

#### **Management Systems Certification**

	QMS ISO 9001:2015 STAGE-1 AUDIT REPORT
Client Ref. No.	
Organization Name	
Address	
Site Address (If any)	
No. of Employees	
E mail id	
Contact Person	
Telephone/Fax	
Scope	
NACE Code	
Audit Team	
Audit Man-days	
Stage of Audit and	
date	
Brief about the	
organization	
Audit Guidance	To evaluate the client's documented system, location & site-specific conditions
	and gather other details through discussions with the client's personnel to
	determine the organization's readiness for the Stage 2 Audit for Certification.
Audit Objective	The purpose of the Stage 1 audit is to evaluate the effective implementation of
	the client's management system. As a minimum you must audit the following
	and your report must show clear audit evidence against these requirements.
	to audit the client's management system documentation and some management
	processes;
	to evaluate the client's location and site-specific conditions and to undertake
	discussions with the client's personnel to determine the preparedness for the
	stage 2 audit;
	to review the client's status and understanding regarding requirements of the
	standard, in particular with respect to the identification of key performance or
	significant aspects, processes, objectives and operation of the management
	system;
	to collect necessary information regarding the scope of the management
	system, processes and any statutory regulations;
	to review the allocation of resources for stage 2 audit and agree with the client
	on the details of the stage 2 audit;
	to evaluate if the internal audits and management review are being planned and
	performed, and that the level of implementation of the management system
	substantiates that the client is ready for the stage 2 audit;
	to verify all information previously supplied is still correct and relevant and any
	changes required to the number of days for stage 2 must be agreed before the audit report is completed;
	audit report is completed,

# AREA OF CONCERN WHICH MAY BE IDENTIFIED AS NON CONFORMITIES DURING STAGE 2 AUDIT



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## **Guidance Note for Opening Meeting Agenda:**

Sl.	Topics	Particular	Completed	
1	Thanks	Give an expression of thanks to the auditee for Choosing BCI.		
2	Attendance	Request attendees to record their attendance		
3	Introduction	Remind timeline to close opening meeting in 15-30 minutes.		
	ļ	Request to give brief introduction with brief roles (participants,		
		observers, guides & Translators)		
4	Scope / Summery	Confirmation of the audit objectives (Assessment for ISO 9001), scope and criteria:		
5	Changes	Changes in documents/Fact to the Application/Stage-1 Audit.		
6	Plan	Confirmation of the audit plan and other relevant arrangements with the		
		auditee, such as the date and time for the closing meeting, any interim		
		meetings between the audit team and the auditee's management, and any late		
7	Method	changes; Methods of Audit: Review of Documents & Records, Interview, Physical		
/	Method	evidence		
8	Sampling	Advise auditee that the audit is sample basis and findings will be based on a		
	Samping	sample of the information selected;		
9	Communication	Confirmation of formal communication channels between the audit team and		
	Channel	the auditee; identify the facilitators.		
10	Language	Confirmation of the language to be used during the audit;		
11	Development	Confirmation that, during the audit, the auditee will be kept informed of audit		
	r	progress;		
12	Resource	Confirmation that the resources and facilities needed by the audit team are		
		available; like Guide, Interpreters, Facility etc		
13	Confidentiality	Confirmation of matters relating to confidentiality and information security;		
14	Safeguard	Confirmation of relevant health and safety, emergency and security		
		procedures for the audit team;		
	Reporting of	NC may be against a clause of the standard i.e. ISO 9001, it's not		
15	Findings	against any person or department.		
	ļ	Method of reporting audit findings & grading (Major, Minor &		
		Observation)		
		Time-span for corrective action (Minor-15 Days, Major-60 Days)		
		Report time: Finding will be discussed at closing meeting and report		
1.5	m : :	will be given within 2 working days.		
16	Termination	Information about conditions under which the audit may be terminated;		
17	Audit Declaration	verify that all members of the organization know what is happening;		
18	Union/Problem	Ascertain union relations or any potential problems;		
19	Confidentiality	Remind the auditee that the audit is confidential.		
20	Closing Meeting	Timing of closing meeting; Participation of the Top Management & where		
		appropriate, those responsible for the functions or processes which have been		
21	Appagla /	audited in the closing meeting.		
21	Appeals / Complaint	information about any system for feedback from the auditee on the findings or conclusions of the audit, including complaints or appeals		
	Compraint	or concrusions of the addit, including complaints or appears	1	



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# **Recording of Attendance**

	AUDIT ATTENDANCE SHEET				ce (by LA)
S. No	Name	Position	Department	Opening	Closing

Sl.	Requirement	Comment	Status* C/N/O/ NA
1	Does the Quality Management system address the key areas of client's business? Quality Manual Reference If any and process description. Context of organization has been understood and addressed by determining external and internal issues. Monitoring and review system and records reference available? Needs and expectations of interested parties understood? Risk based approach adopted? Opportunities identification approach?		
2	Are processes for QMS identified and their sequence & interaction defined?		
3	Is scope of QMS Included in Manual or defined otherwise? Shall determine the boundaries and applicability of the quality management system to establish its scope. Does the Scope is appropriate to the organizational activities?		
4	Do manual or Other documents include Details of exclusions with justifications?		
5	Are all 7(4, 5, 6,7,8,9 and 10) elements of QMS addressed in Quality Manual or addressed in other organizational documents? (Context of organization, leadership, Planning, Support, Operation, Performance Evaluation, Improvement)		



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Sl.	Requirement	Comment	Status* C/N/O/ NA
6	Does the organization have site-specific activities – top level process review		
7	Does organization have identified and complied with the appropriate regulatory and legal requirements; applicable to the product / Services?		
8	Is there a documented statement of Quality Policy? And is appropriate to the purpose and context of the organization and supports its strategic direction Andbe available to relevant interested parties, as appropriate.		
9	Have Quality objectives been established at relevant functions, levels and processes needed for the quality management system.  And objectives shall be: Consistent with Quality Policy, measurable, monitored, communicated and updated as appropriate And records reference of the achievements provided?  Are the 7 mandatory documented clauses		
	and related records references available? (Any other organizational document/ procedure need to be referred?)		
11	Are Internal audits conducted as planned?		
12	Date of Last Internal Audit?  Are Management reviews conducted as planned?		
14	Date of Last MRM?		
15	Are customers complaints recorded? Is		
	there evidence of resolving the same?		
*	Is there any requirement of visit the temporary site(s) in stage-2 audit?		

#### **Guidance Note for Closing Meeting Agenda:**

Sl.	Topics	Particular	Verified
1	Introduction	Particularly if anybody not present at the opening meeting	
2	Thank to company	Thanks to your Team for cooperation during the audit and arrangements for the Audit.	
3	Reaffirmation of Scope	Reconfirm scope of activities assessed	
4	Confirm	Reassure the confidentiality for any information assessed during the audit.	



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	confidentially		
5	Appreciation	Comment on good points within the organisation	
6	Disclaimer	This was audit on sample basis, and it should not mean, that other	
		deficiencies do not exist.	
7	Audit Team	Summary of individual findings from each auditor (if audit team consist	
	Comment	more then 1)	
8	Decision	Significance of categories of non-compliance and summary of	
		findings ,	
		Summary of overall findings and recommendation/Decision	
9	Acknowledgment	Assure that client acknowledge the NCs.	
10	Future Plan	If any NC is identified, Submitting plan for corrective action together with	
		the objective evidences	
11	Follow-up action	Where do we go from here? emphasizing that the final decision regarding	
		certification will be taken by BCI	
	Surveillance Audit	An explanation of the continual Audit (surveillance) procedure and other	
		future actions	
12	Appeal	Explain the Appeal & Complaint option available to the client against any	
		decision of the Audit team.	
13	Invite questions	Invite questions, clarification from company (But no Consultancy)	
14	Signature	Obtain company representative's signature on report to acknowledge	
		receipt.	

# Recommendation for Stage-2 Audit: I have checked, examined and discussed and confirm the following: Mark "X" where applicable.

Sl.	Validation of Critical Points	Yes/No/NA	Comment of the Auditor
1	Relevance of the QMS documentation with activities of the client	es	
2	Scope applied are justified with the present activitie of the Clients	S	
	Any Change in scope needed?		
3	Are temporary sites (i.e installation sites, project locations etc.) available?		
4	Which sites needed to be visited?		
5	will it requires Considerable Travel Time to visit sit	e	
6	is there any Seasonality Factor		
7	Suitability of Audit Timing (Activities at Site)		
8	Process and element of the ISO 9001 in Stage-1 audaddresses?	lit	
Audi	t Duration for Stage 2		
Are o	quoted man-days adequate?		
Any	change in employee detail?		
Any	Change in Scope?		

Any Additional information?

Is there any requirement of verifying the night shift



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in stage-2 audit?					
Audit Summary (Includin	g General Observations/Co.	mments)			
RECOMMENDATION					
	led for Proceeding to Stage				
	nend proceeding to stage 2				
	t the concerns raised by the		rified. A date for stage 2		
	agreed. (within 60 days from				
	nend proceeding without a	further stage 1 Audit due	to the severity of the		
concerns rais	concerns raised by the audit team				
		ı			
		Signature			
Signature					
Name of the Auditor		Name of the			
		Representative			
Date		Date			