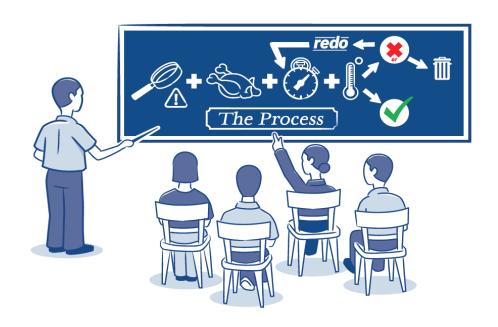
Sample Food Safety Plan MEETS BC REGULATORY REQUIREMENTS

READY TO EAT SAUSAGE





Product Description

Product Description	
1. What is your product name and	Ready to eat sausage (beef)
weight/volume?	Ready to eat sausage (pork)
	(450 g, 10 pieces)
2. What type of product is it (e.g., raw, ready-	Cooked
to-eat, ready-to-cook, or ready for further processing, etc.)?	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?	Wheat
5. What restricted ingredients (preservatives,	Preservative - sodium nitrite
additives, etc.) does your product contain, and in what amounts (e.g., grams)?	(Nitrite in the finished product is between 180 ppm - 200 ppm)
6. What are your food processing steps (e.g., cooking, cooling, pasteurization, etc.)?	Receiving incoming materials, ambient storage, cool refrigerator storage, packaging material storage in a separate location, nitrite weighing, weighing, grinding, mixing, transfer to trolley, refrigerated storage, stuffing, hanging on smoke house rack, hot smoking, cooling, cutting, vacuum packaging, date coding, metal detecting, case packaging and labeling, palletizing, refrigerated storage, shipping.
7. How do you package your product (e.g., vacuum, modified atmosphere, etc.) and what packaging materials do you use?	Vacuum packaging Sausages are vacuum packaged in plastic film. Packaged sausages are packed in corrugated 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 refrigerated. Fresh sausage packages are shipped in a clean, temperature-controlled truck (less than or equal to 4°C)
9. What is the shelf-life of your product under proper storage conditions?	Sausage shelf life is 45 days at refrigerated temperatures (less than or equal to 4°C)

Product Description	
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 product package as YY MM DD. Example: 15 JA 04 (January 04, 2015)
11.Who will consume your product (e.g., the general public, the elderly, the immunocompromised, infants)?	Ready to eat product for the general population. Note: Ready to eat sausage is not suitable for people with wheat allergies or gluten intolerance.
12.How might the consumer mishandle your product, and what safety measures will prevent this?	 Products not stored at correct temperatures can cause illness and can have quality defects – storage and handling instructions are on the label. Products that have passed the best before date can cause illness and can have quality defects – the best before date is printed on the plastic film.
13.Where will the product be sold?	Food service, retail, wholesale and distributor.
14.What information is on your product label?	Individual product package label contains information such as product name, weight, ingredients listing including allergen, 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	
Beef trims	Salt
Pork trims	Black pepper
Natural hog casing	Paprika
Wheat flour	Prague powder
Corn starch	Sodium acetate
Dextrose	Water
Food contact processing aid materials	
Steam	
Food contact packaging materials	
Pre-printed plastic films	
Non-food contact packaging materials	
Corrugated boxes	Tape
Plain labels	Shrink wrap
Ink	Wooden pallets
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 (Regulatory Requirement*)	2. Identifying Critical Control Points (Regulatory Requirement*)	3 Establishing Critical Limits (Regulatory Requirement*)	4 Establishing Monitoring Procedures (Regulatory Requirement*)	5 Establishing Corrective Actions (Regulatory Requirement*)	6 Establishing Verification Procedures (Pending Regulatory Requirement)	7 Keeping Records (Pending Regulatory Requirement)
Biological hazard: Pathogen growth or survival due to inadequate addition of preservative (e.g., Clostridium botulinum, Clostridium perfringens, Listeria monocytogenes) Chemical hazard: More than 200 ppm of nitrite in the finished product due to incorrect weighing	CCP # 1 Nitrite Weighing	1) Nitrite content in the finished product must be between 180 ppm and 200 ppm 2) Nitrite content in the finished product must never be greater than 200 ppm	 Calculate nitrite ppm at each weighing. Calibrate the scale to ensure it is working correctly before weighing the product. Check the correct percentage of sodium nitrite from the curing salt bag label or product specification sheet. Calculate nitrite ppm in finished product using the formula below: Sausage Mix (in kg) = SM Preservative (in g) = P Sodium Nitrite in Preservative (as a %) = SNP Total Emulsion (in kg) = SM + (P x 10⁻³) Total Amount of Sodium Nitrite in Curing Salt (in g) = P x (SNP ÷ 100) Nitrite in Finished Product (in ppm) = (Total Amount of Sodium Nitrite in Curing Salt x 10⁻³ x 10⁶) ÷ Total Emulsion Record the results of the calculation on the "Daily Nitrite Weighing Record," including the date, the time, and initials. 	when the critical limit is not being met 1) Immediately must place all products on hold for this batch. 2) The products must be reprocessed to meet critical limits. If critical limits cannot be met, the products must be destroyed. 3) Investigate the cause of the nonconformance and take necessary corrective actions to prevent reoccurrence. 4) Record all non-conformances and corrective actions taken on the "Daily Nitrite Weighing Record," including the date, the time, and initials.	1) At the end of each production day, review the "Daily Nitrite Weighing Record" to ensure that it has been properly completed. 2) Once per week, ensure that the monitoring of the nitrite weighing check follows the written monitoring procedure. 3) If non-conformance is found during the verification procedure investigate the cause of the non-conformance and take necessary corrective actions to prevent reoccurrence. 4) Record all observations (e.g., nitrite ppm, non-conformances, and corrective actions) on the "Daily Nitrite Weighing Record," including the date, the time, and initials.	Daily Nitrite Weighing Record

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 Requirement*)				(Pending Regulatory Requirement)	(Pending Regulatory
	,					Requirement)
Biological hazard:	CCP # 2	The internal temperature of the	Measure the product's internal	When critical limits are not being	1. At the end of each production	Daily Hot
Pathogen survival due to improper	Hot Smoking	product must be at least 71°C	temperature from different areas of the	met for one or more product	day, review the "Daily Hot	Smoking
temperature distribution and/or		(159.8°F) for at least 15	smoke house rack (top, middle, and	samples	Smoking Record" to ensure that	Record
improper time / temperature		seconds.	bottom) during each smoking session.	1. The product must be smoked for	it has been properly completed.	
applications (e.g., Salmonella spp.,			2. Insert the thermometer into the centre	a longer period of time until the	2. Once per week, ensure that the	
Clostridium botulinum, Clostridium			of the product and wait until the	product's internal temperature	temperature check follows the	
perfringens, Campylobacter jejuni,			thermometer reading is steady.	reaches at least 71°C (159.8°F) for	written monitoring procedure.	
Escherichia coli, Escherichia coli			3. Record the each result on the "Daily Hot	a minimum of 15 seconds. If the	3. If non-conformance is found	
0157:H7, Yersinia spp., Listeria			Smoking Record" including the date, the	limit cannot be met, the product	during the verification	
monocytogenes, Vibrio vulnificus,			time, and initials.	must be destroyed.	procedure, immediately	
Staphylococcus aureus and				1. Immediately investigate the	investigate the cause of the	
enterotoxin, Taenia spp., Cysticercus				cause of the non-conformance	non-conformance and take	
bovis)				and take necessary corrective	necessary corrective actions to	
				actions to prevent reoccurrence	prevent reoccurrence.	
				2. Record all non-conformances and	4. Record all observations on the	
				corrective actions taken on the	"Daily Hot Smoking Record,"	
				"Daily Hot Smoking Record,"	including the date, the time,	
				including the date, the time, and	and the initials.	
				initials.		

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 # 3	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.,		remain between 60°C (140°F)	2.	Calibrate the thermometer to ensure it	1.	Immediately place all products		been properly completed.	
Clostridium perfringens, Listeria		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	
monocytogenes)		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		cooked and re-cooled to meet	3.	If non-conformance is found	
		(40°F) for more than 4 hours.		temperature from different trays of the		the critical limit. If the critical		during the verification	
				trolley (top, middle, and bottom) at		limit is not being met, the		procedure, investigate the	
				each check.		product must be 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.	

1. Identifying Hazards	2. Identifying	3 Establishing Critical Limits	4	Establishing Monitoring Procedures	5	Establishing Corrective Actions		6 Establishing Verification	7 Keeping
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	Points (Regulatory							(Pending Regulatory Requirement)	(Pending
	Requirement*)								Regulatory
									Requirement)
Physical hazard:	CCP # 4	Metal detector must detect 2.5	1.	Test the metal detector at the start,	Α	. When the metal detector fails to	1.	At the end of each production	Daily Metal
Presence of hazardous extraneous	Metal detecting	mm ferrous, 2.5 mm non-		every hour during packaging, and at the	d	etect a metal test sample		day, review the "Daily Metal	Detector Check
metallic material in the finished		ferrous and 3.0 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 2.5 mm ferrous,		passed through a functional		monitoring procedure.	
				2.5 mm non-ferrous and 3.0 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	F	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	C	onformance and take necessary		whether or not the detector is	
				in the product.		orrective actions to prevent		operating effectively, non-	
			5.	·		eoccurrence.		conformances, and corrective	
				detected, the metal detector belt must				actions taken) on the "Daily	
				retract and the rejected product must	R	ecord all non-conformances and		Metal Detector Check Record,"	
								•	
				drop into the rejection box.	C	orrective actions taken on the "Daily		including the date, the time,	

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	Points (Regulatory				(Pending Regulatory Requirement)	(Pending
	Requirement*)					Regulatory
						Requirement)
			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 Nitrite Weighing Record

Critical Control Point # 1 (Biological, Chemical)

Critical Limits:

- 1) Nitrite content in the finished product must be between 180 ppm and 200 ppm
- 2) Nitrite content in the finished product must never be greater than 200 ppm

Calculate nitrite ppm in finished product using the formula below:

Sausage Mix (in kg) = SM

Preservative (in g) = P

Sodium Nitrite in Preservative (as a %) = SNP

Total Emulsion (in kg) = SM + (P x 10^{-3})

Total Amount of Sodium Nitrite in Curing Salt (in g) = $P \times (SNP \div 100)$

Nitrite in Finished Product (in ppm) = (Total Amount of Sodium Nitrite in Curing Salt $\times 10^{-3} \times 10^{6}$) ÷ Total Emulsion

Date	2015/11/02		
Time	9:15		
Batch Number	1		
Sausage Mix Weight (kg)	114		
Curing Salt Weight (g)	350	_	
Total emulsion (kg)	114.350		
Sodium Nitrite in Curing Salt (%)	6.25		
Total Amount of Sodium Nitrite in	21.875		
Curing Salt (g)			
Nitrite in Finished Product (ppm)	191.30		
Initials	CC		

Record non-conformance and corrective actions here:

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

Daily Hot Smoking Record

Critical Control Point # 2 (Biological)

<u>Critical Limits:</u> The internal temperature of the product must be at least 71°C (159.8°F) for a minimum of 15 seconds.

Date	Time	Batch	Product Name	Product'	's Internal	Temperature	Initials
		Number		(Produ	ıct selected	from top,	
				middle	, and botto	om racks of	
					smoke ho	use)	
				Тор	Middle	Bottom	
2015/11/02	12:00	1	Ready to eat sausage	77°C	77°C	76°C	CC
2015/11/02	13:04	2	Ready to eat sausage	68°C	74°C	79°C	CC
2015/11/02	16:00	3	Ready to eat sausage	77°C	79°C	75°C	CC
Record non-confo	rmance ar	nd correctiv	e actions here:				
2015/11/02: Batc	h 2:						
The internal temp	erature of	the sausag	e on the top rack d	id not reac	h 71°C. The	e sausage was p	olaced on
hold and smoked	again until	the interna	al temperature read	ched 71° C	. CC		
Daily verification:				MN	I	Date: 2015/11	/02
Weekly verification	on:			ML		Date: 2015/11	/09

Daily Cooling Record

Critical Control Point #3 (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 Temperature	Middle Tray Temperature	Bottom Tray Temperature	Initials	
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		
16:06	1°C	2°C	1°C	CC	
Record non-conforma	nce and corrective actio	ns here:			
Daily verification: MN			Date: 2015/11/02		
Weekly verification: N	1L		Date: 2015/11/09		

Daily Metal Detector Check Record

Critical Control Point # 4 (Physical)

<u>Critical Limits:</u> Metal detector must detect 2.5 mm ferrous, 2.5 mm non-ferrous, and 3.0 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 Name	2.5 mm	2.5 mm	3.0 mm	Initials
		Number		Ferrous	Non-	Stainless	
					ferrous	Steel	
2015/11/02	12:00	1	Ready to eat sausage				SM
	(start)			✓	✓	√	
	13:05	1	Ready to eat sausage	✓	✓	✓	SM
	14:07	1	Ready to eat sausage	✓	✓	✓	SM
	15:37	1	Ready to eat sausage	✓	✓	✓	SM
	16:04	1	Ready to eat sausage	✓	✓	✓	SM
	17:05	1	Ready to eat sausage	✓	✓	✓	SM
	17:44	1	Ready to eat sausage				SM
	(finish)			✓	✓	√	

Record non-conformance and corrective actions here:

At 16:20, one package was rejected. The product was screened for a metal piece. A small piece (5 mm in size) of metal was found. Upon investigation, it appears that it came from one of the damaged belts. The belt was immediately removed and replaced with a new belt. SM

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

