

CENTER FOR FOOD REGULATION AND RESEARCH EXTERNAL



1. ISSUANCE OF CERTIFICATE OF PRODUCT REGISTRATION (CPR)

1.1. ISSUANCE OF CERTIFICATE OF PRODUCT REGISTRATION (CPR) – AMENDMENT (INITIAL APPLICATION APPROVED FROM MODIFIED E-REGISTRATION (VERSION 2))

'Registration' means the process of approval of an application to register health products prior to engaging in the manufacture, importation, exportation, sale, offer for sale, distribution, transfer, and where applicable, the use, testing, promotion, advertisement, and/or sponsorship of health products. (Republic Act No. 9711)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	In accordance to Administrative Order 50 s. 2001 + Legal Research Fee (LRF).
		Change or Extension of Shelf-life: Php 1,000.00 + 1% LRF Other Types of Amendment: Php 200.00 + 1% LRF

GENERAL GUIDELINES

Please refer to:

A. General Guidelines, B. Specific Guidelines, and C. Procedural Guidelines, IV. GUIDELINES, pages 2-10 of <u>FDA Circular No. 2020-033</u> || Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products"; and

2) III. General Guidelines, and IV. Specific Guidelines of <u>FDA Circular No. 2020-033-A</u> || Addendum to FDA Circular No. 2020-033, "Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products" to include Guidelines for Preassessment and Reiteration of Pre-Assessment Procedures in Applying for Certificate of Product Registration for Food

CHECKLIST OF REQUIREMENTS FOR ALL TYPES OF FOOD PRODUCTS/FOOD CATEGORIZATION: RAW MATERIALS, LOW RISK, MEDIUM RISK AND HIGH RISK FOOD PRODUCTS

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GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE		
☑ Accomplished Application Form as prescribed by FDA regulations	Administrative No. Order 2014-	https://www.fda.gov.ph/		
e.g. E-Registration System	0029			



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☑ Proof of Payment of Fees as pre	scribed by current FDA regulations.	Administrative Order 50 s. 2001	Systems/Means prescribed by FDA
☑ Scanned Application Letter stating the intended changes (indicate ALL the changes/amendments to be made)		Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company
☑ VALID AND APPROPRIATE FDA LICENSE TO OPERATE (LTO) (REQUIRED FOR ALL TYPES OF CPR APPLICATION) *The product being applied must be listed in the FDA approved Product Line/Category.		Administrative No. Order 2014- 0029 Republic Act 9711	FDA Philippines
ADDITIONAL Requirements per An	nendment Type		
AMENDMENT TYPE	☑ ADDITIONAL REQUIREMENTS	BASIS	WHERE TO SECURE
2a. Change in Brand Name	 ☑ Clear and complete loose labels or artworks reflecting the change, as applicable, of all packaging sizes, or equivalents as defined by FDA regulations ☑ Authority from the source or the owner of the brand (imported & local) ☑ IPO registration, if available. 	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company Source/Supplier/Brand Owner IPO/Source/ Supplier
2b. Change in Product Name/Additional Product Description	 ☑ Clear and complete loose labels or artworks reflecting the change, as applicable, of all packaging sizes, or equivalents as defined by FDA regulations *Change in % Alcohol Content and Vintage in Wines as per FDA Circular No. 2020-033-B. 	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2c. Change in Company Name/Business Name	☑ Proof of change in business name (e.g. License to Operate)	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier



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	☑Clear and complete loose labels or artworks reflecting the change, as applicable, of all packaging sizes, or equivalents as defined by FDA regulations		
2d. Change in/Additional Supplier	☑ Any of the following scanned copy of the original documents: Foreign Agency Agreement or Certificate of Distributorship or Appointment letter or Proforma Invoice or Memorandum of Agreement from the new supplier.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2e. Change in Packaging Material and/or Additional Packaging Type	☐ Clear and complete proposed loose labels or artworks, as applicable, of all packaging sizes, or equivalents as defined by FDA regulations ☐ Pictures of the product in all angles and in different packaging sizes, and from at least two different perspectives allowing visual recognition of a product as the same with the one being registered. ☐ Proof of suitability of packaging material for food, including stability of the product in the new packaging.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2f. Change of Packaging in Commercial Presentation (Change/Additional Packaging Size)	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2g. Change or Extension in Shelf- Life	☑ Stability study results with conclusion to support extension or change in shelf-life	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier



2h. Change in/Additional Packaging design	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations *Change in % Alcohol Content and	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
	Vintage in Wines as per <u>FDA Circular</u> <u>No. 2020-033-B</u> .		
2hi. Addition of Claims for Logos	 ☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations. ☑ Valid Certificate (e.g. HALAL, Sangkap pinoy seal, Organic, Kosher, etc.) 	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hii. Change in Label Color	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hiii. Change in Font Size for Product Information	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hiv. Change/Additional Claims for Source of Vitamins/Minerals and Health and Nutrition Claims	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier



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	☑ Certificate of Analysis (duly signed by competent technical staff including the complete name with appropriate parameters and result) or documents to substantiate claims.		
2hv. Change /Update in Nutrition Information (Vitamin and Mineral)	 ☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations. ☑ Certificate of Analysis (duly signed by competent technical staff including the complete name with appropriate parameters and result). 	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hvi. Change/Additional Menu or Serving suggestion (Photograph)	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hvii. Compliance to CPR Remarks	☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations.	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier
2hviii. Declaration of Distributor	 ☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents, reflecting the change/s, as defined by FDA Regulations. ☑ Distributorship Agreement (Notarized, signed by the MAH/ 	Administrative No. Order 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Source/Supplier



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	Applicant Company and distributor		
	reflecting the correct address		
2hix. Change of Manufacturer's	☑ Clear and complete loose labels or	Administrative No. Order 2014-	Applicant Company/
Name	artworks, as applicable, of all packaging	0029	Source/Supplier
	sizes, or equivalents, reflecting the	FDA Circular No. 2020-033	
	change/s, as defined by FDA		
	Regulations.		
	☑ Attestation letter from the		
	manufacturer stating the reason for		
	change in manufacturer's name, and/or		
	ANY of the scanned copy of the original		
	document issued by the Regulatory/		
	Health Authority/Recognized Issuing		
	body/ Attested by recognized		
	Association or duly authenticated by the		
	Philippine Consulate from the country of		
	origin: Certificate of Registration with		
	GMP Compliance or its equivalent or		
	Valid Sanitary Phyto-Sanitary Certificate		
	or Health Certificate or ISO 22000		
	Certificate or FSSC Certificate or		
	HACCP Certificate or Certificate of Free		
01 1 11 5 1 1 11	Sale.	A 1	A 1: 10 /
2hx. Locally Produced with	☑ Clear and complete loose labels or	Administrative No. Order 2014-	Applicant Company/
Additional Activity for Export	artworks, as applicable, of all packaging	0029	Source/Supplier
	sizes, or equivalents, reflecting the	FDA Circular No. 2020-033	
	change/s, as defined by FDA		
	Regulations.		
	☑ LTO as food exporter if the company		
	is not manufacturer.		



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2hxi. Declaration of "Exclusively	☑ Clear and complete loose labels or	Administrative No. Order 2014-	Applicant Company/
Distributed by"	artworks, as applicable, of all packaging	0029	Source/Supplier
	sizes, or equivalents, reflecting the	FDA Circular No. 2020-033	
	change/s, as defined by FDA		
	Regulations.		
	☑ Terms of Agreement/Exclusive		
	Distributorship Agreement.		
2hxii. Declaration of	☑ Clear and complete loose labels or	Administrative No. Order 2014-	Applicant Company/
Manufacturer's Office Address on	artworks, as applicable, of all packaging	0029	Source/Supplier
the Label	sizes, or equivalents, reflecting the	FDA Circular No. 2020-033	
	change/s, as defined by FDA		
	Regulations.		
2i. Transfer of Ownership of a	☑ Proof of Agreement between	Administrative No. Order 2014-	Applicant Company/
Registered Product	previous and current owners of the	0029 FDA Girandan Na 2000 022	Source/Supplier
	product transferring ownership	FDA Circular No. 2020-033	
	☑ Clear and complete loose labels or		
	artworks, as applicable, of all packaging		
	sizes, or equivalents, reflecting the		
	change/s, as defined by FDA		
0: 01	Regulations	A 1 1	A 1: 10 /
2j. Change in	☑ Termination of agreement/Deed of	Administrative No. Order 2014-	Applicant Company/
Importer/Distributor/Trader	assignment	0029 FDA Circular No. 2020-033	Source/Supplier
	☑ Agreement of new	FDA Circulat No. 2020-033	
	manufacturer/importer/distributor or		
	Appointment letter		
	☑ Clear and complete loose labels or		
	artworks, as applicable, of all packaging		
	sizes, or equivalents, reflecting the		
	change/s, as defined by FDA		
	Regulations		



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2k. For Change in	☑ Termination of agreement/Deed of	Administrative No. Order 2014-	Applicant Company/
Importer/Distributor/Trader using a	assignment	0029	Source/Supplier
new user account:	☑ Agreement of new	FDA Circular No. 2020-033	
	manufacturer/importer/distributor or		
	Appointment letter		
	☑ Clear and complete loose labels or		
	artworks, as applicable, of all packaging		
	sizes, or equivalents, reflecting the		
	change/s, as defined by FDA		
	Regulations.		
	☑ Upload ALL INITIAL requirements		
2l. Change in Company	☑ Proof of change in business name	Administrative No. Order 2014-	Applicant Company/
Address/Business Address (Not	(e.g. License to Operate)	0029	Source/Supplier
Applicable to Manufacturer and	☑ Clear and complete loose labels or	FDA Circular No. 2020-033	
Repacker)	artworks reflecting the change, as		
	applicable, of all packaging sizes, or		
	equivalents as defined by FDA		
	regulations		
2m. Change in LTO Number and/or	☑ Copy of updated License to Operate	Administrative No. Order 2014-	Applicant Company/
LTO Validity		0029	Source/Supplier
		FDA Circular No. 2020-033	
2n. Exportation of Previously	☑ Clear and complete loose labels or	Administrative No. Order 2014-	Applicant Company/
Registered Product Initially for	artworks as applicable, of all packaging	0029	Source/Supplier
Local Distribution.	sizes, or equivalents as defined by FDA	FDA Circular No. 2020-033	
	regulations or reflecting compliance to		
	labelling requirements of importing		
	country (if label is different from the		
	approved one)		
	☑ Copy of License to Operate as Food		
	Exporter		



2o. Other Cases as Declared in	e.g. Change in Product Specification	Administrative No. Order 2014-	Applicant Company/
Succeeding FDA Issuances	☑ Copy of updated Product	0029	Source/Supplier
(Examples but not limited to the	Specification Sheet	FDA Circular No. 2020-033	
following; as long as there is no			
change in formulation and no	e.g. Change in Lot Code and		
change in manufacturer's address)	Interpretation		
	☑ Copy of updated Product		
	Specification Sheet		
	☑ Clear and complete loose labels or		
	artworks reflecting the change, as		
	applicable, of all packaging sizes, or		
	equivalents as defined by FDA		
	regulations		

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1.1. Files using the specific product/CASE NUMBER in the INBOX folder, and then accomplishes (including uploading of the COMPLETE documentary requirements) the E-Registration System through the E-Portal https://eportal.fda.gov.ph based on the desired type of application in accordance to current FDA regulation/s on the use of the E-Registration Portal/E-Services. 1.2. Forwards the application to PRE-ASSESSMENT.	Pre-assesses ONLY the completeness of the submitted documents through E-Registration System/E-Portal https://eportal.fda.gov.ph . Result of Pre-assessment will be received by the account holder.	Day 0	Center for Food Regulation and Research (CFRR) PRE-ASSESSOR (e.g. Food-Drug Regulation Officer (FDRO))



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A system generated E-mail notification from FDA will be received by the client upon submission of application for Pre-Assessment.			
2. (If COMPLETE) Receives the Order	2. If found COMPLETE ,	Day 0	CFRR PRE-
of Payment.	Generates Order of Payment through the email of the		ASSESSOR (e.g.
	account holder/client.		FDRO)
(If INCOMPLETE) Receives result of			
Pre-Assessment (Letter of Denial)	If found INCOMPLETE,		
	Generates result of Pre-Assessment.		
3. Pays the assessed fee through	3.1. Receives the payment/Official Receipt (OR)/ proof of	Day 0	Administrative and
Systems/Means prescribed by FDA.	payment through Systems/Means prescribed by FDA, and	Refer to FDA	Finance Services
	then posts the payment.	Cashier 's Citizen Charter	(AFS) STAFF
	3.2. Forwards application to CFRR, once payment is	Citizen Charter	
	posted.		
4. Receives Acknowledgement Receipt with the application and preassessment details.	4.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	8 Working Days	LRD EVALUATOR (e.g. FDRO)
	4.2. Checks application, ALL the submitted documentary requirements, the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	7 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)



	4.3 Reviews the checked application, ALL the submitted documentary requirements, the drafted recommendation of the CHECKER, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, and then finalizes the application by issuing Certificate of Product Registration (CPR) (for APPROVED application) or Letter of Denial (LOD) (for DISAPPROVED application), through the E-Registration System.	5 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
5. If the application is APPROVED , Receives an e-mail notification from FDA indicating that the application is approved, and other pertinent information.	5. Generates electronically signed CPR or LOD.		Information and Communication Technology Management Division (ICTMD) STAFF
If DISAPPROVED , receives a Letter of Denial/Disapproval (LOD) and another e-mail notification containing pertinent information about the application.			
Always as for to the compact FDA as well to	and the E. Dominton King Contains / E. Coming and https://www.fd-	TOTAL: 20 Working Days	
Always refer to the current FDA regulation/s on the us	e of the E-Registration System/E-Services: https://www.fda.gov.ph/		



1.2. ISSUANCE OF CERTIFICATE OF PRODUCT REGISTRATION (CPR) – AUTOMATIC RENEWAL APPLICATION (INITIAL APPLICATION APPROVED FROM MODIFIED E-REGISTRATION (VERSION 2))

'Registration' means the process of approval of an application to register health products prior to engaging in the manufacture, importation, exportation, sale, offer for sale, distribution, transfer, and where applicable, the use, testing, promotion, advertisement, and/or sponsorship of health products. (Republic Act No. 9711)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	In accordance to Administrative Order 50 s. 2001 + Legal Research Fee (LRF).
		Conventional Food (Category 1): Php 1,000.00/5 years of validity + 1% LRF Conventional Food (Category 2): Php 1,250.00/5 year of validity + 1% LRF Food Supplement: Php 5,000.00/5 years of validity + 1% LRF Bottled Water: Php 5,000.00/5 years of validity + 1% LRF

GENERAL GUIDELINES

Please refer to:

- 1) A. General Guidelines, B. Specific Guidelines, and C. Procedural Guidelines, IV. GUIDELINES, pages 2-10 of <u>FDA Circular No. 2020-033</u> || Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products"; and
- 2) III. General Guidelines, and IV. Specific Guidelines of <u>FDA Circular No. 2020-033-A</u> || Addendum to FDA Circular No. 2020-033, "Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products" to include Guidelines for Preassessment and Reiteration of Pre-Assessment Procedures in Applying for Certificate of Product Registration for Food

GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE
☑ Accomplished Application Form as	FDA Circular No.2020-033	https://www.fda.gov.ph/
prescribed by FDA regulations.	FDA Circular No.2020-033-A	
e.g. E-Registration System.		
Select "RENEWAL" as type of application		



using the same case number used in initial application.		
☑ Proof of payment of fees as prescribed by current FDA regulations	Administrative Order 50 s. 2001	Systems/Means prescribed by FDA
☑ Valid and appropriate FDA License to Operate (required for all types of CPR application) *The product being applied must be listed in the FDA approved Product Line/Category.	Administrative No. Order 2014-0029 Republic Act 9711	FDA Philippines
☑ A sworn statement indicating no change in or variation whatsoever in the product is attached to the application.	Implementing Rules and Regulations of Republic Act No. 9711	Applicant Company

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1.1. Files using the specific product/CASE NUMBER in the INBOX folder, and accomplishes (including uploading of the COMPLETE documentary requirements) the E-Registration System through the E-Portal https://eportal.fda.gov.ph based on the desired type of application in accordance to current FDA regulation/s on the use of the E-Registration Portal/E-Services.	Pre-assesses ONLY the completeness of the submitted documents through E-Registration System/E-Portal https://eportal.fda.gov.ph . Result of Pre-assessment will be received by the account holder.	Day 0	Center for Food Regulation and Research (CFRR) PRE-ASSESSOR (e.g. Food-Drug Regulation Officer (FDRO))
1.2. Forwards the application to PRE-ASSESSMENT .			
A system generated E-mail notification from FDA will be received by the client			



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upon submission of application for Pre- Assessment.			
2. (If COMPLETE) Receives the Order of	2. If found COMPLETE,	Day 0	CFRR PRE-
Payment.	Generates Order of Payment through the email of		ASSESSOR (e.g.
. aye	the account holder/client.		FDRO)
(If INCOMPLETE) Receives result of	the descart frontering		1 Bitte)
Pre-Assessment (Letter of Denial)	If found INCOMPLETE,		
Fie-Assessment (Letter of Denial)	Generates result of Pre-Assessment.		
		D 0	A 1
3. Pays the assessed fee through	3.1. Receives the payment/Official Receipt (OR)/	Day 0	Administrative and
Systems/Means prescribed by FDA	proof of payment through Systems/Means	Refer to FDA Cashier	Finance Services
	prescribed by FDA, and then posts the payment.	's	(AFS) STAFF
		Citizen Charter	
	3.2. Forwards application to CFRR, once payment		
	is posted.		
4. Receives Acknowledgement Receipt	4. Finalizes the application by issuing Certificate of	3 Working Days	CFRR APPROVING
with the application and pre-assessment	Product Registration (CPR) (for APPROVED		AUTHORITY
details.	application) or Letter of Denial (LOD) (for		(e.g. DIRECTOR IV)
	DISAPPROVED application), through the E-		(0.9. 220.0)
	Registration System.		
	Tregistration dystem.		
C If the emplication is ADDROVED	5 O		lufa
5. If the application is APPROVED ,	5. Generates electronically signed CPR or LOD.		Information and
Receives an e-mail notification from FDA			Communication
regarding the issuance of Certificate of			Technology
Product Registration (CPR), and other			Management Division
pertinent information.			(ICTMD)
			STAFF
If DISAPPROVED ,			
Receives an e-mail notification from FDA			
regarding the issuance of Letter of			
Denial/Disapproval (LOD), and other			
pertinent information.			
perunent iniornation.			



	TOTAL: 3 Working	
	Days	
Always refer to the current FDA regulation,	/s on the use of the E-Registration System/E-Services: https://www.fda.gov.ph/	



1.3. ISSUANCE OF CERTIFICATE OF PRODUCT REGISTRATION (CPR): AUTOMATIC RENEWAL (DATA CAPTURE)

'Registration' means the process of approval of an application to register health products prior to engaging in the manufacture, importation, exportation, sale, offer for sale, distribution, transfer, and where applicable, the use, testing, promotion, advertisement, and/or sponsorship of health products. (Republic Act No. 9711)

(DATA CAPTURE in the modified e-Registration System/Portal (Version 2) refers to applications processed in the old e-Registration Portal (Version 1) or thru manual registration system)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	In accordance to Administrative Order 50 s. 2001 + Legal Research Fee (LRF).
		Conventional Food (Category 1): Php 1,000.00/5 years of validity + 1% LRF Conventional Food (Category 2): Php 1,250.00/5 year of validity + 1% LRF Food Supplement: Php 5,000.00/5 years of validity + 1% LRF Bottled Water: Php 5,000.00/5 years of validity + 1% LRF

GENERAL GUIDELINES

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- 1) A. General Guidelines, B. Specific Guidelines, and C. Procedural Guidelines, IV. GUIDELINES, pages 2-10 of <u>FDA Circular No. 2020-033</u> | Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products"; and
- 2) III. General Guidelines, and IV. Specific Guidelines of <u>FDA Circular No. 2020-033-A</u> || Addendum to FDA Circular No. 2020-033, "Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products" to include Guidelines for Preassessment and Reiteration of Pre-Assessment Procedures in Applying for Certificate of Product Registration for Food



CHECKLIST OF REQUIREMENTS FOR ALL TYPES OF FOOD PRODUCTS/FOOD CATEGORIZATION: RAW MATERIALS, LOW RISK, MEDIUM RISK AND HIGH RISK FOOD PRODUCTS

RAW MAI ERIALS, LOW RISK, MEDIUM RISK AND HIGH RISK FOOD PRODUCTS					
GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE			
 ☑ Accomplished Application Form as prescribed by FDA regulations. e.g. E-Registration System. 	FDA Circular No.2020-033 FDA Circular No. 2020-033-A	https://www.fda.gov.ph/			
☑ Proof of payment of fees as prescribed by current FDA regulations	Administrative Order 50 s. 2001	Systems/Means prescribed by FDA			
☑ Valid and appropriate FDA License to Operate (LTO) (required for all types of CPR application) *The product being applied must be listed in the FDA approved Product Line/Category.	Administrative No. Order 2014-0029 Republic Act No. 9711	FDA			
☑ A sworn statement indicating no change in or variation whatsoever in the product is attached to the application.	Implementing Rules and Regulations of Republic Act No. 9711	Applicant Company			
☑ Upload ALL INITIAL requirements.	Administrative No. Order 2014-0029 FDA Circular No. 2020-033	Applicant Company/ In reference to the previously filed and approved INITIAL application (The validity of the documents to be submitted must be in accordance to current FDA regulation/s).			

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1.1. Accomplishes (including uploading of	1. Pre-assesses ONLY the completeness of the submitted		Center for Food
the COMPLETE documentary	documents through E-Registration System/E-Portal		Regulation and
requirements) the E-Registration System	https://eportal.fda.gov.ph		Research (CFRR)
through the E-Portal	Result of Pre-assessment will be received by the account		PRE-ASSESSOR
https://eportal.fda.gov.ph based on the	holder.		(e.g. Food-Drug
desired type of application in accordance			Regulation Officer
to current FDA regulation/s on the use of			(FDRO))



			PHILIPPINES
the E-Registration Portal/E-Services.			
1.2. Forwards the application to PRE-ASSESSMENT .			
A system generated E-mail notification from FDA will be received by the client upon submission of application for Pre-Assessment.			
2. (If COMPLETE) Receives the Order of Payment.	2. If found COMPLETE , Generates Order of Payment through the email of the account holder/client.		CFRR PRE- ASSESSOR (e.g. FDRO)
(If INCOMPLETE) Receives result of Pre- Assessment (Letter of Denial)	If found INCOMPLETE, Generates result of Pre-Assessment. To refile, the applicant must start a NEW CASE and upload initially submitted documentary requirements together with the documents for compliance to the deficiencies mentioned.		
3. Pays the assessed fee through Systems/Means prescribed by FDA	 3.1. Receives the payment/Official Receipt (OR)/ proof of payment through Systems/Means prescribed by FDA, and then post the payment. 3.2. Forwards application to CFRR, once payment is posted. 	Refer to FDA Cashier 's Citizen Charter	Administrative and Finance Services (AFS) STAFF
4. The applicant company receives Acknowledgement Receipt with the application and pre-assessment details.	4.1 Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, then drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	3 Working Days	LRD EVALUATOR (e.g. FDRO)



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	4.2 Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, then drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	2 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)
	4.3 Reviews the checked application, ALL the submitted documentary requirements, and the drafted recommendation of the CHECKER, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, and then finalizes the application by issuing Certificate of Product Registration (CPR) (for APPROVED application) or Letter of Denial (LOD) (for DISAPPROVED application), through the E-Registration System.	2 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
5. If the application is APPROVED , Receives an e-mail notification from FDA regarding the issuance of Certificate of Product Registration (CPR), and other pertinent information. If DISAPPROVED , Receives an e-mail notification from FDA regarding the issuance of Letter of Denial/Disapproval (LOD), and other pertinent information.	5. Generates electronically signed CPR or LOD.		Information and Communication Technology Management Division (ICTMD) STAFF
		TOTAL: 7 Working Days	

Always refer to the current FDA regulation/s on the use of the E-Registration System/E-Services: https://www.fda.gov.ph/



1.4. CERTIFICATE OF PRODUCT REGISTRATION (CPR) – INITIAL/ RENEWAL DATA CAPTURE (REGULAR)/ AMENDMENT DATA CAPTURE/ RE-APPLICATION DATA CAPTURE

'Registration' means the process of approval of an application to register health products prior to engaging in the manufacture, importation, exportation, sale, offer for sale, distribution, transfer, and where applicable, the use, testing, promotion, advertisement, and/or sponsorship of health products. (Republic Act No. 9711)

(DATA CAPTURE in the modified e-Registration System/Portal refers to applications processed in the old e-Registration Portal (Version 1) or thru manual registration system).

RENEWAL DATA CAPTURE (REGULAR) in the modified e-Registration System/Portal refers to applications processed in the old e-Registration Portal (Version 1) or thru manual registration system) which is not qualified to the General Guideline/s of AUTOMATIC RENEWAL.

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	In accordance to Administrative Order No. 50 s. 2001 + Legal Research Fee (LRF).
		Conventional Food (Category 1): Php 200.00/year of validity + 1% LRF
		Conventional Food (Category 2): Php 250.00/year of validity + 1% LRF
		Food Supplement: Php 1,000.00/year of validity + 1% LRF
		Bottled Water: Php 1,000.00/year of validity + 1% LRF

GENERAL GUIDELINES

Please refer to:

- 1) A. General Guidelines, B. Specific Guidelines, and C. Procedural Guidelines, IV. GUIDELINES, pages 2-10 of FDA Circular No. 2020-033 || Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products"; and
- 2) III. General Guidelines, and IV. Specific Guidelines of FDA Circular No. 2020-033-A || Addendum to FDA Circular 2020-033, "Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No.



2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products" to include Guidelines for Preassessment and Reiteration of Pre-Assessment Procedures in Applying for Certificate of Product Registration for Food.

CHECKLIST OF REQUIREMENTS

FOR ALL TYPES OF FOOD PRODUCTS/FOOD CATEGORIZATION: RAW MATERIALS, LOW RISK, MEDIUM RISK AND HIGH RISK FOOD PRODUCTS

PRODUCTS			
GENERAL REQUIREMENTS FOR APPLICATION OF CERTIFICATE OF PRODUCT REGISTRATION	BASIS/ISSUANC E	WHERE TO SECURE	
☑ ANNEX D - REQUIREMENTS FOR APPLICATION OF CERTIFICATE OF PRODUCT REGISTRATION	Administrative Order No. 2014- 0029		
☑ Accomplished Initial Application Form as prescribed by current FDA regulations. e.g. E-Registration System	FDA Circular No.2020-033 FDA Circular No.2020-033-A	https://www.fda.gov.ph/	
☑ Proof of Payment of Fees as prescribed by FDA regulations. Please refer to the table <i>Fees to be Paid:</i>	Administrative Order No. 50 s. 2001	Systems/Means prescribed by FDA	
☑ Clear and complete loose labels or artworks, as applicable, of all packaging sizes, or equivalents as defined by FDA regulations.	Administrative Order No. 2014- 0030; and other existing FDA regulation/s with specific labelling requirement/s (e.g. Republic Act No. 8172 Republic Act No. 8976 and its IRR Department Circular No. 2008- 0006 Bureau Circular	Applicant Company/ Manufacturer/Source/Supplie r	



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	No. 2 s. 1999 and etc.)	
☑ Pictures of the product in all angles and in different packaging sizes, and from at least two different perspectives allowing visual recognition of a product as the same with the one being registered, as applicable.	Administrative Order No. 2014- 0029	Applicant Company/ Manufacturer/Source/Supplie r
☑ For FOOD SUPPLEMENT, a sample in actual commercial presentation shall be submitted.	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
☑ As applicable, documents to substantiate claims, such as technical, nutritional or health studies or reports, market-research studies, Certificate of Analysis, quantitative analysis and computations, scientific report or studies published in peer-reviewed scientific journals, certificates or certification to support use of logo/seal on Diamond Sangkap Pinoy Seal, Sangkap Pinoy, Saktong Iodine sa Asin, Halal, Organic, or Kosher food and in compliance with current labeling regulations.	Administrative Order No. 2014- 0029 Administrative Order No. 2014- 0030	Applicant Company/ Manufacturer/Source/Supplie r
☑ VALID AND APPROPRIATE FDA LICENSE TO OPERATE (LTO) (REQUIRED FOR ALL TYPES OF CPR APPLICATION) *The product being applied must be listed in the FDA approved Product Line/Category.	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
For locally produced products: Distributorship Agreement or Contract Agreement signed by duly authorized representative of the establishment or Certificate of Distributorship or Appointment Letter or Memorandum of Agreement from each supplier. e.g. For WHOLESALER:	FDA Circular No. 2020-033 FDA Circular No. 2016-007	Applicant Company/ Manufacturer/Source/Supplie r
 Valid, notarized, and duly signed Distributorship Agreement or Memorandum of Agreement For TRADER: Valid, notarized, and duly signed Toll Manufacturing Agreement 		



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For imported products:	FDA Circular No.	Applicant Company/
☑ Distributorship Agreement or Contract Agreement signed by duly authorized	2020-033	Manufacturer/Source/Supplie
representative of the establishment or Foreign Agency Agreement, Certificate of	FDA Circular No.	r
Distributorship or Appointment Letter or Proforma Invoice or Memorandum of Agreement	2016-007	
from each supplier; and		
☑ Scanned copy of ANY of the following original and valid documents issued to the source		
by the regulatory or health authority from the country of origin per source:		
i) Valid manufacturer's certificate of registration with Good Manufacturing Practices (GMP)		
compliance or its equivalent; or		
ii) Valid Sanitary Phytosanitary Certificate/ Health Certificate; or		
iii) Valid ISO 22000 Certification/FSSC Certificate; or		
iv) Valid Hazard Analysis and Critical Control Point (HACCP) Certificate; or		
v) Certificate of Free Sale (CFS issued by the Regulatory/Health Authority attested by		
recognized Association or duly authenticated by the Philippine Consulate from the country of		
origin)		
*For export market only product, indicate the term FOR EXPORT MARKET ONLY as part of		
the product name in the data entry. Otherwise, your application will be evaluated as for local		
market distribution.		
*For institutional use only products, indicate the term FOR INSTITUTIONAL USE ONLY as		
part of the product name in the data entry. Otherwise, your application will be evaluated as		
conventional food for retail market distribution.		
ADDITIONAL REQUIREMENT/S PER FOOD CATEGORY: RAW MATERIAL, LOW RISK, ME	DIUM RISK AND HIG	H RISK FOOD PRODUCTS
CT ADDITIONAL	BASIS/ISSITANC	

RAW MATERIALS FOOD CATEGORIES	☑ ADDITIONAL REQUIREMENT/S	BASIS/ISSUANC E	WHERE TO SECURE
RAW MATERIALS - all substances that are employed in the processing of a finished product, packed in bulk containers and not labelled as finished product. Raw materials or ingredients would have product specifications that comply with the client requirements and not necessarily a single component.	☑ As applicable, certification to support use of logo/seal on Sangkap Pinoy, Halal, Organic, or Kosher food and in compliance with current labelling regulations.	Administrative Order No. 2014- 0029	Applicant Company/ Manufacturer/Source/Supplie r



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(Source: IV. Definition of terms, No. 36, page 6 of AO			
No. 2014-0029)			
RM01 – Fats, Oils and Fat Emulsions	☑ Valid Certificate of Analysis for	Republic Act No.	Applicant Company/
e.g. Cooking Oils (Coconut, Palm, Soybean	Vitamin A fortificant used for	8976	Manufacturer/Source/Supplie
and Corn)	COOKING OILS (e.g. Coconut,	Implementing	r
	Palm, Soybean and Corn)	Rules and	
		Regulation of	
	*Finished food products in bulk	Republic Act No.	
	intended for further processing	8976	
	shall conform with the applicable		
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM02 - Processed Fruits, Vegetable and Edible	*Finished food products in bulk	Administrative	Applicant Company/
Fungi, Seaweeds and Nuts	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM03 - Confectionery	*Finished food products in bulk	Administrative	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		



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	E-Registration data entry (e.g. under Product Specifications).		
RM04 - Cereals	*Finished food products in bulk intended for further processing shall conform with the applicable Administrative Orders set forth for Low-Risk, Medium-Risk and High-Risk Food Products. Compliance must be through declaration in the E-Registration data entry (e.g. under Product Specifications).	Administrative Order No. 2014- 0029	Applicant Company/ Manufacturer/Source/Supplie r
RM05 - Bakery Wares and Bakery Related Products e.g. Wheat Flour	✓ Valid Certificate of Analysis for Vitamin A and Iron fortificant used for WHEAT FLOUR *Finished food products in bulk intended for further processing shall conform with the applicable Administrative Orders set forth for Low-Risk, Medium-Risk and High-Risk Food Products. Compliance must be through declaration in the E-Registration data entry (e.g. under Product Specifications).	Republic Act No. 8976 Implementing Rules and Regulation of Republic Act No. 8976	Applicant Company/ Manufacturer/Source/Supplie r
RM06 - Sweeteners including Honey e.g. Refined Sugar, Brown Sugar, Cane Sugar	 ✓ Valid Certificate of Analysis for Vitamin A fortificant used for REFINED SUGAR *Finished food products in bulk intended for further processing shall conform with the applicable Administrative Orders set forth for 	Republic Act No. 8976 Implementing Rules and Regulation of Republic Act No. 8976	Applicant Company/ Manufacturer/Source/Supplie r



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	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM07 - Salt, Spices, Soups, Sauces, Salads and	☑ Valid Certificate of Analysis for	Republic Act No.	Applicant Company/
Protein Products	lodine Content used for IODIZED	8172	Manufacturer/Source/Supplie
e.g. lodized Salt, Soy Sauce	SALT	FDA Circular No.	r
	07.121	2013-007	
	*Finished food products in bulk		
	intended for further processing		
	shall conform with the applicable		
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
	,	FDA Memorandum	Applicant Company/
	☑ Valid Certificate of Analysis for	No. 2011-028	Manufacturer/Source/Supplie
	3MCPD content of SOY SAUCE	140. 2011 020	r
RM08 - Beverages (excluding Dairy Products) Non-	*Finished food products in bulk	Administrative	Applicant Company/
Alcoholic	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for	0020	•
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
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RM09 - Beverages (excluding Dairy Products)	*Finished food products in bulk	Administrative	Applicant Company/
Alcoholic	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM10- Dairy products and Analogues	*Finished food products in bulk	<u>Administrative</u>	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM11- Frozen Desserts	*Finished food products in bulk	Administrative	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
DM40 Proceed Fish and Fish Products Including	under Product Specifications).	A -1::	A
RM12 - Processed Fish and Fish Products Including	*Finished food products in bulk	Administrative	Applicant Company/
Molluscs, Crustaceans and Echinoderms	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable Administrative Orders set forth for	0029	1
	_		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		



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	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM13 - Herbal Products	*Finished food products in bulk	<u>Administrative</u>	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM14 - Vitamins and Minerals	*Finished food products in bulk	<u>Administrative</u>	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM15 - Products with Nutritional Substances	*Finished food products in bulk	<u>Administrative</u>	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM16 - Food Additives	*Finished food products in bulk	Administrative	Applicant Company/
	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	<u>0029</u>	r



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	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM17 - Edible Casings (except natural casings from	*Finished food products in bulk	<u>Administrative</u>	Applicant Company/
animal sources)	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
RM18 - Processed Meat and Meat Products,	*Finished food products in bulk	Administrative	Applicant Company/
including poultry and game	intended for further processing	Order No. 2014-	Manufacturer/Source/Supplie
	shall conform with the applicable	0029	r
	Administrative Orders set forth for		
	Low-Risk, Medium-Risk and High-		
	Risk Food Products. Compliance		
	must be through declaration in the		
	E-Registration data entry (e.g.		
	under Product Specifications).		
LOW RISK FOOD PRODUCTS	☑ ADDITIONAL REQUIREMENT/S	BASIS/ISSUANC E	WHERE TO SECURE
LOW RISK FOOD PRODUCTS - foods that are			
unlikely to contain pathogenic microorganisms and			
will not normally support their growth because of			
food characteristics and foods that are unlikely to contain harmful chemicals.			
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A1 - Butter oil, anhydrous milkfat, ghee	☑ In the Electronic Registration	Administrative	Applicant Company/
"The milkfat products anhydrous milkfat, anhydrous	Data Entry – Product	<u>Order 132 s. 1970</u>	Manufacturer/Source/Supplie
butter oil and butter oil are products derived exclusively	Specifications		r
from milk and/or products obtained from milk by a	Physical/Chemical/Microbiological		
process that almost completely removes water and	, declare the results (under		
nonfat solids. Ghee is a product obtained exclusively	specification) for the following		
from milk, cream or butter by a process that almost	Parameters: %Milk Fat by weight;		
completely removes water and nonfat solids; it has a	% Milk Solids not fat by weight; %		
specially developed flavour and physical structure"	water by weight; Salt (optional) for		
(Source URL:	BUTTER (Whipped, Pasteurized)		
https://www.fao.org/gsfaonline/foods/details.html?id=41)	☑ In the Electronic Registration	Administrative	Applicant Company/
	Data Entry – Product	Order 132 s. 1970	Manufacturer/Source/Supplie
	Specifications		r
	Physical/Chemical/Microbiological		
	, declare the results (under		
	specification) for the following		
	Parameters: %Milk Fat by weight;		
	% Milk Solids not fat by weight; %		
	water by weight; Salt (optional) for		
	WHEY BUTTER		
	☑ In the Electronic Registration	Administrative	Applicant Company/
	Data Entry – Product	Order No. 232 s.	Manufacturer/Source/Supplie
	Specifications	1974	r
	Physical/Chemical/Microbiological		
	, declare the results (under		
	specification) for the following		
	Parameters: % Fat; % Moisture		
	for MARGARINE		
	*The product shall conform with		
	the standards for optional		



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	ingredients and additional label declaration for MARGARINE.		
A2 - Vegetable Oils and Fats e.g. Coconut, Palm, Soybean and Corn "Edible fats and oils obtained from edible plant sources. Products may be from a single plant source or marketed and used as blended oils that are generally designated as edible, cooking, frying, table or salad oils. Virgin oils are obtained by mechanical means (e.g., pressing or expelling), with application of heat only so as not to alter the natural composition of the oil. Virgin oils are suitable for consumption in the natural state. Cold pressed oils are obtained by mechanical means without application of heat. Examples include: virgin olive oil, cottonseed oil, peanut oil, and vanaspati." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=42)	☑ Valid Certificate of Analysis for Vitamin A fortificant (in mg RE/L) used for COOKING OILS (e.g.Coconut, Palm, Soybean and Corn) *The specific form of Vitamin A fortificant used (e.g. Retinol Palmitate) shall be declared in the Electronic Registration Data Entry.	Republic Act No. 8976 Implementing Rules and Regulation of Republic Act No. 8976	Applicant Company/ Manufacturer/Source/Supplie r
A3 - Animal Fats "All animal fats and oils should be derived from animals in good health at the time of slaughter and intended for human consumption. Lard is fat rendered from the fatty tissue of swine. Edible beef fat is obtained from fresh bovine fatty tissue covering the abdominal cavity and surrounding the kidney and heart, and from other compact, undamaged fat tissues. Such fresh fat obtained at the time of slaughter is the "killing fat." Prime beef fat (premiere jus or oleo stock) is obtained by lowheat rendering (50-55C) of killing fat and selected fat trimmings (cutting fat). Secunda beef fat is a product with typical beef fat odor and taste obtained by rendering (60-65C) and purifying beef fat. Rendered pork fat is fat obtained from the tissue and bones of swine. Edible tallow (dripping) is produced by the	☑ In the Electronic Registration Data Entry – Product Specifications Physical/Chemical/Microbiological , declare the results (under specification) for the following Parameters: Saponification Value; lodine Value for LARD	Administrative Order No. 231 s. 1974	Applicant Company/ Manufacturer/Source/Supplie r



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rendering of fatty tissue (excluding trimmings and cutting fat), attached muscles and bones of bovine animals or sheep. Fish oils are derived from suitable sources such as herring, sardines, sprat, and anchovies. Other examples include: tallow and partially defatted beef or pork fatty tissue." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=43)			
A4 - Fat emulsions mainly of type oil-in-water "Includes fat-based counterparts of dairy-based foods excluding dessert products." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=50) e.g. Imitation milk - a fat-substituted milk produced from nonfat milk solids by addition of vegetable fats (coconut, safflower or corn oil), non-dairy whipped cream, non- dairy toppings and vegetable cream	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A5 - Fat emulsions mainly of type water-in-oil "Include all emulsified products excluding fat-based counterparts of dairy products and dairy desserts." (Source URL: https://www.fao.org/qsfaonline/foods/details.html?id=44) e.g., Margarine, reduced-fat based desserts	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A6 - Fat-based desserts excluding dairy-based desserts "Includes fat-based counterparts of dairy-based desserts. Includes ready-to-eat products and their mixes. Also includes non-dairy fillings for desserts." e.g., ice cream-like product made with vegetable fats (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=51)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
B1 - Dehydrated fruits or vegetables, including candied fruits	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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"Includes glazed fruits (fruit treated with a sugar so and dried), candied fruit (dried glazed fruit immerse sugar solution and dried so that the fruit is covered candy-like sugar shell), and crystallized fruit is prep (dried glazed fruit rolled in icing or granulated sugar dried). Examples include: cocktail (maraschino) che candied citrus peel, candied citrons (e.g. used in his fruitcakes), and mostarda di frutta." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?ice.page.granulades	ed in a by a pared r and erries, pliday	Administrative	Applicant Company/
"Jams, preserves and conserves are thick, spread products prepared by boiling whole fruit or pieces of fruit, fruit pulp or puree, with or without fruit juice of concentrated fruit juice, and sugar to thicken, and which pectin and fruit pieces may be added. Jelly it clear spreadable product prepared similarly to jame except that it has a smoother consistency and doe contain fruit pieces. Marmalade is a thick spreadable fruit slurry prepared from whole fruit, fruit pulp or product prepared to thicken, to which pectin and fruit pieces and fruit peel pieces in the be added. 38,40 Includes dietetic counterparts made.	Specifications Physical/Chemical/Microbiological , declare the results (under specification) for the following Parameters: Soluble Solids for JELLY/JELLIES *The product shall conform with the standard of quality and additional label declaration for	Administrative Order No. 239 s. 1975	Applicant Company/ Manufacturer/Source/Supplie r
non-nutritive high-intensity sweeteners. Examples include: orange marmalade, grape jelly, and strawl jam" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id	Specifications Physical/Chemical/Microbiological	Administrative Order No. 238 s. 1975	Applicant Company/ Manufacturer/Source/Supplie r



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	*The product shall conform with the standard of quality and additional label declaration for PRESERVES OR JAMS.		
B3 - Dehydrated vegetable protein products e.g., Textured Vegetable Protein	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
B4 - Fruits or Vegetables in vinegar, oil or brine "Products prepared by treating raw vegetables with salt solution excluding fermented soybean products." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=80) Note: Fruits or vegetables in vinegar, oil or brine in canned, bottled or hermetically sealed containers must be file under Medium Risk Food Product - MRC3	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
B5 - Fruit-based spreads excluding jams, jellies and marmalades "Includes all other fruit-based spreads, such as apple butter and lemon curd. Also includes condiment-type fruit products such as mango chutney and raisin chutney." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=65)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
e.g. fruit pulp, purees, fruit toppings, fruit sauce, fruit syrup, coconut milk and cream "Fruit pulp is not usually intended for direct consumption. It is a slurry of lightly steamed and strained fresh fruit, with or without added preservatives. Fruit puree (e.g., mango puree, prune puree) is produced in the same way, but has a smoother, finer texture, and may be used as fillings for pastries, but is not limited to this use. Fruit sauce (e.g., pineapple sauce or strawberry sauce) is made from boiled fruit pulp with or without added	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



sweeteners and may contain fruit pieces. Fruit sauce may be used as toppings for fine bakery wares and ice cream sundaes. Fruit syrup (e.g., blueberry syrup) is a more liquid form of fruit sauce that may be used as a topping e.g., for pancakes.Non-fruit toppings are included in category 05.4 (sugar- and chocolate-based toppings) and sugar syrups (e.g. maple syrup) are included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or comminuted fresh endosperm (kernel) of coconut palm
cream sundaes. Fruit syrup (e.g., blueberry syrup) is a more liquid form of fruit sauce that may be used as a topping e.g., for pancakes.Non-fruit toppings are included in category 05.4 (sugar- and chocolate-based toppings) and sugar syrups (e.g. maple syrup) are included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
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topping e.g., for pancakes.Non-fruit toppings are included in category 05.4 (sugar- and chocolate-based toppings) and sugar syrups (e.g. maple syrup) are included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
included in category 05.4 (sugar- and chocolate-based toppings) and sugar syrups (e.g. maple syrup) are included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
toppings) and sugar syrups (e.g. maple syrup) are included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
included in category 11.4. Coconut milk and coconut cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
cream are products prepared using a significant amount of separated, whole, disintegrated macerated or
of separated, whole, disintegrated macerated or
comminuted fresh endosperm (kernel) of coconut palm
and expelled, where most filterable fibers and residues
are excluded, with or without coconut water, and/or with
additional water. Coconut milk and coconut cream are
treated by heat pasteurization, sterilization or ultrahigh
temperature (UHT) processes. Coconut milk and
coconut cream may also be produced in concentrated or
skim (or "light") forms. Examples of traditional foods in
this sub-category are: tamarind concentrate (clean
extract of tamarind fruit with not less than 65% total
soluble solids), tamarind powder (tamarind paste mixed
with tapioca starch), tamarind toffee (mixture of tamarind
pulp, sugar, milk solids, antioxidants, flavours, stabilizers and preservatives), and fruit bars (a mixture of fruit
(mango, pineapple, or guava) pulp mixed with sugar,
flavours and preservatives, dried into a sheet)."
(Source URL:
https://www.fao.org/gsfaonline/foods/details.html?id=67)
B7 - Cooked fruits
"Fruit that is steamed hoiled haked or fried with or
without a coating, for presentation to the consumer. NOT APPLICABLE NOT APPLICABLE APPLICABLE NOT APPLICABLE
Examples include: baked apples, fried apple rings, and



			PHILIPPINES
peach dumplings (baked peaches with a sweet dough			
covering."			
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=71)			
B8 - Frozen vegetables, seaweeds, and nuts and			
seeds			
"Fresh vegetables are usually blanched and frozen.			
Examples include: quick-frozen corn, quick-frozen	NOT APPLICABLE	NOT	NOT APPLICABLE
French-fried potatoes, quick frozen peas, and quick	NOTALLECABLE	APPLICABLE	NOTALLECABLE
frozen whole processed tomatoes."			
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=78)			
B9 - Vegetable seaweeds, nut and seed in pulps and		NOT	
preparations other than food in HR Letter B2	NOT APPLICABLE	APPLICABLE	NOT APPLICABLE
e.g. Aloe extract, potato pulp, horseradish pulp		ALLEGABLE	
B10 - Cooked or fried vegetables and seaweeds			
"Vegetables that are steamed, boiled, baked, or fried,			
with or without a coating, for presentation to the			
consumer. Examples include: simmered beans, pre-fried	NOT APPLICABLE	NOT	NOT APPLICABLE
potatoes, fried okra, and vegetables boiled down in soy	1401741 EIG/ABEE	APPLICABLE	1401741 LIONBLE
sauce (tsukudani)."			
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=85)			
C1 - Confectionery			
"Includes all types of products that mainly contain sugar			
and other dietetic counterparts and may or may not		NOT	
contain cocoa (e.g. Hard candy, soft candy, nougats and	NOT APPLICABLE	APPLICABLE	NOT APPLICABLE
marzipans"		7 TEIO/IDEE	
Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=93			
C2 - Chewing gum		NOT	
"Product made from natural or synthetic gum base	NOT APPLICABLE	APPLICABLE	NOT APPLICABLE
containing flavours, sweeteners (nutritive or non-		, ar Elonbee	



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nutritive), aroma compounds, and other additives. Includes bubble gum and breath-freshener gum products." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=97)			
C3 - Decorations, toppings (non-fruit), and sweet			
sauces			
"Includes ready-to-eat icings and frostings for cakes, cookies, pies and bread and flour confectionery, as well as mixes for these products." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=98) e.g., Ready-to-eat icings and frostings for cakes, cookies etc, maple, caramel and flavoured syrups	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
D1 - Flour, starches (including soybean powder) and	☑ Valid Certificate of Analysis for	Republic Act No.	Applicant Company/
flour mixes	Vitamin A fortificant (in mg/kg as	8976	Manufacturer/Source/Supplie
"The basic milled products of cereal grains, roots,	retinol) and Iron fortificant (in mg	Implementing	r
tubers, pulses, pith or soft core of palm tree or legumes	Fe/kg) used for WHEAT FLOUR	Rules and	
sold as such or used as ingredients (e.g. in baked goods)."	*TI :: (Regulation of Republic Act No.	
(Source URL:	*The specific form of Vitamin A	8976	
https://www.fao.org/gsfaonline/foods/details.html?id=101	fortificant used (e.g. Retinol Palmitate) and Iron fortificant	0070	
)	used (e.g. Elemental Iron, Ferrous		
e.g. Wheat flour, corn flour, bran	Sulfate, Ferrous Fumarate) shall		
	be declared in the Electronic		
	Registration Data Entry.		
D2 - Breakfast cereals including rolled oats			
"Includes all ready-to-eat, instant, and regular hot			
breakfast cereal products. Examples include: granola-	NOT APPLICABLE	NOT	NOT APPLICABLE
type breakfast cereals, instant oatmeal, farina, corn	NOTALLOADLE	APPLICABLE	NOTALLEGABLE
flakes, puffed wheat or rice, multi-grain (e.g. rice, wheat			
and corn) breakfast cereals, breakfast cereals made			



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from soy or bran, and extruded-type breakfast cereals made from grain flour or powder." (Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=104			
)			
e.g. granola type breakfast cereals, corn flakes,			
multi-grain			
D3a - Fresh pastas and noodles and like products			
"Products that are untreated (i.e. not heated, boiled,			
steamed, cooked, pre-gelatinized or frozen) and are not			
dehydrated. These products are intended to be			
consumed soon after preparation. Examples include:	NOT APPLICABLE	NOT	NOT APPLICABLE
unboiled noodles, and "skins" or crusts for spring rolls, wontons, and shuo mai."		APPLICABLE	
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=106			
)			
D3b - Dried pastas and noodles and like products			
"Products that are untreated (i.e. not heated, boiled,			
steamed, cooked, pre-gelatinized or frozen) and are			
dehydrated."		NOT	
(Source URL:	NOT APPLICABLE	APPLICABLE	NOT APPLICABLE
https://www.fao.org/gsfaonline/foods/details.html?id=107			
e.g. spaghetti pasta, bean vermicelli, rice			
vermicelli, macaroni, rice noodles			
D3c - Pre-cooked pastas and noodles and like			
products			
"Products that are treated (i.e. heated, boiled, steamed,		NOT	
cooked, pre-gelatinized or frozen). These products may	NOT APPLICABLE	APPLICABLE	NOT APPLICABLE
be sold directly to the consumer (e.g. pre-cooked, chilled		AFFLICABLE	
gnocchi to be heated prior to consumption), or may be			
the starch component of prepared meals (e.g., heat-and-			



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serve frozen dinner entrees containing spaghetti, macaroni or noodles; canned spaghetti and meatballs entrée). Also includes instant noodles (sokuseki-men; e.g. pre-cooked ramen, udon, rice noodles), that are pre-gelatinized, heated and dried prior to sale to the consumer." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=108) e.g. Instant noodles			
D4 - Cereal and starch-based desserts "Dessert products containing cereal, starch or grain as the main ingredient. Also includes cereal- or starch based fillings for desserts. Examples include: rice pudding, semolina pudding, tapioca pudding, rice flour dumplings (dango), a steamed yeast-fermented wheat flour dough dessert (musipan), and a starchy pudding based dessert (namagashi)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=109)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
"Products containing flaked or ground cereal or grain that when combined with other ingredients (e.g., egg, water, milk) are used as a coating for fish or poultry. Products are usually sold as dry mix of the cereal or grain component. Examples include breading for tempura batter." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=110) e.g. for breading or batters for fish or poultry	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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D6 - Pre-cooked or processed rice products	☑ Valid Certificate of Analysis for	Republic Act No.	Applicant Company/
e.g. Prepackaged Rice in Retail Size, Iron Rice	Iron fortificant (in mg Fe/kg) used	8976	Manufacturer/Source/Supplie
Premix	for RICE	Implementing	r
		Rules and	
	*The specific form of Iron	Regulation of	
	fortificant used (e.g. Ferrous	Republic Act No.	
	Sulfate) shall be declared in the	8976	
	Electronic Registration Data		
	Entry.	EDAO: L N	A 1: 10 /
	☑ In the Electronic Registration	FDA Circular No.	Applicant Company/
	Data Entry – Product	<u>2007-010-A</u>	Manufacturer/Source/Supplie
	Specifications		I
	Physical/Chemical/Microbiological		
	, declare the results (under specification) for the following		
	Parameters: Iron Content (in mg		
	Iron (Fe)/100g and Moisture		
	Content for IRON RICE PREMIX		
	Contone for Internal Internal		
	*The specific form of Iron		
	fortificant used (e.g. Ferrous		
	Sulfate) shall be declared in the		
	Electronic Registration Data		
	Entry.		
	**The product shall conform with		
	the Composition and Quality		
	Factors for Iron Rice Premix		
D7a - Soybean based beverages			
"Products prepared from dried soybeans that are soaked	NOT APPLICABLE	NOT	NOT APPLICABLE
in water, pureed, boiled and strained, or prepared from		APPLICABLE	
soybean flour, soybean concentrate, or soybean isolate."			



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(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=269			
D7b - Soybean based film "Film formed on the surface of boiling soybean-based beverage that is dried." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=270) e.g. Fuzhu - asian food which is a protein–lipid film isolated from soymilk surface through high-temperature incubation	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
D7c - Soybean curd (tofu) "Soybean curd is prepared from dried soybeans that are soaked in water, pureed, and strained to produce soybean-based beverage, which is then made into a curd with a coagulant, and placed in a mould. Soybean curds may be of a variety of textures (e.g. soft, semi-firm, firm)" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=271)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
D7d - Semi-dehydrated soybean curd "Soybean curd that has been pressed while being moulded into blocks so that some moisture has been removed, but so that it is not completely dried. Semi- dehydrated soybean curd typically contains 62% water, and has a chewy texture." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=272)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
D7e - Dehydrated soybean curd	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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"Soybean curd from which all moisture has been removed through the process of freezing, aging, and dehydrating. It may be reconstituted with water or sauce for consumption, or is used directly in prepared dishes. It may also be deep-fried or simmered in sauce." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=276)			
D7f - Other soybean protein products "Other products from soybeans composed mainly of soybean protein such as extruded, textured, concentrated, and isolated soybean protein." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=279) e.g. Soy-based "chicken" meat	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1a - Breads and rolls - yeast leavened breads and specialty breads, soda breads "Includes yeast-leavened and specialty breads and soda bread." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=115 e.g. White bread, raisin bread, whole wheat bread, hamburger rolls, hotdog buns	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1b - Crackers excluding sweet crackers "The term "cracker" refers to a thin, crisp wafer, usually of unsweetened dough." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=118 e.g. Soda Crackers, Rye Crisps, Matzohs	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1c - Other ordinary bakery products "Includes all other ordinary bakery wares, such as	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



			PHILIPPINES
cornbread and biscuits. The term "biscuit" in this category refers to a small cake of shortened bread, leavened with baking powder or baking soda. It does not refer to the British "biscuit," which is a "cookie" or "sweet cracker" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=119) e.g. Bagels, pita, English muffins			
F1d - Bread-type products, including bread stuffing and bread crumbs "Includes bread-based products such as croutons, bread stuffing and stuffing mixes, and prepared doughs (e.g. for biscuits)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=120) e.g. Croutons	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1e - Steamed bread and buns "Oriental-style leavened wheat or rice products that are cooked in a steamer. Products may be made with or without filling. In China, products without filling are called steamed bread (mantou), and those with filling are called steamed buns (baozi or bao). Twisted rolls of various shapes (huajuan) may also be prepared. Examples include: filled dumplings and steamed bun with meat, jam or other filling (manjyu)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=121) Other e.g., Siopao	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1f - Mixes for bread and ordinary bakery wares "Includes all the mixes containing the dry ingredients to	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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which wet ingredients (e.g., water, milk, oil, butter, eggs) are added to prepare a dough for baked goods." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=122)			
F2 - Fine bakery wares and mixes - Mixes for fine bakery wares "Mixes containing the dry ingredients to which wet ingredients (e.g. water, milk, oil, butter, eggs) are added to prepare a dough for fine baked goods." e.g. cake mix, flour confectionery mix, pancake mix, pie mix, and waffle mix (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=126)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
G1 - Refined and raw sugars "Nutritive sweeteners, such as fully or partially purified sucrose (derived from sugar beet and sugar cane), glucose (derived from starch), or fructose." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=174) e.g. Refined Sugar, Raw Cane Sugar	☑ Valid Certificate of Analysis for Vitamin A fortificant used for REFINED SUGAR	Republic Act No. 8976 Implementing Rules and Regulation of Republic Act No. 8976	Applicant Company/ Manufacturer/Source/Supplie r
G2 - Brown Sugar "Includes large-grain, brown or yellow lump sugars, such as Demerara sugar" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=182)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
G3 - Sugar solutions and syrups "Includes co-products of the sugar refining process (e.g. treacle and molasses), invert sugar (equimolar mixture of glucose and fructose produced from the hydrolysis of	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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sucrose), and other sweeteners, such as high fructose corn syrup, high fructose inulin syrup and corn sugar." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=183) e.g. Maple Syrup, Vanilla Syrupm Flavoured Syrups			
G4 - Other sugars and syrups including coconut sugar e.g. Coloured sugar crystals for cookies	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
"Honey is the natural sweet substance produced by honeybees from the nectar of blossoms or secretions of plants. The honeybees collect the nectar or secretions, transform it by combination with specific substances of the bees' own, and store it in a honeycomb to ripen and mature." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=185) e.g. Wildflower Honey and Clover Honey	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
G6- Table-top sweeteners, including those containing high-intensity sweeteners "Includes products that are preparations of high-intensity sweeteners (e.g. acesulfame potassium) and/or of polyols (e.g. sorbitol) which may contain other additives and/or nutritive ingredients, such as carbohydrates. These products, which are sold to the final consumer, may be in powder, solid (e.g. tablets or cubes), or liquid form." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=186)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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In - Salt and Salt substitutes Salt - "Primarily food-grade sodium chloride. Includes table salt, iodized and fluoride iodized salt, and dendritic salt." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=189) "Salt substitutes are seasonings with reduced sodium content intended to be used on food in place of salt." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=190)	✓ Valid Certificate of Analysis for lodine Content for SALT, ROCK SALT, SEA SALT (Excluding Himalayan Pink Salt, Gourmet Salt) * "All food manufacturers processors using food-grade salt are also required to use iodized salt in the processing of their products and must comply with the provisions of this Act not later than one (1) year from its effectivity. Provided, That the use of iodized salt shall not prejudice the quality and safety of their food products: Provided, however, That the burden of proof and testing for any prejudicial effects due to iodized salt fortification lies on the said food manufacturers/processor." – RA No. 8172	Republic Act No. 8172 FDA Circular No. 2013-007	Applicant Company/ Manufacturer/Source/Supplie r
I2 - Herbs, spices, seasonings and condiments "Herbs and spices are usually derived from botanical sources, and may be dehydrated, and either ground or whole. Examples of herbs include basil, oregano and thyme. Examples of spices include cumin and caraway seeds. Spices may also be found as blends in powder or paste form." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=192)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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"Condiments and seasonings are mixtures of herbs and spices together with other food ingredients (such as salt, vinegar, lemon juice, molasses, honey or sugar, and sweeteners). Examples include meat tenderizers, onion salt, garlic salt, Oriental seasoning mix (dashi), topping to sprinkle on rice (furikake, containing, e.g. dried seaweed flakes, sesame seeds and seasoning), and seasoning for noodles. The term "condiments" as used in the Food Category System does not include condiment sauces (e.g. ketchup, mayonnaise, mustard) or relishes." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=193)			
"Liquid produced from fermentation of ethanol from a suitable source (e.g. wine, cider). Examples include, cider vinegar, wine vinegar, malt vinegar, spirit vinegar, grain vinegar, raisin vinegar, and fruit (wine) vinegar." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=194)	☑ In the Electronic Registration Data Entry – Product Specifications Physical/Chemical/Microbiological , declare the results (under specification) for the following Parameters: % Acidity; % Total Solids; % Ash; Lead Content; Copper Content and Arsenic Content; *Additional for Malt Vinegar: Phosphorus Pentoxide and Nitrogen Contents for VINEGAR	Administrative Order No. 134 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
I4 – Mustards "Condiment sauce prepared from ground, often defatted mustard seed that is mixed into a slurry with water, vinegar, salt, oil and other spices and refined. Examples include Dijon mustard, and "hot" mustard (prepared from seeds with hulls)."	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



			PHILIPPINES
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=195			
I5 - Soups and broths "Concentrated soup to be reconstituted with water and/or milk, with or without addition of other optional ingredients (e.g. vegetables, meat, noodles). Examples include: bouillon powders and cubes; powdered and condensed soups (e.g. mentsuyu); and stock cubes and powders." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=198	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
IGA - Mixes for sauces and gravies "Concentrated product, usually in powdered form, to be mixed with water, milk, oil or other liquid to prepare a finished sauce or gravy. Examples include mixes for cheese sauce, hollandaise sauce, and salad dressing (e.g. Italian or ranch dressing)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=202)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
If the control of the	☑ In the Electronic Registration Data Entry – Product Specifications Physical/Chemical/Microbiological , declare the results (under specification) for the following Parameters: Specific Gravity; Total Solids; Salt Content; Protein Content for PATIS	Administrative Order No. 325 s. 1977	Applicant Company/ Manufacturer/Source/Supplie r



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I7 - Yeast and like products "Includes baker's yeast and leaven used in the manufacture of baked goods. Includes the Oriental products koji (rice or wheat malted with A. oryzae) used in the production of alcoholic beverages." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=205)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
Isa - Fermented Soybean Paste (e.g. Miso) "The product is made of soybeans, salt, water and other ingredients, using the process of fermentation. The product includes dou jiang (China), doenjang (Republic of Korea), or miso (Japan), which maybe used in the preparation of soups or dressings, or as a seasoning." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=207)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
I8b- Soybean Sauce "A liquid seasoning obtained by fermentation of soybeans, non-fermentation (e.g. hydrolysis) of	☑ Valid Certificate of Analysis for 3-MCPD for SOY SAUCE	FDA Memorandum 2011-028	Applicant Company/ Manufacturer/Source/Supplie r
soybeans, or by hydrolysis of vegetable protein" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=211)			
I9- Protein products other than from soybeans, marinades "Includes, for example, milk protein, cereal protein and vegetable protein analogues or substitutes for standard products, such as meat, fish or milk. Examples include: vegetable protein analogues, fu (a mixture of gluten (vegetable protein) and flour that is sold dried (baked) or raw, and is used as an ingredient, e.g. in miso soup) and proteinaceous meat and fish substitutes."	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=218			
e.g. Vegetable Protein Analogues			
J1a - Non-alcoholic (soft) beverages without herbal	☑ In the Electronic Registration	Administrative	Applicant Company/
ingredients	Data Entry – Product	Order No. 136-As.	Manufacturer/Source/Supplie
e.g. Roasted coffee beans, coffee grounds, Freeze-dried	Specifications	1985	r
coffee	Physical/Chemical/Microbiological	1000	·
	, declare the results (under		
	specification) for the following		
	Parameters: Moisture Content		
	(%w/w); Caffeine (%w/w dry		
	, , ,		
	basis); Ash (%w/w dry basis;		
	Water-insoluble Solids (%w/w, dry		
	basis); pH; Solubility; Sensory		
	Attributes; Arsenic Content; Lead		
	Content for INSTANT COFFEE	A 1 1 1 1 1 1 1	10 /
	☑ In the Electronic Registration	Administrative	Applicant Company/
	Data Entry – Product	Order No. 136-B s.	Manufacturer/Source/Supplie
	Specifications	<u>1985</u>	r
	Physical/Chemical/Microbiological		
	, declare the results (under		
	specification) for the following		
	Parameters: Moisture Content		
	(%w/w); Caffeine (%w/w, dry		
	basis); Ash (%w/w, dry basis;		
	Water-insoluble Solids (%w/w, dry		
	basis); Carbohydrates (% w/w, dry		
	basis); pH; Solubility; Sensory		
	Attributes; Arsenic Content; Lead		
	Content for SOLUBLE COFFEE		
	•		



	140000	T	PHILIPPINES
	WITH ADDED CARBOHYDRATES		
J1b - Non-alcoholic (soft) beverages with herbal ingredients e.g. Green Tea, Chamomile Tea	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
J2a - Beer and Malt Beverages "Alcoholic beverages brewed from germinated barley (malt), hops, yeast, and water. Examples include: ale, brown beer, weiss beer, pilsner, lager beer, oud bruin beer, Obergariges Einfachbier, light beer, table beer, malt liquor, porter, stout, and barleywine." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=254)	☑ For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including methanol content); b) a certificate of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b)	Memorandum Circular No. 13 s. 1989	Applicant Company/ Manufacturer/Source/Supplie r
	✓ For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages	Memorandum Circular No. 13 s. 1989	Applicant Company/ Manufacturer/Source/Supplie r
J2b - Cider and Perry "Fruit wines made from apples (cider) and pears (perry). Also includes cidre bouche." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=255)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



"Alcoholic beverage obtained exclusively from the partial or complete alcoholic fermentation of fresh grapes, whether crushed or not, or of grape must (juice)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=256) e.g. Still grape wine, sparkling and semi-sparkling grape wines, fortified grape wine, grape liquor wine, sweet grape wine, red wine, white wine, rose wine ### For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) a certificate of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) ### For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages ### JZd - Wines other than grape ### Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or ### Or Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or				PHILIPPINES
specifications of raw materials and finished product (including methanol content); b) a certificate of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) If predictions of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape Includes wines made from fruit other than grapes, and from other agricultural products, including grain (e.g. rice). These wines may be still or	J2c - Grape Wines	☑ For IMPORTED ALCOHOLIC	<u>Memorandum</u>	Applicant Company/
whether crushed or not, or of grape must (juice)." (Source URL: https://www.fao.org/qsfaonline/foods/details.html?id=256) e.g. Still grape wine, sparkling and semi-sparkling grape wines, fortified grape wine, grape liquor wine, sweet grape wine, red wine, white wine, rose wine We for LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or Applicant Company/ Manufacturer/Source/Supplie For INCOALLY Memorandum Circular No. 13 s. 1989 Memorandum Circular No. 13 s. 1989 Applicant Company/ Manufacturer/Source/Supplie For INPORTED ALCOHOLIC Beverages For INCOALLY Memorandum Circular No. 13 s. 1989 Applicant Company/ Manufacturer/Source/Supplie Circular No. 13 s. Sepecifications of raw materials 1989 Circular No. 13 s. 1989	"Alcoholic beverage obtained exclusively from the partial	BEVERAGES: a) Technical	Circular No. 13 s.	Manufacturer/Source/Supplie
(Source URL: https://www.fao.org/qsfaonline/foods/details.html?id=256) e.g. Still grape wine, sparkling and semi-sparkling grape wines, fortified grape wine, grape liquor wine, sweet grape wine, red wine, white wine, rose wine For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape For IMPORTED ALCOHOLIC BEVERAGEs: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages For IMPORTED ALCOHOLIC BEVERAGEs: a) Technical specifications of raw materials and finished product (including grain (e.g. rice). These wines may be still or These wines wine wine wine with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) Applicant Company/ Memorandum These wines wines wines wines and regulation stated in (b) These wines	or complete alcoholic fermentation of fresh grapes,	specifications of raw materials	<u>1989</u>	r
of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) If or LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) If or LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages If or IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including grain (e.g. rice). These wines may be still or	whether crushed or not, or of grape must (juice)."	and finished product (including		
origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) Memorandum Circular No. 13 s. 1989 Applicant Company/ Manufacturer/Source/Supplie or Imported National specifications of raw materials and finished product (including grain (e.g. rice). These wines may be still or	(Source URL:	methanol content); b) a certificate		
origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b) Memorandum Circular No. 13 s. 1989 Applicant Company/ Manufacturer/Source/Supplie r For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including grain (e.g. rice). These wines may be still or	https://www.fao.org/gsfaonline/foods/details.html?id=256	of compliance with the country of		
e.g. Still grape wine, sparkling and semi-sparkling grape wines, fortified grape wine, grape liquor wine, sweet grape wine, red wine, white wine, rose wine For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw material for compounded alcoholic beverages J2d - Wines other than grape For IMPORTED ALCOHOLIC BEVERAGEs: a) Technical specifications of raw materials and finished products, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or Memorandum Circular No. 13 s. Memor		origin's standards and regulation		
(b) For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials specifications of raw materials and finished product (including specifications of raw materials and finished product (including specifications of raw materials specifications of raw	e.g. Still grape wine, sparkling and semi-sparkling	for alcoholic beverages; c) copy of		
## For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages ### J2d - Wines other than grape Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or For LOCALLY Memorandum Circular No. 13 s. 1989 Memorandum Circular No. 13 s. Memorandum Circular No. 13	grape wines, fortified grape wine, grape liquor wine,	standards and regulation stated in		
MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or Manufacturer/Source/Supplie Circular No. 13 s. Manufacturer/Source/Supplie The product (including Memorandum Circular No. 13 s. Memorandum Circular No. 13 s. Memorandum Circular No. 13 s. Manufacturer/Source/Supplie The product (including and finished	sweet grape wine, red wine, white wine, rose wine	(b)		
BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or BEVERAGE: a) Technical specifications of raw materials and finished product (including and fi		☑ For LOCALLY	<u>Memorandum</u>	Applicant Company/
specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or J2d - Wines other than grape Specifications of raw materials and finished product (including and finished and fi		MANUFACTURED ALCOHOLIC	Circular No. 13 s.	Manufacturer/Source/Supplie
specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or specifications of raw materials and finished product (including specifications of raw materials and finished specifica		BEVERAGE: a) Technical	<u>1989</u>	r
and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or Applicant Company/ Memorandum Circular No. 13 s. 1989 Applicant Company/ Manufacturer/Source/Supplie r		· · · · · · · · · · · · · · · · · · ·		
ethyl alcohol, used as raw material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or ethyl alcohol, used as raw material for compounded alcoholic beverages Image: For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including) Applicant Company/ Manufacturer/Source/Supplie r		and finished product (including		
material for compounded alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or material for compounded alcoholic beverages ✓ For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including		methanol content); b) source of		
alcoholic beverages J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or alcoholic beverages ✓ For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including		ethyl alcohol, used as raw		
J2d - Wines other than grape "Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or □ For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including serior including seri		material for compounded		
"Includes wines made from fruit other than grapes, apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or BEVERAGES: a) Technical specifications of raw materials and finished product (including		alcoholic beverages		
apples and pears, and from other agricultural products, including grain (e.g. rice). These wines may be still or and finished product (including	J2d - Wines other than grape	☑ For IMPORTED ALCOHOLIC	Memorandum	Applicant Company/
including grain (e.g. rice). These wines may be still or and finished product (including	"Includes wines made from fruit other than grapes,	BEVERAGES: a) Technical	Circular No. 13 s.	Manufacturer/Source/Supplie
	apples and pears, and from other agricultural products,	specifications of raw materials	<u>1989</u>	r
	including grain (e.g. rice). These wines may be still or	and finished product (including		
sparkling. Examples include: rice wine (sake), and methanol content); b) a certificate	sparkling. Examples include: rice wine (sake), and	methanol content); b) a certificate		
	sparkling and still fruit wines."	of compliance with the country of		
t origin o standardo and rogalation	(Source URL:	origin's standards and regulation		
https://www.fao.org/gsfaonline/foods/details.html?id=260 for alcoholic beverages; c) copy of	https://www.fao.org/gsfaonline/foods/details.html?id=260	for alcoholic beverages; c) copy of		
standards and regulation stated in		standards and regulation stated in		
e.g. Fruit wine, rice wine (b)	e.g. Fruit wine, rice wine	(b)		
☑ For LOCALLY Memorandum Applicant Company/		☑ For LOCALLY		
MANUFACTURED ALCOHOLIC Circular No. 13 s. Manufacturer/Source/Supplie		MANUFACTURED ALCOHOLIC		Manufacturer/Source/Supplie
BEVERAGE: a) Technical 1989 r		BEVERAGE: a) Technical	<u>1989</u>	r



			PHILIPPINES
	specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages		
J2e - Mead "Alcoholic liquor made from fermented honey, malt and spices, or just of honey. Includes honey wine." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=261) e.g. Honey wine	☑For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including methanol content); b) a certificate of compliance with the country of origin's standards and regulation for alcoholic beverages; c) copy of standards and regulation stated in (b)	Memorandum Circular No. 13 s. 1989	Applicant Company/ Manufacturer/Source/Supplie r
	☑For LOCALLY MANUFACTURED ALCOHOLIC BEVERAGE: a) Technical specifications of raw materials and finished product (including methanol content); b) source of ethyl alcohol, used as raw material for compounded alcoholic beverages	Memorandum Circular No. 13 s. 1989	Applicant Company/ Manufacturer/Source/Supplie r
J2f - Distilled spirituous beverages (>15%alcohol) "Includes all distilled spirituous beverages derived from grain (e.g. corn, barley, rye, wheat), tubers (e.g. potato), fruit (e.g. grapes, berries) or sugar cane that contain greater than 15% alcohol."	☑ For IMPORTED ALCOHOLIC BEVERAGES: a) Technical specifications of raw materials and finished product (including methanol content); b) a certificate of compliance with the country of origin's standards and regulation	Memorandum Circular No. 13 s. 1989	Applicant Company/ Manufacturer/Source/Supplie r



(0)			PHILIPPINES
(Source URL:	for alcoholic beverages; c) copy of		
https://www.fao.org/gsfaonline/foods/details.html?id=262	standards and regulation stated in		
	(b)		
e.g. Brandy, whisky, rhum, tequila, vodka	☑ For LOCALLY	<u>Memorandum</u>	Applicant Company/
	MANUFACTURED ALCOHOLIC	Circular No. 13 s.	Manufacturer/Source/Supplie
	BEVERAGE: a) Technical	<u>1989</u>	r
	specifications of raw materials		
	and finished product (including		
	methanol content); b) source of		
	ethyl alcohol, used as raw		
	material for compounded		
	alcoholic beverages		
J2g - Aromatized alcoholic beverages	☑ For IMPORTED ALCOHOLIC	<u>Memorandum</u>	Applicant Company/
"Includes all non-standardized alcoholic beverage	BEVERAGES: a) Technical	Circular No. 13 s.	Manufacturer/Source/Supplie
products."	specifications of raw materials	<u>1989</u>	r
(Source URL:	and finished product (including		
https://www.fao.org/gsfaonline/foods/details.html?id=263	methanol content); b) a certificate		
	of compliance with the country of		
e.g. Aperitif wine	origin's standards and regulation		
	for alcoholic beverages; c) copy of		
	standards and regulation stated in		
	(b)		
	☑ For LOCALLY	Memorandum	Applicant Company/
	MANUFACTURED ALCOHOLIC	Circular No. 13 s.	Manufacturer/Source/Supplie
	BEVERAGE: a) Technical	1989	r
	specifications of raw materials		
	and finished product (including		
	methanol content); b) source of		
	ethyl alcohol, used as raw		
	material for compounded		
	alcoholic beverages		



			PHILIPPINES
K1 - Snacks - potato - cereal - or starch-based (from roots and tubers, pulses and legumes) "Includes all savoury snacks, with or without added flavourings, but excludes unsweetened crackers. Examples include potato chips, popcorn, pretzels, rice crackers (senbei), flavoured crackers (e.g. cheese-flavoured crackers), bhujia (namkeen; snack made of a mixture of flours, maize, potatoes, salt, dried fruit, peanuts, spices, colours, flavours, and antioxidants), and papads (prepared from soaked rice flour or from black gram or cow pea flour, mixed with salt and spices, and formed into balls or flat cakes)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=265)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
K2 - Chicharon e.g. Pork chicharon, mushroom chicharon	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
K3 - Snacks - fish-based "This describes savoury crackers with fish, fish products or fish flavouring." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=267) e.g. Fish Crackers, dried fish chips	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
MEDIUM RISK FOOD PRODUCTS	☑ ADDITIONAL REQUIREMENT/S	BASIS/ISSUANC E	WHERE TO SECURE
MEDIUM RISK FOOD PRODUCTS - foods that may contain pathogenic micro-organisms but will not normally support their growth because of food characteristics; or food that is unlikely to contain pathogenic micro-organisms because of food type or processing, but may support the formation of toxins or			



the growth of pathogenic micro-organisms. (AO No. 2014-0029)			PHILIPPINES
A1a - Condensed milk (plain) "Condensed milk is obtained by partial removal of water from milk to which sugar may have been added. For evaporated milk, the water removal may be accomplished by heating." "Includes partially dehydrated milk, evaporated milk, sweetened condensed milk, and khoa (cow or buffalo milk concentrated by boiling)." (Source URL:	✓ Valid Certificate of Analysis for % Total Milk Solids and % Milk Fat for EVAPORATED MILK, EVAPORATED WHOLE MILK, EVAPORATED FULL CREAM MILK, UNSWEETENED CONDENSED WHOLE MILK, UNSWEETENED FULL CREAM CONDENSED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
https://www.fao.org/gsfaonline/foods/details.html?id=13)	✓ Valid Certificate of Analysis for % Total Milk Solids and % Milk Fat for SWEETENED CONDENSED MILK, SWEETENED CONDENSED WHOLE MILK, SWEETENED FULL CREAM CONDENSED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
	*The product shall conform with the standards for optional ingredients and additional label declaration for Sweetened Condensed Milk, Sweetened Condensed Whole Milk, Sweetened Full Cream Condensed Milk.		
	✓ Valid Certificate of Analysis for % Milk Solids for EVAPORATED SKIMMED MILK,	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r



		PHILIPPINES
UNSWEETENED CONDENSED SKIMMED MILK		
✓ Valid Certificate of Analysis for% Milk Solids for SWEETENEDCONDENSED SKIMMED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
☑ Valid Certificate of Analysis for %Milk Fat and % Solids-Not-Fat for RECONSTITUTED, RECONSTRUCTED OR RECOMBINED EVAPORATED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the standards for optional ingredients and additional label declaration for Reconstituted, Reconstructed or Recombined Evaporated Milk.		
☑ Valid Certificate of Analysis for % Milk Fat and % Solids-Not-Fat for RECONSTITUTED, RECONSTRUCTED OR RECOMBINED SWEETENED CONDENSED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
✓ Valid Certificate of Analysis for % Milk Solids for RECONSTITUTED, RECONSTRUCTED OR RECOMBINED EVAPORATED SKIMMED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r



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☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
% Non-Fat Milk Solids, Vitamin A	Order No. 132 s.	Manufacturer/Source/Supplie
and Vitamin D (if added) for	<u>1970</u>	r
EVAPORATED FILLED MILK		
*The % Total Oil Content shall be		
declared in the Electronic		
Registration Data Entry.		
**The product shall conform with		
the identity, standards for optional		
ingredients and additional label		
declaration for Evaporated Filled		
Milk.		
☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
· · · · · · · · · · · · · · · · · · ·	Order No. 132 s.	Manufacturer/Source/Supplie
% Non-Fat Milk Solids, Vitamin A		
and Vitamin D (if added) for	<u>1970</u>	
SWEETENED CONDENSED		
FILLED MILK		
*The % Total Oil Content shall be		
declared in the Electronic		
Registration Data Entry.		
**The product shall conform with		
the identity, standards for optional		
ingredients and additional label		
declaration for Sweetened		
Condensed Filled Milk.	ED 4 0: 1 1:	<u> </u>
☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
SWEETENED CONDENSED		r
MILK: Coliforms CFU/g, Yeast &		
Mold Count CFU/g & Aerobic		
Plate Count CFU/g		
riale coulii cru/y		



	☑ Valid Certificate of Analysis for Microbiological parameters for LIQUID MILK (EVAPORATED): Commercial Sterility	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A1b - Beverage whiteners "Milk or cream substitute consisting of a vegetable fatwater emulsion in water with milk protein and lactose or vegetable proteins for use in beverages such as coffee and tea. Also includes the same type of products in powdered form." "Includes condensed milk analogues, blends of evaporated skimmed milk and vegetable fat and blends of sweetened condensed skimmed milk and vegetable fat." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=14) e.g. Condensed creamer	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A2 - Milk powder and cream powder and powder analogues (plain) "Includes plain milk powders, cream powders, or combination of the two, and their analogues. Includes products based on skim, part-skim, low-fat and whole milk." (Source URL:	☑ Valid Certificate of Analysis for % Milk Solids, % Milk Fat and % Water for WHOLE MILK POWDER (DRIED FULL CREAM MILK, FULL CREAM MILK POWDER, DRY WHOLE MILK, MILK POWDER, DRIED MILK)	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
https://www.fao.org/gsfaonline/foods/details.html?id=20) "Milk cream powder analogues are products based on a fat-water emulsion and dried for use other than as a beverage whitener. Examples include imitation dry cream mix and blends of skimmed milk and vegetable fat in powdered form."	✓ Valid Certificate of Analysis for % Solids, % Fat and % Water for SKIMMED MILK POWDER ✓ Valid Certificate of Analysis for % Milk Solids, % Milk Fat and % Water for PARTLY SKIMMED MILK POWDER	Administrative Order No. 132 s. 1970 Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
(Source URL:	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
https://www.fao.org/gsfaonline/foods/details.html?id=22)	% Milk Fat and Moisture Content	Order No. 132 s.	Manufacturer/Source/Supplie
	for MALTED MILK POWDER	<u>1970</u>	
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	% Butterfat, % Total Milk Solids	Order No. 132 s.	Manufacturer/Source/Supplie
	and Moisture Content for	1970	r
	BUTTERMILK POWDER (DRIED		
	BUTTERMILK)		
	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	MILK POWDER (e.g. WHOLE,		r
	NONFAT, FILLED MILK,		
	BUTTERMILK, WHEY & WHEY		
	PROTEIN AND MILK INTENDED FOR CHILDREN MORE THAN		
	36 MONTHS OF AGE AND		
	ADULTS): Salmonella/25g		
A3 - Milk products for specific age groups or target	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
population	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
e.g. Powdered milk for children above 3 years and	MILK POWDER (e.g. WHOLE,		r
pregnant women	NONFAT, FILLED MILK,		
	BUTTERMILK, WHEY & WHEY		
	PROTEIN AND MILK INTENDED		
	FOR CHILDREN MORE THAN		
	36 MONTHS OF AGE AND ADULTS): Salmonella/25g		
	,	Administrative	Applicant Company/
	☑ Valid Certificate of Analysis to support Nutrition Information	Order No. 2014-	Manufacturer/Source/Supplie
	declaration on the label	0029	r
		Administrative	
			•



			PHILIPPINES
		Order No. 2014- 0030	
B1 - Non-Dairy based frozen desserts "Includes fat-based counterparts of dairy-based desserts. Includes ready-to-eat products and their mixes. Also includes non-dairy fillings for desserts. An example is an ice cream-like product made with vegetable fats." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=51)	✓ Valid Certificate of Analysis for Microbiological parameters for ICE CREAM & SHERBET (PLAIN AND FLAVORED): Coliforms CFU/g, Listeria monocytogenes/25g, Salmonella/25g, Aerobic Plate Count CFU/g & S. aureus CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	✓ Valid Certificate of Analysis for Microbiological parameters for ICE CREAM WITH ADDED INGREDIENTS (NUTS, FRUITS, COCOA ETC.): Coliforms CFU/g, S. aureus CFU/g, Salmonella/25g, Aerobic Plate Count CFU/g & Listeria monocytogenes/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
B2 - Edible ices - popsicles "This category includes water-based frozen desserts, confections and novelties, such as "Italian"-style ice, and flavoured ice." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=52) e.g. Ice candy, ice popsicles	☑ Valid Certificate of Analysis for Microbiological parameters for FLAVORED ICE: Aerobic Plate Count CFU/g, Coliforms MPN/g or CFU/g or /25g, Yeast and Mold Count CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
C1 - Tomato products e.g. Tomato Catsup, tomato sauce, tomato paste	☑ Valid Certificate of Analysis for Total Soluble Solids, Specific Gravity, Total Acidity in terms of acetic acid, Arsenic Content, Lead Content, Copper Content, Zinc Content and Tin Content for TOMATO CATSUP	Administrative Order No. 233 s. 1974	Applicant Company/ Manufacturer/Source/Supplie r



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C2 - Frozen fruits "Fruit that may or may not be blanched prior to freezing. The product may be frozen in a juice or sugar syrup. Examples include frozen fruit salad and frozen strawberries." (Source URL:	*The product shall conform with the identity and standard of quality of Tomato Catsup. ☑ Valid Certificate of Analysis for Microbiological parameters for FROZEN FRUITS (pH >4.5): E. coli CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
C3 - Canned or bottled (pasteurized) or retort pouch fruit and vegetable preserve in juice, syrup, brine "Fully preserved product in which fresh fruit is cleaned and placed in cans or jars with natural juice or sugar syrup (including artificially sweetened syrup) and heat-sterilized or pasteurized. Includes products processed in retort pouches. Examples include: canned fruit salad, and applesauce in jars." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=63) "Fully preserved product in which fresh vegetables are cleaned, blanched, and placed in cans or jars in liquid (e.g. brine, water, oil or sauce), and heat-sterilized or pasteurized. Examples include: canned chestnuts, canned chestnut puree, asparagus packed in glass jars, canned and cooked pink beans, canned tomato paste (low acid), and canned tomatoes (pieces, wedges or whole)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=81)	✓ Valid Certificate of Analysis for Microbiological parameters for FRUITS AND VEGETABLE PRODUCTS IN HERMETICALLY SEALED CONTAINERS: Commercial Sterility	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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☑ Valid Certificate of Analysis for		Applicant Company/
Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
FERMENTED VEGETABLE		r
(READY TO EAT) (e.g. KIMCHI):		
Yeast and Mold Count CFU/g,		
Coliforms MPN/g or CFU/g or		
/25g, E. coli MPN/g or CFU/g or		
/25g, Salmonella/25g & S. aureus		
cfu/g		
	☑ Valid Certificate of Analysis for Microbiological parameters for FERMENTED VEGETABLE (READY TO EAT) (e.g. KIMCHI): Yeast and Mold Count CFU/g, Coliforms MPN/g or CFU/g or /25g, E. coli MPN/g or CFU/g or /25g, Salmonella/25g & S. aureus	NOT APPLICABLE PLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE NOT APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE STATE OF APPLICABLE NOT APPLICABLE STATE OF APPL



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Chinese cabbage and vegetable preparation), and sauerkraut (fermented cabbage)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=84)			
C8 - Vegetable protein products (canned and frozen)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
D - Cocoa products and chocolate products "Cocoa Mixes (powders) and cocoa mass/cake: Includes a variety of products that are used in the manufacture of other chocolate products or in the preparation of cocoa-based beverages." (Source URL:	☑ Valid Certificate of Analysis for Microbiological parameters for COCOA POWDER: Molds CFU/g, Salmonella/25g, Coliforms, MPN/g or CFU/g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
Cocoa Mixes (syrups) "Products that may be produced by adding a bacterial amylase to cocoa liquor. The enzyme prevents the syrup from thickening or setting by solubilizing and dextrinizing cocoa starch. Includes products such as chocolate syrup used to prepare chocolate milk or hot chocolate." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=89) Cocoa-based spreads, including fillings "Products in which cocoa is mixed with other ingredients (usually fat-based) to prepare a spreadable paste that is used as a spread for bread or as a filling for fine bakery wares. Examples include: cocoa butter, fillings for bonbons and chocolates, chocolate pie filling, and nut-chocolate based spreads for bread (Nutella-type product)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=90)	✓ Valid Certificate of Analysis for Microbiological parameters for CHOCOLATE PRODUCTS: Molds CFU/g, Salmonella/25g, Coliforms MPN/g or CFU/g & Aerobic Plate Count CFU/g. ✓ Valid Certificate of Analysis for Microbiological parameters for CHOCOLATE CONFECTIONARIES: Molds CFU/g, Osmophilic yeast CFU/g, Salmonella/25g, Coliforms MPN/g or CFU/g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



Cocoa and Chocolate Products Chocolate

"Chocolate is produced from cocoa nibs, mass, press cake, powder, or liquor with or without addition of sugar, cocoa butter, aroma or flavouring substances, and optional ingredients (e.g. nuts). This category is for chocolate as defined in the Standard for Chocolate and Chocolate Products (CODEX STAN 87-1981) and for confectionery that meet the standard and may contain other contain other ingredients, for example chocolatecovered nuts and fruit (e.g. raisins). This category includes only the chocolate portion of any confectionery within the scope of food category 05.2. Examples include: bonbons, cocoa butter confectionery (composed of cocoa butter, milk solids and sugar), white chocolate, chocolate chips (e.g. for baking), milk chocolate, cream chocolate, sweet chocolate, bitter chocolate, enrobing chocolate, chocolate covered in a sugar-based "shell" or with coloured decorations, filled chocolate (chocolate with a texturally distinct center and external coating, and chocolate with added edible ingredients." (Source URL:

https://www.fao.org/gsfaonline/foods/details.html?id=91)

Imitation Chocolate, Chocolate substitute products
"Includes chocolate-like products that may or may not be
cocoa-based, but have similar organoleptic properties as
chocolate, such as carob chips, and cocoa-based
products that contain greater than 5% vegetable fat.
These chocolate-like products may contain additional
optional ingredients and may include filled confectionery.
Examples include: compound chocolate, flavoured and
coloured compound chocolate, compound chocolate



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coatings, and imitation chocolate covered nuts and fruit (e.g. raisins)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=92)			
E1 - Fermented soybeans "The product is prepared from soybeans that have been steamed and fermented with certain fungi or bacteria (starter). The soft, whole beans have a distinctive aroma and taste. It includes products such as dou chi (China), natto (Japan), and tempe (Indonesia)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=277)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
E2 - Fermented soybean curd "The product is prepared by forming soybean curd into a loaf during the fermentation process. It is a soft, flavoured product, either in red, rice-yellow, or grey-green." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=278)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
F1ai - Cured (including salted) non-heat treated processed meat, poultry and game products in whole pieces or cuts "Salted products are treated with sodium chloride. Dry cured (dry pickled) products are prepared by rubbing salt directly on the meat surface. Wet pickle cured products are prepared by submerging the meat in a brine solution. Pump cured products are prepared by	☑ Valid Certificate of Analysis for Microbiological parameters for PACKAGED COOKED, CURED/SALTED MEAT (HAM, BACON): S. aureus CFU/g, Salmonella/25g & Listeria monocytogenes/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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injecting brine into the meat. Curing may also be achieved by addition of additives. Smoked products are also included here. Examples include: bacon (cured, dry-cured, immersion-cured, pump-cured); side bacon;	☑ Valid Certificate of Analysis for Microbiological parameters for CURED/SMOKED POULTRY: S. aureus CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
corned beef; marinaded beef; and different types of Oriental pickled products: miso-pickled meat (miso-zuke), koji-pickled meat (koji-zuke), and soy sauce-pickled meat (shoyu-zuke)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=133	☑ Valid Certificate of Analysis for Nitrate and/or Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	Applicant Company/ Manufacturer/Source/Supplie r
F1aii - Cured (including salted) and dried non-heat treated processed meat, poultry and game products in whole pieces or cuts "The meat cuts may be cured or salted and then dried, or they may only be dried. Drying is achieved either in hot air or in vacuum. Examples include: dried salt pork, dehydrated meat, stuffed loin, Iberian ham, and	☑ Valid Certificate of Analysis for Microbiological parameters for PACKAGED COOKED, CURED/SALTED MEAT (HAM, BACON): S. aureus CFU/g, Salmonella/25g & Listeria monocytogenes/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
proscuitto-type ham." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=134)	☑ Valid Certificate of Analysis for Microbiological parameters for CURED/SMOKED POULTRY: S. aureus CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for Nitrate and/or Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	Applicant Company/ Manufacturer/Source/Supplie r
F1aiii - Fermented non-heat treated processed meat, poultry and game products - processed meat in whole pieces or cuts "Fermented products are a type of pickled product produced by the action of lactic acid bacteria in the	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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presence of salt. Examples include: potted beef and pickled (fermented) pig's feet." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=135)			
F2ai - Cured (including salted) non-heat treated processed comminuted meat, poultry and game products "Salted products are treated with sodium chloride. Dry cured (dry pickled) products are prepared by rubbing salt directly on the meat surface. Wet pickle cured products are prepared by submerging the meat in a brine solution. Pump cured products are prepared by injecting brine into the meat. Curing may also be achieved by addition of additives. Smoked products are also included here. Examples include: bacon (cured, dry-cured, immersion-cured, pump-cured); side bacon; corned beef; marinaded beef; and different types of Oriental pickled products: miso-pickled meat (miso-zuke), koji-pickled meat (koji-zuke), and soy sauce-pickled meat (shoyu-zuke)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=133) e.g. chorizos (spicy pork sausages), salami-type	✓ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM POULTRY MEAT INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g, Salmonellas/25g ✓ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM SPECIES OTHER THAN POULTRY INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g, Salmonellas/25g Aerobic Plate Count, CFU/g, Salmonellas/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
products, salchichon, tocino (fresh, cured sausage), pepperoni, and smoked sausage.	☑ Valid Certificate of Analysis for Nitrate and/or Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	



F2aii - Cured (including salted) and dried non-heat treated processed comminuted meat, poultry and game products (jerky, shredded beef/pork) "The comminuted or mechanically deboned products may be cured or salted as described for category 08.3.1.1, and then dried, or they may only be dried. Drying is achieved either in hot air or in vacuum. Examples include: pasturmas, dried sausages, cured and dried sausages, beef jerky, Chinese sausages (including traditional cured or smoked pork sausage), and sobrasada." (Source URL; https://www.fao.org/gsfaonline/foods/details.html?id=141)	☑ Valid Certificate of Analysis for Microbiological parameters for DRIED ANIMAL PRODUCTS: S. aureus CFU/g, Clostridium perfringens CFU/g and Salmonella/25 ☑ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM POULTRY MEAT INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g, Salmonellas/25g ☑ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM SPECIES OTHER THAN POULTRY INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g, Salmonellas/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	Valid Certificate of Analysis for Nitrate and/or Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	



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F2aiii - Fermented non-heat treated processed	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
comminuted meat, poultry and game products	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
"Fermented products are a type of pickled product	FERMENTED, COMMINUTED		r
produced by the action of lactic acid bacteria in the	MEAT, NOT COOKED (DRY &		
presence of salt. Certain types of sausages may be	SEMI-DRY FERMENTED		
fermented."	SAUSAGES): E. coli MPN/g, S.		
(Source URL:	aureus CFU/g &		
https://www.fao.org/gsfaonline/foods/details.html?id=142	Salmonella/25gCFU/g &		
)	Salmonella/25g		
e.g., pre-grilled beef patties; foie gras and pates; brawn	Valid Certificate of Analysis for	Administrative	Applicant Company/
and head cheese; cooked, cured chopped meat;	Nitrate and/or Nitrite Content (if	Order No. 154 s.	Manufacturer/Source/Supplie
chopped meat boiled in soy sauce (tsukudani); canned	utilized)	1971 and Bureau	r
corned beef; luncheon meats; meat pastes; cooked		Circular No. 2006-	
meat patties; cooked salami-type products; cooked		<u>016</u>	
meatballs; saucises de strasbourg; breakfast sausages;			
brown-and-serve sausages; and terrines (a cooked			
chopped meat mixture).		EDA O: I N	A 1: 10 /
H1a - Smoked, dried, fermented, and/or salted fish	✓ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
and fish products, including molluscs, crustaceans	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
and echinoderms	ETHNIC FOOD PRODUCTS -		r
"Smoked fish are usually prepared from fresh deep	DRIED, SALTED FISH: Aerobic		
frozen or frozen fish that are dried directly or after	Plate Count CFU/g, Yeast and		
boiling, with or without salting, by exposing the fish to	Mold Count CFU/g, Coliforms		
freshly-generated sawdust smoke. Dried fish are prepared by exposing the fish to sunlight or drying	MPN/g, E. coli MPN/g and S.		
directly or after boiling in a special installation; the fish	aureus MPN/g		
may be salted prior to drying. Salted fish are either			
rubbed with salt or placed in a salt solution. This	☑ Valid Certificate of Analysis for		
manufacturing process is different from that described in	Microbiological parameters for		
food category 09.3 for marinated and pickled fish. Cured	SMOKED FISH: Aerobic Plate		
fish is prepared by salting and then smoking fish.	Count CFU/g, Salmonella/25g, E.		
Examples include: salted anchovies, shrimp, and shad;	coli MPN/g and S. aureus CFU/g		
smoked chub, cuttlefish and octopus; fish ham; dried			
and a contract and a compact, non-right, arrow	1	l	



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and salted species of the Gadidae species; smoked or salted fish paste and fish roe; cured and smoked sablefish, shad, and salmon; dried shellfish, dried bonito (katsuobushi), and boiled, dried fish (niboshi)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=158	☑ Valid Certificate of Analysis for Microbiological parameters for SALT FERMENTED FISH AND SHRIMPS (BAGOONG): Aerobic Plate Count CFU/g and Coliforms CFU/g		
H2a - Fish and fish products, includings molluscs, crustaceans and echinoderms - marinated and/or in jelly "Marinated products are manufactured by soaking the fish in vinegar or wine with or without added salt and spices. They are packaged in jars or cans and have a limited shelf life. Products in jelly may be manufactured by tenderizing fish products by cooking or steaming, adding vinegar or wine, salt and preservatives, and solidifying in a jelly. Examples include: "rollmops" (a type of marinated herring), sea eel (dogfish) in jelly and fish aspic." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=160)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
H2b - Fish and fish products, includings molluscs, crustaceans and echinoderms - pickled and/or in MH2brine "Pickled products are sometimes considered a type of marinaded product. Pickling results from the treatment of the fish with with a salt and vinegar or alcohol (e.g., wine) solution. Examples include: different types of Oriental pickled products: koji-pickled fish (koji-zuke), lees-pickled fish (kasu-zuke), miso-pickled fish (miso-zuke), soy sauce-pickled fish (shoyu-zuke), and vinegar-	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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pickled fish (su-zuke); pickled whale meat; and pickled herring and sprat." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=161)			
H2c - Salmon substitutes, caviar and other fish roe products "Roe is usually produced by washing, salting and allowing to ripen until transparent. The roe is then packaged in glass or other suitable containers. The term "caviar" refers only to the roe of the sturgeon species (e.g. beluga). Caviar substitues are made of roe of various sea and freshwater fish (e.g., cod and herring) that are salted, spiced, dyed and may be treated with a preservative. Examples include: salted salmon roe (sujiko), processed, salted salmon roe (ikura), cod roe, salted cod roe (tarako) and lumpfish caviar." Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=162)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
H2d - Semi-preserved fish and fish products, including molluscs, crustaceans and echinoderms, excluding products under MR Letter H.1 a to c. "Examples include fish or crustacean pates and traditional Oriental fish paste. The latter is produced from fresh fish or the residue from fish sauce production, which is combined with other ingredients such as wheat flour, bran, rice or soybeans. The product may be further fermented." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=163)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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e.g. fish or crustacean pates and traditional Oriental fish paste			
I1 - Preserved eggs, including alkaline, salted and canned eggs (salted eggs, century eggs) "Includes traditional Oriental preserved products, such as salt-cured duck eggs (Hueidan), and alkaline treated "thousand-year-old-eggs" (pidan)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=171) e.g. salt-cured duck eggs (Hueidan), and alkaline treated "thousand-year-old-eggs" (pidan)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
I2 - Egg-based desserts "Includes ready-to-eat products and products to be prepared from a dry mix. Examples include: flan and egg custard. Also includes custard fillings for fine bakery wares (e.g. pies)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=172	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
Ja - Cakes, cookies, pies pastries, doughnuts, sweet rolls, scones, muffins, waffles - plain/without filling e.g. pancakes, waffles, filled sweet buns (anpan), Danish pastry, wafers or cones for ice cream, flour confectionery, and trifles	✓ Valid Certificate of Analysis for Microbiological parameters for BAKED GOODS: Yeast CFU/g, Mold CFU/g, Aerobic Plate Count, CFU/g, Coliforms CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
Jb - Frozen dough	☑ Valid Certificate of Analysis for Microbiological parameters for FROZEN AND REFRIGERATED DOUGHS: Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
K1 - Soups and broths "Water- or milk-based products consisting of vegetable,	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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meat or fish broth with or without other ingredients (e.g. vegetables, meat, noodles). Examples include: bouillon, broths, consommés, water- and cream-based soups, chowders, and bisques." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=197)			
K2a - Emulsified sauces and dips "Sauces, gravies, dressings, and dips based, at least in part, on a fat- or oil-in water emulsion. Examples include: salad dressing (e.g., French, Italian, Greek, ranch style), fat-based sandwich spreads (e.g., mayonnaise with mustard), salad cream, fatty sauces and snack dips (e.g., bacon and cheddar dip, onion dip)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=200)	 ☑ In the Electronic Registration Data Entry – under Complete List of Ingredients, declare the % by weight of edible vegetable oil content of the finished product for MAYONNAISE ☑ Valid Certificate of Analysis for calcium disodium EDTA (calcium disodium ethylenediaminetetraacetate) or disodium EDTA (disodium ethylenediaminetetraacetate) content, IF ADDED in MAYONNAISE *The product shall conform with the identity, standards for optional ingredients and additional label declaration for MAYONNAISE. ☑ Valid Certificate of Analysis for Microbiological parameters for EMULSIFIED SAUCE PH ≤ 4.6 	Administrative Order No. 235 s. 1975 FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r Applicant Company/ Manufacturer/Source/Supplie
	(E.G. MAYONNAISE, THOUSAND ISLAND, RANCH,		



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K2b - Non-emulsified sauces (ketchup, cheese sauce, cream sauce, brown gravy) "Include water-, coconut milk-, and milk-based sauces, gravies and dressings. Examples include: barbecue sauce, tomato ketchup, cheese sauce, Worcestershire sauce, Oriental thick Worcestershire sauce (tonkatsu sauce), chili sauce, sweet and sour dipping sauce, and white (cream-based) sauce (sauce consisting primarily of milk or cream, with little added fat (e.g. butter) and flour, with or without seasoning or spices)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=201	FRENCH): Aerobic Plate Count CFU/g, Yeast and Mold Count CFU/g, Salmonella/25g & Listeria monocytogenes/25g ☑ Valid Certificate of Analysis for Microbiological parameters for SALADS AND SANDWICH SPREADS (excluding cocoa milk based sadwich spreads): Aerobic Plate Count CFU/g, Salmonella/25g & Listeria monocytogenes/25g ☑ Valid Certificate of Analysis for Total Solids; Titratable Acidity (as acetic acid); pH for BANANA SAUCE/BANANA CATSUP *The product shall conform with the standards for the identity, essential composition, quality factors and label declaration for BANANA SAUCE/BANANA CATSUP.	Administrative Order No. 123-As. 1985	Applicant Company/ Manufacturer/Source/Supplie r
K3 - Salads (e.g. macaroni salad, potato salad) and sandwich spreads excluding cocoa- and nut-based spreads under HR Letter B.8 (peanut butter) and MR D.1.c (cocoa-based spreads) "Includes prepared salads, milk-based sandwich	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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spreads, non-standardized mayonnaise-like sandwich spreads, and dressing for coleslaw (cabbage salad)" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=204)			
L1a - Fruit and vegetable juices - (fruit juice, vegetable juice, concentrates for fruit juice, concentrates for vegetable juice) FRUIT JUICE "Fruit juice is the unfermented but fermentable liquid obtained from the edible part of sound, appropriately mature and fresh fruit or of fruit maintained in sound condition by suitable means. The juice is prepared by	☑ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
suitable processes, which maintain the essential physical, chemical, organoleptical and nutritional characteristics of the juices of the fruit from which it comes. The juice may be cloudy or clear, and may have restored (to the normal level attained in the same kind of fruit) aromatic substances and volatile flavour	☑ Valid Certificate of Analysis for Microbiological parameters for FROZEN JUICE CONCENTRATES: Aerobic Plate Count CFU/ml & Yeast and Mold Count CFU/ml	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
components, all of which must be obtained by suitable physical means, and all of which must have been recovered from the same kind of fruit. Pulp and cells obtained by suitable physical means from the same kind of fruit may be added. A single juice is obtained from one kind of fruit. A mixed juice is obtained by blending	✓ Valid Certificate of Analysis for Microbiological parameters for JUICES IN HERMETICALLY SEALED CONTAINERS (TETRA PACK ETC.): Commercial Sterility	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
two or more juices or juices and purees, from different kinds of fruit. Fruit juice may be obtained, e.g. by directly expressing the juice by mechanical extraction processes, by reconstituting concentrated fruit juice	☑ Valid Certificate for Microbiological parameters for POWDERED BEVERAGES (e.g. ICED TEA, POWDERED JUICES/MIXES): Aerobic Plate	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



(food category 14.1.2.3) with water, or in limited
situations by water extraction of the whole fruit (e.g.,
prune juice from dried prunes). Examples include:
orange juice, apple juice, black currant juice, lemon
juice, orange-mango juice, and coconut water."
(Source URL:

https://www.fao.org/gsfaonline/foods/details.html?id=239, you may also refer to AO No. 90-A s. 1980)

VEGETABLE JUICE

"Vegetable juice is the liquid unfermented but fermentable product intended for direct consumption obtained by mechanical expression, crushing, grinding, and/or sieving of one or more sound fresh vegetables or vegetables preserved exclusively by physical means. The juice may be clear, turbid, or pulpy. It may have been concentrated and reconstituted with water. Products may be based on a single vegetable (e.g. carrot) or blends of vegetables (e.g. carrots, celery)." (Source URL:

https://www.fao.org/gsfaonline/foods/details.html?id=240

CONCENTRATES FOR FRUIT JUICE

"It is prepared by the physical removal of water from fruit juice in an amount to increase the Brix level to a value at least 50% greater than that established for reconstituted juice from the same fruit. In the production of juice that is to be concentrated, suitable processes are used, and may be combined, with simultaneous diffusion of the pulp cells or fruit pulp by water, provided that the water-extracted soluble fruit solids are added in-line to the primary juice, before the concentration procedure. Fruit

	Count CFU/g & Coliforms CFU/g		
<u> </u>	✓ Valid Certificate of Analysis for Microbiological parameters for FRUIT BEVERAGE PRODUCTS: Aerobic Plate Count CFU/ml, Yeast and Mold Count CFU/ml, Coliforms CFU/ml & E.coli CFU/ml.	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
<u>)</u>			
t t			

Count CFU/g, Yeast and Mold



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juice concentrates may have restored (to the normal level attained in the same kind of fruit) aromatic substances and volatile flavour components, all of which must be obtained by suitable physical means, and all of which must be recovered from the same kind of fruit. Pulp and cells obtained by suitable physical means from the same kind of fruit may be added. Sold in liquid, syrup and frozen forms for the preparation of a ready-to-drink juice by addition of water. Examples include: frozen orange juice concentrate, and lemon juice concentrate." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=241)			
CONCENTRATES FOR VEGETABLE JUICE "Prepared by the physical removal of water from vegetable juice. Sold in liquid, syrup and frozen forms for the preparation of a ready-to-drink juice by addition of water. Includes carrot juice concentrate." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=242)			
L1b - Fruit and vegetable nectars (fruit nectar, vegetable nectar, concentrates for fruit nectar, concentrates for vegetable nectar) FRUIT NECTAR "Fruit nectar is the unfermented but fermentable product obtained by adding water with or without the addition of sugar, honey, syrups, and/or sweeteners to fruit juice, concentrated fruit juice, fruit purees or concentrated fruit	☑ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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purees, or a mixture of those products. Aromatic	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
substances, volatile flavour components, pulp and cells,	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
all of which must have been recovered from the same	FROZEN JUICE		r
kind of fruit and obtained by suitable physical means,	CONCENTRATES : Aerobic Plate		
may be added. Products may be based on a single fruit	Count CFU/ml & Yeast and Mold		
or on fruit blends. Examples include: pear nectar and	Count CFU/ml		
peach nectar."	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
(Source URL:	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
https://www.fao.org/gsfaonline/foods/details.html?id=244	JUICES IN HERMETICALLY		r
)	SEALED CONTAINERS (TETRA		
	PACK ETC.): Commercial Sterility		
VEGETABLE NECTAR	☑ Valid Certificate for	FDA Circular No.	Applicant Company/
"Product obtained by adding water with or without the	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
addition of sugar, honey, syrups, and/or sweeteners to	POWDERED BEVERAGES (e.g.		r
vegetable juice or concentrated vegetable juice, or a	ICED TEA, POWDERED		
mixture of those products. Products may be based on a	JUICES/MIXES): Aerobic Plate		
single vegetable or on a blend of vegetables."	Count CFU/g, Yeast and Mold		
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=245	Count CFU/g & Coliforms CFU/g		
nttps://www.rao.org/gsraomine/roods/details.ntmi?id=245	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
/	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
CONCENTRATES FOR FRUIT NECTAR	FRUIT BEVERAGE PRODUCTS:		r ···
"Prepared by the physical removal of water from fruit	Aerobic Plate Count CFU/ml,		
nectar or its starting materials. Sold in liquid, syrup and	Yeast and Mold Count CFU/ml,		
frozen forms for the preparation of a ready-to-drink	Coliforms CFU/ml & E.coli		
nectar by addition of water. Examples: pear nectar	CFU/ml.		
concentrate and peach nectar concentrate."			
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=246			
)			
/			

CONCENTRATES FOR VEGETABLE NECTAR



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"Prepared by the physical removal of water from vegetable nectar. Sold in liquid, syrup and frozen forms for the preparation of ready-to-drink nectars by addition of water." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=247)			
L1c - "Sport," "energy", or "electrolyte drinks" "Includes so-called "energy" drinks that are carbonated and contain high levels of nutrients and other ingredients (e.g. caffeine, taurine, carnitine)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=249)	☑ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	Valid Certificate of Analysis for Caffeine and Vitamin B and/or mineral/s (whichever is applicable) content	Administrative No. Order 2014-0029	Applicant Company/ Manufacturer/Source/Supplie r
	Label bearing the Precaution Statement: "Excessive intake of caffeine may cause sleeplessness, palpitation and other similar side effects. Not recommended for children, pregnant and lactating women, people who may have heart problems and/or those sensitive to caffeine."	Administrative Order No. 2014- 0030	Applicant Company/ Manufacturer/Source/Supplie r



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L1ci - Carbonated water-based flavored drinks "Includes water-based flavored drinks with added carbon dioxide with nutritive, non-nutritive and/or intense sweeteners and other permitted food additives. Includes gaseosa (water-based drinks with added carbon dioxide, sweetener, and flavour), and sodas such as colas, pepper-types, root beer, lemon-lime, and citrus types, both diet/light and regular types. These beverages may be clear, cloudy, or may contain particulated matter (e.g. fruit pieces). Includes so-called "energy" drinks that are carbonated and contain high levels of nutrients and other ingredients (e.g. caffeine, taurine, carnitine).	☑ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL Valid Certificate of Analysis for Caffeine Content for COLA-TYPE BEVERAGE	Administrative Order 88-A s.	Applicant Company/ Manufacturer/Source/Supplie r Applicant Company/ Manufacturer/Source/Supplie
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=249) e.g. colas, pepper-types, root beer, lemon-lime, and citrus types, both diet/light and regular types)		1984	r
L1cii - Non-carbonated water-based flavored drinks "Include water-based flavoured drinks without added carbon dioxided, fruit and vegetable juice-based drinks (e.g. almond, aniseed, coconut-based drinks, and ginseng drink), fruit flavoured ades (e.g. lemonade, orangeade), squashes (citrus-based soft drinks), capile groselha, lactic acid beverage, ready-to-drink coffee and tea drinks with or without milk or milk solids, and herbal- based drinks (e.g. iced tea, fruit-flavoured iced tea, chilled canned cappucino drinks) and "sports" drinks containing electrolytes. These beverages may be clear or contain particulated matter (e.g. fruit pieces), and may be unsweetened or sweetened with sugar or a non- nutritive high-intensity sweetener. Includes so-called "energy" drinks that are non-carbonated and contain	✓ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL ✓ Valid Certificate of Analysis for Microbiological parameters for CHILLED YOUNG COCONUT WATER (BUKO JUICE): Aerobic	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



high levels of nutrients and other ingredients (e.g. caffeine, taurine, carnitine)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=250)	Plate Count CFU/mL, Yeast and Mold Count CFU/mL and Coliforms CFU/mL		
L1ciii - Concentrates (liquid or solid) for water-based flavored drinks "Include powder, syrup, liquid and frozen concentrates for the preparation of carbonated or non-carbonated water-based non-alcoholic beverages by addition of water or carbonated water. Examples include: fountain syrups (e.g. cola syrup), fruit syrups for soft drinks, frozen or powdered concentrate for lemonade and iced tea mixes." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=251)	 ☑ Valid Certificate of Analysis for Microbiological parameters for FROZEN JUICE CONCENTRATES: Aerobic Plate Count CFU/ml & Yeast and Mold Count CFU/ml ☑ Valid Certificate for Microbiological parameters for POWDERED BEVERAGES (e.g. ICED TEA, POWDERED JUICES/MIXES): Aerobic Plate Count CFU/g, Yeast and Mold Count CFU/g & Coliforms CFU/g 	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
L1d - Powdered cocoa drink mixes (cocoa) "Examples include: drinking chocolate powder; breakfast cocoa; cocoa dust (fines), nibs, mass, press cake; chocolate liquor; cocoa mixes (powders for preparing the hot beverage); cocoa-sugar mixture; and dry mixes for sugar-cocoa confectionery." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=88)	✓ Valid Certificate for Microbiological parameters for POWDERED BEVERAGES (e.g. ICED TEA, POWDERED JUICES/MIXES): Aerobic Plate Count CFU/g, Yeast and Mold Count CFU/g & Coliforms CFU/g ✓ Valid Certificate of Analysis for Microbiological parameters for COCOA POWDER: Molds CFU/g, Salmonella/25g, Coliforms,	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



	MPN/g or CFU/g & Aerobic Plate Count CFU/g		PHILIPPINES
M1 - Vitamins and minerals as Food Supplement "Includes vitamin and mineral supplements in unit dose forms such as capsules, tablets, powders, solutions etc., where national jurisdictions regulate these products as food" (Source URL:	☑ Valid Shelf life study with stability data containing relevant information on the critical parameters of the finished product, period conducted and conclusion	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
"means a processed food product intended to supplement the diet that bears or contains one or more of the following dietary ingredients: vitamin, mineral, herb, or other botanical, amino acid, and dietary substance to increase the total daily intake in amounts conforming to the latest Philippine recommended energy and nutrient intakes or internationally agreed minimum	✓ Valid Certificate of Analysis of the physico-chemical (Vitamins or Minerals or Amino Acids Assays) and/or microbiological parameters of the finished product (whichever is applicable) *The amount of Vitamins shall conform with the prescribed level of Office Order No. 22 s 1991	Administrative Order No. 2014- 0029 Office Order No. 22 s 1991 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
daily requirements. It usually is in the form of capsules, tablets, liquids, gels, powders or pills and not represented for use as a conventional food or as the sole item of a meal or diet or replacement of drugs and medicines." (Source URL:	☑ Clear and complete loose labels or artworks declaring the term "Food Supplement" and the phrase "NO APPROVED THERAPEUTIC CLAIMS" based on	Bureau Circular No. 2 s 1999	Applicant Company/ Manufacturer/Source/Supplie r
https://www.officialgazette.gov.ph/2009/08/18/republic-act-no-9711/) e.g. Vitamin C + Zinc Food Supplement Capsule	☑ Sample in actual commercial presentation *for the procedure on submission, please refer to: IV. Guidelines, C. Procedural Guidelines 2. L. ii., Pages 7-8 of FDA Circular No. 2020-033	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r



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M2 - Amino acids as Food Supplement "Includes vitamin and mineral supplements in unit dose forms such as capsules, tablets, powders, solutions etc., where national jurisdictions regulate these products as food" (Source URL:	☑ Valid Shelf life study with stability data containing relevant information on the critical parameters of the finished product, period conducted and conclusion	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
"means a processed food product intended to supplement the diet that bears or contains one or more of the following dietary ingredients: vitamin, mineral, herb, or other botanical, amino acid, and dietary substance to increase the total daily intake in amounts conforming to the latest Philippine recommended energy and nutrient intakes or internationally agreed minimum	✓ Valid Certificate of Analysis of the physico-chemical (Vitamins or Minerals or Amino Acids Assays) and/or microbiological parameters of the finished product (whichever is applicable) *The amount of Vitamins shall conform with the prescribed level of Office Order No. 22 s 1991	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
daily requirements. It usually is in the form of capsules, tablets, liquids, gels, powders or pills and not represented for use as a conventional food or as the sole item of a meal or diet or replacement of drugs and medicines." (Source URL:	☑ Clear and complete loose labels or artworks declaring the term "Food Supplement" and the phrase "NO APPROVED THERAPEUTIC CLAIMS" based on	Bureau Circular No. 2 s 1999	Applicant Company/ Manufacturer/Source/Supplie r
https://www.officialgazette.gov.ph/2009/08/18/republic-act-no-9711/) e.g. Branched-Chain Amino Acids (BCAA) Food Supplement Powder	☑ Sample in actual commercial presentation *for the procedure on submission, please refer to: IV. Guidelines, C. Procedural Guidelines 2. L. ii., Pages 7-8 of FDA Circular No. 2020-033	Administrative Order No. 2014- 0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplie r
N - Processed nuts, including coated nuts and nut mixtures (with e.g. dried fruits)	☑ Valid Certificate of Analysis for Microbiological parameters for SNACK FOODS: Molds, CFU/g,	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



		1	PHILIPPINES
e.g. Yoghurt-, cereal-, and honey-covered nuts, and dried fruit-nut-and-cereal snacks (e.g. "trail mixes")	Yeast & Yeast-like fungi, CFU/g, Coliforms, CFU/g, Aerobic Plate Count, CFU/g.		
HIGH RISK FOOD PRODUCTS	☑ ADDITIONAL REQUIREMENTS	BASIS/ISSUANC E	WHERE TO SECURE
HIGH RISK FOOD PRODUCTS - foods that may contain pathogenic microorganisms and will support the formation of toxins and or the growth or pathogenic microorganisms and foods that may contain harmful chemicals. (AO No. 2014-0029)			
A1a - Milk (plain) and buttermilk (plain) "Plain fluid milk obtained from milking animals (e.g., cows, sheep, goats, buffalo) that has been processed. Includes pasteurized, ultra-high temperature (UHT) treated, sterilized, homogenized, or fat adjusted milk.	☑ Valid Certificate of Analysis for % Milk Solids Not Fat and % Milk Fat for MILK, CARABAO'S AND/OR BUFFALO'S MILK AND GOAT'S (NATIVE) MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
Includes, but is not limited to, skim, part-skim, low-fat and whole milk." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=3)	☑ Valid Certificate of Analysis for % Milk Solids Not Fat and % Milk Fat for SKIM MILK OR SKIMMED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
"Includes plain recombined fluid milks, plain reconstituted fluid milks, plain composite milks, non-flavoured vitamin and mineral fortified fluid milks, protein adjusted milks, lactose reduced milk, and plain milkbased beverages."	☑ Valid Certificate of Analysis for % Milk Solids Not Fat and % Milk Fat for RECONSTITUTED, RECONSTRUCTED OR RECOMBINED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=4)	☑ Valid Certificate of Analysis for % Milk Solids Not Fat for RECONSTITUTED, RECONSTRUCTED OR RECOMBINED SKIMMED MILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for % Milk Solids Not Fat for BUTTERMILK	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r



		PHILIPPINES
☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
% Milk Fat for LOWFAT MILK	Order No. 132 s.	Manufacturer/Source/Supplie
AND RECONSTITUTED,	<u>1970</u>	I
RECONSTRUCTED OR		
RECOMBINED LOWFAT MILK	A also is intentions	Annicant Cananany
☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
% Non-Fat Milk Solids, Vitamin A	Order No. 132 s.	Manufacturer/Source/Supplie
and Vitamin D (if added) for	<u>1970</u>	ľ
FILLED MILK		
*The 0/ Total Oil Content shall be		
*The % Total Oil Content shall be declared in the Electronic		
Registration Data Entry. **The product shall conform with		
the identity, standards for optional		
ingredients and additional label		
declaration for Filled Milk.		
*PASTEURIZED MILK AND	Administrative	Applicant Company/
STERILISED MILK shall conform	Order No. 132 s.	Manufacturer/Source/Supplie
with the prescribed standard of	1970	r
identity and quality	1970	1
✓ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	2022-012	Manufacturer/Source/Supplie
Microbiological parameters for LIQUID MILK (EVAPORATED &	2022-012	r
READY TO DRINK)-		
UHT/STERILIZED: Commercial		
Sterility		
,	FDA Circular No.	Applicant Company/
☑ Valid Certificate of Analysis for	2022-012	Manufacturer/Source/Supplie
Microbiological parameters for	<u> </u>	r
PASTEURIZED MILK: Coliforms		1
CFU/mL, Salmonella/25mL,		
Listeria monocytogenes/25mL,		



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	Psychrotrophic bacteria cfu/mL & Aerobic Plate Count CFU/g (Plain/Flavored)		
A1b - Dairy-based drinks, flavored and/or fermented "Includes all mixes and ready-to-drink fermented or not fermented milk-based drinks with flavourings and/or food ingredients that intentionally impart flavour, excluding mixes for cocoa. Examples, include but are not limited to, chocolate milk, chocolate malt drinks, strawberry-flavoured yoghurt drink, lactic acid bacteria drinks, whey-based drinks, and lassi (liquid obtained by whipping curd from the lactic acid fermentation of milk,	*FLAVORED MILK, FLAVORED RECONSTITUTED MILK, FLAVORED DRINK OR FLAVORED DAIRY DRINK, AND CHOCOLATE DRINK OR CHOCOLATE FLAVORED DRINK shall conform with the prescribed standard of identity and quality	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
and mixing with sugar or intense sweetener)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=6)	✓ Valid Certificate of Analysis for Microbiological parameters for LIQUID MILK (READY TO DRINK)-UHT/STERILIZED: Commercial Sterility	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	✓ Valid Certificate of Analysis for Microbiological parameters for NON-ALCOHOLIC BEVERAGES (e.g. READY TO DRINK, SOFTDRINKS, ICED TEA, ENERGY DRINKS, JELLY DRINKS): Yeast and Mold Count CFU/mL, Coliforms CFU/mL & Aerobic Plate Count CFU/mL	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A2ai - Fermented milk (plain), non heat-treated after fermentation "Includes fluid and non-fluid plain products, such as yoghurt and plain drinks based on fermented milk." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=9)	☑ Valid Certificate of Analysis for % Milk Fat Content by weight; % Milk Solids Non-Fat Content by weight; Acidity of the product when solid for YOGURT AND FLAVORED YOGURT	Administrative Order No. 132 s. 1970	



			PHILIPPINES
			Applicant Company/ Manufacturer/Source/Supplie
	*Toned Milk shall conform with the prescribed standard of identity and quality	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie
	✓ Valid Certificate of Analysis for Microbiological parameters for YOGURT AND OTHER FERMENTED MILK: S. aureus CFU/mL, Coliforms CFU/mL, Salmonella/25mL & Lactic acid CFU/mL (required minimum level: ≥10 ⁶ CFU/mL)	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A2aii - Fermented milks (plain), heat-treated after fermentation Includes fluid and non-fluid plain products, such as yoghurt and plain drinks based on fermented milk "except that they have been heat-treated (e.g. sterilized or pasteurized) after fermentation."	✓ Valid Certificate of Analysis for % Milk Fat Content by weight; % Milk Solids Non-Fat Content by weight; Acidity of the product when solid for YOGURT AND FLAVORED YOGURT	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=10)	*Toned Milk shall conform with the prescribed standard of identity and quality	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for Microbiological parameters for HEAT TREATED, FERMENTED MILK (STERILIZED, UHT): Commercial Sterility	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A2b - Renneted milk (plain) "Plain, coagulated milk produced by the action of milk coagulating enzymes. Includes curdled milk."	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=11)			
A3a - Pasteurized cream (plain) "Cream subjected to pasteurization by appropriate heat treatment or made from pasteurized milk. Includes milk cream and "half-and-half."" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=16)	✓ Valid Certificate of Analysis for Microbiological parameters for PASTEURIZED CREAM : Coliforms CFU/g, Salmonella/25g, Listeria monocytogenes/25g, Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A3b - Sterilized and UHT creams, whipping and whipped creams, and reduced fat creams (plain) "Includes every cream, regardless of fat content, which	☑ Valid Certificate of Analysis for % Butterfat for CREAM	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
has undergone a higher heat-treatment than pasteurization. Also includes pasteurized creams with a reduced fat content, as well as every cream intended for whipping or being whipped. Sterilized cream is subjected to appropriate heat-treatment in the container in which it is presented to the consumer. Ultra-heat treated (UHT) or ultrapasteurized cream is subjected to	☑ Valid Certificate of Analysis for % Butterfat for LIGHT CREAM TABLE CREAM OR COFFEE CREAM	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for % Milk Fat for WHIPPING CREAM	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
the appropriate heat treatment (UHT or ultrapasteurization) in a continuous flow process and aseptically packaged. Cream may also be packaged under pressure (whipped cream). Includes whipping	☑ Valid Certificate of Analysis for % Butterfat for LIGHT WHIPPING CREAM	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
cream, heavy cream, whipped pasteurized cream, and whipped cream-type dairy toppings and fillings." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=17)	☑ Valid Certificate of Analysis for % Milk Fat for HEAVY CREAM OR HEAVY WHIPPING CREAM	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
intpo.//www.ido.org/goldonii/io/ioodo/doldiio.ntinii:id=1//	☑ Valid Certificate of Analysis for % Milk Fat for HALF-AND HALF	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for Microbiological parameters for	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



	CREAM (UHT/STERILIZED): Commercial Sterility		PHILIPPINES
A3c - Clotted cream (plain) "Thickened, viscous cream formed from the action of milk coagulating enzymes. Includes sour cream (cream subjected to lactic acid fermentation." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=18)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A3d - Cream analogues "Cream substitute consisting of a vegetable fat-water emulsion in liquid or powdered form for use other than as a beverage whitener. Includes instant whipped cream toppings and sour cream substitutes." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=19)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A4a - Unripened cheese "Unripened cheese, including fresh cheese, is ready for consumption soon after manufacture. Examples include cottage cheese (a soft, unripened, coagulated curd cheese), creamed cottage cheese (cottage cheese covered with a creaming mixture), cream cheese (rahmfrischkase, an uncured, soft spreadable cheese), mozzarella and scamorza cheeses. Includes the whole	✓ Valid Certificate of Analysis for % Milk Fat and % Moisture for CREAM CHEESE *The product shall conform with the identity and standards for optional ingredients for Cream Cheese.	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
unripened cheese and unripened cheese rind (for those unripened cheeses with a "skin" such as mozzarella). Most products are plain, however, some, such as cottage cheese and cream cheese, may be flavoured or contain ingredients such as fruit, vegetables or meat." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=24)	✓ Valid Certificate of Analysis for % Milk Fat and % Moisture for COTTAGE CHEESE DRY CURD or DRY CURD COTTAGE CHEESE *The product shall conform with the identity, standards for optional ingredients and additional label	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r



		PHILIPPINE2
declaration for Cottage Cheese Dry Curd or Dry Curd Cottage Cheese.		
☑ Valid Certificate of Analysis for % Milk Fat and % Moisture for COTTAGE CHEESE	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Cottage Cheese.		
✓ Valid Certificate of Analysis for % Milk Fat and % Moisture for LOW FAT COTTAGE CHEESE	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Low Fat Cottage Cheese.		
☑ Valid Certificate of Analysis for % Milk Fat for SKIM MILK CHEESE	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the identity for Skim Milk Cheese.		
✓ Valid Certificate of Analysis for Microbiological parameters for SOFT CHEESE (FROM PASTEURIZED MILK): Enterobacteriaceae CFU/g, E.coli	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
CFU/g, Salmonella/ 25g, Listeria		



			PHILIPPINES
	monocytogenes/ 25g & S. aureus CFU/g		
	✓ Valid Certificate of Analysis for Microbiological parameters for HARD AND SEMI-HARD CHEESE: Enterobacteriaceae CFU/g, E.coli CFU/g, Salmonella/ 25g, Listeria monocytogenes/ 25g & S. aureus CFU/g		
	☑ Valid Certificate of Analysis for Microbiological parameters for CREAM CHEESE PRODUCTS: Coliforms CFU/g or MPN/g or /25g. E. coli CFU/g or MPN/g or /25g and Yeast and Molds CFU/g		
	☑ Valid Certificate of Analysis for Microbiological parameters for ALL RAW MILK CHEESE; RAW MILK UN-RIPENED CHEESE W/ MOISTURE > 50%, pH > 5.0: Campylobacter/25g, Listeria monocytogenes/25g, Salmonella/25g & S. aureus CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A4bi - Ripened cheese, includes rind "Refers to ripened (including mould-ripened) cheese, including rind, or any part thereof, such as cut, shredded, grated or sliced cheese. Examples of ripened cheese include: blue cheese, brie, gouda, havarti, hard grating cheese, and Swiss cheese."	☑ Valid Certificate of Analysis for % Moisture and % Milk Fat (of solids) for CHEDDAR CHEESE *The product shall conform with the identity and standards for	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
· ·	optional ingredients for Cheddar		
https://www.fao.org/gsfaonline/foods/details.html?id=26)	Cheese.		
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	% Moisture and % Milk Fat (of	Order No. 200-As.	Manufacturer/Source/Supplie
	solids) for WASHED CURD	1973	r
		1070	'
	CHEESE (SOAKED CURD		
	CHEESE)		
	*The product shall conform with		
	the identity and standards for		
	Washed Curd Cheese (Soaked		
	Curd Cheese).		
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	% Moisture and % Milk Fat (of	Order No. 200-As.	Manufacturer/Source/Supplie
	solids) for COLBY CHEESE	1973	r
	solids) for GOLDT GITLEGE	1010	·
	*The product shall conform with		
	•		
	the identity and standards for		
<u> </u>	Colby Cheese.		
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	% Moisture and % Milk Fat (of	Order No. 200-As.	Manufacturer/Source/Supplie
	solids) for GRANULAR CHEESE	<u>1973</u>	r
	(STIRRED CURD CHEESE)		
	,		
	*The product shall conform with		
	the identity and standards for		
	Granular Cheese (Stirred Curd		
	Cheese).		
	/	Administrative	Applicant Company
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	% Moisture and % Milk Fat (of	Order No. 200-As.	Manufacturer/Source/Supplie
	solids) for BRICK CHEESE	<u>1973</u>	r



*The product shall conform with the identity and standards for optional ingredients Swiss Cheese. Valid Certificate of Analysis for % Moisture and % Milk Fat (of Order No. 200-A s. Manufacturer/Source/Supplied			<u> </u>
% Moisture and % Milk Fat (of solids) for SWISS CHEESE *The product shall conform with the identity and standards for optional ingredients Swiss Cheese. ☑ Valid Certificate of Analysis for % Moisture and % Milk Fat (of Solids) Manufacturer/Source/Supplied and Solids (of SWISS CHEESE) Manufacturer/Source/Supplied and Solids (of SWISS Cheese) Administrative Order No. 200-As. Manufacturer/Source/Supplied (of Swiss Cheese)	the identity and standards for optional ingredients for Brick		
the identity and standards for optional ingredients Swiss Cheese. ☑ Valid Certificate of Analysis for % Moisture and % Milk Fat (of Order No. 200-As. Manufacturer/Source/Supplied	% Moisture and % Milk Fat (of	Order No. 200-As.	
% Moisture and % Milk Fat (of Order No. 200-A s. Manufacturer/Source/Supplied	the identity and standards for optional ingredients Swiss	1	
solids) for GRUYERS CHEESE 1973	· · · · · · · · · · · · · · · · · · ·	Order No. 200-As.	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the identity and standards for optional ingredients Gruyers Cheese.	the identity and standards for optional ingredients Gruyers	1	
✓ Valid Certificate of Analysis for % Moisture and % Milk Fat (of solids) for EDAM CHEESE Applicant Company/ Manufacturer/Source/Supplier r	% Moisture and % Milk Fat (of	Order No. 200-As.	Applicant Company/ Manufacturer/Source/Supplie r
*The product shall conform with the identity and standards for optional ingredients Edam Cheese.	the identity and standards for optional ingredients Edam		
✓ Valid Certificate of Analysis for % Moisture and % Milk Fat (of solids) for PARMESAN CHEESE Applicant Company/ Manufacturer/Source/Supplier	% Moisture and % Milk Fat (of	Order No. 200-As.	Applicant Company/ Manufacturer/Source/Supplie r



	1	PHILIPPINES
*The product shall conform with the identity and standards for optional ingredients Parmesan Cheese.		
☑ Valid Certificate of Analysis for Microbiological parameters for SOFT CHEESE (FROM PASTEURIZED MILK): Enterobacteriaceae CFU/g, E.coli CFU/g, Salmonella/ 25g, Listeria monocytogenes/ 25g & S. aureus CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
☑ Valid Certificate of Analysis for Microbiological parameters for HARD AND SEMI-HARD CHEESE: Enterobacteriaceae CFU/g, E.coli CFU/g, Salmonella/ 25g, Listeria monocytogenes/ 25g & S. aureus CFU/g		
✓ Valid Certificate of Analysis for Microbiological parameters for CREAM CHEESE PRODUCTS: Coliforms CFU/g or MPN/g or /25g. E. coli CFU/g or MPN/g or /25g and Yeast and Molds CFU/g		
✓ Valid Certificate of Analysis for Microbiological parameters for ALL RAW MILK CHEESE; RAW MILK UN-RIPENED CHEESE W/ MOISTURE > 50%, pH > 5.0:	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
	Campylobacter/25g, Listeria monocytogenes/25g, Salmonella/25g & S. aureus CFU/g		
A4bii - Rind of ripened cheese "Refers to the rind only of the cheese. The rind of the cheese is the exterior portion of the cheese mass that initially has the same composition as the interior portion of the cheese, but which may dry after brining and ripening." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=27)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A4biii - Cheese powder (for reconstitution) "Dehydrated product prepared from a variety or processed cheese. Product is intended either to be reconstituted with milk or water to prepare a sauce, or used as-is as an ingredient (e.g. with cooked macaroni, milk and butter to prepare a macaroni and cheese casserole). Includes spray-dried cheese." Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=28)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A4c - Whey cheese "A solid or semi-solid product obtained by concentration of whey with or without the addition of milk, cream or other materials of milk origin, and moulding of the concentrated product. Includes the whole cheese and the rind of the cheese." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=29)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A4di - Plain processed cheese	☑ Valid Certificate of Analysis for % Moisture Content, % Fat Content in Dry Matter and %	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r



	T	T	PHILIPPINES
"Processed cheese product that does not contain added flavours, seasonings, fruit, vegetables and/or meat. Examples include: American cheese, requeson."	PROCESS CHEESE		
(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=31)	*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized		
	Process Cheese.		
	✓ Valid Certificate of Analysis for % Moisture Content, % Fat Content and % Milk Fat (when the food contains other foodstuffs) for PASTEURIZED PROCESS CHEESE FOOD	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
	*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized Process Cheese Food.		
	☑ Valid Certificate of Analysis for % Moisture Content and % Fat Content for PASTEURIZED PROCESS CHEESE SPREAD	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
	*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized Process Cheese Spread.		



		PHILIPPINES
☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
SOFT CHĔESE (FROM		r
PASTEURIZED MILK):		
Enterobacteriaceae CFU/g, E.coli		
CFU/g, Salmonella/ 25g, Listeria		
monocytogenes/ 25g & S. aureus		
CFU/q		
CF0/g		
☑ Valid Certificate of Analysis for		
Microbiological parameters for		
HARD AND SEMI-HARD		
CHEESE: Enterobacteriaceae		
CFU/g, E.coli CFU/g, Salmonella/		
25g, Listeria monocytogenes/ 25g		
& S. aureus CFU/g		
☑ Valid Certificate of Analysis for		
Microbiological parameters for		
CREAM CHEESE PRODUCTS:		
Coliforms CFU/g or MPN/g or		
/25g. E. coli CFU/g or MPN/g or		
/25g and Yeast and Molds CFU/g		
✓ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
ALL RAW MILK CHEESE; RAW	ZOZZ OTZ	r
MILK UN-RIPENED CHEESE W/		'
MOISTURE > 50%, pH > 5.0:		
Campylobacter/25g, Listeria		
monocytogenes/25g,		
Salmonella/25g & S. aureus		
CFU/g		



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	✓ Valid Certificate of Analysis for Microbiological parameters for PROCESSED CHEESE SPREAD: Coliforms CFU/g, S. aureus CFU/g & Aerobic Plate	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	Count CFU/g		
A4dii - Flavored processed cheese "Processed cheese product that contains added flavours, seasonings, fruit, vegetables and/or meat. Examples include: neufchatel cheese spread with vegetables, pepper jack cheese, cheddar cheese spread with wine, and cheese balls (formed processed cheese coated in nuts, herbs or spices)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=32)	✓ Valid Certificate of Analysis for % Moisture Content, % Fat Content in Dry Matter and % Lactose for PASTEURIZED PROCESS CHEESE *The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
	Process Cheese.		
	☑ Valid Certificate of Analysis for % Moisture Content, % Fat Content and % Milk Fat (when the food contains other foodstuffs) for PASTEURIZED PROCESS CHEESE FOOD	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r
	*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized Process Cheese Food.		
	☑ Valid Certificate of Analysis for % Moisture Content and % Fat	Administrative Order No. 200-As. 1973	Applicant Company/ Manufacturer/Source/Supplie r



		PHILIPPINES
Content for PASTEURIZED PROCESS CHEESE SPREAD		
*The product shall conform with the identity, standards for optional ingredients and additional label declaration for Pasteurized Process Cheese Spread.		
☑ Valid Certificate of Analysis for Microbiological parameters for SOFT CHEESE (FROM PASTEURIZED MILK): Enterobacteriaceae CFU/g, E.coli CFU/g, Salmonella/ 25g, Listeria monocytogenes/ 25g & S. aureus CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
☑ Valid Certificate of Analysis for Microbiological parameters for HARD AND SEMI-HARD CHEESE: Enterobacteriaceae CFU/g, E.coli CFU/g, Salmonella/ 25g, Listeria monocytogenes/ 25g & S. aureus CFU/g		
☑ Valid Certificate of Analysis for Microbiological parameters for CREAM CHEESE PRODUCTS: Coliforms CFU/g or MPN/g or /25g. E. coli CFU/g or MPN/g or /25g and Yeast and Molds CFU/g		



			PHILIPPINES
	☑ Valid Certificate of Analysis for Microbiological parameters for ALL RAW MILK CHEESE; RAW MILK UN-RIPENED CHEESE W/ MOISTURE > 50%, pH > 5.0: Campylobacter/25g, Listeria monocytogenes/25g, Salmonella/25g & S. aureus CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Valid Certificate of Analysis for Microbiological parameters for PROCESSED CHEESE SPREAD: Coliforms CFU/g, S. aureus CFU/g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A4e - Cheese analogues "Products that look like cheese, but in which milkfat has been partly or completely replaced by other fats. Includes imitation cheese, imitation cheese mixes, and imitation cheese powders." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=33)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A4f - Whey protein cheese "Product containing the protein extracted from the whey component of milk. These products are principally made by coagulation of whey proteins. Example: ricotta cheese." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=34)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A5 - Dairy-based desserts "Includes ready-to-eat flavoured dairy dessert products and dessert mixes."	☑ Valid Certificate of Analysis for % Milk Fat Content by weight; % Milk Solids Non-Fat Content by	Administrative Order No. 132 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
"Includes flavoured yoghurt (a milk product obtained by fermentation of milk and milk products to which flavours and ingredients (e.g., fruit, cocoa, coffee) have been added) that may or may not be heat-treated after fermentation." Other examples include: "jellied milk, frozen flavoured yoghurt, junket (sweet custard-like dessert made from flavoured milk set with rennet), dulce de leche (cooked milk with sugar and added ingredients such as coconut or chocolate), butterscotch pudding and chocolate mousse. Includes traditional milk-based sweets prepared from milk concentrated partially, from khoa (cow or buffalo milk concentrated by boiling), or chhena (cow or buffalo milk, heat coagulated aided by acids like citric acid, lactic acid, malic acid, etc), sugar or synthetic sweetener, and other ingredients (e.g. maida (refined wheat flour), flavours and colours (e.g. peda, burfee, milk cake, gulab jamun, rasgulla, rasmalai, basundi)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=35)	weight; Acidity of the product when solid for YOGURT AND FLAVORED YOGURT ☑ Valid Certificate of Analysis for Microbiological parameters for YOGURT AND FERMENTED MILK: S. aureus CFU/mL, Coliforms CFU/mL, Salmonella/25mL & Lactic acid CFU/mL (required minimum level: ≥10^6 CFU/mL) ☑ Valid Certificate of Analysis for Microbiological parameters for ETHNIC MILK-BASED CONFECTIONERIES (e.g. PASTILLAS and YEMA): Yeast and Mold Count CFU/g, Salmonella/25, Coliforms MPN/g or CFU/g and Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
A6a - Liquid whey and whey products "Whey is the fluid separated from the curd after coagulation of milk, cream, skimmed milk or buttermilk with milk coagulating enzymes during the manufacture of cheese, casein or similar products. Acid whey is obtained after the coagulation of milk, cream, skimmed milk or buttermilk, mainly with acids of the type used for the manufacture of fresh cheese." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=37)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



			PHILIPPINES
A6b - Dried whey and whey products "Whey powders are prepared by spray- or roller-drying whey or acid whey from which the major portion of the milkfat has been removed." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=38)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A7 - Milk for manufacture	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
A8 - Dairy-based frozen desserts "Includes frozen dairy confections and novelties, and dairy-based fillings." "Other examples include: ice cream (frozen dessert that may contain whole milk, skim milk products, cream or butter, sugar, vegetable oil, egg products, and fruit, cocoa, or coffee), ice milk (product similar to ice cream with reduced whole or skim milk content, or made with nonfat milk)" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=35)	✓ Valid Certificate of Analysis for Microbiological parameters for ICE CREAM & SHERBET (PLAIN AND FLAVORED): Coliforms CFU/g, Listeria monocytogenes/25g, Salmonella/25g, Aerobic Plate Count CFU/g & S. aureus CFU/g ✓ Valid Certificate of Analysis for Microbiological parameters for ICE CREAM WITH ADDED INGREDIENTS (NUTS, FRUITS, COCOA ETC.): Coliforms CFU/g, S. aureus CFU/g, Salmonella/25g, Salmonella/25g CFU/g & Listeria monocytogenes/25g	FDA Circular No. 2022-012 FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r Applicant Company/ Manufacturer/Source/Supplie r
B1 - Dried fruits and vegetable - plain/sun-dried seaweeds, and nuts and seeds "Products in which the natural water content has been reduced below that critical for growth for microorganisms without affecting the important nutrients. The product may or may not be intended for rehydration prior to	✓ Valid Certificate of Analysis for Microbiological parameters for SUN DRIED FRUITS: Mold CFU/g, Osmophilic Yeasts CFU/g & E. coli MPN/g	FDA Circular No. 2022-012 FDA Circular No.	Applicant Company/ Manufacturer/Source/Supplie r Applicant Company/
consumption. Includes vegetable powders that are obtained from drying the juice, such as tomato powder	☑ Valid Certificate of Analysis for Microbiological parameters for	2022-012	Manufacturer/Source/Supplie



			PHILIPPINES
and beet powder. Examples include: dried potato flakes and dried lentil. Examples of Oriental dried products include: dried sea tangle (kelp; kombu), dried sea tangle with seasoning (shio-kombu), dried seaweed (tororo-kombu), dried gourd strips (kampyo), dried laver (nori), and dried laminariales (wakame)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=79)	DRIED VEGETABLE: E. coli MPN/g		
B2 - Vegetable seaweed, and nut and seed - purees, spreads "Vegetable purees are finely dispersed slurries prepared from the concentration of vegetables, which may have been previously heat-treated (e.g., steamed). The slurries may be filtered prior to packaging. Examples include: tomato puree, peanut butter (a spreadable paste made from roasted and ground peanuts by the addition of peanut oil), other nut butters (e.g., cashew	Valid Certificate of Analysis for % Fat Content and % Water Insoluble Inorganic Residue for Peanut Butter *The product shall conform with the identity and label statement for optional ingredients for Peanut Butter.	Administrative Order No. 228 s. 1974	Applicant Company/ Manufacturer/Source/Supplie r
butter), and pumpkin butter." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=82)	✓ Valid Certificate of Analysis for Microbiological parameters for PEANUT BUTTER & OTHER NUT BUTTERS: Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
D - Chocolate with nuts	 ✓ Valid Certificate of Analysis for Microbiological parameters for CHOCOLATE PRODUCTS: Molds CFU/g, Salmonella/25g, Coliforms MPN/g or CFU/g & Aerobic Plate Count CFU/g. ✓ Valid Certificate of Analysis for Microbiological parameters for 	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	CHOCOLATE CONFECTIONARIES: Molds		



			PHILIPPINES
	CFU/g, Osmophilic yeast CFU/g, Salmonella/25g, Coliforms MPN/g or CFU/g & Aerobic Plate Count CFU/g		
F1 - Fine bakery products with fillings: meat, milk, poultry, cream, and other perishable foods; icings and coatings "The term "sweet cracker" or "sweet biscuit" used in this category refers to a cookie-like product that may be eaten as a dessert. Examples include: butter cake,	☑ Valid Certificate of Analysis for Microbiological parameters for BAKED GOODS: Yeast CFU/g, Mold CFU/g, Aerobic Plate Count CFU/g, Coliforms CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
cheesecake, fruit-filled cereal bars, pound cake (including kasutera), moist cake (type of starchy dessert (namagashi)), western cakes, moon cakes, sponge cake, fruit-filled pies (e.g. apple pie), oatmeal cookies, sugar cookies and British "biscuits" (cookies or sweet crackers)." (Source URL: https://fao.org/gsfaonline/foods/details.html?id=124)	✓ Valid Certificate of Analysis for Microbiological parameters for COATED OR FILLED, DRIED SHELF-STABLE BISCUITS: Coliforms MPN/g & Salmonella/25g ✓ Valid Certificate of Analysis for	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	Microbiological parameters for ETHNIC FLOUR-BASED CONFECTIONERIES e.g. PIAYA): Yeast and Mold Count CFU/g and Coliforms CFU/g		
	✓ Valid Certificate of Analysis for Microbiological parameters for FROZEN BAKERY PRODUCTS (READY TO EAT) WITH LOW ACID OR HIGH Aw FILLINGS OR TOPPINGS: S. aureus CFU/g & Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	FROZEN BAKERY PRODUCTS		r
	(TO BE COOKED) WITH LOW		
	ACID OR HIGH AW FILLINGS		
	OR TOPPINGS: S. aureus CFU/g		
FO Cooking with muto	& Salmonella/25g	EDA Cinavilar Na	Analisant Cananany
F2 - Cookies with nuts	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
	BAKED GOODS: Yeast CFU/g,		r
	Mold CFU/g, Aerobic Plate Count		
	CFU/g, Coliforms CFU/g &		
C4a Haat translad musaaas ad marat maraktura sad	Salmonella/25g	EDA Cinavilan Na	Applicant Cansus and
G1a - Heat-treated processed meat, poultry and	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
game products in whole pieces or cuts (canned) "Includes cooked (including cured and cooked, and dried	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
and cooked), heat-treated (including sterilized) and	MEAT PRODUCTS IN		'
canned meat cuts. Examples include: cured, cooked	HERMETICALLY SEALED		
ham; cured, cooked pork shoulder; canned chicken	CONTAINERS: Commercial		
meat; and meat pieces boiled in soy sauce (tsukudani)."	Sterility		
(Source URL:			
https://www.fao.org/gsfaonline/foods/details.html?id=136	☑ Valid Certificate of Analysis for		
)	Microbiological parameters for PACKAGED COOKED		
,	CURED/SALTED MEAT: S.		
	aureus, CFU/g, Salmonella/25g,		
	Listeria Monocytogenes/25g		
	Listeria Monocytogenes/20g		
	☑ Valid Certificate of Analysis for		
	Microbiological parameters for		
	MARINATED MEAT PRODUCTS:		
	Salmonella/25g, Listeria		
	Loannondia/209, Libitina		



			PHILIPPINES
	monocytogenes/25g, S. aureus, CFU/g		
	☑ Valid Certificate of Analysis for Nitrate or Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	Applicant Company/ Manufacturer/Source/Supplie r
G1b - Frozen processed meat, poultry and game products in whole pieces or cuts (marinated pork/beef/chicken cuts) "Includes raw and cooked meat cuts that have been frozen. Examples include: frozen whole chickens, frozen chicken parts, and frozen beef steaks." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=137)	✓ Valid Certificate of Analysis for Microbiological parameters for MARINATED MEAT PRODUCTS: Salmonella/25g, Listeria monocytogenes/25g, S. aureus, CFU/g ✓ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM POULTRY MEAT INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g, Salmonella/25g ✓ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM SPECIES OTHER THAN POULTRY INTENDED TO BE EATEN COOKED: Aerobic Plate Count, CFU/g, E.coli, CFU/g,	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	Salmonella/25g		1



✓ Valid Certificate of Analysis for Microbiological parameters for FOODS COOKED IMMEDIATELY PRIOR TO SALE OR CONSUMPTION (e.g. Takeaway food, burgers, kebabs, sausages, pizza, ready meals (cook/chill and cook/freeze) after regeneration): Aerobic Plate Count CFU/g, Enterobacteriaceae CFU/g, E. coli CFU/g, S. aureus (coagulase +) CFU/g, Salmonella/25g and Listeria monocytogenes/25g

☑ Valid Certificate of Analysis for Microbiological parameters for COOKED POULTRY MEAT, FROZEN TO BE REHEATED BEFORE EATING: Aerobic Plate Count, CFU/g, S. aureus, CFU/g, Listeria monocytogenes/25g, Salmonella/25g, Campylobacter Jejuni/25g

✓ Valid Certificate of Analysis for Microbiological parameters for COLD CUTS, FROZEN & CHILLED HOTDOGS: E. coli MPN/g or CFU/g or /25g, Salmonella/25g, S. aureus CFU/g,





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Count, CFU/g, E.coli, CFU/g, Salmonella/25g	
✓ Valid Certificate of Analysis for Microbiological parameters for FOODS COOKED IMMEDIATELY PRIOR TO SALE OR CONSUMPTION (e.g. Takeaway food, burgers, kebabs, sausages, pizza, ready meals (cook/chill and cook/freeze) after regeneration): Aerobic Plate Count CFU/g, Enterobacteriaceae CFU/g, E. coli CFU/g, S. aureus (coagulase +) CFU/g, Salmonella/25g and Listeria monocytogenes/25g	
✓ Valid Certificate of Analysis for Microbiological parameters for COOKED POULTRY MEAT, FROZEN TO BE REHEATED BEFORE EATING: Aerobic Plate Count, CFU/g, S. aureus, CFU/g, Listeria monocytogenes/25g, Salmonella/25g, Campylobacter Jejuni/25g	
☑ Valid Certificate of Analysis for Microbiological parameters for COLD CUTS, FROZEN & CHILLED HOTDOGS: E. coli	



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	MPN/g or CFU/g or /25g, Salmonella/25g, S. aureus CFU/g, L. monocytogenes/25g & Aerobic Plate Count CFU/g ☑ Valid Certificate of Analysis for Nitrate and Nitrite Content (if utilized)	Administrative Order No. 154 s. 1971 and Bureau Circular No. 2006- 016	Applicant Company/ Manufacturer/Source/Supplie r
G2b - Frozen processed comminuted meat, poultry and game products (nuggets, patties, dumplings salami, meat loaf, hotdog) "Includes raw, partially cooked and fully cooked comminuted or mechanically deboned meat products that have been frozen. Examples include: frozen hamburger patties; frozen breaded or battered chicken fingers." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=144)	✓ Valid Certificate of Analysis for Microbiological parameters for COLD CUTS, FROZEN & CHILLED HOTDOGS: E. coli MPN/g or CFU/g or /25g, Salmonella/25g, S. aureus CFU/g, L. monocytogenes/25g & Aerobic Plate Count CFU/g ✓ Valid Certificate of Analysis for Microbiological parameters for COOKED POULTRY MEAT, FROZEN TO BE REHEATED BEFORE EATING (e.g. prepared frozen meals chicken burgers, chicken turkey rolls, chicken nuggets, other breaded poultry meat products): Aerobic Plate Count CFU/g, S. aureus CFU/g, Listeria monocytogenes/25g, Salmonella/25 and Campylobacter jejuni/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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✓ Valid Certificate of Analysis for Microbiological parameters for MARINATED MEAT PRODUCTS (e.g. Marinated meat and meat preparations (tapa, sisig, etc.), - Marinated poultry, Dim sum made from meat (siomai)): Salmonella/25g, Listeria monocytogenes/25g and S. aureus CFU/g		
☑ Valid Certificate of Analysis for Microbiological parameters for FOODS COOKED IMMEDIATELY PRIOR TO SALE OR CONSUMPTION (e.g. Takeaway food, burgers, kebabs, sausages, pizza, ready meals (cook/chill and cook/freeze) after regeneration): Aerobic Plate Count CFU/g, Enterobacteriaceae CFU/g, E. coli CFU/g, S. aureus (coagulase +) CFU/g, Salmonella/25g and Listeria monocytogenes/25g	EDA Circular No.	Applicant Company
✓ Valid Certificate of Analysis for Microbiological parameters for MINCED MEAT AND MEAT PREPARATIONS MADE FROM SPECIES OTHER THAN POULTRY INTENDED TO BE	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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	EATEN COOKED:		
	Salmonella/25g, Aerobic Plate		
	Count CFU/g and E. coli CFU/g		
	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	MEAT PASTE & PATE:		r
	Salmonella/25g, Clostridium		•
	J .		
	perfringens CFU/g, S. aureus		
	CFU/g, Coliforms CFU/g &		
	Aerobic Plate Count CFU/g		
	☑ Valid Certificate of Analysis for	Administrative	Applicant Company/
	Nitrate and Nitrite Content (if	Order No. 154 s.	Manufacturer/Source/Supplie
	utilized)	<u>1971</u> and <u>Bureau</u>	r
	,	Circular No. 2006-	
		<u>016</u>	
H1a - Frozen fish, fish fillets and fish products	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
"Fresh, including partially cooked, fish subjected to	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
freezing or quick-freezing at sea and on land for further	FRESH FROZEN FISH: E. coli		r
processing. Examples include: frozen or deep frozen	MPN/g, S. aureus CFU/g, V.		
clams, cod fillets, crab, finfish, haddock, hake, lobster,	parahaemolyticus MPN/g,		
minced fish, prawns and shrimp; frozen fish roe; frozen	Salmonella/25g & Aerobic Plate		
surimi; and frozen whale meat."	Count CFU/q		
(Source URL:	9	FDA Circular No.	Applicant Company/
https://www.fao.org/gsfaonline/foods/details.html?id=151	☑ Valid Certificate of Analysis for		
1 Titips://www.rao.org/gsraoriiine/100ds/details.html: id=101	Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
/	FROZEN RAW CRUSTACEANS:		r
	E. coli MPN/g, S. aureus CFU/g,		
	Salmonella/25g, V.		
	parahaemolyticus MPN/g, Aerobic		
	Plate Count CFU/g		
	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	FRESH & FROZEN BIVALVE		l r
		l	



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H1b - Frozen battered fish, fish fillets and fish products, including molluscs, crustaceans and echinoderms "Uncooked product prepared from fish or fish portions, with dressing in eggs and bread crumbs or batter. Examples include: frozen raw breaded or batter-coated shrimp; and frozen or quick-frozen breaded or batter-coated fish fillets, fish portions and fish sticks (fish fingers)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=152	MOLLUSCS: E. coli MPN/g, Salmonella/25g, V. parahaemolyticus MPN/g & Aerobic Plate Count CFU/g ☑ Valid Certificate of Analysis for Microbiological parameters for FISH AND CRUSTACEAN BASED PROCESSED MEAT (e.g. fish ball, squid ball): Aerobic Plate Count CFU/g, S. aureus CFU/g, V. parahaemolyticus MPN/g and E. coli MPN/g.	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
H1c - Frozen minced and creamed fish products "Uncooked product prepared from minced fish pieces in cream-type sauce" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=153)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
H1di - Cooked fish and fish products "Cooked products include steamed, boiled or any other cooking method except frying. The fish may be whole, in portions or comminuted. Examples include: fish sausage; cooked fish products boiled down in soy sauce (tsukudani); cooked surimi product (kamaboko); crabflavoured cooked kamaboko product (kanikama); cooked fish roe; cooked surimi; cooked, tube-shaped surimi product (chikuwa); and cooked fish and lobster paste (surimi-like products."	 ✓ Valid Certificate of Analysis for Microbiological parameters for AQUATIC PRODUCTS: Salmonella/25g, V. parahaemolyticus MPN/g and S. aureus CFU/g ✓ Valid Certificate of Analysis for Microbiological parameters for 	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



(Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=155) H1dii - Cooked molluscs, crustaceans and echinoderms "Cooked products include steamed, boiled or any other cooking method except frying. Examples include: cooked crangon crangon and crangon vulgaris (brown shrimp; cooked shrimp, clams and crabs." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=156)	PRE-COOKED BREADED FISH: E.coli, MPN/g, S. aureus, CFU/g, Aerobic Plate Count, CFU/g ☑ Valid Certificate of Analysis for Microbiological parameters for FROZEN COOKED CRUSTACEANS: E. coli MPN/g, S. aureus CFU/g, V. parahaemolyticus CFU/g, Salmonella/25g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
)	✓ Valid Certificate of Analysis for Microbiological parameters for COOKED, CHILLED & FROZEN CRABMEAT: E. coli MPN/g, S. aureus CFU/g, V. parahaemolyticus MPN/g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
H1diii - Fried fish and fish products "Ready-to-eat products prepared from fish or fish portions, with or without further dressing in eggs and bread crumbs or batter, that are fried, baked, roasted or barbecued, and then packaged or canned with or without sauce or oil. Examples include: ready-to-eat fried surimi, fried calamari, and fried soft-shell crabs." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=157)	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
H2 - Fully preserved including canned or fermented fish and fish products "Products with extended shelf life, manufactured by pasteurizing or steam retorting and packaging in	☑ Valid Certificate of Analysis for Microbiological parameters for FISH & SHELLFISH PRODUCTS, COOKED	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



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vacuum-sealed air-tight containers to ensure sterility. Products may be packed in their own juice or in added oil or sauce. Examples include: canned tuna, clams, crab, fish roe and sardines; gefilte fish balls; and surimi (heat-pasteurized)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=164)	CRUSTACEANS IN HERMETICALLY SEALED CONTAINERS (THERMALLY PROCESSED) EG. COOKED BAGOONG/SHRIMP PASTE: Commercial Sterility Valid Certificate of Analysis for Total Solids, Protein and NaCl for BAGOONG (FISH AND SHRIMP)	Administrative Order No. 128 s. 1970	Applicant Company/ Manufacturer/Source/Supplie r
la - Liquid egg products "The purified whole egg, egg yolk or egg white is pasteurized and chemically preserved (e.g. by addition of salt)." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=168)	✓ Valid Certificate of Analysis for Microbiological parameters for PASTEURIZED EGG PRODUCTS (SMOKED LIQUID, FROZEN, DRIED): Coliforms CFU/g, Salmonella/25g, Yeast and Mold Count CFU/g (for dried products) & SPC/APC CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
Ib - Frozen egg products "The purified whole egg, egg yolk or egg white is pasteurized and frozen." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=169)	✓ Valid Certificate of Analysis for Microbiological parameters for PASTEURIZED EGG PRODUCTS (SMOKED LIQUID, FROZEN, DRIED): Coliforms CFU/g, Salmonella/25g, Yeast and Mold Count CFU/g (for dried products) & SPC/APC CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
Ic - Dried and/or heat coagulated egg products "Sugars are removed from the purified whole egg, egg yolk or egg white, which is then pasteurized and dried." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=170)	☑ Valid Certificate of Analysis for Microbiological parameters for PASTEURIZED EGG PRODUCTS (SMOKED LIQUID, FROZEN, DRIED): Coliforms CFU/g, Salmonella/25g, Yeast	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r



			PHILIPPINES
	and Mold Count CFU/g (for dried		
	products) & SPC/APC CFU/g		
J1 - Infant formula, follow-on formula and formula	INFÁNT FORMULA & FORM	ULAS FOR SPECIA	L MEDICAL PURPOSES
for special medical purposes for infants		NDED FOR INFANT	
Tor oposiar modical purposes for intante		Codex Stan 72-	Applicant Company/
INFANT FORMULA	☑ Valid Certificate of Analysis for	1981 Rev. 2007	Manufacturer/Source/Supplie
	Energy, Protein, Total Fat,	1901 Rev. 2007	iviariuracturer/Source/Supplie
"A human milk substitute for infants (aged no more than	Linolenic Acid, Total		r
12 months) that is specifically formulated to provide the	Carbohydrates per 100g, Vitamins		
sole source of nutrition during the first months of life up	and Minerals, Trace Minerals and		
to the introduction of appropriate complementary	Other Substances,		
feeding. Product is in a liquid form, either as a ready-to-	Lauric/Mystiric/Trans Fatty Acids,		
eat product, or is reconstituted from a powder."	Optional Ingredients- Taurine,		
(Source URL:	DHA and Contaminants		
https://www.fao.org/gsfaonline/foods/details.html?id=225	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	POWDERED INFANT FORMULA	LOLL OIL	r
FOLLOW-UP FORMULA	WITH OR WITHOUT ADDED		1
"Food intended for use as a liquid part of the			
complementary feeding of infants (aged at least 6	LACTIC ACID PRODUCING		
months) and for young children (aged 1-3 years). They	CULTURES (INTENDED FOR 0		
may be ready-to-eat or in a powdered form to be	TO 6 MONTHS OLD):		
reconstituted with water."	Cronobacter spp./10g,		
	Salmonella/25g, Aerobic Plate		
(Source URL:	Count CFU/g &		
https://www.fao.org/gsfaonline/foods/details.html?id=226	Enterobacteriaceae/10g		
)	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
FORMULA FOR SPECIAL MEDICAL PURPOSES FOR	INFANT FORMULA- LIQUID		l r
INFANTS	(UHT/STERILIZED): Commercial		
"Foods for special dietary use that are specially	Sterility		
processed or formulated and presented for the dietary	,	Department	Applicant Company/
management of infants and may be used only under	☑ Clear and complete loose		
medical supervision. They are intended for the exclusive	labels or artworks compliant with	Circular No. 2008-	Manufacturer/Source/Supplie
	Department Circular 2008-0006	0006	Γ



			PHILIPPINES
or partial feeding of infants with limited or impaired	☑ For FSMP: Scientific Studies	Codex Stan 72-	Applicant Company/
capacity to take, digest, absorb or metabolize ordinary	indicating safety and benefits of	1981 Rev. 2007	Manufacturer/Source/Supplie
infant formulae or certain nutrients contained therein, or	the product for intended medical	and Administrative	r
who have other special medically-determined nutrient	condition	Order No. 2014-	
requirement, whose dietary management cannot be		0029	
achieved only by modification of the normal diet, by	FOLLOW-UP F	ORMULA/MILK SUF	PLEMENT
other foods for special dietary uses, or by a combination	☑ Valid Certificate of Analysis for	Codex Stan 156-	Applicant Company/
of the two."	Energy, Protein, Total Fat,	1987	Manufacturer/Source/Supplie
(Source URL:	Linolenic Acid, Total		r
https://www.fao.org/gsfaonline/foods/details.html?id=227	Carbohydrates per 100g, Vitamins		
)	and Minerals, Trace Minerals and		
	Other Substances,		
	Lauric/Mystiric/Trans Fatty Acids,		
	Optional Ingredients- suitable for		
	6 months onwards and		
	scientifically proven.		
	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
	FOLLOW-UP FORMULA/MILK		r
	SUPPLEMENT (FROM 6		
	MONTHS INFANTS TO 36		
	MONTHS YOUNG CHILDREN);		
	FORMULA FOR SPECIAL		
	MEDICAL PURPOSES FOR		
	YOUNG CHILDREN:		
	Salmonella/25g, Aerobic Plate		
	Count CFU/g &		
	Enterobacteriaceae/10g		
	☑ Clear and complete loose	<u>Department</u>	Applicant Company/
	labels or artworks compliant with	Circular No. 2008-	Manufacturer/Source/Supplie
	Department Circular 2008-0006.	0006	r
	CEREAL-BASED FOOD	S FOR INFANTS &	YOUNG CHILDREN



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	☑ Valid Certificate of Analysis for Energy, Protein, Carbohydrates, Lipids, Minerals and Vitamins per 100 kcal or 100 kJ	Codex Stan 074- 1981, Rev 1-2006	Applicant Company/ Manufacturer/Source/Supplie r
J2 - Complementary foods for infants and young children "Foods that are intended for infants 6 months of age and	✓ Valid Certificate of Analysis for Microbiological parameters for CEREAL-BASED FOODS FOR INFANTS: Bacillus cereus CFU/g, Clostridium perfringes CFU/g, Aerobic Plate Count CFU/g, Salmonella/25g & Coliforms MPN/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
older, and for progressive adaptation of infants and children to ordinary food. Products may be ready-to-eat or in powder form to be reconstituted with water, milk, or other suitable liquid. Examples include: cereal-, fruit-, vegetable-, and meat-based "baby foods" for infants, "toddler foods," and "junior foods"; lactea flour, biscuits and rusks for children." (Source URL:	✓ Valid Certificate of Analysis for Microbiological parameters for DRIED AND INSTANT PRODUCTS REQUIRING RECONSTITUTION: Coliforms MPN/g, Aerobic Plate Count CFU/g, Salmonella/25g & Listeria monocytogenes/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
https://www.fao.org/gsfaonline/foods/details.html?id=228	✓ Valid Certificate of Analysis for Microbiological parameters for DRIED PRODUCTS REQUIRING RECONSTITUTION AND BOILING BEFORE CONSUMPTION: Coliforms MPN/g, Salmonella/25g & Aerobic Plate Count CFU/g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	☑ Clear and complete loose labels or artworks declaring the statement "Infants six months onwards should be given fresh,	Department Circular No. 2008- 0006	Applicant Company/ Manufacturer/Source/Supplie r



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	indigenous and natural food in combination with continued breastfeeding based on		
	Department Circular 2008-0006.		
	CAN	NNED BABY FOODS	
	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 73- 1981 amended 1989	
	☑ Valid Certificate of Analysis for Microbiological parameters for BABY FOODS IN	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
	HERMETICALLY SEALED CONTAINERS: Commercial Sterility		
	☑ Clear and complete loose labels or artworks declaring the statement "Infants six months onwards should be given fresh, indigenous and natural food in combination with continued breastfeeding based on Department Circular 2008-0006.	Department Circular No. 2008- 0006	Applicant Company/ Manufacturer/Source/Supplie r
J3. Dietetic foods intended for special medical purposes (excluding products under HR Letter J.1.) "Foods for special dietary use that are specially processed or formulated and presented for the dietary	☑ Scientific Studies indicating safety and benefits of the product for intended medical condition	Codex Stan 180- 1991 and Administrative No. Order 2014-0029	Applicant Company/ Manufacturer/Source/Supplie r
management of patients and may be used only under medical supervision. They are intended for the exclusive or partial feeding of patients with limited or impaired	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 180- 1991	Applicant Company/ Manufacturer/Source/Supplie r
capacity to take, digest, absorb or metabolize ordinary foods or certain nutrients contained therein, or who have other special medically-determined nutrient requirement,	☑ Clear and complete loose labels or artworks compliant with Codex Stan 180-1991.	Codex Stan 180- 1991	Applicant Company/ Manufacturer/Source/Supplie r



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whose dietary management cannot be achieved only by modification of the normal diet, by other foods for special dietary uses, or by a combination of the two." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=229)			
J4 - Dietetic formula for slimming purposes and weight reduction "Formula foods that when presented as "ready-to-eat" or	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 181- 1991	Applicant Company/ Manufacturer/Source/Supplie r
when prepared in conformity with the directions for use are specifically presented as replacements for all or part of the total daily diet. Includes products with reduced caloric content such as those that are low in sugar and/or fat, sugar- or fat-free, or contain sugar- and/or fat-substititues." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=230)	☑ Clear and complete loose labels or artworks compliant with Codex Stan 181-1991	Codex Stan 181- 1991	Applicant Company/ Manufacturer/Source/Supplie r
J5 - Dietetic foods (e.g. supplementary foods for dietary use) excluding products under HR Letter J.1 to 4 and Letter K, Food Supplements) "Products of high nutritional content, in liquid or solid	☑ Scientific Studies indicating safety and suitability of the product to specific disease and disorder to which it is intended	Codex Stan 146- 1985 and Administrative No. Order 2014-0029	Applicant Company/ Manufacturer/Source/Supplie r
form (e.g. protein bars), to be used by individuals as part of a balanced diet to provide supplemental nutrition. Products are not intended to be used for purposes of	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 146- 1985	Applicant Company/ Manufacturer/Source/Supplie r
weight loss or as part of a medical regimen." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=231)	☑ Clear and complete loose labels or artworks compliant with Codex Stan146-1985	Codex Stan 146- 1985	Applicant Company/ Manufacturer/Source/Supplie r
J6 - Weaning foods for infants and growing children	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE



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J7 - Dietetic foods for special medical purpose "Foods for special dietary use that are specially processed or formulated and presented for the dietary management of patients and may be used only under	☑ Scientific Studies indicating safety and benefits of the product for intended medical condition	Codex Stan 180- 1991 and Administrative No. Order 2014-0029	Applicant Company/ Manufacturer/Source/Supplie r
medical supervision. They are intended for the exclusive or partial feeding of patients with limited or impaired capacity to take, digest, absorb or metabolize ordinary	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 180- 1991	Applicant Company/ Manufacturer/Source/Supplie r
foods or certain nutrients contained therein, or who have other special medically-determined nutrient requirement, whose dietary management cannot be achieved only by	☑ Clear and complete loose labels or artworks compliant with Codex Stan 180-1991	Codex Stan 180- 1991	Applicant Company/ Manufacturer/Source/Supplie r
modification of the normal diet, by other foods for special dietary uses, or by a combination of the two" (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=229)	✓ Valid Certificate of Analysis for Microbiological parameters for READY-TO-USE THERAPEUTIC FOODS (RUTF) AND READY-TO-USE-SUPPLEMENTARY FOODS (RUFS), 6-59 MONTHS OF AGE: Salmonella/25g	FDA Circular No. 2022-012	Applicant Company/ Manufacturer/Source/Supplie r
J8 - Dietetic formulas for weight control "Formula foods that when presented as "ready-to-eat" or when prepared in conformity with the directions for use	☑ Valid Certificate of Analysis to support Nutrition Information	Codex Stan 181- 1991	Applicant Company/ Manufacturer/Source/Supplie r
are specifically presented as replacements for all or part of the total daily diet. Includes products with reduced caloric content such as those that are low in sugar and/or fat, sugar- or fat-free, or contain sugar- and/or fat-substitutes." (Source URL: https://www.fao.org/gsfaonline/foods/details.html?id=230)	☑ Clear and complete loose labels or artworks compliant with Codex Stan 181-1991	Codex Stan 181- 1991	Applicant Company/ Manufacturer/Source/Supplie r
J - Bottled Water "means water that is placed in a sealed container or packaged and is offered for sale for human consumption as drinking water."	☑ Valid Certificate of Analysis for Physico-Chemical Properties (Turbidity, Color, Odor, Taste, pH, TDS, Conductivity, Calcium.	Administrative Order No. 18-A s. 1993	Applicant Company/ Manufacturer/Source/Supplie r



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(Source: AO No. 18-A s. 1993)	Magnesium, Sodium, Potassium, Chloride, Sulfate), Contaminants (Nitrates, Nitrites, Iron,		
	,		
	manganese, Copper, Zinc,		
	Aluminum, Fluoride, organic		
	Matter, Surfactants), Toxic		
	Contaminants (Arsenic,		
	Cadmium, Cyanide, Chromium,		
	Lead, Mercury, Selenium,		
	Phenolic Substances), Volatile		
	Organic Compounds (Carbon		
	tetrachloride, Benzene,		
	Trihalomethanes), Pesticides &		
	Related Substances		
	(Carbamates, Organochlorines,		
	Organophosphates, Herbicides,		
	Fungicides, PCB), Radionuclides		
	(Gross Alpha Activity, Gross Beta		
	Activity) and Microbiological		
	Parameters (Coliforms, Fecal		
	Strepcocci, Pseudomonas		
	Aeruginosa, HPC)		
	Clear and complete loose labels	Administrative	Applicant Company/
	or artworks compliant with	Order No. 39 s.	Manufacturer/Source/Supplie
	Administrative Order No. 39 s.	<u>1996</u> and	r
	1996 and Administrative Order	Administrative	
	No. 18-A s. 1993.	Order No. 18-As.	
		1993	
K1 - Herbs and botanicals and/or Products with	Shelf-life study with stability data	Administrative	Applicant Company/
other nutritional substances and/or combination as	containing relevant information on	Order No. 2014-	Manufacturer/Source/Supplie
Food Supplement	the critical parameters of the	0029	r
"Includes vitamin and mineral supplements in unit dose	finished product, period		
forms such as capsules, tablets, powders, solutions etc.,	conducted and conclusion		



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where national jurisdictions regulate these products as	Valid Certificate of Analysis of the	<u>Administrative</u>	Applicant Company/
food"	physico-chemical (Vitamins or	Order No. 2014-	Manufacturer/Source/Supplie
(Source URL:	Minerals or Amino Acids or	0029	r
https://www.fao.org/gsfaonline/foods/details.html?id=232	Ingredient Assays) and/or		
	microbiological parameters of the		
,	finished product (whichever is		
"means a processed food product intended to	applicable)		
supplement the diet that bears or contains one or more			
of the following dietary ingredients: vitamin, mineral,	*The amount of Vitamins shall		
herb, or other botanical, amino acid, and dietary	conform with the prescribed level		
substance to increase the total daily intake in amounts	of Office Order No. 22 s 1991		
conforming to the latest Philippine recommended energy	Clear and complete loose labels	Bureau Circular	Applicant Company/
and nutrient intakes or internationally agreed minimum	or artworks declaring the term	No. 2 s 1999	Manufacturer/Source/Supplie
daily requirements. It usually is in the form of capsules,		<u>140. 2 5 1999</u>	wariulacturer/Source/Supplie
1	"Food Supplement" and the		
tablets, liquids, gels, powders or pills and not	phrase "NO APPROVED		
represented for use as a conventional food or as the	THERAPEUTIC CLAIMS" based		
sole item of a meal or diet or replacement of drugs and	on <u>BC 2 S. 1999</u>		
medicines."	Sample in actual commercial	Administrative	Applicant Company/
(Source URL:	presentation	Order No. 2014-	Manufacturer/Source/Supplie
https://www.officialgazette.gov.ph/2009/08/18/republic-		0029	r
<u>act-no-9711/</u>)	*for the procedure on submission,		
	please refer to: IV. Guidelines, C.		
	Procedural Guidelines 2. L. ii.,		
	Pages 7-8 of FDA Circular 2020-		
	033		
	For VIRGIN COCONUT OIL	Bureau Circular	Applicant Company/
	FOOD SUPPLEMENT WITH	2006-018	Manufacturer/Source/Supplie
	FLAVOR:		l r
	1) That the raw material (virgin		
	coconut oil) used conforms with		
	the Philippine National Standards		
	for Virgin Coconut Oil;		
	Tion virgini occornation,	1	



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2) That the flavoring added should be generally recognized as safe and suitable for human consumption as evidenced by a certification from the supplier. The nature of flavor used (natural, nature-identical, artificial) shall be indicated in the list of ingredients; 3) No other food additive shall be allowed except the flavor; 4) The label shall conform with BC 2 s. 1999; 5) The term "Food Supplement" shall be part of the product name ☑ Valid Certificate of Analysis for Microbiological parameters for VIRGIN COCONUT OIL: Aerobic Plate Count CFU/ml, Coliform MPN/ml or CFU/ml, Yeast and Mold Count CFU/ml, Salmonella spp. /25ml and E. coli MPN/ml or CFU/ml For GINKGO BILOBA: 1.) Valid Certificate of Analysis for	FDA Circular No. 2022-012 Bureau Circular No. 02 s. 2004	Applicant Company/ Manufacturer/Source/Supplie
Mold Count CFU/ml, Salmonella spp. /25ml and E. coli MPN/ml or		
For GINKGO BILOBA :		



For TAUFFRO / Day diagon /	Duragu Circular	Applicant Company
For TAHEEBO / Pau d'arco /	Bureau Circular	Applicant Company/
Lapacho:	No. 17 s. 2004	Manufacturer/Source/Supplie
Clear and complete label		r
declaring the precautions:		
1. "This product is not intended to		
diagnose, treat, cure, and prevent		
disease"		
2. "Maximum daily intake up to 3		
cups per day only"		
3. "should not be taken with		
aspirin, ticlopidine, ginkgo biloba,		
ginseng, warfarin & heparin"		
4. "should not be taken by		
pregnant or breast-feeding		
mother"		
5. "should not be taken at least		
one week before contemplated		
operation"		
6. Stop intake of this product in		
the event of nausea, vomiting,		
diarrhea, skin pallor, bruises and		
nose bleeding.		
For PROBIOTICS WHICH	Bureau Circular	Applicant Company/
BACTERIAL STRAINS NOT	No. 16 s. 2004	Manufacturer/Source/Supplie
FOUND IN THE ACCEPTABLE		r
LIST shall be subject to (1)		
demonstration of evidence of safe		
use as food supplement and (2)		
analysis of the bacterial species		
found in formulation. Likewise,		
BFAD shall use as reference:		
WHO-FAO "Guidelines for the		



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Evaluation of Probiotics in Food" (2002). A. The BFAD also would like to inform everyone concerned that, for a Probiotic to the effective, the following properties should be demonstrated: a. beneficial effect on the host organism b. should be able to survive in the digestive tract c. should adhere to the mucosal epithelial cells d. should exhibit enhancement		
•		
d. Epidemiological surveillance of adverse incidents in consumers (post-market)		



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	e. If the strain under evaluation		
	belongs to a species that is a		
	known mammalian toxin producer,		
	it must be tested for toxin		
	production. One possible scheme		
	for testing toxin production has		
	been recommended by the EU		
	Scientific Committee on Animal		
	Nutrition (SCAN, 2000)		
	f. If the strain under evaluation		
	belongs to a species with known		
	hemolytic potential,		
	determination of hemolytic activity		
	is required.		
K2 - Herbs and botanicals and/or Products with	☑ Valid Certificate of Analysis for	FDA Circular No.	Applicant Company/
other nutritional substances and/or combination as	Microbiological parameters for	<u>2022-012</u>	Manufacturer/Source/Supplie
Conventional Food Product	NON-ALCOHOLIC BEVERAGES		r
e.g. Powdered Juice with marine collagen, coffee	(e.g. READY TO DRINK,		
powder with barley grass, tongkat ali and royal jelly	SOFTDRINKS, ICED TEA,		
	ENERGY DRINKS, JELLY		
	DRINKS) : Yeast and Mold Count		
	CFU/mL, Coliforms CFU/mL &		
	Aerobic Plate Count CFU/mL		
	☑ Valid Certificate for	FDA Circular No.	Applicant Company/
	Microbiological parameters for	2022-012	Manufacturer/Source/Supplie
	POWDERED BEVERAGES (e.g.		r
	ICED TEA, POWDERED		
	JUICES/MIXES): Aerobic Plate		
	Count CFU/g, Yeast and Mold		
	Count CFU/g & Coliforms CFU/g		



L. New in the international or local market/Other New Products/Unclassified or Unlisted in A.O. 2014-0029 Annex A	NOT APPLICABLE	NOT APPLICABLE	NOT APPLICABLE
FOOD PRODUCTS CONTAINING TRANS-FATTY ACIDS (TFA) FDA Circular 2021-028, FDA Circular No.2021-028-A	 ☑ technical specifications of raw materials indicating specific oil(s) and/or fat(s) used and the processing it underwent; ☑ recent (within 12 months from date of application) certificate of analysis of the finished product from an accredited laboratory of the FDA and Philippine Accreditation Board/Office (PAB/PAO) or from the country of origin (for imported products), reflecting the TFA content per 100g or ml, validated reference methods of analysis, and the limit of detection for the method used in the analysis of TFA; and ☑ for prepackaged processed food containing naturally-occurring TFA of more than 2g TFA per 100g or ml of the total fat, recent (within 12 months from date of application) certificate of analysis showing that the TFA is naturally-occurring and/or obtained from ruminant animal, from an accredited laboratory of 	FDA Circular No. 2021-028 FDA Circular No. 2021-028-A FDA Circular No. 2020-033-B Administrative Order No. 2021- 0039	Applicant Company/ Manufacturer/Source/Supplie r



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the FDA and Philippine	
Accreditation Board/Office	
(PAB/PAO) or from the country of	
origin, with validated reference	
method of analysis and the limit of	
detection for the method used in	
the analysis.	
	Accreditation Board/Office (PAB/PAO) or from the country of origin, with validated reference method of analysis and the limit of detection for the method used in

FOR AMENDMENT DATA CAPTURE

DATA CAPTURE in the modified E-Registration System refers to applications processed in the old E-Registration System (Version 1) or thru manual registration system.

manual registration system.				
GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE		
 ☑ Accomplished Application Form as prescribed by FDA regulations e.g. E-Registration System 	FDA Circular No. 2020-033 FDA Circular No. 2020-033-A	https://www.fda.gov.ph/		
☑ Proof of Payment of Fees as prescribed by current FDA regulations.	Administrative Order No. 50 s. 2001	Systems/Means prescribed by FDA		
☑ Scanned Application Letter stating the intended changes (indicate ALL the changes/amendments to be made)	Administrative Order No. 2014-0029 FDA Circular No. 2020-033 FDA Circular No. 2020-033-A	Applicant Company/ Manufacturer/Source/Supplier		
✓ VALID AND APPROPRIATE FDA LICENSE TO OPERATE (LTO) (REQUIRED FOR ALL TYPES OF CPR APPLICATION) *The product being applied must be listed in the FDA approved Product Line/Category.	Administrative Order No. 2014-0029 FDA Circular No. 2020-033	Applicant Company/ Manufacturer/Source/Supplier		
☑ Upload ALL INITIAL requirements if previously approved application is in the old E-Registration System (Version 1) or thru manual registration system	FDA Circular No. 2020-033 FDA Circular No. 2020-033-A	Applicant Company/ Manufacturer/Source/Supplier		
☑ Additional Requirements per Amendment Type. Please refer to TITLE OF	Administrative Order No. 2014-0029 FDA Circular No. 2020-033 FDA Circular No. 2020-033-A	Applicant Company/ Manufacturer/Source/Supplier		



CERTIFICATION/PERMIT: CERTIFICATE OF PRODUCT REGISTRATION (CPR) – AMENDMENT (INITIAL APPLICATION	
APPROVED FROM MODIFIED E- REGISTRATION (VERSION 2) - III.	
ADDITIONAL Requirements per Amendment	
Туре.	

FOR RE-APPLICATION DATA CAPTURE

DATA CAPTURE in the modified E-Registration System refers to applications processed in the old E-Registration System (Version 1) or thru manual registration system.

manual registration system.				
GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE		
☑ Accomplished Application Form as	FDA Circular No. 2020-033	https://www.fda.gov.ph/		
prescribed by FDA regulations	FDA Circular No. 2020-033-A			
e.g. E-Registration System				
☑ Upload ALL INITIAL requirements AND	Administrative Order No. 2014-0029	Applicant Company/ In reference to the		
compliance to the deficiencies stated in the	FDA Circular No. 2020-033	previously filed and disapproved INITIAL		
previously issued Letter of Denial (LOD) within	FDA Circular No. 2020-033-A	application		
6 months upon receipt of LOD.				
☑ Proof of Payment of Fees as prescribed by	Administrative Order No. 50 s. 2001	Systems/Means prescribed by FDA		
current FDA regulations.				

FOR RENEWAL DATA CAPTURE (REGULAR)

DATA CAPTURE in the modified E-Registration System refers to applications processed in the old E-Registration System (Version 1) or thru manual registration system.

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GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE	
 ☑ Accomplished Application Form as prescribed by FDA regulations e.g. E-Registration System 	FDA Circular No. 2020-033 FDA Circular No. 2020-033-A	https://www.fda.gov.ph/	



☑ VALID AND APPROPRIATE FDA LICENSE	Administrative Order No. 2014-0029	Applicant Company/
TO OPERATE (LTO) (REQUIRED FOR ALL	FDA Circular No. 2020-033	Manufacturer/Source/Supplier
TYPES OF CPR APPLICATION)		
*The product being applied must be listed in		
the FDA approved Product Line/Category.		
☑ Upload ALL INITIAL requirements if	Administrative Order No. 2014-0029	Applicant Company/
previously approved application is in the old E-	FDA Circular No. 2020-033	Manufacturer/Source/Supplier
Registration System (Version 1) or thru manual	FDA Circular No. 2020-033-A	
registration system		
☑ Proof of Payment of Fees as prescribed by	Administrative Order No. 50 s. 2001	Systems/Means prescribed by FDA
current FDA regulations.		

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1.1. Accomplishes (including uploading of the COMPLETE	Pre-assesses ONLY the completeness of the submitted documents through E-Registration		Center for Food Regulation and Research (CFRR) PRE-
documentary requirements) the E- Registration System through the E-	System/E-Portal https://eportal.fda.gov.ph .		ASSESSOR (e.g. Food-Drug Regulation Officer
Portal https://eportal.fda.gov.ph based on the desired type of	Result of Pre-assessment will be received by the account holder.		(FDRO))
application in accordance to current FDA regulation/s on the use of the			
E-Registration Portal/E-Services.			
1.2. Forwards the application to PRE-ASSESSMENT .			
A system generated E-mail notification from FDA will be			
received by the client upon submission of application for Pre-Assessment.			



(If COMPLETE) Receives the Order of Payment. (If INCOMPLETE) Receives result of Pre-Assessment (Letter of Denial)	2. If found COMPLETE Generates Order of Payment through the email of the account holder/client. If found INCOMPLETE , Generates result of Pre-Assessment. To refile, the applicant must start a NEW CASE and upload initially submitted documentary requirements together with the documents for compliance to the deficiencies mentioned. For Food Supplement application, the proof of submission of sample can be re-uploaded to the new application.		CFRR PRE-ASSESSOR (e.g. FDRO)
3. Pays the assessed fee through Systems/Means prescribed by FDA.	 3.1. Receives the payment/Official Receipt (OR)/ proof of payment through Systems/Means prescribed by FDA, and then posts the payment. 3.2. Forwards application to CFRR, once payment is posted. 	Refer to FDA Cashier 's Citizen Charter	Administrative and Finance Services (AFS) STAFF
4. Receives Acknowledgement Receipt with the application and preassessment details.	4.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	8 Working Days	LRD EVALUATOR (e.g. FDRO)



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	4.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	7 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)		
	4.3. Reviews the checked application, ALL the submitted documentary requirements, and the drafted recommendation of the CHECKER, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, and then finalizes the application by issuing Certificate of Product Registration (CPR) (for APPROVED application) or Letter of Denial (LOD) (for DISAPPROVED application), through the E-Registration System.	5 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)		
5. If the application is APPROVED , receives Certificate of Product Registration (CPR), and other pertinent information. If DISAPPROVED , receives an email notification from FDA regarding the issuance of Letter of Denial/Disapproval (LOD), and other pertinent information.	5. Generates electronically signed CPR or LOD.		Information and Communication Technology Management Division (ICTMD) STAFF		
		TOTAL: 20			
Always refer to the current EDA regula	 htion/s on the use of the E-Registration System/E S	Working Days	fda gov ph/		
Always refer to the current FDA regulation/s on the use of the E-Registration System/E-Services: https://www.fda.gov.ph/					



1.5. ISSUANCE OF CERTIFICATE OF PRODUCT REGISTRATION (CPR) – RE-APPLICATION (INITIAL APPLICATION DISAPPROVED FROM MODIFIED E-REGISTRATION)

'Registration' means the process of approval of an application to register health products prior to engaging in the manufacture, importation, exportation, sale, offer for sale, distribution, transfer, and where applicable, the use, testing, promotion, advertisement, and/or sponsorship of health products. (Republic Act No. 9711)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction		Highly Technical
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	In accordance to Administrative Order 50 s. 2001 + Legal Research Fee (LRF).
		Re-application Fee PhP 200.00 + 1% LRF

GENERAL GUIDELINES

Please refer to:

- 1) A. General Guidelines, B. Specific Guidelines, and C. Procedural Guidelines, IV. GUIDELINES, pages 2-10 of <u>FDA Circular No. 2020-033</u> || Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products"; and
- 2) III. General Guidelines, and IV. Specific Guidelines of <u>FDA Circular No. 2020-033-A</u> || Addendum to FDA Circular No. 2020-033, "Procedure for the Use of the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged Processed Food Products" to include Guidelines for Preassessment and Reiteration of Pre-Assessment Procedures in Applying for Certificate of Product Registration for Food

CHECKLIST OF REQUIREMENTS FOR ALL TYPES OF FOOD PRODUCTS/FOOD CATEGORIZATION:

RAW MATERIALS, LOW RISK, MEDIUM RISK AND HIGH RISK FOOD PRODUCTS

GENERAL REQUIREMENTS	BASIS/ISSUANCE	WHERE TO SECURE
☑ Accomplished Application Form as prescribed by current regulations.		https://www.fda.gov.ph/ 1) For the Certificate of Analysis: a) Applicant Company/



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Through the E-Registration System, upload/attach the compliance to the deficiencies stated in the previously issued		Manufacturer/Source/Supplier; or b) Laboratory analysis issued/conducted by FDA
Letter of Denial (LOD) within 6 months upon receipt of LOD, using the same case number.		accredited laboratories.
receipt of LOD, using the same case number.		2) For other technical document(s): a) Applicant Company/ Manufacturer/Source/Supplier
☑ Proof of payment of fees as prescribed by current FDA regulations.	Administrative Order 50 s. 2001	Systems/Means prescribed by FDA
☑ Valid and appropriate FDA License to Operate (required for all types of CPR application) *The product being applied must be listed in the FDA approved Product Line/Category.	Administrative No. Order 2014-0029 Republic Act No. 9711	FDA Philippines

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1.1. Files using the specific product/CASE NUMBER in the INBOX folder, and then accomplishes (including uploading of the COMPLETE documentary requirements) the E-Registration System through the E-Portal https://eportal.fda.gov.ph based on the desired type of application in accordance to current FDA regulation/s on the use of the E-Registration Portal/E-Services. 1.2. Forwards the application to PRE-ASSESSMENT.	Pre-assesses ONLY the completeness of the submitted documents through E-Registration System/E-Portal https://eportal.fda.gov.ph . Result of Pre-assessment will be received by the account holder.	Day 0	Center for Food Regulation and Research (CFRR) PRE- ASSESSOR (e.g. Food-Drug Regulation Officer (FDRO))



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A system generated E-mail notification from FDA will be received by the client upon submission of application for Pre-Assessment.			
2. (If COMPLETE) Receives the Order of Payment.	2. If found COMPLETE , Generates Order of Payment through the email of the account holder/client.	Day 0	CFRR PRE- ASSESSOR (e.g. FDRO)
(If INCOMPLETE) Receives result of Pre- Assessment (Letter of Denial)	If found INCOMPLETE, Generates result of Pre-Assessment. To refile, the applicant must start a NEW CASE and upload initially submitted documentary requirements together with the documents for compliance to the deficiencies mentioned.		
3. Pays the assessed fee through Systems/Means prescribed by FDA	3.1. Receives the payment/Official Receipt (OR)/ proof of payment through Systems/Means prescribed by FDA, and then post the payment.3.2. Forwards application to CFRR, once payment is	Day 0 Refer to FDA Cashier 's Citizen Charter	Administrative and Finance Services (AFS) STAFF
	posted.		
4. The applicant company receives Acknowledgement Receipt with the application and pre-assessment details.	4. 1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then will forward the same to the CHECKER.	Day 0 8 Working Days	LRD EVALUATOR (e.g. FDRO)
	4.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is	7 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)



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	for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.		
	4.3. Reviews the checked application, ALL the submitted documentary requirements, drafted recommendation of the CHECKER, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, and then finalizes the application by issuing Certificate of Product Registration (CPR) (for APPROVED application) or Letter of Denial (LOD) (for DISAPPROVED application), through the E-Registration System.	5 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
5. If the application is APPROVED , Receives an e-mail notification from FDA indicating that the application is approved, and other pertinent information. If DISAPPROVED , receives a Letter of Denial/Disapproval (LOD), and other pertinent information.	5. Generates electronically signed CPR or LOD.		Information and Communication Technology Management Division (ICTMD) STAFF
		TOTAL: 20 Working Days	
Always refer to the current FDA regulation/	」 s on the use of the E-Registration System/E-Services: <u>https:</u> ,		



2. ISSUANCE OF DIAMOND SANGKAP PINOY SEAL

Diamond Sangkap Pinoy Seal – refer to the seal of good nutrition quality that will be awarded as an incentive to BFAD (FDA) registered staple manufacturer who will fortify their products according to standards. (Administrative Order No. 82 s. 2003)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	••	Government to Business
Type of Transaction	••	Highly Technical Transaction
Who May Avail	••	All FOOD Manufacturers of Fortified Products
Fees to be Paid		P8,000.00 non-refundable fee for the use of the seal (Regular Seal)
rees to be raid	•	P500.00 processing fee for every application (Regular Seal or Diamond Seal) + 1% LRF

CHECKLIST OF REQUIREMENTS Submit ONE (1) scanned copy of the required document.	WHERE TO SECURE
☑ Basic Requirements based on RA No. 8976 (Food Fortification Law of 2000), RA No. 8172 (ASIN Law), the Sangkap Pinoy Seal Program Manual of Operations (December 2000) and Administrative Order No. 82 s. 2003 (Guidelines on the Granting of Diamond Sangkap Pinoy Seal to Manufacturers of Fortified Products):	https://www.fda.gov.ph/
☑ Duly accomplished application forms	FDA Philippines
☑ Copy of valid and appropriate LTO issued by the FDA	FDA Philippines
☑ Results of product analysis for vitamin A, Iron and Iodine from an FDA recognized laboratory.	For the Certificate of Analysis: Laboratory analysis issued/conducted by FDA accredited laboratories.
☑ Sample label with Diamond Sangkap Pinoy Seal	Applicant Company/ Manufacturer/Source/Supplier
☑ Proof of payment	Systems/Means prescribed by FDA
☑ Inspection report with Certificate of Compliance	FDA Regional Field Office



CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
	Receives document requirements from FDA Regional Field Office and decks the same to CFRR technical evaluators.	1 Working Day	LICENSING AND REGISTRATION DIVISION (LRD) EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	8 Working Days	LRD EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	5 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)
	2.3. Prints the authorization.	1 Working Day	CFRR ADMINISTRATIVE STAFF
	2.4. Signs the Certification/Authorization.	4 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
	3. Forwards the Certificate/Authorization to the Office of Director General, for signature.	1 Working Day	CFRR ADMINISTRATIVE STAFF
		TOTAL: 20 Working Days	



3. ISSUANCE OF E-REGISTRATION PORTAL USER ACCOUNT

The applicant shall be assigned an FDA account to apply through the E-Registration System.

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification		Government to Business
Type of Transaction		Simple
Who May Avail		All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters)
Fees to be Paid	:	NONE

CHECKLIST OF REQUIREMENTS	WHERE TO SECURE
GENERAL GUIDELINES	https://www.fda.gov.ph/
Please refer to:	
C. Procedural Guidelines, IV. GUIDELINES, pages 5-6 of FDA Circular No. 2020-033 Procedure for the Use of	
the Modified Electronic Registration System for Raw Materials and Prepackaged Processed Food Products	
Repealing FDA Circular No. 2016-014 "Procedure for the Use of Electronic Registration System for Prepackaged	
Processed Food Products"	
ISSUANCE OF CFRR E-REGISTRATION USER ACCOUNT	
☑ Send a request for a user account to cfrr@fda.gov.ph	Applicant Company
SUBJECT: CFRR: E-Registration	
BODY:	
Email Address:	
Last Name:	
First Name:	
Middle Name:	
Company Name:	
LTO No.:	
LTO validity:	



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☑ The email must contain an attached scanned copy of notarized authorization letter (please see Annex B of FDA	Applicant Company
Circular No. 2020-033) from a company with a valid License-to-Operate (LTO).	
CHANGE IN THE APPLICANT COMPANY'S REPRESENTATIVE	Applicant Company
☑ Send a request for change in credentials of the CFRR E-Registration User Account to cfrr@fda.gov.ph	Applicant Company
SUBJECT: CFRR: E-Registration	
COBCCOTT OF TAX. E Programation	
BODY:	
Email Address:	
Last Name:	
First Name:	
Middle Name:	
Company Name:	
LTO No.:	
LTO validity:	
☑ The email must contain an attached scanned copy of notarized Affidavit of Undertaking (please see Annex C of	Applicant Company
FDA Circular No. 2020-033) from a company with a valid License-to-Operate (LTO).	
RENEWAL OF USER ACCOUNT AT LEAST 90 DAYS PRIOR TO EXPIRATION	Applicant Company
☑ Send a request for renewal of user account to cfrr@fda.gov.ph	Applicant Company
SUBJECT: CFRR: E-Registration	
BODY:	
Email Address:	
Last Name:	
First Name:	
Middle Name:	
Company Name:	



	PHILIPPINES
LTO No.:	
LTO validity:	
ISSUED USER ACCOUNT BY THE FDAC FOR E-LTO CAN BE REVALIDATED TO ACCESS E-REGISTRATION	Applicant Company
☑ Send a request for revalidation of user account to cfrr@fda.gov.ph	Applicant Company
SUBJECT: CFRR: E-Registration	
BODY:	
Email Address:	
Last Name:	
First Name:	
Middle Name:	
Company Name:	
LTO No.:	
LTO validity:	
RETRIEVAL OF USER NAME AND/OR PASSWORD OF E-REGISTRATION ACCOUNT (IN CASES OF	Applicant Company
PROBLEMS WITH USER NAME AND/OR PASSWORD)	
☑ Send a request for retrieval of user name and/or password to cfrr@fda.gov.ph	Applicant Company
SUBJECT: CFRR: E-Registration	
BODY:	
BOD1.	
Email Address:	
Last Name:	
First Name:	
Middle Name:	
Company Name:	
LTO No.:	



LTO validity:

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1. Submits required documents/information to the above-mentioned e-mail address.	 1.1. Checks the e-mail request. 1.2. If compliant, user name and password will be issued to the client, via e-mail. Otherwise, the personnel will send an e-mail to the applicant company/authorized representative to request for lacking document(s) or clarify information. 	3 Working Days	Food Drug Action Center (FDAC) or Center for Food Regulation and Research (CFRR) STAFF



4. ISSUANCE OF GOOD MANUFACTURING PRACTICES (GMP) CERTIFICATE

Good manufacturing practices refer to a quality assurance system aimed at ensuring that products are consistently manufactured, packed, repacked or held to quality standards appropriate for the intended use. It is thus concerned with both manufacturing and quality control procedure. (Republic Act No. 10611)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	••	Highly Technical Transaction
Who May Avail	••	All FOOD Manufacturers (Importer of raw material for own use/Exporters)
Fees to be Paid		GMP – Php 500.00 + LRF per year

CHECKLIST OF REQUIREMENTS Submit ONE (1) scanned copy of the required document.	WHERE TO SECURE
☑ Inspection report with certificate of Compliance/ Recommendation Letter from RFO	FDA Regional Office
☑ Copy of valid and appropriate LTO issued by the FDA	FDA Philippines
☑ Proof of payment	Systems/Means prescribed by FDA

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
	Receives document requirements from FDA Regional Field Office and decks the same to CFRR technical evaluators.	1 Working Day	LICENSING AND REGISTRATION DIVISION (LRD) EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or	8 Working Days	LRD EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))



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	Disapproval, and then forwards the same to the CHECKER.		
	2.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	5 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)
	2.3. Prints the authorization.	1 Working Day	CFRR ADMINISTRATIVE STAFF
	2.4. Signs the Certification/Authorization.	4 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
3. Receives the Certificate/Authorization.	3. Forwards the Certificate/Authorization to Food and Drug Action Center (FDAC) for release of Records Section.	1 Working Day	CFRR ADMINISTRATIVE STAFF
		TOTAL: 20 Working Days	



5. ISSUANCE OF HAZARD ANALYSIS AND CRITICAL CONTROL POINTS (HACCP) CERTIFICATE

Hazard Analyses at Critical Control Points (HACCP) refer to a science-based system which identities, evaluates and controls hazards which are significant for food safety at critical points during a given stage in the food supply chain. (Republic Act No. 10611)

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical Transaction
Who May Avail	:	All FOOD Manufacturers (Importer of raw material for own use/Exporters)
Fees to be Paid	:	HACCP – Php1,000.00 + LRF per year

CHECKLIST OF REQUIREMENTS Submit ONE (1) scanned copy of the required document.	WHERE TO SECURE
☑ Inspection report with certificate of Compliance/ Recommendation Letter from RFO	FDA Regional Office
☑ Copy of valid and appropriate LTO issued by the FDA	FDA Philippines
☑ Proof of payment	Systems/Means prescribed by FDA

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
	Receives document requirements from FDA Regional Field Office and decks the same to CFRR technical evaluators.	1 Working Day	LICENSING AND REGISTRATION DIVISION (LRD) EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))



	2.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	8 Working Days	LRD EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	5 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)
	2.3. Prints the authorization.	1 Working Day	CFRR ADMINISTRATIVE STAFF
	2.4. Signs the Certification/Authorization.	4 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
3. Receives the Certificate/Authorization.	3. Forwards the Certificate/Authorization to Food and Drug Action Center (FDAC) for release of Records Section.	1 Working Day	CFRR ADMINISTRATIVE STAFF
		TOTAL: 20 Working Days	



6. ISSUANCE OF IMPORT PERMIT

Import permit is the authorization issued by the FDA to an establishment to import a prepackaged processed food, bulk food and raw materials in the Philippines for the purpose of research and development and shall not be intended for market testing purposes and donated food products.

Center/Office/Division	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Simple Transaction
Who May Avail	:	All FOOD Manufacturers/Traders/Distributors (Importers/ Wholesalers/ Exporters) and Donee/Consignee
Fees to be Paid	:	In accordance with Administrative Order No. 50 s. 2001 Import Permit: Php 500.00/invoice + 1% LRF

CHECKLIST OF REQUIREMENTS	WHERE TO SECURE
Submit ONE (1) scanned copy of the required document.	
FOR RELEASE OF SAMPLES:	
	No specific format, this document is initiated by applicant
☑ Application Letter	company
☑ Notarized Affidavit of Undertaking	See sample template (Annex A)
☑ Certificate of Analysis/ Certificate of Free Sale	Country of Origin or Source of Product to be imported
☑ Pro Forma Invoice	Product Source/company
☑ Packing List	Product Source/company
☑ Bill of Lading/Airway Bill (if available)	Courier or Shipping company
☑ Valid License to Operate	FDA Issued
	FDA Cashier/Other FDA Authorized Payment Portals or
□ Payment (Php 510.00/inclusive of 1% LRF)	Banks
FOR RELEASE OF DONATED FOOD:	
☑ BIHC Endorsement Letter	BIHC of DOH (The Director)



	* Please refer to DOH Administrative Order 2020-0001) for
	the requirement to secure BHIC endorsement.
☑ Letter request from Donee	From Donee
☑ Certificate of Quality (should reflect the expiration or last recommended date of consumption) / Certificate of Free Sale	Product Source/Company
☑ Certificate of Donation	From Donor
☑ Deed of Acceptance	From Donee
☑ Invoice Packing List	From product source/company
☑ Bill of Lading/Airway Bill (if available)	Courier or shipping company
	FDA Cashier/Other FDA Authorized Payment Portals or
☑ Payment (Php 510.00/inclusive of 1% LRF)	Banks

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1. Submits documents through email to the Food and Drug Action Center (FDAC).	Receives the submitted documents.	Day 0	Food Drug Action Center (FDAC) (e.g. Information Officer I or II)
Receives the Document Tracking number as reference for payment.	2. Issues an Acknowledgement Receipt and 14-digit Document Tracking Number (DTN) as reference of the applicant.	Day 0	Food Drug Action Center (FDAC) (e.g. Information Officer I or II)
3. Pays the assessed fee through any FDA Authorized means (e.g. Landbank LinkBiz). (Php 510.00/Invoice).	3. Receives the complete documents and proof of payment through automated transaction.	Day 0	Food Drug Action Center (FDAC) (e.g. Information Officer I or II)
4. Sends the proof of payment to FDAC through email.	4.1. Receives the proof of payment and updates the FDA FIS.	Day 0	Food Drug Action Center (FDAC) (e.g. Information Officer I or II)



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	4.2. Verifies and posts the payment through FDA	Day 0	Administrative and
	FIS.	4.11	Finance Services (AFS)
	4.3. Forwards the application to CFRR receiving,	4 Hours	Food Drug Action Center
	and also updates the FIS indicating that the		(FDAC) (e.g. Information
	application is transmitted to CFRR.	4.11	Officer I or II)
	4.4. Receives application and updates the FIS	4 Hours	Center for Food
	indicating that the application is forwarded to		Regulation and Research
	assigned CFRR evaluator.		(e.g. Administrative
	4.5. 5	4.1.1	Assistant III)
	4.5. Evaluates the correctness of documents and	4 Hours	Center for Food
	updates the FIS indicating that the application is		Regulation and Research
	forwarded to checker for quality assurance.		(e.g. Food-Drug
			Regulation Officer (FDRO
	4.0.01 1:60	4.11	II or III)
	4.6. Checks if the recommendation is	4 Hours	Center for Food
	appropriate/accurate, and updates the FIS indicating		Regulation and Research
	that the application is forwarded to the Center		(e.g. Senior FDRO or
	Director.	4.1.1	Division Chief)
	4.7. Renders the final decision on the	4 Hours	Center for Food
	recommendation and updates the FIS.		Regulation and Research
			Approving Authority
5 D : U MADORT REPAIR	5.5	4.1.1	(e.g. Director IV)
5. Receives the IMPORT PERMIT.	5. Forwards the Permit/Authorization to Records	4 Hours	Center for Food
	section for release and updates the FIS indicating		Regulation and Research
	the same.		(e.g. Administrative Aide
		TOTAL OVAL	VI)
		TOTAL: 3 Working	
		Days	



7. ISSUANCE OF LAW ENFORCEMENT AGENCY (LEA) REQUEST FOR PRODUCT/ LICENSE-TO-OPERATE VERIFICATION THROUGH THE REGULATORY ENFORCEMENT UNIT

Verification of the authorization (i.e., License-to-Operate and Certificate of Product Registration) of the establishment and products as requested by the Law Enforcement Agency in line with an ongoing investigation.

Center/Office/Division	:	Center for Food Regulation and Research (CFRR) – Food Safety Unit (FSU)
Classification	:	Government to Government (G2G)
Type of Transaction	••	Highly Technical Transaction
Who May Avail	••	FDA Center - Regulatory Enforcement Unit (REU)
Fees to be Paid		None

CHECKLIST OF REQUIREMENTS Submit ONE (1) scanned copy of the required document.	WHERE TO SECURE
☑ Letter Request for Product Verification/ License-to-Operate Verification/ Food Products from Law Enforcement Agencies	Requesting Party
☑ Output Documents (Verification Report)	Food Safety Technical Staff
☑ Technology (Internet, Printer, Computer)	Office

INTERNAL CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1. Creates referral for the received Product Verification/ License-to- Operate Verification Letter Request from Law Enforcement Agencies (LEAs)		Day 0	Regulatory Enforcement Unit Staff (Requesting Office)



1.1 Receives, double-checks the completeness of the documents/ samples referred and decks referral.	1 Working Day	Food Safety Unit (FSU) Administrative Staff
1.2 Verifies the status of License to Operate of the Establishment / Registration of the Food Product and/or Food Supplement.	15 Working Days	FSU Evaluator (e.g. Food-Drug Regulation Officer (FDRO))
1.3 Reviews the Information and Recommendation of the Evaluator, and forwards the Referral Report to the OIC, Food Safety Unit for Quality assurance.	2 Working Days	FSU Checker (e.g. Senior Food-Drug Regulation Officer)
1.4 Checks the Referral Report for Quality Assurance, and then forwards the Referral to the CFRR Director.	1 Working Day	FSU, Officer In-Charge (OIC)
1.5 Checks and signs the Final Referral for release.	1 Working Day	Center for Food Regulation and Research (CFRR) Approving Authority (e.g. DIRECTOR IV)
1.6 Mails the final referral to the requesting Law Enforcement Agency		CFRR STAFF
	TOTAL: 20 Working Days	



8. ISSUANCE OF SALES PROMO PERMIT (INITIAL AND AMENDMENT APPLICATION)

Authorization issued for activities conducted by the companies which is intended for broad consumer participation which contains promises of gain such as prizes, in cash or in kind, as reward for the purchase of a product, security, service or winning in a contest, game, tournament, and other similar competitions which involves determination of winner/s and which utilize mass media or other widespread means of information. It is also issued for activities purely intended to increase the sales, patronage and/or goodwill of a product.

Center/Office/Di vision	:	Center for Food Regulation and Research (CFRR)
Classification	••	Government to Business
Type of Transaction	:	Complex Transaction
Who May Avail	:	Food Manufacturers, Importers, Exporters, Wholesalers/Distributors and Third Party Marketing Agencies
Fees to be Paid		In accordance to DTI-DOH JAO NO. 1 s. 2000 Amount of Prizes: (Fees) Php 150,000.00- below Php 300,000.00: Php 1,000.00.00 + 1% LRF Php 300, 001.00-Php 500,000.00: Php 2,000.00 + 1% LRF Php 500,001.00- Php 1,000,000.00: Php 3,000.00 + 1% LRF Above Php 1,000.000.00: Php 5,000.00 + 1% LRF Coverage: (Fees) NCR only or in several regions in NCR and Nationwide: Php 1,000.00.00 + 1% LRF More than one (1) region in NCR and Nationwide: Php 750.00 + 1% LRF Several provinces/cities/municipalities within a single region: Php 500.00 + 1% LRF Single province/city/municipality: Php 250.00 + 1% LRF Amendment/Extension: Php 300.00 + 1% LRF

CHECKLIST OF REQUIREMENTS	WHERE TO SECURE
Submit ONE (1) scanned copy of the required document.	



INITIAL APPLICATION	
☑ Integrated Application Form	https://www.fda.gov.ph/
☑ Completely and accurately filled-up Information Sheet and Mechanics of Sales Promotion	https://www.fda.gov.ph/
☑ Photocopy of valid Certificate of Product Registration (CPR) and Cosmetic Notification (NN)of the company	FDA Issued
☑ Advertising/Collateral Materials to be used in the promotion, if any	Applicant Company
	FDA Cashier/Other FDA Authorized Payment Portals or Banks (where
☑ Proof of Payment of Fees	payment was made)
AMENDMENT APPLICATION	
☑ Integrated Application Form	https://www.fda.gov.ph/
☑ Letter of Intent stating the desired changes	Applicant Company
☑ Photocopy of Approved Permit	FDA Issued
☑ Additional Advertising/Collateral Materials to be used in Promotion if any	Applicant Company
☑ Proof of Payment of Fees	FDA Cashier/Other FDA Authorized Payment Portals or Banks (where payment was made)



SALES PROMO PERMIT (INITIAL AND AMENDMENT APPLICATION) PROCESS FLOW based on <u>FDA Circular No.2021-013</u>: Interim Guidelines of the Center for Food Regulation and Research (CFRR) for the Application and Receiving of Sales Promo Permit Applications in Compliance to the Republic Act No. 11032 otherwise known as The Ease of Doing Business and Efficient Government Service Delivery Act Of 2018 or current FDA regulation.

CLIENT STEPS	AGENCY ACTION	PROCESSING TIME	PERSON RESPONSIBLE
1. Requests for DTN and schedule of submission for pre-assessment to Food and Drug Action Center (FDAC) through email.	1. Provides the DTN and schedule of submission for preassessment through email to the client.	Day 0	Food Drug Action Center (FDAC)
2. Submits documents for pre-assessment through email to Center for Food Regulation and Research (CFRR) on their assigned schedule.	2. Pre-assesses the completeness and correctness of the submitted documents.	Day 0	CFRR EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO)
3. Receives an email to proceed with the payment and must pay through any FDA Authorized means (e.g. Landbank LinkBiz) or an email stating the deficiency/ies noted on the documents for the client to comply.	3. If complete and correct, Sends an email stating that the company can proceed with the payment will be sent to the email address of the authorized representative. A CFRR pre-assessment slip will also be attached on the email. Otherwise, an email stating the deficiency/ies noted on the documents for the client to comply and they will be advice to secure another DTN and schedule.	Day 0	CFRR STAFF
4. Pays the indicated fee as per Integrated Application Form through any applicable payment system prescribed by FDA.	4.1 Verifies and posts the payment through updating the FDA FIS.	Refer FDA Cashier Citizen's Charter	Administrative and Finance Services (AFS) STAFF



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	4.2. Forwards the application to CFRR and updates the FIS indicating the same.	1 Working Day	FDAC STAFF
	4.3. Receives the Sales Promo Permit Application, decks the application to the assigned evaluator, and updates the FIS indicating the same.	1 Working Day	CFRR STAFF
	4.4. Evaluates the consistency of the documents submitted during the pre-assessment stage and the documents received from FDAC, and then forwards the application to the Checker and updates the FIS indicating the same.	1 Working Day	CFRR EVALUATOR (e.g. FDRO)
	4.5. Checks if the recommendation is appropriate and updates the FIS indicating that the application is forwarded to the Center Director.	1 Working Day	CFRR CHECKER (e.g. SENIOR FDRO or DIVISION CHIEF)
	4.6. Renders the final decision on the recommendation and updates the FIS.	1 Working Day	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
	4.7. Forwards the Sales Promotion Permit to FDA Records section for release, and updates the FIS indicating the same	1 Working Day	CFRR STAFF
5. Receives the Certificate/Authorization through courier or pick-up.	5. Updates the status via FIS and release the Certificate/Authorization through courier or pick-up	1 Working Day	Releasing Section Staff
		TOTAL: 7 working days	



9. ISSUANCE OF SANGKAP PINOY SEAL

Sangkap Pinoy Seal Program (SPSP) - a strategy to encourage food manufacturers to fortify processed foods or food products with essential nutrients at levels approved by the DOH. The fundamental concept of the program is to authorize food manufacturers to use the DOH seal of acceptance for processed foods or food products, after these products passed a set of defined criteria. The seal is a guide used by consumers in selecting nutritious foods. (Republic Act No. 8976)

Center/Office/Di vision	:	Center for Food Regulation and Research (CFRR)
Classification	:	Government to Business
Type of Transaction	:	Highly Technical Transaction
Who May Avail	:	All FOOD Manufacturers of Fortified Products
Fees to be Paid	:	P8,000.00 non-refundable fee for the use of the seal (Regular Seal) P500.00 processing fee for every application (Regular Seal or Diamond Seal) + 1% LRF

CHECKLIST OF REQUIREMENTS Submit ONE (1) scanned copy of the required document.	WHERE TO SECURE	
☑ Basic Requirements based on RA No. 8976 (Food Fortification Law of 2000), RA No. 8172 (ASIN Law), the Sangkap Pinoy Seal Program Manual of Operations (December 2000) and Administrative Order No. 82 s. 2003 (Guidelines on the Granting of Diamond Sangkap Pinoy Seal to Manufacturers of Fortified Products):	https://www.fda.gov.ph/	
☑ Duly accomplished application forms	FDA Philippines	
☑ Copy of valid and appropriate LTO issued by the FDA	FDA Philippines	
☑ Results of product analysis for vitamin A, Iron and Iodine from an FDA recognized laboratory.	For the Certificate of Analysis: Laboratory analysis issued/conducted by FDA accredited laboratories.	
☑ Sample label with Sangkap Pinoy Seal	Applicant Company/ Manufacturer/Source/Supplier	
☑ Proof of payment	Systems/Means prescribed by FDA	
☑ Inspection report with Certificate of Compliance	FDA Regional Field Office	



CLIENT	AGENCY ACTION	PROCESSING	PERSON RESPONSIBLE
STEPS	Receives document requirements from FDA Regional Field Office and decks the same to CFRR technical evaluators.	TIME 1 Working Day	LICENSING AND REGISTRATION DIVISION (LRD) EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.1. Evaluates the application and ALL the submitted documentary requirements in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the CHECKER.	8 Working Days	LRD EVALUATOR (e.g. Food-Drug Regulation Officer (FDRO))
	2.2. Reviews the evaluated application, ALL the submitted documentary requirements, and the drafted recommendation of the EVALUATOR, in accordance to existing FDA regulation/s and Quality Work/Standard Procedures, drafts recommendation if the application is for Approval or Disapproval, and then forwards the same to the APPROVING AUTHORITY.	5 Working Days	LRD CHECKER (e.g. SENIOR FDRO or SUPERVISOR or DIVISION CHIEF)
	2.3. Prints the authorization.	1 Working Day	CFRR ADMINISTRATIVE STAFF
	2.4. Signs the Certification/Authorization.	4 Working Days	CFRR APPROVING AUTHORITY (e.g. DIRECTOR IV)
	3. Forwards the Certificate/Authorization to the Office of Director General, for signature.	1 Working Day	CFRR ADMINISTRATIVE STAFF
		TOTAL: 20 Working Days	