

Corrective Action Report

Tracking Number:	Reported By:	Date Found:	Status

1. Describe the problem

Priority Classification					
Legal or compliance issues?	<input type="checkbox"/> Concerns must be forwarded to Top management for review				
Safety concerns?	<input type="checkbox"/> Concerns must be forwarded to the H&S Department for review				
Repeated nonconformities in:	<input type="checkbox"/> Repeated issues must be sent to the QMS Manager for review				
Select	Products/services?	<input checked="" type="checkbox"/> High (3)	<input type="checkbox"/> Medium (2)	<input type="checkbox"/> Low (1)	<input type="checkbox"/> OFI (0)
	Processes?	<input checked="" type="checkbox"/> High (3)	<input type="checkbox"/> Medium (2)	<input type="checkbox"/> Low (1)	<input type="checkbox"/> OFI (0)
	Management system?	<input checked="" type="checkbox"/> High (3)	<input type="checkbox"/> Medium (2)	<input type="checkbox"/> Low (1)	<input type="checkbox"/> OFI (0)

How was the nonconformance detected?			
Source:	<input type="checkbox"/> Feedback - Supplier	<input type="checkbox"/> Audit - Client	<input type="checkbox"/> In-process inspection
	<input type="checkbox"/> Feedback - Customer	<input type="checkbox"/> Review - Management	<input type="checkbox"/> First article inspection
	<input type="checkbox"/> Feedback - Stakeholder	<input type="checkbox"/> Review - Department	<input type="checkbox"/> Receiving inspection
	<input type="checkbox"/> Feedback - Employee	<input type="checkbox"/> Review - Design	<input type="checkbox"/> Final inspection
	<input type="checkbox"/> Audit - Internal	<input type="checkbox"/> Nonconformity - Process	<input type="checkbox"/> Monitoring
	<input type="checkbox"/> Audit - Registrar	<input type="checkbox"/> Nonconformity - Product	<input type="checkbox"/> Customer complaint
	<input type="checkbox"/> Audit - Supplier	<input type="checkbox"/> Nonconformity - Service	<input type="checkbox"/> Customer feedback
	<input type="checkbox"/> Other:		

Define the process name in which the problem arose	
Location:	1.
	2.
	3.

Include a description of requirements (reference drawing, specification, standard as appropriate). A detailed description of the nonconformance. Sketches, drawings, specifications, etc., should be used to help the understanding of the nonconformity.

1. Problem Statement;
2. Failure analysis;
3. Is/Is Not analysis;
4. Process flow with control points identified;
5. Problem Description.

Description of the Issue:
Describe the issue using plain language, and list any relevant documents, names of others who may be aware. Need to have correct problem description to identify causes. Use terms that are understood by all.

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3. Analyze the causes

Establish a small group of people with the process/product knowledge, allocated time, authority and skill in the required technical disciplines to solve the problem.

Corrective Action Team			
Name	Position/Dept	Skills	Responsibility

Root-cause Analysis
Describe how the root-cause was established and attach copies of the chosen root-cause analysis method: A3, 5-Y, 8D, Fishbone, or FMEA. Update Part A of the Corrective Action Tracker (Col G).

4. Propose correction action

Description of Corrective Action:
Describe the proposed method for permanent corrective action. Update Part B of the Corrective Action Tracker (Col P).

5. Implement corrective action

Establish a small group of people with the process/product knowledge, allocated time, authority and skill in the required technical disciplines to implement corrective actions. Update Part B of the Corrective Action Tracker (Cols O, R & S).

Corrective Action Team			
Name	Position/Dept	Date Assigned	Completion Date

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Corrective Action Implemented By:			
Implemented By:	Signature	Position	Date
Reviewed By:	Signature	Position	Date

6. Verify effectiveness

Verification:
Describe how the corrective action addressed the root cause, list relevant evidence. Update Part B of the Corrective Action Tracker (Col Y).

Feedback & Acknowledgments:
Mention pertinent feedback from the personnel involved and acknowledge those that contributed to resolving the corrective action.

7. Close Out

Corrective Action Close Out:
Verify that the planned actions were taken as scheduled and assess their effectiveness in permanently preventing the undesirable condition, situation, non-conformity, or failure from recurring. Update Part B of the Corrective Action Tracker (Col Z).

Corrective Action Completed By:			
Actioned By	Signature	Position	Date
Verification By	Signature	Position	Date