

Supplier's Audit Form (Second - Party) (F.021)

A. General:

Company name:		
Address:		
City/Town:	_ Province:	Country:
Postal/Zip code:	Tel. ()	_ Fax.()
Head of Quality Deptt:		Title:
Head of quality deptt. Reports t	to:	Title:
Other contacts:		Title:
		Title:
Product or service:		
To which standard(s) does your	quality system conforms to:	

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В.	Quality	/ Management System:
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Do you have a quality Management System Policy Manual in use?
which includes: scope of quality management system, documented procedures.
Do you have documented procedures and records required by this International Standard?
Are these quality management system documents controlled?
Do you have a process to assure the timely review, distribution and implementation of all customer engineering specifications and changes on customer required schedule?
When do you plan to upgrade your QMS (i.e. ISO to TS to IATF)?
Management Responsibility:
Do you have a company Quality Policy?
Do procedures describe the authority of those responsible for managing, performing, and verifying work effecting quality?
Is the quality system reviewed at appropriate intervals by your company's management to ensure continuing suitability and effectiveness?
Resource Management:
Are verification resources such as inspection, test and quality system audits independent of those responsible for work being performed?
Do you have personnel performing work affecting product requirement confirmity competent enough?

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_	Product Realization:
	Do you determine quality objectives and requirements for the product?
	How do you determine required verification, validation, monitoring, measurement, inspection and test activities specific to the product?
	Do you ensure confidentiality of customer-contracted products and projects under development?
-	
	Do you determine and implement effective arrangements for communicating to the customers related to product information, customer feed back including complaints?
	Do you have written procedures assuring that all drawings, procedures and specifications used are of the latest revision?
	How are all affected personnel made aware of the latest revisions?
	Does your Quality Asssurance deptt. Review all the drawings, specifications and procedures and their revisions?
	Do your purchase orders ensure that all of your product purchase specific requirements are mentioned on them?
	Do you ensure that purchased product conforms to specified purchase requirements?
	Do you maintain a quality history on received parts?
	Do you perform sampling inspection?

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12	State briefly how your receiving inspector performs his/her function?
13	How are the suppliers selected?
11	How is supplier avaluation and to avaluation done?
14	How is supplier evaluation and re-evaluation done?
15	Do you use any of customer- approved sources? If yes what is the criteria
	for ensuring quality product?
16	
	Does it include all product requirements?
17	Do you have quality inspection plan for products?
10	Do you have a corrective action program for defective surchased material
18	Do you have a corrective action program for defective purchased material

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and parts? Please describe:



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19	Do you identify inspection status of received product? Please explain
	your system for keeping non-conforming material segreegated:
20	Do you have work instructions available as required?
21	Do your work instructions show use of suitable equipment, monitoring
	and measuring equipment, product release implementation, delivery &
	post delivery activities?
22	How do you verify job set ups such as initial run of a job, material change
23	How is preventive and predictive maintenance carried out?
	e.g. identifying key process equipment, planning maintenance activities,
	avaialability of replacement parts, improving maintenance objectives.
24	How do you maintain production tooling?
	e.g. maintenance and repair, storage, set-up, tool change etc.
25	How do you manage inventory?

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e.g. inventory turns, assuring stock rotation as FIFO, controlling obsolete product.



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26	How do you maintain instruments caliberation? e.g. verifying at specified intervals, have identification, be protected from damage during handling, maintenace and storage.
F.	Measurement, Analysis and Improvement:
1	How do you monitor information as to customer satisfaction for all product requirments? e.g. customer satisfaction survey etc.
2	Do you have a process in place for performing internal audits within your organization?
3	Do you maintain an in-process inspection history for manufactured products?
4	Are these records traceable to a specific lot?
5	Do you perform sampling inspection?
6	State briefly, how do you perform your in-process inspection?

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7	Do you have a corrective action program for in-process defectives?
8	Describe your system for identifying the inspection status of in-process material?
9	Do you maintain final inspection history for manufactured products?
0	Are these records traceable to a specific lot?
.1	Do you perform sampling inspection?
2	State briefly, how do you perform your final inspection?
.3	Do you have a corrective action program for final inspection defectives?
.4	Describe your system for identifying the inspection status of final inspection material?
.5	Describe your system for keeping non-conforming product segregated?

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provement through use of quality
a analysis, corrective and preventive
es in order to prevent recurrence.
uding customer complaints), determining cause,
nese do not recur, determining and implementing
tion taken and reviewing effectiveness of corrective
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