Internal Audits & Inspections

**Definition**

Internal audits are an in-house check to identify gaps or deficiencies in the SQF System.

**Applicable Code Requirements**

1. 2.5.4
2. 2.1.2.1 iv.
3. 2.5.3.1

**Review Glossary Terms**

1. Audit Checklist
2. Competence
3. Corrective Action
4. Exempt
5. Non-conformance
6. N/A
7. SQF System
8. System Elements

**Implementation & Audit Guidance**

**What does it mean?**

This element requires the site to audit the activities in their System on a regular basis to ensure that everything is running as intended. The internal audit program should include all aspects of the system elements and the good manufacturing modules at least annually. The internal audit can be broken down into multiple pieces and conducted throughout the year to meet the annual requirement. However, all elements must have been audited at least once. It may be necessary to update the audit schedule based on customer complaints, internal or external food safety events such as recalls or regulatory warning letters, or internal or external audit results.

This program requires a document(s) that outlines the methods and responsibilities, schedule, and process for completing the internal audit all the way through to verification of corrective actions.

For internal audits to be effective, staff conducting internal audits must be trained in internal auditing techniques, information gathering and objective observation. This training need not be “formal” training provided by an external source. Internal auditor training covers internal audit procedures, including the planning and scheduling of internal audits, preparing internal audit reports, and initiating and following up on audit findings. Internal audits should combine several information gathering techniques, including interview of personnel, review of records and observation of current conditions.

To ensure the objectiveness of the internal audit the site is required, where possible, to use personnel who are separate from the area being audited to conduct internal audits. The inclusion of the words “where possible” illustrates that in the case of some very small suppliers this may not be possible. In such cases, the site is required to demonstrate that the alternative internal audit arrangement meets the objective of this requirement and is communicated to all affected parties.

Finally, the outcomes of all internal audits, including any corrective actions taken, must be recorded. Any audit tool that is developed by the supplier can be utilized to perform the internal
Audits provided it covers the required areas and programs. Records of internal audits should be included in your document control and record policy.

Internal audit results are communicated and included as a part of the management review process.

**Why is it in the Code & why is it important?**

**This is a mandatory clause.**

Internal audits help the site to identify faults in their System so that it can be improved. The internal audit program is a verification method and when used properly, can identify risk, areas for improvement and/or needed corrective and preventative actions prior to an external audit or the occurrence of a food safety event.

See RIO Chart on following page.
### RIO Road to Audits (Records, Interviews, and Observations)

<table>
<thead>
<tr>
<th>Records</th>
<th>Interviews</th>
<th>Observations</th>
</tr>
</thead>
<tbody>
<tr>
<td>The following are examples of records and/or documents to assist in the implementation and review of this topic:</td>
<td>The following are examples of people to interview to assist in the implementation and review of this topic:</td>
<td>The following are examples of observations to assist in the implementation and review of this topic:</td>
</tr>
<tr>
<td>- Internal Audit Procedure</td>
<td>- Internal Audit Team</td>
<td>- An internal audit or facility inspection being conducted.</td>
</tr>
<tr>
<td>- Audit Schedule</td>
<td>- SQF Practitioner</td>
<td>- Corrective actions from internal audits in practice.</td>
</tr>
<tr>
<td>- Audit Checklist or similar tool</td>
<td>- Senior Site Management</td>
<td></td>
</tr>
<tr>
<td>- Audit Report</td>
<td>- Corrective Action Reports</td>
<td></td>
</tr>
<tr>
<td>- Objective Evidence Gathered</td>
<td>- Training Records</td>
<td></td>
</tr>
<tr>
<td>- Corrective Action Reports</td>
<td>- GMP Inspection Reports</td>
<td></td>
</tr>
<tr>
<td>- Training Records</td>
<td>- Management Review</td>
<td></td>
</tr>
<tr>
<td>- GMP Inspection Reports</td>
<td>Meeting minutes</td>
<td></td>
</tr>
</tbody>
</table>

### Additional References

- SQFI Checklist (Excel):