

Auditing Certifications and Requirements

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Outline

- What and Who
- How to Start
- Qualification Audit
- Facility Audits





What and Who

- An auditing agency is an organization that is independent of any facilities that manufacturer products for consumer interest.
- Auditing agencies will consist of inspectors/auditors that perform unannounced inspections of facilities that have registered in a standard program.
- Agencies are qualified and accredited by a certification body that ensures the agency is performing inspections per the standard program.
- Qualified agencies are typically found on the certifications body website.



How to Start

- Log onto the certifications' body website to determine which agencies have been qualified to perform these inspections.
- Contact agency and discuss prices, contracts, etc.
- Once relationship has been established with an accredited agency, a qualification audit will be scheduled.
- Auditors will be performing inspections in accordance with regulations set forth by the Pellet Fuel Institute Residential/Commercial Densified Fuel QA/QC Handbook



Qualification Audit

- Once the auditing agency and manufacturer have agreed upon a contract, a qualification audit will be scheduled.
- During this audit, the inspector will review all relevant documentation and equipment to be sure the facility is in compliance with the established regulations.
- Inspector will also collect samples to be sent to an accredited independent laboratory.
- Once this audit is complete, the certification body will be informed by the agency that the facility has been qualified.



Audits

- Inspector/Auditor is required to perform 12 unannounced audits in a calendar year. (Typically once a month)
- During the audit, the manufacturing facility is required to have all relevant documentation easily accessible for the auditor to inspect.
- This will include production since the last audit and all QC records.
- There will also be an inspection of bags and labels to make sure they are in compliance.



Audits

- Once the inspector/auditor has completed the records check, they will then inspect the facility grounds for properly labeling of bags and storage of material.
- Sample bags (40 lbs.) will then be randomly selected from the inventory at a rate of 1 bags for every 1,000 tons produced and submitted to an accredited laboratory for analysis. One bag must be submitted on each visit.
- If during the audit, any corrective actions are necessary than the auditor will communicate with the facility these corrective actions before leaving the plant.



Corrective Actions

- There are two measures where corrective actions maybe necessary.
 - During the audit, the inspector finds documents or materials (bags, labels, etc) that does not meet the regulations.
 - After the agency has received the laboratory analysis from the samples submitted by the auditor.
- Both corrective actions require the agency to increase the frequency of the inspections until the facility has satisfied the requirements of the regulations.
- If the corrective actions are not resolved during follow up inspection, than the facility can be subjected to Warning, Operational restrictions or Suspensions.



Please understand that the auditors are not at the facility to "Police" or to control the way a facility does business. Their only purpose is to ensure the certifications and proper production of material to the clients. Most auditors and manufacturing facilities have a very good working relationship.



THANK YOU FOR YOUR TIME

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