


FIRST QUALITY CERTIFICATION		Doc. Ref. No.:	QPR-014-SMK-F004
	Document Title: Application Form	Page No.	Page 1 of 5
		Rev. No.	Rev.00
		Date:	01-Apr-18
		Issue No.	01

COMPANY INFORMATION

Company Name:	
Corporate Address:	
Tel. No(s).:	Fax No.:
Company Email:	Web:
Authorized Contact Person Name:	Mobile:
Designation:	Email:
Org. Head:	Email:

CERTIFICATION STANDARD(S)

<input type="checkbox"/> ISO 9001	<input type="checkbox"/> ISO 14001	<input type="checkbox"/> OHSAS 18001	<input type="checkbox"/> ISO 22000	<input type="checkbox"/> HACCP	<input type="checkbox"/> Others _____
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NACE Code(s):

Scope (Activity):

No. of Permanent Employees: ()	No. of Temporary Employees: ()	TOTAL No. of Employees: ()
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No. of Permanent Site(s):	No. of Temporary Site(s), if any:
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Address (1)	Address (1)
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Address (2)	Address (2)
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Address (3)	Address (3)
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Address (4)	Address (4)
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Address (5)	Address (5)
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Guidance number of auditor days (based on effective number of employees:	Standard(s):		
	Initial Assessment:	<input type="checkbox"/> YES	<input type="checkbox"/> NO
	Surveillance: F/\$	<input type="checkbox"/> YES	<input type="checkbox"/> NO
	Re-assessment	<input type="checkbox"/> YES	<input type="checkbox"/> NO

CONSULTANTS DETAILS (IF ANY)

Company Name:

Standards(s):

<input type="checkbox"/> ISO 9001	<input type="checkbox"/> ISO 14001	<input type="checkbox"/> OHSAS 18001	<input type="checkbox"/> ISO 22000	<input type="checkbox"/> HACCP	<input type="checkbox"/> Others _____
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Quotation to be issued:	Client Stamp:
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<input type="checkbox"/> Transfer	<input type="checkbox"/> Application for Certification	<input type="checkbox"/> Audit Planning	<input type="checkbox"/> Document Review
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<input type="checkbox"/> Stage 1 Assessment	<input type="checkbox"/> Stage 2 Main Assessment	<input type="checkbox"/> Combined Stage 1 & 2 Assessment	<input type="checkbox"/> Re-assessment
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<input type="checkbox"/> Surveillance Year 1	<input type="checkbox"/> Surveillance Year 2	<input type="checkbox"/> _____	<input type="checkbox"/> _____
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TIME ALLOCATED		
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Addition or Reduction:	Audit Planning:	Document Review:
Stage 1 Initial Assessment:	Stage 2 Main Assessment:	Re-assessment:
Clear Corrective Actions:	Certification Review:	Annual Surveillance:
Site Visit Time:	Travel Time (Duration):	
Scope review form suitable? [] YES [] NO	Other Comments/ Specific Competence Requirements:	
State of implementation of standards? [] YES [] NO		

Statutory and Regulatory Requirements (Related to the nature of work and Management System Certification):

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Outsourced Process: (if any, which effects the conformity of the product/service)
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In case of outsourced process, what type and extent of controls have been applied in order to ensure that the externally provided functions or processes do not adversely affect the effectiveness of your management system (MS)?


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In case of outsourced process, how have you evaluated and determined organization's ability to meet your requirement and legal compliances?
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STANDARD REQUIREMENTS (v ONLY WHERE APPLICABLE)		
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QMS	DOCUMENTED?	
i. Is the category "design & development", included in the activities to be certified?	[] YES	[] NO
ii. Is there any process outsourced, which affects product conformity?	[] YES	[] NO
iii. Exclusions, if any?	[] YES	[] NO
1. Policy	[] YES	[] NO
2. Identification and methodology of Risk	[] YES	[] NO
3. Risk Treatment Plan	[] YES	[] NO
4. Risk Assessment Report	[] YES	[] NO
5. Statement of Applicability with regard to the standards	[] YES	[] NO
6. Records required by the standards	[] YES	[] NO
7. Internal Audit	[] YES	[] NO
8. Detail of the service outsourced	[] YES	[] NO

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9. Assessment of Risks and Opportunities defined	<input type="checkbox"/> YES	<input type="checkbox"/> NO
10. Records of internal audits conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
11. Records of Management Review conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
12. Defined scope of the Quality Management System	<input type="checkbox"/> YES	<input type="checkbox"/> NO
13. Process Flow diagrams indicating interactions of all steps in the operation	<input type="checkbox"/> YES	<input type="checkbox"/> NO
14. List all functions/departments of your organization	<input type="checkbox"/> YES	<input type="checkbox"/> NO
15. List of all applicable regulatory and statutory legislations	<input type="checkbox"/> YES	<input type="checkbox"/> NO
EMS ONLY	DOCUMENTED?	
1. Policy	<input type="checkbox"/> YES	<input type="checkbox"/> NO
2. Is there any process outsourced, which affects product conformity?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
3. Details of the site(s); your company managing at the same time?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
4. A Register of Significant Environment Aspects?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
5. An Environmental Management Manual?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
6. An Internal Environmental Audit Programme?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
7. Has the Internal Environmental Audit Programme been implemented?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
8. Kindly provide list of Significant Aspects & License required?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
9. Storage condition & permitted quantities of hazardous material?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
10. Records of internal audits conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
11. Records of Management Review conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
12. Defined scope of the Environmental Management System	<input type="checkbox"/> YES	<input type="checkbox"/> NO
13. Process Flow diagrams indicating interactions of all steps in the operation	<input type="checkbox"/> YES	<input type="checkbox"/> NO
14. List all functions/departments of your organization	<input type="checkbox"/> YES	<input type="checkbox"/> NO
15. List of all applicable regulatory and statutory legislations	<input type="checkbox"/> YES	<input type="checkbox"/> NO
OHSAS ONLY	DOCUMENTED?	
1. Policy	<input type="checkbox"/> YES	<input type="checkbox"/> NO
2. Applicable Legal & Other Requirements?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
3. Hazards Identified?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
4. Please detail any critical Occupational Health & Safety Risks Identified?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
5. Eliminating Hazards & Reducing OH&S Risks?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
6. Emergency Preparedness and Response Plan or Process?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
7. Incident Investigation Process?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
8. Records of internal audits conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
9. Records of Management Review conducted	<input type="checkbox"/> YES	<input type="checkbox"/> NO
10. Defined scope of the OHSAS Management System	<input type="checkbox"/> YES	<input type="checkbox"/> NO
11. Process Flow diagrams indicating interactions of all steps in the operation	<input type="checkbox"/> YES	<input type="checkbox"/> NO
12. List all functions/departments of your organization	<input type="checkbox"/> YES	<input type="checkbox"/> NO
13. List of all applicable regulatory and statutory legislations	<input type="checkbox"/> YES	<input type="checkbox"/> NO
FSMS ONLY	DOCUMENTED?	
1. Policy	<input type="checkbox"/> YES	<input type="checkbox"/> NO
2. Applicable Legal & Other Requirements?	<input type="checkbox"/> YES	<input type="checkbox"/> NO
3. Food Hazards Identified?	<input type="checkbox"/> YES	<input type="checkbox"/> NO


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4. Food Safety Team	[] YES	[] NO
5. Records of internal audits conducted	[] YES	[] NO
6. Records of Management Review conducted	[] YES	[] NO
7. Defined scope of the Food Safety Management System	[] YES	[] NO
8. Process Flow diagrams indicating interactions of all steps in the operation	[] YES	[] NO
9. List all functions/departments of your organization	[] YES	[] NO
10. List of all applicable regulatory and statutory legislations	[] YES	[] NO
11. List of pre-requisite programmes implemented	[] YES	[] NO
12. List of finished products and their intended use	[] YES	[] NO
13. Product recall	[] YES	[] NO

HACCP ONLY	DOCUMENTED?	
1. Policy	[] YES	[] NO
2. Applicable Legal & Other Requirements?	[] YES	[] NO
3. Hazards Identified?	[] YES	[] NO
4. Recipe(s)/List of ingredients	[] YES	[] NO
5. List of Equipment and Materials	[] YES	[] NO
6. Hazard Worksheet/list of hazards	[] YES	[] NO
7. Process Flow Diagram	[] YES	[] NO
8. HACCP Worksheet	[] YES	[] NO
9. Cooking Potentially Hazardous Foods SOP	[] YES	[] NO
10. Cooling Potentially Hazardous Foods SOP	[] YES	[] NO
11. Eliminating Bare Hand Contact SOP	[] YES	[] NO
12. Handwashing SOP	[] YES	[] NO
13. Hot and Cold Holding of Potentially Hazardous Foods SOP	[] YES	[] NO
14. Personal Hygiene SOP	[] YES	[] NO
15. Time As a Control SOP	[] YES	[] NO
16. Using and Calibrating Thermometer SOP	[] YES	[] NO
17. Cooking and Reheating Log	[] YES	[] NO
18. Cooling Log	[] YES	[] NO
19. Refrigeration Log	[] YES	[] NO
20. Thermometer Calibration Log	[] YES	[] NO
21. Training Log & Food Safety Checklist	[] YES	[] NO

REVIEWED BY:	CLIENT REPRESENTATIVE:
Name:	Name:
Designation:	Designation:
Date:	Date:
Remarks:	

APPLICATION REVIEW BY FIRST QUALITY CERTIFICATION (FQC)		
1. Does accreditation request available with FQC? (Refer accreditation letter)	[] YES	[] NO
2. Does territory of the application in active list? (Refer accreditation letter)	[] YES	[] NO
3. Does Scope demand available with FQC? (Refer accreditation letter)	[] YES	[] NO

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4. Does MS certification request available with FQC? (Refer accreditation letter)	[] YES	[] NO
5. Does the above information complete?	[] YES	[] NO

REVIEWED BY:		
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Name:	Signature:	Date:
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1. Can the application be further processed?	[] YES	[] NO
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REMARKS (IF ANY):