



INTERAGENCY MANUFACTURER QUALITY QUESTIONNAIRE

Ref MSF: MSF-NFOS-F1.1-2

Revision: 10

14/02/2013

Scope

This questionnaire applies to all specialized food (ready to use or not) suppliers. It aims to get more details/information about the quality management system in place at the factory, the production means, and the controls implemented in the factory to prevent final products from main microbiological, chemical and physical risks.

Manufacturer/supplier information

	MANUFACTURER:	SUPPLIER (IF DIFFERENT THAN MANUFACTURER)
NAME:		
Address:		
Phone & email		

This questionnaire has been completed by:

Name(s)	
Function(s)	
e-mail(s)	

List of documents to provide

Mandatory:

- This Manufacturer Quality Questionnaire filled
- Annex 1: List of products manufactured in the site (see paragraph 2.3)
- Annex 2 – Environmental monitoring program (see paragraph 4.3)
- Map of the manufacturing site, detailing the zoning system, flow of material, personnel, waste... site (see paragraph 4.3)
- Copy of the procedure for cleaning operations, including cleaning of the production zone. (see paragraph 4.8)

If applicable:

- Food manufacturing license (see paragraph 1)
- Copy of the certifications (quality system/environment... ie : ISO 22 000 / ISO 14 001...)

Outcome (to be filled by MSF / UNICEF / WFP)

Date sent	
Date returned	

Manufacturer confidence level Low Medium High Very high



CONTENTS

RECOMMENDATIONS TO THE MANUFACTURERS..... 3

1. COMPANY IDENTIFICATION..... 4

2. GENERAL INFORMATION 4

 2.1. ORGANISATION..... 4

 2.2. PLANT WORKFORCE..... 4

 2.3. PRODUCTS 4

 2.4. SUB-CONTRACTING..... 5

3. QUALITY MANAGEMENT SYSTEM..... 5

 3.1. QUALITY MANAGEMENT SYSTEM 5

 3.2. TECHNICAL DOCUMENTATION..... 5

 3.3. HACCP..... 5

4. GOOD MANUFACTURING PRACTICES 6

 4.1. PERSONNEL 6

 4.2. TRAINING..... 6

 4.3. PREMISES..... 6

 4.4. EQUIPMENT 7

 4.5. SECURITY..... 7

 4.6. PEST CONTROL..... 7

 4.7. PROTECTION AGAINST FOREIGN BODIES 7

 4.8. CLEANING..... 8

 4.9. ENVIRONMENTAL MONITORING PROGRAM..... 8

 4.10. FLUIDS 8

 4.11. MAINTENANCE..... 9

 4.12. ENVIRONMENT 9

5. PRODUCTION AND QUALITY CONTROL..... 9

 5.1. MONITORING OF INCOMING MATERIALS..... 9

 5.2. TRACEABILITY & RECALL 10

 5.3. NON-CONFORMING PRODUCTS 11

6. LABORATORY 11

 6.1. LABORATORY..... 11

 6.2. CALIBRATION 11

7. STORAGE AND TRANSPORT 11

 7.1. STORAGE 11

 7.2. DESTRUCTION..... 12

 7.3. TRANSPORT..... 12

ANNEX 1 – LIST OF PRODUCTS MANUFACTURED ON THE SITE 13

ANNEX 2 – ENVIRONMENTAL MONITORING PROGRAM..... 14

RECOMMENDATIONS TO THE MANUFACTURERS

- Codex alimentarius (http://www.codexalimentarius.net/web/index_en.jsp)
 - Recommended International Code of Practice. General Principles of Food Hygiene CAC/RCP 1-1969, Rev. 4-2003
 - Code of Hygienic Practice for Powdered Formulae for Infants and Young Children (CAC/RCP 66 – 2008)
 - All standards linked to specific products for ingredients/raw materials and final products (ex : aflatoxins levels in peanuts, peroxide levels in vegetable oils, radioactive elements in milks, etc.) are detailed in the applicable Product Specifications Sheet (ref QA-NFOS-T:PSS+).
- Iso <http://www.iso.org/iso/en/ISOOnline.frontpage>
 - ISO 22000:2005: Food safety management systems – Requirements for any organization in the food chain
 - ISO/TS 22004 – Guidance on the application of ISO 22000:2005
 - ISO 9001:2000

- - -

MSF/UNICEF/WFP highly recommends the manufacturer to read the technical guidance:

- “Control of salmonella in low-moisture foods” and its annex, published by The GMA Association of Food, Beverage and Consumer Products Companies, in February 4, 2009.
<http://www.gmaonline.org/downloads/technical-guidance-and-tools/SalmonellaControlGuidance.pdf>
- General Guidelines on sampling CAC/GL 50-2004, codex alimentarius

- - -

MSF/UNICEF/WFP highly recommends the manufacturer to read the following documents:

- US FDA – ‘The Changing Science of Peanut Butter’
http://www.google.com/url?sa=t&rct=j&q=&esrc=s&source=web&cd=10&ved=0CIABEBYwCQ&url=http%3A%2F%2Fwww.ncagr.gov%2Fncfoodsafetyforum%2Fpresentations%2FDonald%2520Zink%2520-%25202009%2520Food%2520Safety%2520Forum.ppt&ei=eNUQUdfnLem00QXI0YHOCA&usq=AFQjCNHuf_TaCO34yz6961RBxeoY6gvdvg&sig2=JL9_YJO7XEFutx9YW19faw&bvm=bv.41867550.d.d2k
- Thermal inactivation of Salmonella in peanut Butter Li Ma et al., 2009. J. Food Protect. 72:1596 – 1601



2.4. SUB-CONTRACTING

Do you sub-contract?

- Some of the production? yes no

If yes, which part and where?

- Some of the services (maintenance, sanitation, cleaning of premises, storage...)? yes no

If yes, which one?

3. QUALITY MANAGEMENT SYSTEM

What is the number of persons employed in the quality department?

3.1. QUALITY MANAGEMENT SYSTEM

Do you have a documented Quality Management system (Quality Manual)? yes no

Is your Quality Assurance system certified? yes no

If yes What is the reference standard?

Who is the certifying organisation?

What is the date of the last certification?

What is the title of the certification (application field)?

Please provide a copy of the last certificate

Do you organise formal internal audits? yes no

If yes How many per year?

Do you organise formal management reviews? yes no

If yes How many per year?

Do you have a procedure for the management of the customer specifications? yes no

Do you have a customer complaint management system? yes no

3.2. TECHNICAL DOCUMENTATION

Do you have a technical file on each finished product ? yes no

Do you have a Quality Plan (QP) for each finished product (technical documentation of process parameters, control criteria...)? yes no

3.3. HACCP

Do you have a HACCP plan for the production of **each product**? yes no

Have you set up a team for HACCP? yes no

If yes Name & position of the coordinating person:

Do all employees know what the HACCP concept is? yes no

Are all employees able to locate the CCPs pertaining to their area of responsibility? yes no

Is your HACCP system regularly audited? yes no

If yes What is the date of the latest audit?



4. GOOD MANUFACTURING PRACTICES

4.1. PERSONNEL

Is there a clear organization chart? yes no

Do you have documentation for job description? yes no

Do you have the curriculum vitae of your key employees? yes no

Is health screening regularly carried out on all employees and prior to employment? yes no

Do you provide protective clothing? yes no
If yes, What is the frequency of change?

Are the following articles mandatory: Head covers Masks Special shoes Other:

Is hand washing part of hygiene regulations? yes no

Is it forbidden to:

- Smoke in the workshops and warehouses? yes no
- Eat in the workshops and warehouses? yes no
- Bring personal effects (bags...) into the workshops? yes no
- Wear jewellery? yes no

Are employees personnel items stored in a special place? yes no

4.2. TRAINING

Are there signs supporting GMP's posted? yes no

Do you have a training programme? yes no
If yes Does it include quality assurance (GMPs, HACCP, hygiene...) for everyone? yes no

Do you have training for new, temporary and seasonal personnel? yes no
If yes Does it include quality assurance (GMPs, HACCP, hygiene...) for everyone, including technical department? yes no

Do you evaluate training needs ? yes no

Do you have documented training records? yes no

Do you evaluate the effectiveness of training? yes no

4.3. PREMISES

Are zoning principles applied in the plant? yes no

Is there any specific identification (on a map) of: Standard hygiene areas Sensitive areas
Are access restrictions for these areas defined and materialized? yes no

Are different flows (material, personnel, waste) defined to avoid cross contamination? yes no
Are different flows written on a plan ? yes no

Please provide a copy of a map including the zoning system, flow of material, personnel, waste...

Are change rooms and bathrooms available and logically located? yes no

Is ventilation adequate and sufficient? yes no

Is lightning adequate and sufficient? yes no



4.4. EQUIPMENT

- Is equipment suitable for the intended use? yes no
- Does equipment design and condition (smooth, surface...) facilitate effective cleaning? yes no
- Are instructions on how to use the equipment available? yes no
- Are the labels for calibration and maintenance available on the equipment? yes no

4.5. SECURITY

- Is access to the plant secured? yes no
- Are visitors required to sign in and sign out ? yes no
- Are Emergency exits adequate in number and location? yes no
- Are fire extinguishers adequate in number and location? yes no
- Are first-aid procedures and equipment available? yes no

4.6. PEST CONTROL

- Is there a control plan (documented) against the following:
 Rodents Birds Insects Other:.....
- Are pest control devices (including rodent traps and electrical fly killers) adequate in number & location, and located away from exposed food products? yes no
- Do you use a specialised external company for pest control? yes no
 If yes Which one?
 What is the frequency of the interventions?
- Is there a written report after each inspection? yes no
- Do you know which treatment products are used? yes no
- Are corrective actions implemented in case of regular detection of pest activity? yes no

4.7. PROTECTION AGAINST FOREIGN BODIES

Foreign bodies are any element which is not part of the product (glass, metal, insect, plastic, stone, wood, hair,...), as well as all elements which may come from the materials being processed (shells, stones, pips, leaf...) and which should have been eliminated during processing (cleaning, washing, sorting out, etc.).

- Have the areas where one might find glass objects or materials been identified? yes no
 If yes Are these areas checked in any particular way? yes no
- Are the light bulbs and fluorescent tubes protected?
 - Throughout the whole site? yes no
 - Only in the areas where the product is exposed? yes no
- Does a procedure exist to specify what to do after glass breakage occurs? yes no
- Are the lines used for MSF/UNICEF/WFP equipped with metallic foreign body detectors? yes no
 If yes Is there a procedure for checking the metal detectors? yes no
 Is the metal detector frequently calibrated ? yes no
 Are personnel issued with metal detectable plasters for cuts/grazes? yes no



Has the facility eliminated the use of wooden items or surfaces? yes no

4.8. CLEANING

Have you set up a cleaning plan? yes no
 If yes, does it include

- The roofs? yes no
- The waste storing areas? yes no
- The dustbins and waste containers? yes no
- The equipment? yes no

Have you implemented dry cleaning for the production zone?
 If yes How often? yes no

Do you sometimes have wet cleaning?
 If yes How often? yes no

Do you keep records of all the cleaning operations? yes no

Do you check after cleaning operations?
 If yes What inspection methods do you use? yes no

Do you keep a list of cleaning products used? yes no

Do you use a specialized external company/contractor for cleaning?
 If yes Which one and for what type of operations? yes no

Please provide a copy of the detailed procedure for cleaning operation, including cleaning of the production zone.

4.9. ENVIRONMENTAL MONITORING PROGRAM

Have you set up a surveillance plan for contamination usual pathogens (Salmonella, Enterobacteriaceae ...) on product contact surfaces? yes no

Have you set up a surveillance plan for contamination usual pathogens (Salmonella, Enterobacteriaceae ..) on non-product contact surfaces, equipment and in the vicinity of the production lines? yes no

Please detail the environmental monitoring program in the annex 2.

4.10. FLUIDS

What is the volume of water used per day?

What is the origin of the water? town well surface
 Do you use other types of water (softened, industrial, recycled, ...)? yes no

Is the process for making the water drinkable under your responsibility? yes no

Do you have a water monitoring plan? yes no
 If yes, What is the kind of control?
 What is the frequency of control?

Have arrangement been made with local health officials to ensure immediate notification of the plan if potability of public water supply is compromised? yes no



Do you have a mapping of the various water circuits? yes no
Is each circuit physically identified inside the plant? yes no

Does steam ever come in contact with the product during the process? yes no

Is steam used for equipment sanitation? yes no

Are inspections made on the steam? yes no NA
If yes Please list the different inspections and their frequency

4.11. MAINTENANCE

Is there a maintenance plan in place? yes no
If yes Does it include preventive maintenance? yes no

Do you keep records of maintenance operations and equipment ? yes no

Have you identified the chemicals (sanitizers, detergents, lubricants...) used in the product's immediate vicinity? yes no
If yes Are they surveyed in any particular way? yes no
Are they stored securely? yes no
Do you keep an up-to-date list of the chemicals used? yes no
Are all chemicals labeled "for food contact" (ex USDAH1 class)? yes no
Are they labeled correctly? yes no

4.12. ENVIRONMENT

Do you have someone in charge of environment? yes no

Do you have a defined policy and measurable goals in terms of environment? yes no

Do you have precise actions plan to reach these objectives? yes no

Does the site comply with all applicable national/regional legislation? (e.g. licenses/permits, emissions into the air, waste water discharge...)
yes no

Is the site audited for environmental performance, impact and compliance? yes no

Have you undertaken an ISO 14001 certification procedure? yes no
If yes Who is the certifying organization?
What is the date of the last certification?
 Please provide a copy of the last certificate

Have the latest inspections by the authorities shown any cases of non-conformity? yes no NA

Do you have a procedure of environmental crisis management? yes no

5. PRODUCTION AND QUALITY CONTROL

5.1. MONITORING OF INCOMING MATERIALS

Do you know the geographical origin of all raw materials used ? yes no



What are your supply networks and channels?

- Producers on contract Intermediary agency Spot purchase Distribution
 Other:

How do you approve your suppliers?

.....

Do you audit your suppliers?

- yes no

If yes How often?

What is the date of the last audit?

Do your raw and packaging materials have to meet specifications?

- yes no

Do you buy any materials from non approved suppliers?

- yes no

If yes please specify materials and reasons

.....

Do you keep samples of material?

- yes no

If yes How long?

5.2. TRACEABILITY & RECALL

With respect to upstream traceability:

With a batch number, can you find all the history of the finished products (composition, processing parameters, analytical results, raw materials used...)?

- yes no

How long does it take you to find all the information?

Has this timing been verified through real tests?

- yes no

Do you require your suppliers to set up a traceability system?

- yes no

Do you audit raw material traceability at your suppliers?

- yes no

Concerning raw materials, how far can you go back?

Do you follow traceability of recycled/rework product?

- yes no NA

With respect to downstream traceability, for each batch, are you able to find:

- All customer sites delivered?

- yes no

- Quantity delivered for each site?

- yes no

- Date of deliveries?

- yes no

How long does it take you to find all the information?

Has this timing been verified through real tests?

- yes no



Do you require your customers to set up a traceability system? yes no NA

Do you have a procedure for recall? yes no

Do you have a designated person(s) responsible for recall? yes no

Do you audit the traceability system regularly? yes no
If yes What is the date of the last audit?
What part of the traceability system was audited?

5.3. NON-CONFORMING PRODUCTS

Are they physically identified? yes no

Do you keep a record of them? yes no

Who decides what to do with non-conforming products?

6. LABORATORY

6.1. LABORATORY

Does the plant have an internal laboratory? yes no
If yes, kind of analysis? Organoleptic Physical-Chemical Microbiological Others:

Is there a validation system for the laboratory methods? yes no NA
Are in-house methods documented and approved by a suitably qualified person? yes no NA

Do you use outside laboratories for some analyses? yes no
If yes Do you use accredited laboratories for those analyses? yes no
How do you select/approve external laboratories?
.....
.....

6.2. CALIBRATION

Is there a written procedure for the calibration of all equipment and instruments, including new equipment and instrument prior to use? yes no

How long are calibration records kept?

7. STORAGE AND TRANSPORT

7.1. STORAGE

Do you have an outside storage location? yes no
If yes How far is it from the factory?
Does the warehouse belongs to your company? yes no
Is there staff permanently present at this warehouse? yes no

Do you have specific premises for storing the following:
- Raw materials? yes no
- Packaging materials? yes no
- Chemicals? yes no
- Finished products? yes no



Are there specific storage conditions (temperature, humidity) for materials? yes no

If yes Please describe:
.....
.....

Are those parameters recorded? yes no

If yes, how often?

Are the persons authorized to change those parameters clearly defined? yes no

How do you protect finished products on the pallets:

- Between layers? yes no

- Plastic film cover? yes no

Are procedures in place for stock rotation? yes no

Are materials properly marked with rotation codes (receipt dates, manufacture dates...)? yes no

Are periodic stock reconciliations performed by comparing the actual and recorded stocks (inventory)? yes no

Are significant stock discrepancies investigated? yes no

Is there sufficient space along all walls (<30cm) to permit proper cleaning and inspection for pest activity? yes no

7.2. DESTRUCTION

Do you have a procedure for destruction of raw material/finished products? yes no

7.3. TRANSPORT

Do you have set procedures for inspecting the equipment before unloading and loading (cleanliness, absence of odor, absence of suspicious products...)? yes no

Do you keep a record of these checks? yes no

Are there specific transport conditions (temperature, humidity) for the ingredients based on stability studies? yes no

If yes, Are those parameters recorded? yes no



ANNEX 1 – LIST OF PRODUCTS MANUFACTURED ON THE SITE

MSF products	UNICEF products	WFP products	Products	Reference of production line	Filling conditions (packaging type)
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
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<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			



ANNEX 2 – ENVIRONMENTAL MONITORING PROGRAM

Where	bacteria	Method	Frequency	Number of sampling points	target	% of out of target results within the last year	Laboratory used (internal/external)	
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
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							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External
							<input type="checkbox"/> Internal	<input type="checkbox"/> External

Name and location (country) of the external laboratory(ies):