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HOW INTERNAL AUDITING ADDRESSES YOUR ORGANISATIONAL COMPLIANCE CONCERNS

Dr Charles Beacroft 30th April 2025

Agenda

- Overview of ISO management systems
- Understanding what is internal auditing
- Benefits of internal auditing from a compliance perspective
- Q&A



— OUR — PURPOSE

IS TO HELP CUSTOMERS DELIVER PRODUCTS THE WORLD CAN TRUST

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NQA specialises in certification in high technology and engineering sectors.





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Certification body in **Aerospace** sector

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Certification body in **Automotive** sector

UK'S NO.2

Certification body in **Aerospace** sector



Dr Charles Beacroft NQA Training Manager



Your Presenter

- Completed 1 year as the Training Manager for NQA Certification Ltd
 - Responsible for growth in the NQA Training Business Unit
 - Managing the delivery of training courses for all NQA customers
- Lead Auditor in ISO 9001 and ISO 27001, Internal Auditor in ISO 14001 and ISO 45001
- Completed a PhD at the University East Anglia (UEA), with significant university lecturing and teaching experience at University College London, the London School of Economics, and UEA
- 4 years at Pearson Education, delivering postgraduate online distance learning for thousands of students at seven UK universities



OVERVIEW TO ISO MANAGEMENT SYSTEMS



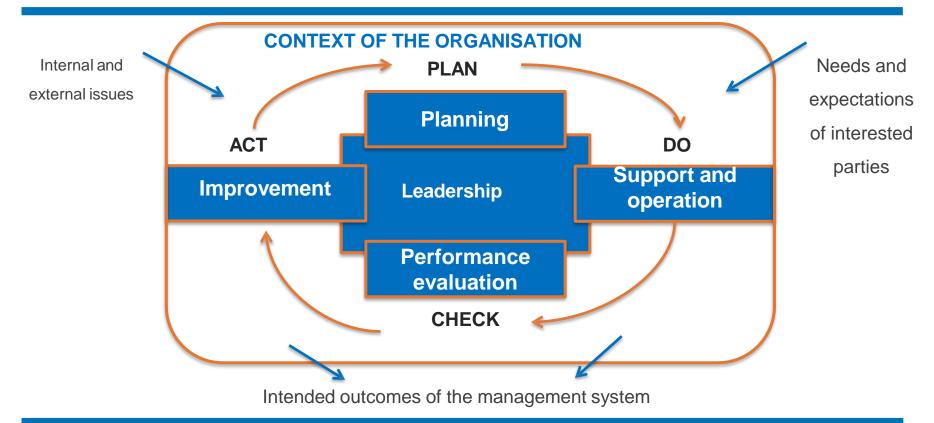
PROCESS MODEL APPROACH

- Determine the processes within the management system
- Identify inputs and expected outputs
- Determine their sequence and interactions
- Ensure effective controls
- Address risks and opportunities





PLAN-DO-CHECK-ACT





ANNEX SL HIGH-LEVEL STRUCTURE

- 1. Scope
- 2. Normative References
- Terms and Definitions
- 4. Context of the Organization
- 5. Leadership
- 6. Planning
- 7. Support
- 8. Operation
- Performance Evaluation
- 10. Improvement

Establishes the basis for the management system

Aligned to nearly all ISO management system structures





CLAUSE 9 & 10 OF THE ISO STANDARD

Clause 9 – Performance Evaluation

- 9.1 Monitoring, measurement, analysis and evaluation
- 9.2 Internal audit
- 9.3 Management review

Check

Clause 10 – Improvement

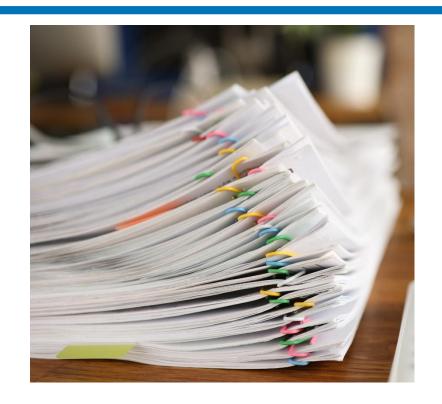
- 10.1 General
- 10.2 Nonconformity and corrective actions
- 10.3 Continual improvement

Act



PROCESSES AND DOCUMENTED INFORMATION

- A key part of a management system is the emphasis on the use of processes, including organisational, supporting, and operational processes
- All such processes are part of the documented information, which allows for the management system to be monitored
- This fits the requirements set out in Clause 7.5
 These documents are essential for conducting internal audits





UNDERSTANDING WHAT IS INTERNAL AUDITING



AUDIT

Audit: Systematic, independent and documented process for obtaining objective evidence and evaluating it objectively to determine the extent to which the audit criteria are fulfilled.

(ISO 14001:2015)



TYPES OF AUDIT

1st party - internal audits

Audits carried out by an organisation on their own systems.

2nd party - supplier/vendor audits

Audits carried out by an organisation on their suppliers / sub-contractors.

3rd party – external audits

Audits performed by an independent body, usually for certification purposes.



INTERNAL AUDIT OBJECTIVES

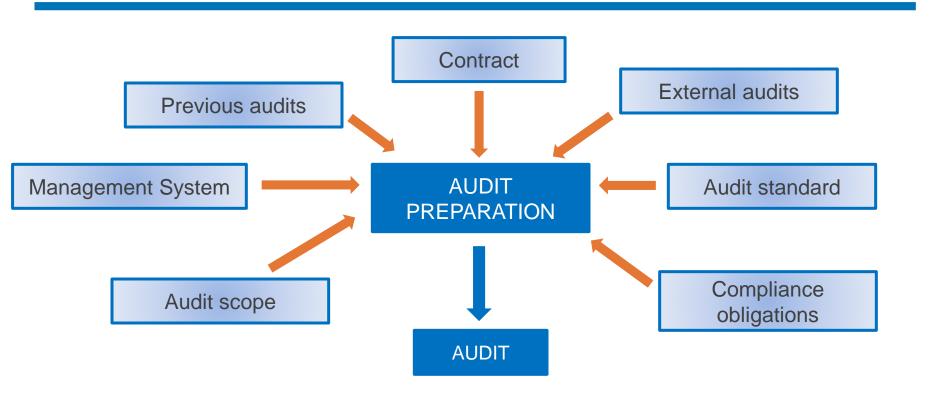


Audit objectives include:

- Conformance to the standard
- Conformance with their own management system
- Conformance to contract requirements
- Conformance to certification body requirements
- Conformance with compliance obligations



AUDIT PREPARATION – INPUTS





AUDITOR ACTIVITIES

- Conducting an opening meeting
- Following the audit plan
- Asking questions and recording answers
- Collecting evidence
- Assessing evidence against criteria
- Taking notes
- Conducting a closing meeting





AUDIT CHECKLIST

OI 1	
Sheet	of

Area/Procedure(s) audited Auditor	Date
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Question/Topic	Ref	Response	Evidence



AUDIT CHECKLIST (CONT.)

Question/topic	Ref	Response	Evidence
Overview of waste processing	(-)		
Review process	14001 8.1		
Method of receipt and storage			
Stored so as to prevent pollution, litter, spills?	14001 6.1.2		
Tour all waste storage areas and check			
Documentation –	14001 6.1.3		
Evidence of haz/non hazardous			
Detailed descriptions, SIC, EWC codes clear?			
Statutory and regulatory requirements			
Method of disposal clear/specified?	14001 6.2		
Any evidence of recycling in line with obj?	14001 6.2		
Identification of waste	14001 6.1.2		
Communication to other departments			
Identified – who and how?			
Communication	14001 7.4		
Who? How? Evidence?	14001 7.4		
Handling, storage, identification of waste	14001 8.1		
Competence and training of staff	14001 7.2		
Competence of contractor	14001 7.3		
Waste carrier and disposal permit	14001 6.1.3		
Process in event of loss of control of waste?	S. C. Carrier St.		



AUDIT CHECKLIST (CONT.)

Area/processes audited: Waste Management Auditor: C.

Moore Date: 17 July

Question/topic	Ref	Response	Evidence
Overview of waste processing	-	Toured the waste storage areas and discussed with Waste Manager.	Generally good housekeeping - Minor littering and oil spills.
Review process Method of receipt and storage Stored so as to prevent pollution, litter, spills?	14001 8.1	Production deliver waste to relevant areas Segregate to mixed general, haz, liquid, etc.	Observed areas for wood, plastic, oil and chemicals, etc. See photos. Some minor spills of
Tour all waste storage areas and check Documentation –	14001 6.1.2	Some minor spills observed in oil storage!	oil not cleaned up! Some uncovered skips, i.e., plastic!
Evidence of haz/non hazardous Detailed descriptions, SIC, EWC codes clear?	14001 6.1.3	Clear segregation in place in most areas Procedure makes these clear as does	Chemical and oil waste separated General mixed recyclables skip
Statutory and regulatory requirements Method of disposal clear/specified?		signage, i.e., for oil, wood, plastic, etc. Waste matrix provides waste routes	Wood skip Plastic skips
Any evidence of recycling in line with obj? Identification of waste	14001 6.2	Ongoing segregation and recycling figures available for 2015/16.	All segregated waste is recycled. Monthly figures seen for 2015/16
Communication to other departments Identified – who and how?	14001 6.1.2	Emails, posters, memos in place to	Email seen re: oil spills from WM
Communication Who? How? Evidence? Handling, storage, identification of waste	14001 7.4	production displayed to notice board. Toolbox talk on waste delivered February	to Production Manager. Topic segregation of waste - 7 pax Incl. D Wood, S Holmes, D
A 8 68	14001 8.1	Training is given at induction, in regular	Watson.
Competence and training of staff Competence of contractor Waste carrier and disposal permit	14001 7.2 14001 7.3	toolbox talks and following incidents, such as spills. Reviewed waste Carriers Permit for	TBT talk register seen for spills March - 8 personnel in attendance.
Process in event of loss of control of waste?	14001 6.1.3		Last collection April note number 15879 EWC and SIC clear - OK



REASONS TO RAISE NON-CONFORMITIES

'Non-fulfilment of a requirement'

Ref: ISO 9000:2015

- The audit criteria are not being met e.g.
 - System does not comply with the standard
 - Practice does not comply with their own management system
 - A specific customer contractual requirement is not being complied with
 - Regulatory requirements not being met



CONTENTS OF A NON-CONFORMITY REPORT

Appendix 1: Nonconformity Report

Report No:

ABC17/8/1

Sheet No 1 of 2

Department:

Auditor(s):

Stores

C Moore

Auditee manager: S

Keeper

Date of audit:

15 August

Standard ref: ISO 14001

Category

8.5.2

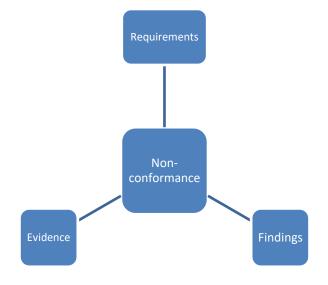
MAJOR MINOR (Delete one)

NONCONFORMITY/OFI (Delete one) - To be completed by the auditor

Not all waste held in stores is clearly identified

Oil waste in Area C e.g.

Chemical Waste in Area B



Signed (For department): S Keeper

Signed (Auditor): C Moore

CORRECTIVE ACTION

Action taken to eliminate the cause of a nonconformity and to prevent recurrence.

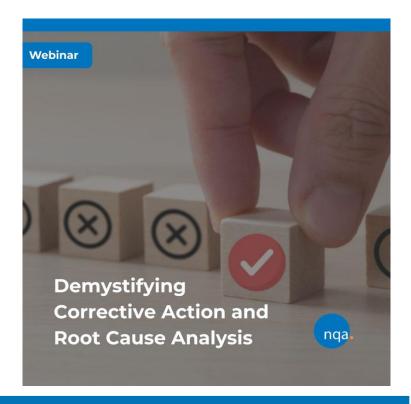
(ISO 9000:2015)



ROOT CAUSE ANALYSIS

- Eliminate the cause of the problem
- Prevent problems from recurring
- Address causes not symptoms

e.g. Re-train workers
 Mistake-proof
the process





CORRECTIVE ACTION

CORRECTIVE ACTION (To be completed by auditee and to include root cause analysis where applicable)

Identify all waste as required.

Review waste labelling system to establish reason for errors and amend as required.

Signed (For department): S Keeper

Signed (Auditor): C Moore

To be completed by: (Date): end of August

Follow-up date: end of Sept



VERIFICATION OF CORRECTIVE ACTION

VERIFICATION (To be performed by auditor, giving details of checks made)

Bar code labelling now introduced and procedure amended to reflect new system.

Operators have been given a toolbox talk on the need for and regulatory requirements related to waste and the importance of storing correctly and clearly identifying such.

All waste checked and found OK, for example batches 1379, 1568 and 1782

Corrective action complete and satisfactory: C Moore Date: 30 Sept



INTERNAL AUDITS – COMPLIANCE BENEFITS FOR YOUR ORGANISATION



Health Check

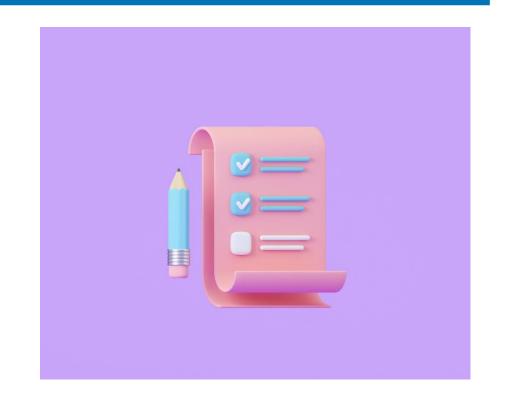
- Effective means of conducting a 'health check' on your organisation
- If done correctly, they are an objective, evidence-based review of your organisation's adherence to the relevant ISO standard and the compliance obligations





Compliance

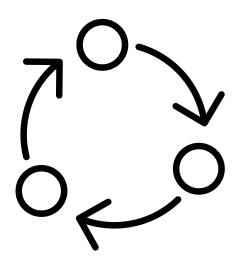
- Ensures that the system complies with both the requirements of the organisation, the ISO standard, and other relevant regulations
- If internal audits are not completed to the required standard, the organisation may lose its ISO standard certification which would be a significant risk to the organisation's model and aims





Continual Improvement

 Internal audits help organisations create a culture of continual improvement through reviewing processes, assessing nonconformance and suggesting improvements to the performance of the management system





Change Management

Internal audits allow for positive, productive change to occur with an organisation,
 especially if it is needed to improve a process to meet compliance requirements





Risk Management

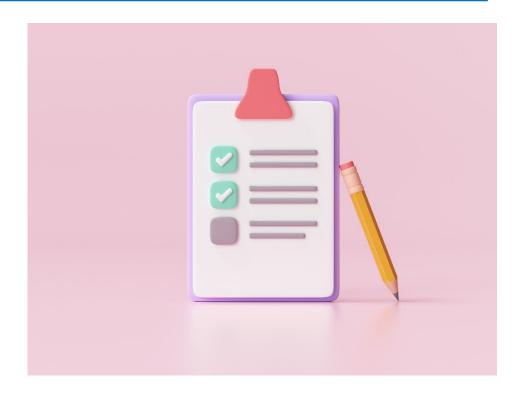
 Internal audits ensure that organisations are recording, assessing, and sufficiently mitigating risks within an organisation, with effective risk management essential for the success of an organisation's management system





Process vs. Individuals

 ISO requirements focus on processes over individuals, with internal audits focusing on the establishment of robust processes which remove single points of failure within an organisation





Internal vs. External

- Internal audits are usually completed annually (dependent on the risk level), as opposed to once every two to three years for external audits
- Internal audits provide a more accurate account of an organisation's compliance to the standard and its own processes, delving in more deeply than an external audit
- Effective means of improving the management system,
 with the results of internal audits essential for pushing
 the business forward on a regular basis





ORGANISATIONAL REQUIREMENTS FOR THE EFFECTIVE USE OF INTERNAL AUDITORS

 To ensure that internal auditors can bring the most value to an organisation, it is essential that the following steps are taken by organisation to ensure they have the most impact.

Formal training of Internal Auditors

Experienced and Knowledgeable Internal Auditors

Supported by Top Management Collaboration with all Employees of an Organisation



CONCLUSION



CERTIFICATION AND TRAINING SERVICES

We specialize in management systems certification for:



QUALITY



AEROSPACE (QUALITY)



AUTOMOTIVE (QUALITY)



SUSTAINABILITY



ENERGY



HEALTH AND SAFETY



INFORMATION RESILIENCE



FOOD SAFETY



RISK MANAGEMENT



MEDICAL DEVICES



NQA INTERNAL AUDITOR TRAINING



CQI and IRCA ISO 9001 Internal Auditor Training (A17931)

Virtual - Tutor Led (Quality)

O 2 Days III Level 2 - Intermediate



NQA ISO 14001 Internal Auditor Training

Virtual - Tutor Led (Environmental)

(3 1 Day III Level 2 - Intermediate



CQI and IRCA ISO 27001 Internal Auditor Training (A2574)

Virtual - Tutor Led (Information Security)

O 2 Days II Level 2 - Intermediate



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