Re	ecord all non-conformances and		Metal Detector Check Record,"	

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	Points (Regulatory				(Pending Regulatory Requirement)	(Pending
	Requirement*)					Regulatory
						Requirement)
			drop into the rejection box.	corrective actions taken on the "Daily	including the date, the time,	
			6. Record the metal sample check as	Metal Detector Check Record,"	and initials.	
			acceptable (" \checkmark ") (i.e., the metal	including the date, the time, and		
			detector is operating correctly) or not	initials.		
			acceptable ("X") (i.e., the metal			
			detector is not operating correctly) on			
			the "Daily Metal Detector Check			
			Record," including the date, the time,			
			and initials.			

Daily Baking Record

Critical Control Point # 1 (Biological)

<u>Critical Limits:</u> The internal temperature of the product must be at least 74°C (165°F) for at least 15 seconds.

Date	Time	Batch	Product Name	Product's Internal Temperature			Initials
		Number			(Product selected from top, middle, and bottom racks of		
				mudic	oven)	ii racks or	
				Тор	Middle	Bottom	
2015/11/02	12:00	1	Duck pot pie	77°C	77°C	76°C	СС
2015/11/02	13:04	2	Duck pot pie	76°C	72°C	78°C	СС
2015/11/02	16:00	3	Duck pot pie	77°C	79°C	75°C	CC
December 2016		d					
Record non-confo		<u>ia correctiv</u>	<u>'e actions nere:</u>				
2015/11/02: Batcl	h 2:						
The internal temp	erature of	the pie on	the middle rack did	not reach	74°C. Pies w	ere baked aga	ain until
the internal tempo	erature rea	ached 74°C	. CC				
Daily verification:			MN Date: 2015/11		/02		
Weekly verificatio	n:		ML Date: 2015/11/			/09	

Daily Cooling Record

Critical Control Point # 2 (Biological)

<u>Critical Limits:</u> During cooling, the product's internal temperature must not remain between 60°C (140°F) and 20°C (70°F) for more than 2 hours. The product's internal temperature must not remain between 60°C (140°F) and 4°C (40°F) for more than 4 hours.

Production Date: 2015/11/02 Batch Number 1

Time	Top Tray	Middle Tray	Bottom Tray Temperature	Initials			
	Temperature	Temperature	remperature				
12:30	66°C	66°C	64°C	CC			
13:04	36°C	38°C	36°C	CC			
14:05	19°C	18°C	16°C	CC			
15:04	9°C	8°C	6°C	CC			
16:06	1°C	2°C	1°C	CC			
Record non-conformar	nce and corrective action	ons here:					
Dell effective and			D. L. 2045/44/02				
Daily verification: MN Date: 2015/11/02							
Weekly verification: M	L		Date: 2015/11/09				

Daily Metal Detector Check Record

Critical Control Point #3 (Physical)

<u>Critical Limits:</u> Metal detector must detect 3.0 mm ferrous, 3.0 mm non-ferrous, and 3.5 mm stainless steel test samples when the test samples are passed through the detector with the product. The metal detector must reject the product.

Record the metal sample check as acceptable (" \checkmark ") (i.e., the metal detector is operating correctly) or not acceptable ("X") (i.e., the metal detector is not operating correctly)

Date	Time	Batch	Product	3.0 mm	3.0 mm	3.5 mm	Initials
		Number	Name	Ferrous	Non-	Stainless Steel	
					ferrous		
2015/11/02	12:00	1	Duck pot pie	,	,	,	SM
	(start)			√	√	√	
	13:05	1	Duck pot pie	✓	✓	√	SM
	14:07	1	Duck pot pie	Х	✓	✓	SM
	15:37	1	Duck pot pie	✓	✓	✓	SM
	16:04	1	Duck pot pie	✓	✓	✓	SM
	17:05	1	Duck pot pie	✓	✓	✓	SM
	17:44	1	Duck pot pie	,	,	,	SM
	(finish)			√	√	√	

Record non-conformance and corrective actions here:

At 14:07, a 3.0 mm ferrous test sample was not detected by the metal detector. The line was stopped. Products were placed on hold since last successful check at 13:05. At 15:30, the metal detector was repaired and calibrated. SM

Daily verification:	MN	Date: 2015/11/02
Weekly verification:	ML	Date: 2015/11/09

