

GETTING THE MOST OUT OF INTERNAL AUDITS



OUR — **PURPOSE**

IS TO HELP **CUSTOMERS** DELIVER PRODUCTS
THE WORLD CAN **TRUST**

NQA is a world leading certification body with global operations.

NQA specialises in certification in high technology and engineering sectors.





AMERICA'S NO.1

Certification body in Aerospace sector

TOP 3 IN THE UK

ISO 9001, ISO 14001, ISO 45001, ISO 27001

CHINA'S NO.1

Certification body in **Automotive** sector

GLOBAL NO.1

Certification body in telecommunications and **Automotive** sector

GLOBAL NO.3

Certification body in **Aerospace** sector

UK'S NO.2

Certification body in Aerospace sector



CERTIFICATION AND TRAINING SERVICES

We specialize in management systems certification for:



QUALITY



AEROSPACE (QUALITY)



AUTOMOTIVE (QUALITY)



ENVIRONMENT



ENERGY



HEALTH AND SAFETY



INFORMATION RESILIENCE



FOOD SAFETY



RISK MANAGEMENT



MEDICAL DEVICES

NATIONWIDE TRAINING SERVICES

ACCREDITED COURSES



Virtual Learning



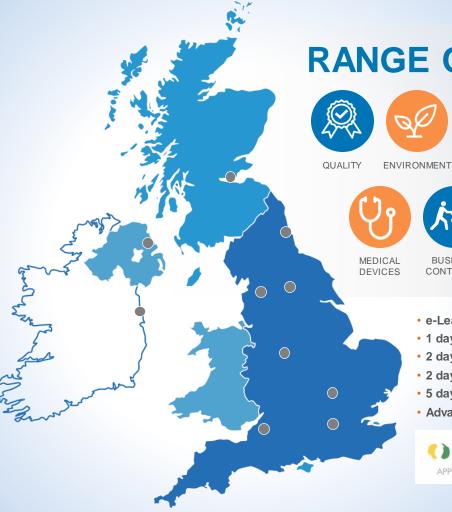
e-Learning / Live Webinars



In-house **Training**



Public Training Nationwide Locations















ENERGY

HEALTH AND SAFETY

SECURITY



BUSINESS CONTINUITY





INTEGRATED MANAGEMENT

• e-Learning Introduction

- 1 day Introduction Courses
- 2 day Implementation Courses
- 2 day Internal Auditor NQA or IRCA
- 5 day Lead Auditor NQA or IRCA
- Advanced Training







KEY INFO

- 45 minute webinar
- Questions in the chat box
- Q&A at the end
- Recording of webinar circulated shortly

YOUR PRESENTER



Judith Hargreaves

ISO 9001, ISO 14001, ISO 37001, ISO 45001, ISO 50001, PAS 43:2018, NHSS 17, ISO 17021-1



Technical and Projects Manager

Judith has been a Regional Assessor with NQA for 4.5 years covering multiple industries and key clients. She now sits within the Risk and Regulatory team reporting directly to the Global Accreditation Director.

Judith is also an Internal Auditor for NQA covering our global offices and occasionally joins our field team on client audits in order to keep up-to-date and involved with core Standards and industry developments.

She is currently responsible for gaining training and accreditation to so that she can help NQA to continually offer our clients the latest new standards, aid with transitions for updated Standards, and liaise between NQA and UKAS and other Regulatory bodies.



OBJECTIVES OF TODAY'S WEBINAR

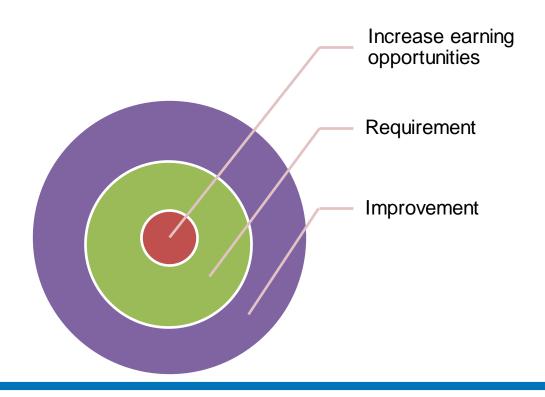
- Understand how to get the most out of the mandatory element of internal audits
- Plan effectively
- Improve and use



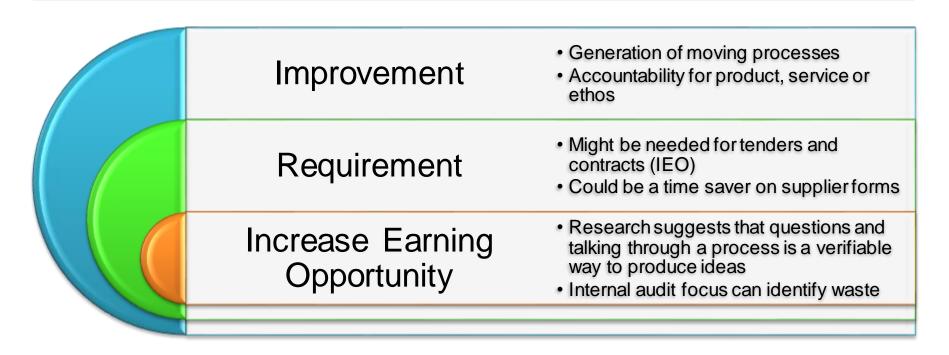


Why do you want the Standard you are certified for?











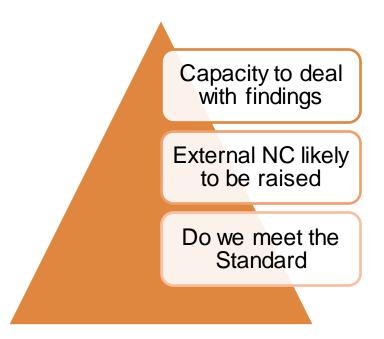
Increased earning opportunity:

What do we want to know to optimise this?





GET THE MOST OUT OF IT – REQUIREMENT





IMPROVEMENT:

Better place to work

More robust processes

Structure

Step to next level

Manage change



PLAN EFFECTIVELY

Now we know why we want the certificate we plan our audits already with some key objectives:

- What the Standards require
- Information which may be useful to you
- What core processes you want to test
- Highest risk

Who do you need involved?

- Process owner
- Process users



PLAN EFFECTIVELY

The Process Owner is accountable for that process and daily implementation

The Approver is the person that takes accountability for the content of the process

BOTH should be aware of their responsibility and what is in their own process.

Oversight of the use of the process is a good measure as to how it is working, but internal audit should question it, challenge it and find any aspect non-compliant

- On occasions where the anticipated timeframe for completion is exceeded, the complainant shall be kept updated in respect of progress by the nominated investigator(s). Progress reports (as with the outcome of the investigation) shall be formally communicated to the complainant and records retained. If issues are experienced with timeframes this may be managed by the Technical and Projects Manager or escalated to the Global Accreditation Director.
- 3.5 The individual(s) responsible for the validation, investigation and provisional decision shall be demonstrably independent from the subject of the complaint. For example; not those who conducted the audit or made the certification decision in question.
- 3.6 The individual(s) responsible for the investigation and provisional decision shall meet all competence criteria defined by NQA pertaining to the role of decision maker in respect of the scheme or schemes to which the complaint relates.
- 3.7 The outcome of the investigation and provisional decision shall be reviewed and approved by an applicable and demonstrably independent senior manager prior to communication to the complainant by the assigned investigator.
- 3.8 The Senior Manager with authority for the process subject to complaint shall be responsible for the initiation, monitoring and close out of corrective actions identified as being incumbent upon NQA. This shall be concluded in a timely manner.

Process Owner: J Hargreaves Approved: Steve Russell Issue 8 Feb 2023



PLAN EFFECTIVLEY

The plan itself should documented and a live document. Getting people out of the business to take part in an audit will always be met with 'haven't got time' however.... This can be seen as a risk potentially and can help form your plan!

Risk Register Audit Plan v1					
Risk	Because	Effect			
	Don't always				
	understand				
	improvements, can be	Waste of time, no further			
Lack of buy in from	seen as a negative	forward with			
auditee	exercise	improvements			
People needed not		Not able to discuss full			
available	Time factor, holidays	process or deficiency			
Not finding a major		Close out costs, potential			
problem which the	Not confident,	cost of special visit,			
external auditor	material not available,	becoming reactive rather			
does	not enough training	than proactive			
		Close out costs, potential			
Not knowing the		cost of special visit,			
process well	Not the area familiar	becoming reactive rather			
enough to audit it	with	than proactive			
	Risk Lack of buy in from auditee People needed not available Not finding a major problem which the external auditor does Not knowing the process well	Risk Because Don't always understand improvements, can be seen as a negative exercise People needed not available Not finding a major problem which the external auditor does Not knowing the process well Don't always understand improvements, can be seen as a negative exercise Time factor, holidays Not confident, material not available, not enough training			



PLAN EFFECTIVLEY

Assuming we have assessed risk we can put communications out to help people understand, plan around holidays and factor in when you can target busiest departments.

Your plan must be documented but it doesn't have to be pretty!

Audit areas	Needed	Auditor	Date
	Process owner,		
	Process		3rd
	approver,		March
Sales process	process user	Jude	2023
	Process owner,		
	Process		18th
	approver,		April
Operations	process user	Jude	2023
	Process owner,		
	Process		2nd
Management	approver,		Septemb
Processes	process user	Jade	er 2023



THE AUDIT ITSELF

Gold	Waste of Time	
Explain the process as you understand it and show me how it works	Do we have a process? Yes, anything changed? No – job done	
S.3 of the process says we have a Form32 to complete x where are these and show me the last 5 used	I need to do an audit – has anything unusual happened since last time? No – Job done	
Can you show me an example of where you might have had to work outside of the process and tell me why?	Can you give me 5 order numbers so I can put them in my report	
What parts of the process could be improved?	I have found that 3 out 5 examples were incorrect – I won't write it down just fix it	
What do we do when x happens	If you can't find it now just put something together for me and email it	



THE AUDIT ITSELF

- s.3.4 what is the timeframe? Show me how this is met and monitored show me emails of the complainant being updated show me the records of investigation for xxx
- s.3.5 show me how you verified independence
- s.3.6 show me the competence records in relation to this specific complaint
- s.3.7 show me the verification of the senior manager

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Risk Register Audit Findings v1						
Reference	Risk	Because	Effect			
	Document control					
	was not seen to be					
	effective in all areas		Puts us on radar for			
	- Old contract used,	Can be requested by	external agencies, costs			
	out of date H&S	HSE or EA, contract	may be incurred. Old			
	policy seen, waste	may be causing	contracts may haveold			
	transfer notes	detriment hence	clauses making void or			
R1	missing off system	change	costing us			
			Loss of business,			
	Despatch dates not	shift work - might be	reputation, unknown scle			
R2	clearly identified	missed by everyone	of problem until complaint			
		The issues were not				
		classed as NC's as they				
		were dealt with at the				
	Non-conformances	time. Process not	Cannot monitor how many			
	not raised and put	deemed worthwhile	errors are made, if there is			
R3	into the system	by staff	a training issue, costs			
			An incident may occur and			
	Daily checklists	Responsible person	documentation shows a			
	were seen to be	says never a problem	check that did not happen			
	ticked even on	and is trying to save	placing all documents into			
R4	dates not yet been	time.	quetsion.			



1

Consider assigning an educated guess to the risks eg HSE visit cost

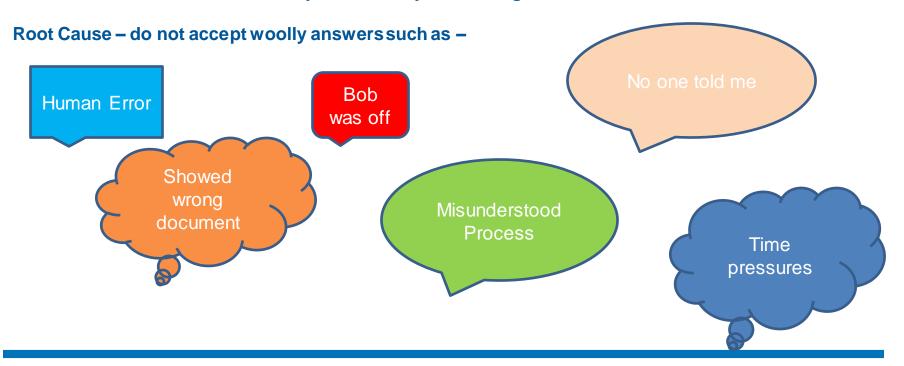
2

Put a mitigation plan in place to reduce the risk of this reoccurring 3

Delegate these risks to a specific owner to manage and review



The corrective action will be closely linked with your risk register:





Whilst these are all reasonable – they tell us nothing!

Human error – what can the process do to prevent human error –software? reminders? Quality checks?

Showed wrong document – how was that accessible and what can be done to prevent this as incorrect documents should not be in the system. Is it a training issue – did the not know what was being asked?

Bob was off – who can do Bob's job when he is off?

No one told me – what process do we have in place to make sure ALL employees are aware, is the current one failing?



How much has root cause and corrective action cost?

- Time spent
- Any rework
- Chasing people for paperwork and redoing root cause

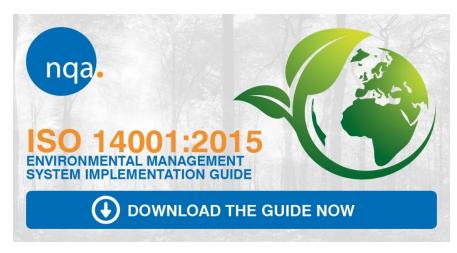


- Find positives!
- What is working well that can be replicated elsewhere in the business
- Have we found any rising stars?



NQA RESOURCES

- Training
- Regular news, legislative updates and blogs via our website. Or sign up to InTouch <u>here</u>.
- Implementation guides >>>
- Annex SL Comparison Tool download
- NQA Associate Partner Programme







ADVANCED TRAINING COURSES

- Leadership within ISO
- Effective management of change
- Managing your supply chain relationships
- Effective evaluation of compliance
- How to identify risks & opportunities
- Participation & consultation of workers
- Demonstrating customer satisfaction
- Managing information security remotely
- GDPR How to think like a DPO
- Operational resilience planning
- Risk / process based auditing
- Effective root cause analysis

www.nqa.com/en-gb/training/advanced





Q8A



THANK YOU

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