

Sample Submission Form

Client Details			(For Office Use: Registration Details)			
Company Name				CAIQTEST Pacific Uniqu	e ID	
Address				Temperature/Condition	1	
Contact Person			Received by			
Phone				Date/Time Received		
Email			Due Date			
Client Reference			Second Checked by			
Quote #		Order#		Reported		Invoiced
Submitter (if not client)	Additional Comments		nents			
Submitter email						
Invoice To (if not client)						
	Sample's Information					
Name of Sample				Batch No.		
Production Date				Expiration Date		
Manufacturer				Approval No.		
Place of Origin				Importer		
Quantity				Weight/Units		
□Initial Import □Non-Initial Import						
Sample Description	Color Powder Liquid Solid					
Modes of packing	□Pre-Packed: □Sealed □Unsealed □Bulk					
Storage Requirement	□Room Temperature □Cold Storage □Freeze □Other					
Product Categories						
	①Infant Formula ②Milk Power ③Liquid Milk ④Whey Powder ⑤Cheese ⑥Honey ⑦Yoghurt					



	Sample Submission Details	
Test Item	Test Method	Additional Notes



	Test Code						
1	Protein	27	Potassium	53□	Whey Protein/ Total Protein	Package	
2	Fat	28	Phosphorus				
3	Vitamin A	29□	Selenium		Total Trotein	73□	UHT Milk
4	Vitamin D3	30□	Sodium	Sodium 54		74	UHT cream
5	Vitamin E α-TE	31	Tin	55□	Sensory (Color, Taste, Texture,Odor,Recon)	75 🗆	Milk Powder
6	Vitamin K1	32□	Zinc	56□	Acidity	76□	Infant Formula (Non-Initial)
7	Vitamin B1 (as thiamine)	33□	Ca:P Ratio	57□	OPO (Fat)	77 🗆	Pasteurised Milk
8	Vitamin B2	34□	Iodine	58□	Non-Fat Total Milk Solids	78□	Honey Suite 1
9	Vitamin B6	35□	Nitrate	59□	Energy		
10	Vitamin B12	36□	Nitrite		Carbohydrate:		
11□	Folic acid (FA)	37□	Aflatoxin M1	60□	(Protein, Fat, Ash, Moisture)		
12	Pantothenic acid	38□	Mercury		Microbiology		
13	Biotin	39□	Arsenic		wiiciobiology		
14	Choline (Choline Hydroxide)	40□	Chromium	Chromium 61			
15	Inositol	41	Fatty Acids *		(APC)		
16□	Taurine	42□	Trans Fatty Acids*/** 62		Total Coliforms (MPN/Plate Count)		
17	L-carnitine	43□	Vitamin C	63□	S. aureus (P/A or Plate Count)		
18	Impurities	44 🗌	Niacin 64		E. sakazakii (MPN)		
19	Ash	45□	Lutein	65□	Salmonella		
20□	Moisture	46□	Chloride	66□	L.monocytogenese (P/A)		
21	Calcium	47 🗌	Lactose	67□	Bacillus cereus		
22	Copper	48□	Lactose/ Total Carbohydrate Ratio	68□	Yeast and Mould		
23 🗌	Iron	49□	FOS (Polyfructose)	69□	Total Yeast		
24	Lead	50□	GOS**	70 🗆	Total Mould		
25□	Magnesium	51	Nucleotide	71	Osmophillic Yeast		
26□	Manganese	52□	Lactoferrin	72 □	Total Activity/ Non Peroxide Activity (TA/ NPA)		

Note

- 1) * For Fatty acids and Trans fatty acids please specify which results required on the test report
- 2) ** GOS test same as Trans fatty acids please provide 50g raw material with purity and 50g base powder



		Package C	ption		
Infant Formula (Repeat Import or Non - Initial Import)	Honey Suite 1	Milk Powder	Pasteurised Milk	UHT Milk	UHT cream
Protein Fat Vitamin A Vitamin D3 Vitamin E α-TE Vitamin K1 Vitamin B1 Vitamin B2 Vitamin B6 Vitamin B12 Folic acid (FA) Pantothenic acid Biotin Choline (Choline Hydroxide) Inositol Taurine L-Carnitine Calcium Phosphorus Iodine Selenium Iron Copper Zinc Aerobic Plate Count Total Coliforms (E. Sakazaki): Stage 1 Salmonellas (S. aureus): Stage 1 Nitrate Nitrite Lead Aflatoxin M1	Aerobic Plate Count (APC) Total Coliforms Salmonellas S. aureus Total Yeast Total Mould	Protein Arsenic Chromium Lead Nitrate Nitrite Aerobic Plate Count (APC) Total Coliforms Salmonellas S. aureus	Aerobic Plate Count (APC) Total Coliforms Salmonellas S. aureus Nitrite	Mercury Lead Chromium Arsenic Aflatoxin M1 Commercial - Sterility	Commercial - Sterility Lead



Sample requirements:

1. Sampling quantities to meet inspection requirements.

Infant Formula and Milk Powers: For products with retail packaging >= 400g (mL), randomly sample at least 6 retail packages. For products with retail packaging less than 400g (mL), randomly sample at least 6 units where 1 unit should be no less than 800g. (e.g. if the retail package is 200g, 4 packages are 1 unit).

Liquid Milk: For products with retail packaging >= 0.75 L, randomly sample at least 4 complete packages. For products with retail packaging less than 0.75 L, randomly sample at least 8 retail packages.

Whey Power: For products with retail packaging >= 400g, randomly sample at least 6 retail packages. For products with retail packaging less than 400g, randomly sample at least 6 units where 1 unit should be no less than 800g. (e.g. if the retail package is 200g, 4 packages are 1 unit).

Others (butter, cream, etc.): For products with retail packaging >=400 g, randomly sample at least 6 retail packages. For products with retail packaging less than 400g, randomly sample at least 6 units where 1 unit should be no less than 800g (e.g. if the retail package is 200g, 4 packages are 1 unit).

* If any microbiological test involved, please send an extra individual packaging sample.

(For Five points microbiology test please send five individual packaging samples.)

2. Guarantee the sample is the same batch with customs clearance.

Terms and Conditions:

- 1. The customer is responsible for the accuracy of sample information when submitting samples for testing at CAIQTEST(Pacific). CAIQTEST(Pacific) are only responsible for the testing results as they relate to the samples received.
- 2. If no test method is selected, the laboratory will choose the default testing for the sample type received. CAIQTEST(Pacific) will not make interpretations or offer opinions on test results. Any comment made against a test result is for reference only. CAIQTEST(Pacific) takes no responsibility for questions arising due to comments against test results.
- 3. CAIQTEST(Pacific) complies with relevant laws and regulations. The laboratory will ensure the impartiality, accuracy and authenticity of the test results. All client information and testing data will be kept strictly confidential. Details of Quotations and testing applications will not be disclosed to any third party without written permission, except where required by law, judicial or administrative regulatory requirements.
- 4. If there are any changes to the testing or the time frame of contracted work, either the "Application for Amendment" (where changes are requested by the client) or the "Executive Deviation Record" (where CAIQTEST(Pacific) makes changes) will be filled out and must be mutually agreed on.
- 5. If there are any objections or queries relating to the test results they must be received within 15 days from receipt of the Test Report. Where objections or queries are received outside the 15 days period they will be deemed as no objection. If changes to the test report must be made due to error on CAIQTEST(Pacific)'s part, the laboratory will cover the cost of re-issue. If the need to change the test report is the result of client error or request, the client shall submit an "Application for Amendment" and wait for approval from CAIQTEST(Pacific) for the change. The client will be responsible for fees and charges incurred in changing the test report. If a report is changed and re-issued, the original must be returned.
- 6. Test reports cannot be used for advertising purposes, court evidence, arbitration or other relevant activities unless approval from CAIQTEST(Pacific) has been given.
- 7. Reports cannot be modified by clients after issue. If modifications are required which do not violate the relevant provisions, the client must fill in the "Report Change Request" form. The change request form must be received within 60 days of the report issue date, and if the change is accepted, the client is responsible for paying any incurred cost. After 60 days CAIQTEST(Pacific) will not accept requests to modify reports.
- 8. If inaccuracy in the test report is the result of fault or negligence on the part of CAIQTEST(Pacific) and causes direct economic losses, CAIQTEST(Pacific) will provide compensation, where compensation does not exceed twice the cost of detection. Where indirect economic losses, mental damage, unforeseen losses and third parties' losses occur, CAIQTEST(Pacific) will not assume liability.
- 9. CAIQTEST(Pacific) will not take responsibility or provide compensation for damage to the integrity of the test sample or loss/damage to the report resulting from Force Majeure, natural disasters, national laws/policies or administrative enforcement action.
- 10. The Application Agreement and Quotation will take effect immediately upon signing of this form by the client or their authorized representative. Invalidation for any reason of any individual provision does not affect the validity of the other provisions laid out in this Agreement.

Signature of Applicant:

Application Date: