Management System Lead Auditor Training

Module 8. Audit Reporting and Documentation Processes



Learning Outcome



Able to effectively document and report audit findings in accordance with management system

management system Develop the skills to accurately assess audit evidence, identify nonconformities, and prepare clear and concise audit reports

proparo orean any concise audit reports Gain knowledge to ensure the completeness, traceability, and confidentiality of audit records

confidentiality of audit records



Module 8

Audit Reporting and Documentation Processes







ISO 19011:2018 -Guidelines for auditing management systems

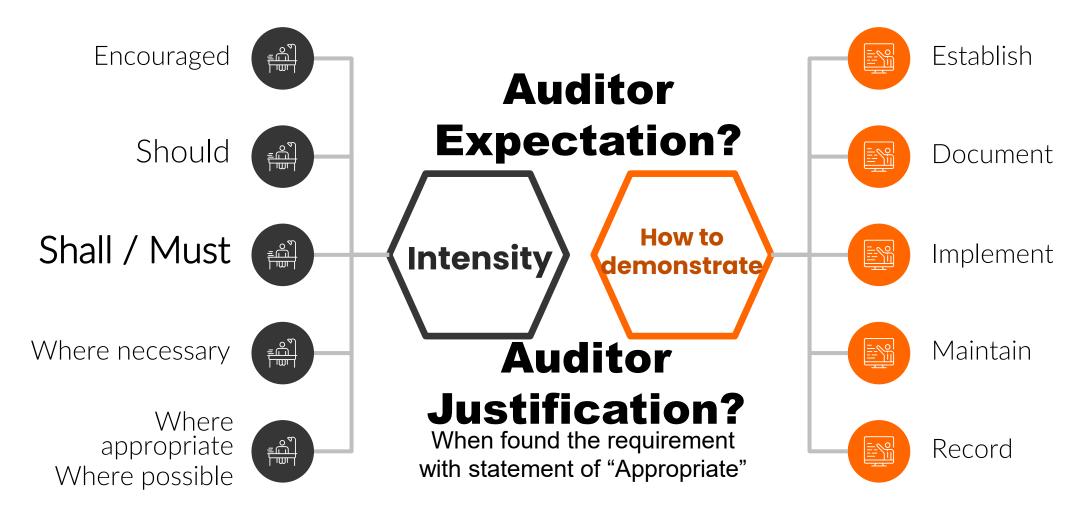


ISO/IEC 17021-1:2015 - Conformity assessment -Requirements for bodies providing audit





Justifying Requirements



Recording the findings of an audit involves capturing evidence from various sources, including observation, interviews, and records. By integrating evidence from observation, interviews, and paper records, auditors can ensure a holistic and well-documented representation of their findings, enabling accurate analysis, recommendations, and the delivery of a comprehensive audit report.





Findings Classification

FEATURES	STRENGTH	CONFORMANCE	0.	F.I	n	nn.No	C	Mj.N	NC
LEGAL REQUIREMENT	~	~	•	✓	✓	~	✓	~	×
STANDARD REQUIREMENT	✓	✓	~	~	~	~	×	Repeated case	•
INTERNAL REQUIREMENT	✓	~	~	×	~	×	~	~	~
RISK	×	×	~	×	With evidence	~	~	~	~
ADD VALUE	✓	×	×	×	×	×	×	×	×
	OUTSTANDING	POSITIVE I	RATHER NEGATIVE		/E NEGATIVE		NEGATIVE		



Conforming to requirements

Legal is higher requirement

Justification of RISK: a) Penalties b) Company's image c) Cost d) Financial



Conforming to requirements

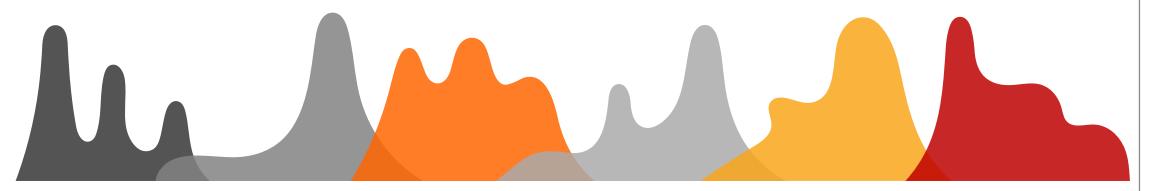
Standard is higher requirement

Justification of RISK: a) Effectiveness

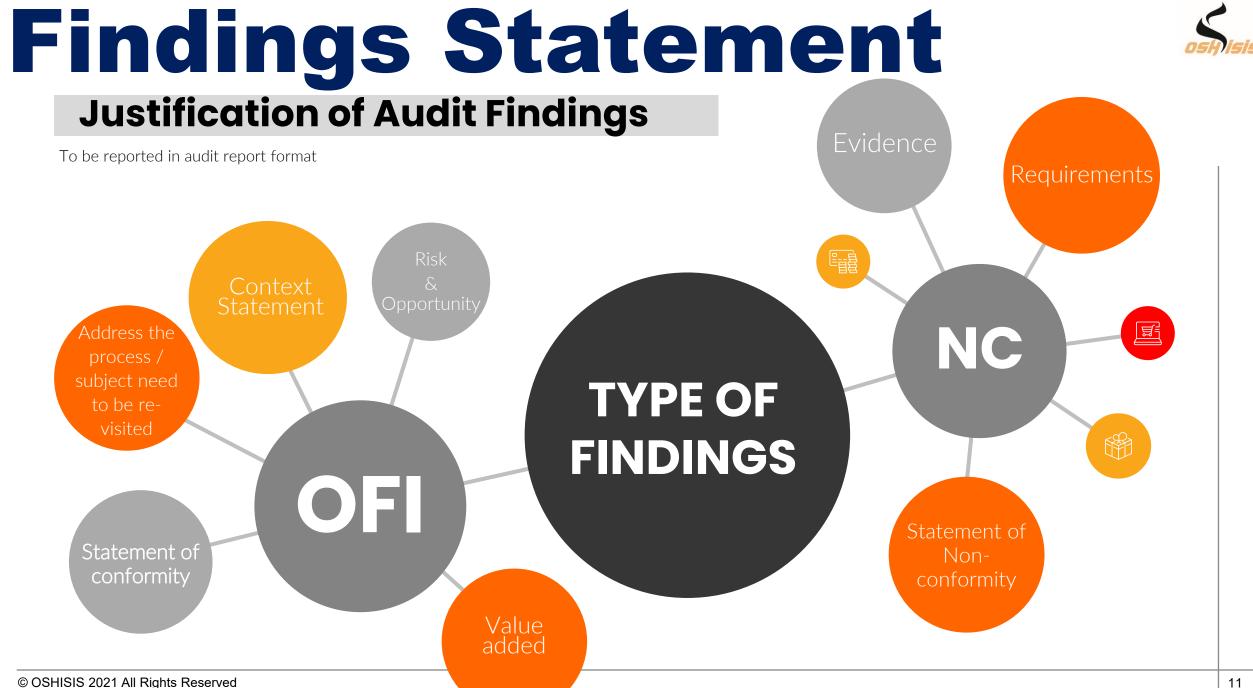


Conforming to requirements

Internal SOP is higher requirement



Justification of RISK: a) Effectiveness b) System integrity



Findings Statement



To be reported in audit report format



Statement of conformity

Statement of process / subject to be relooked

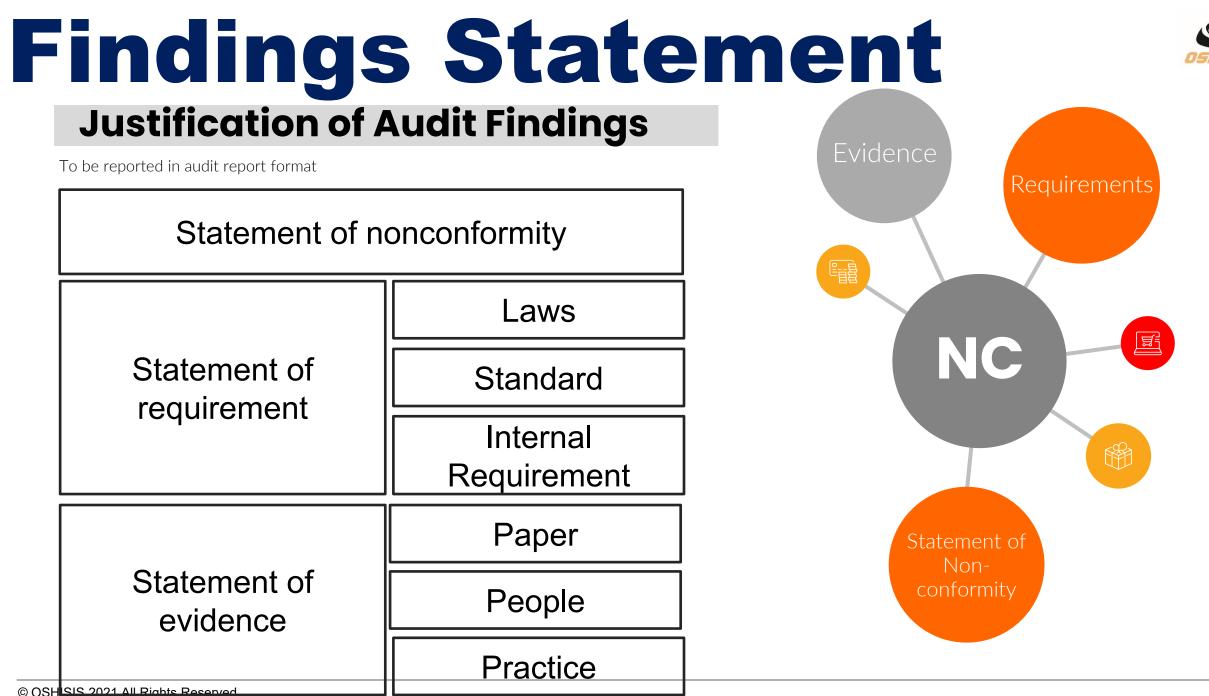
Specific context to emphasize

and address..

Risk

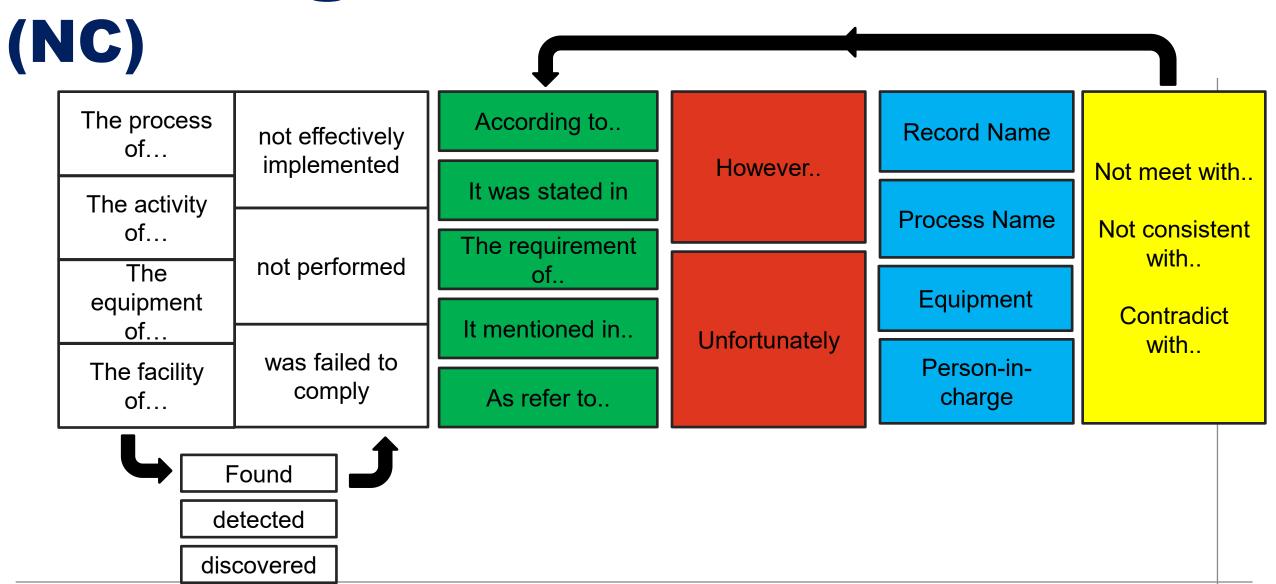
or

Benefit(s)



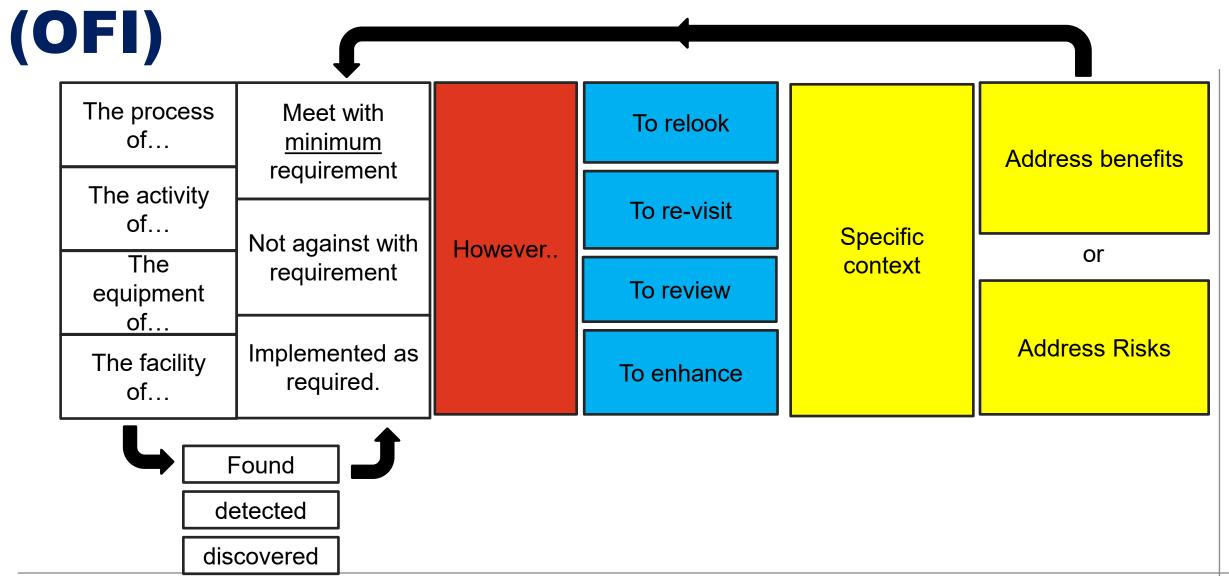
Findings Statement





Findings Statement





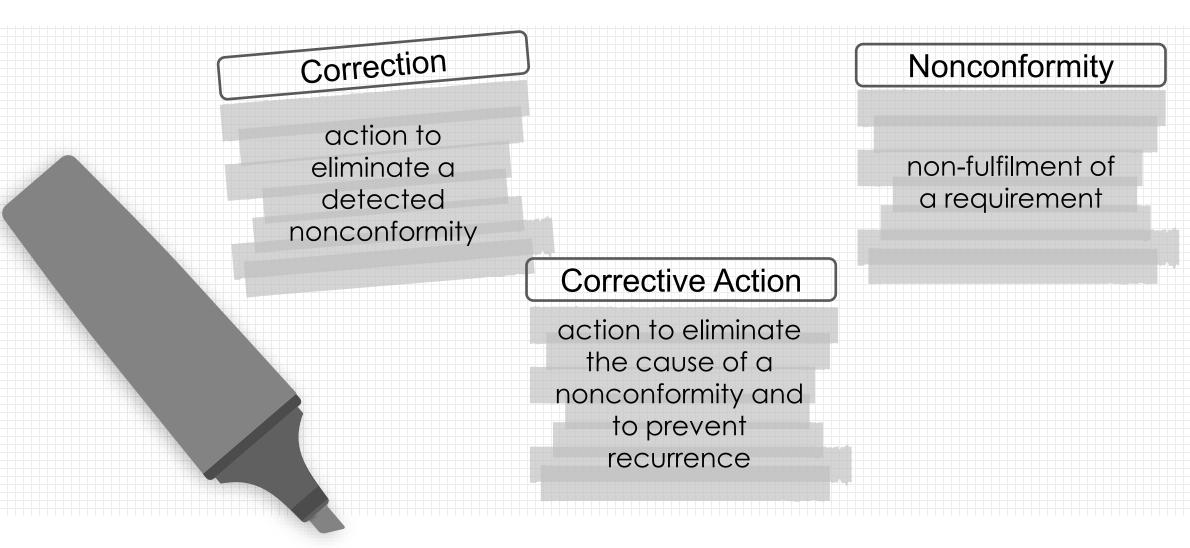


Module 9

Nonconformities, Corrective Action and Follow-Up



DEFINITION – ISO 9000:2015





NONCONFORMITY FORM (NCR Form Sample)

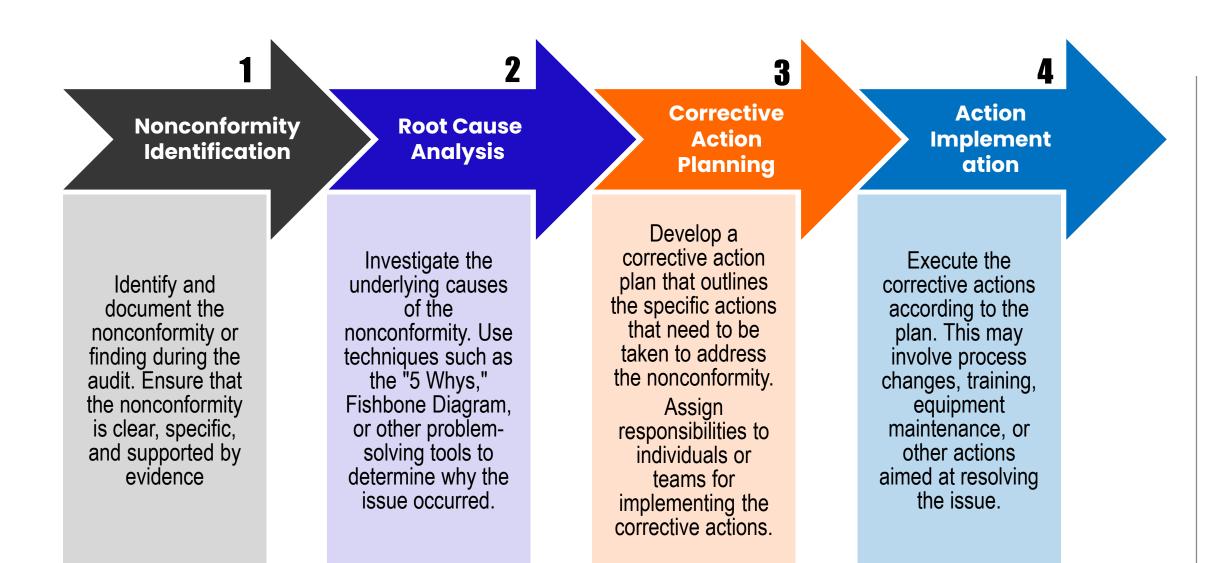
Date	Dept		Standard Clause	
NCR No.	Audited Area Auditor	Statement of No		
Correction		Corrective Action	[Provide a detailed of including what was requirements, and a Objec [Attach or reference identification of the documents, record	
[Describe any immediate actions taken to address the nonconformity, including containment actions.]		[Specify the corrective actions that will be taken to eliminate the nonconformity, including responsible parties and timelines.] [Describe any preventive actions that will be implemented to prevent recurrence of similar		
Root Cause Analysis		nonconformities.]	Approval and Review	
[Describe the root cause(s) of the nonconformity, if known. Use techniques like the "5 Whys" or "Fishbone Diagram" if applicable.]		Verification of Corrective Action [Outline how the effectiveness of corrective actions will be verified and who will be responsible for this verification.] [Signatures of the auditors involved in identifying the nonconformity.]	[Specify the names and signatures of individuals responsible for approving the corrective and preventive actions, and the date of approval.]	

Description of Nonconformity

Statement of Nonconformity						
cluding what was	description of the nor observed, discrepand any relevant evidence	cies from				
Attach or referentidentification of th	tive Evidence ce any evidence supp e nonconformity, such ds, or photographs.]					
proval and Review	Closure Status	Follow-up Action				
becify the names d signatures of ividuals ponsible for proving the rective and	[Indicate whether the nonconformity has been closed and the date of closure.]	[Document any additional actions taken during follow-up audits or reviews.]				
eventive actions, d the date of proval 1	[List any additional documents or evidence related to the nonconformity.]					

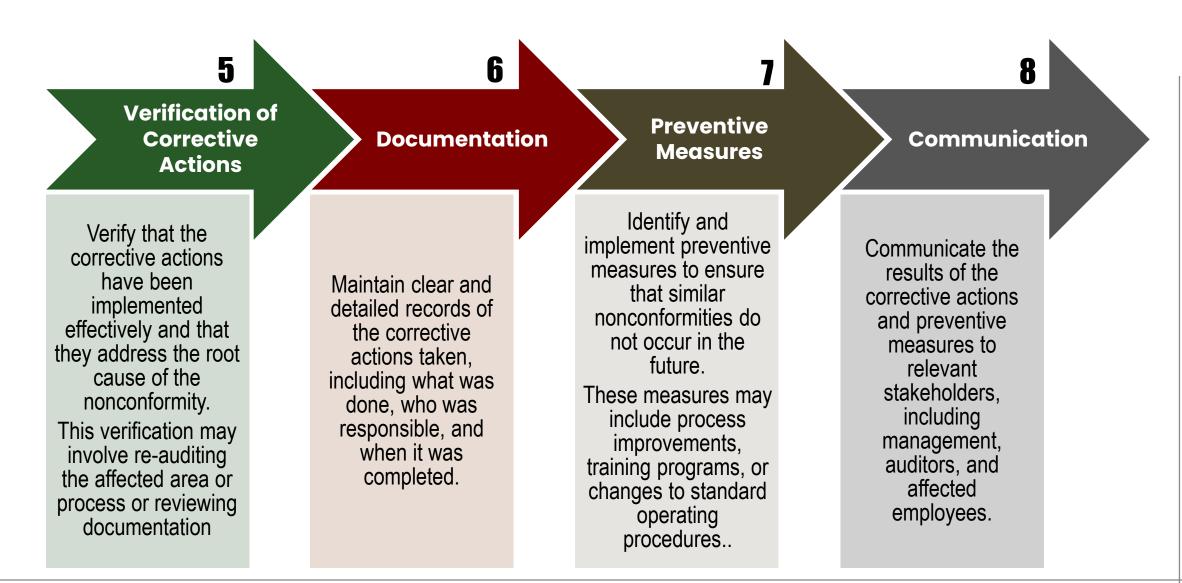
STEPS FOR IMPLEMENTING CORRECTIVE ACTIONS





STEPS FOR IMPLEMENTING CORRECTIVE ACTIONS





STEPS FOR IMPLEMENTING CORRECTIVE ACTIONS





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Review the effectiveness of the corrective actions and preventive measures. Ensure that they have resolved the nonconformity and prevented recurrence. Close out the nonconformity or audit finding once it has been satisfactorily addressed.

Follow-Up Audits

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Documentation and Reporting

Continuous Improvement

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Conduct follow-up audits to verify the sustained effectiveness of the corrective actions and preventive measures over time. Continue monitoring the area or process to ensure long-term

compliance.

Document the entire corrective action process, including the analysis, planning, implementation, and verification stages. Report on the status and effectiveness of corrective actions and preventive measures to management and relevant stakeholder

Use the lessons learned from the corrective action process to drive continuous improvement in the organization's processes, systems, and overall performance