

# Internal Audit Summary Page

Date of Audit: \_\_\_\_\_

<p><b>Step 1 – Select one process only</b></p> <p> <input type="checkbox"/> Governance Processes  <input type="checkbox"/> QMS Processes  <input type="checkbox"/> Business / Support Processes  <input type="checkbox"/> Sample Processes  <input type="checkbox"/> Testing/Inspection Processes  <input type="checkbox"/> Review / Decision Making Processes  <input type="checkbox"/> Evaluation / Certification Processes         </p>	<p><b>Name of Process</b></p> <p>_____</p> <p>(Comments)</p> <p style="text-align: right;"><i>(See Summary of Findings Pages)</i></p>	
<p><b>Step 2 – Document Review</b></p> <p>Is system documented?   <input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> N/A</p> <p>Is system implemented?   <input type="checkbox"/> Yes   <input type="checkbox"/> No   <input type="checkbox"/> N/A</p> <p>Note: Any “No” must result in a finding</p>	<p><i>(Review Documents – prepare Checklist Page)</i></p>	
	<b>Are These Things Available?</b>	<b>Are Records Maintained?</b>
Documents that govern the process	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Supporting procedures	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Necessary space and environment	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Staff training	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Staff qualification	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Supporting QMS Procedures <ul style="list-style-type: none"> <li>• Feedback, disputes/appeals</li> <li>• Identification of NCs, PNCs, OFIs</li> <li>• Determining impact and root cause if needed</li> <li>• Corrective / preventive action if needed</li> <li>• Follow up for effectiveness if needed</li> </ul>	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
<p><b>Step 5 - General Comments</b></p> <ul style="list-style-type: none"> <li>• Findings overleaf have been agreed</li> <li>• Recommendation for next audit:   <input type="checkbox"/> As regularly scheduled</li> <li style="margin-left: 100px;"><input type="checkbox"/> On _____ (date)</li> </ul>		

\_\_\_\_\_  
(Lead Auditor)

\_\_\_\_\_  
(Date)



# Internal Audit Summary of Findings

Step 4 – Complete one Finding Form for each entry

Finding No.		Detail of Finding <i>(cite procedure/document/record)</i>	Recommend Level	
Int Aud #	IDR Log #			
			OK <input type="checkbox"/>	OFI <input type="checkbox"/>
			PNC <input type="checkbox"/>	N/C <input type="checkbox"/>
			OK <input type="checkbox"/>	OFI <input type="checkbox"/>
			PNC <input type="checkbox"/>	N/C <input type="checkbox"/>
			OK <input type="checkbox"/>	OFI <input type="checkbox"/>
			PNC <input type="checkbox"/>	N/C <input type="checkbox"/>
			OK <input type="checkbox"/>	OFI <input type="checkbox"/>
			PNC <input type="checkbox"/>	N/C <input type="checkbox"/>

**Incident and Deviation Report**

Date: \_\_\_\_\_  
 Serial # \_\_\_\_\_  
 Number of pages attached \_\_\_\_\_

**Note:** Only one incident or deviation per report.

Deviation  Potential Deviation  Opportunity for Improvement

(Select one ref only)  MOTIVA QMS: \_\_\_\_\_  External: \_\_\_\_\_  
 →

1. Description of the incident or deviation  
 \_\_\_\_\_  
 \_\_\_\_\_

2. Description of the immediate remedial action (remediation) taken, including any correction or prevention  
 \_\_\_\_\_  
 \_\_\_\_\_  

QM review (initials)	Investigation assigned to	Date:
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3. Is full Corrective/Preventive Action Required? Yes if there are any "Yes" boxes checked.

	Yes	No	
Is there an unacceptable risk to ABC?	<input type="checkbox"/>	<input type="checkbox"/>	<i>If all answers are "No" then only remediation is required.</i>
Is the technical validity of ABC results affected?	<input type="checkbox"/>	<input type="checkbox"/>	
Is it easier to effect permanent resolution than many little remediations?	<input type="checkbox"/>	<input type="checkbox"/>	

4. Proposed Solution (and Investigation of Root Cause if required) Date Due: \_\_\_\_\_

Root Cause(s) of condition:	<b>Not required (eg: remediation only)</b> <input type="checkbox"/>
Proposed solution:	Corrective Action <input type="checkbox"/> Preventive Action <input type="checkbox"/> Remediation Only <input type="checkbox"/>
Investigator's Signature and Date _____, _____	

5. Confirmation of Solution Implementation

Condition resolved (root cause eliminated/opportunity exploited) <input type="checkbox"/>	Date implemented _____
Supervisor/Manager Initials _____	QM closure (Initials) _____

6. Follow up Date Due: \_\_\_\_\_

Follow up required? Yes - <input type="checkbox"/> No - <input type="checkbox"/>	If not, why not? _____
Monitoring of condition assigned to: _____	Date Completed _____
"Solution is deemed EFFECTIVE." <input type="checkbox"/>	QM review (Initials) _____

Date: \_\_\_\_\_

Serial # \_\_\_\_\_

Number of pages attached \_\_\_\_\_

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Potential Deviation

Opportunity for Improvement

**(Select one ref only)**

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MOTIVA

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Date: \_\_\_\_\_

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MOTIVA

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