

AS9100 Issues

Challenges Pitfalls Internal Auditing Continual Improvement

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Challenge: Aerospace Wants Improvements

Aerospace Supplier Improvements

- Better monitoring and measuring of processes
- Improved Quality & Delivery
- Better Internal control of processes, paperwork, traceability, schedules
- Utilizing Nonconforming Product info for Data Analysis, CA, & PA
- Systematic Root Cause Analysis
- Proactive Preventive Actions (# CA's vs PA's)
- More effective Internal Audits with Findings (you should be finding your problems)
- More effective Corrective Actions directed toward Process not Product
- Documentation requirements for certs, test results, etc.
- Improved Coordination with subcontractors
- Clear information in a Traveler Package

Registrar Improvements

- Tougher guidelines for auditors
- Registration/Surveillance Audit results consistent with performance to the primes
- Measuring # Findings
- Site Audits at Registrar & Witnessed Audits at Suppliers Registration or Surveillance
- Tracking individual auditors and # of OFI's vs # of NC's
- Primes do not want OFI's when a NC is warranted

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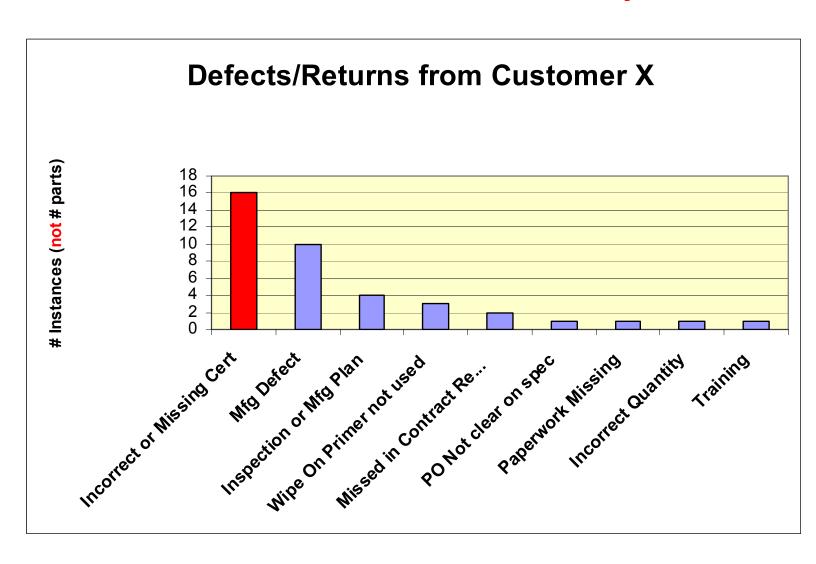
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Effective Corrective Action

Aerospace is not getting better products from Suppliers through corrective action process

There are repeat problems

Points to Ineffective Root Cause Analysis



Basic Root Cause Analysis

- 1. Identify the problem
- 2. List the potential root causes.
 - → Materials → People → Environment
 - → Methods (Process) → Machines
- 3. Are there sub-causes? List them.
- 4. Pick the most likely causes.
- 5. What action is needed to fix/prevent them.

Data Collection?

Quick fixes?

Next Steps?

6. Action implemented and monitored?

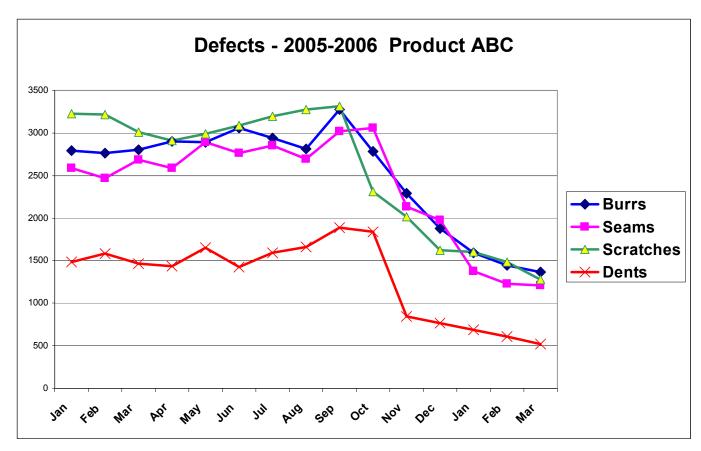
Process Change

Clear Roles & Responsibilities

Define what to monitor to see if fix is effective

- 7. Did it work?
 - → No, go back to line 4.
 - → Yes, document it, standardize it, and celebrate!

Link Metrics to Process Changes



Process Log - Product ABC		
Date	Change	Comments
9/15/2005	Started Preventive Maintainance	Machines 1 and 2
10/12/2005	Expanded PM	Machines 3,4,5,6
10/20/2005	Training - Mat'l Handling	Receiving & Whse
11/5/2005	Replaced all needlenose pliers	
12/18/2005	New Setup Fixtures at Wct 15	Operators trained
1/5/2006	Training - Defect Identification	Raw Mat'l
1/30/2006	Changed Suppliers	Housings and glass

Internal Auditing

The Purpose of Internal Auditing

- Determine Compliance
- Check the effectiveness of business processes
- Ensure Customer requirements are met
- Identify Opportunities for Improvement
- Voice of the process

This is the power of the QMS for improving the business

Make the data & metrics talk to you!

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Pitfalls, Potholes, & Speedbumps Taking Internal Auditing to Next Level

- Missing Signatures on Operations
- Changes on Traveler w/o Initials & Dates
- Unidentified material, bottles, cans, etc.
- Expired items
- Epoxy or gluing Pot Life not identified
- Poor configuration control
- Quantities on Traveler do not add up
- "As Found" and "As Left" not on Cal History
- Comparison of Certs to Spec on Receiving
- Not flowing down all requirements on PO
- Flow down CA to suppliers
- Follow Up on CA's when results not achieved
- Clear & Complete Design Plan with task sequence, significant stages, roles, resp, safety, key characteristics
- Design Review "authorize progression to next stage"
- Clear & Documented Verification & Validation Testing

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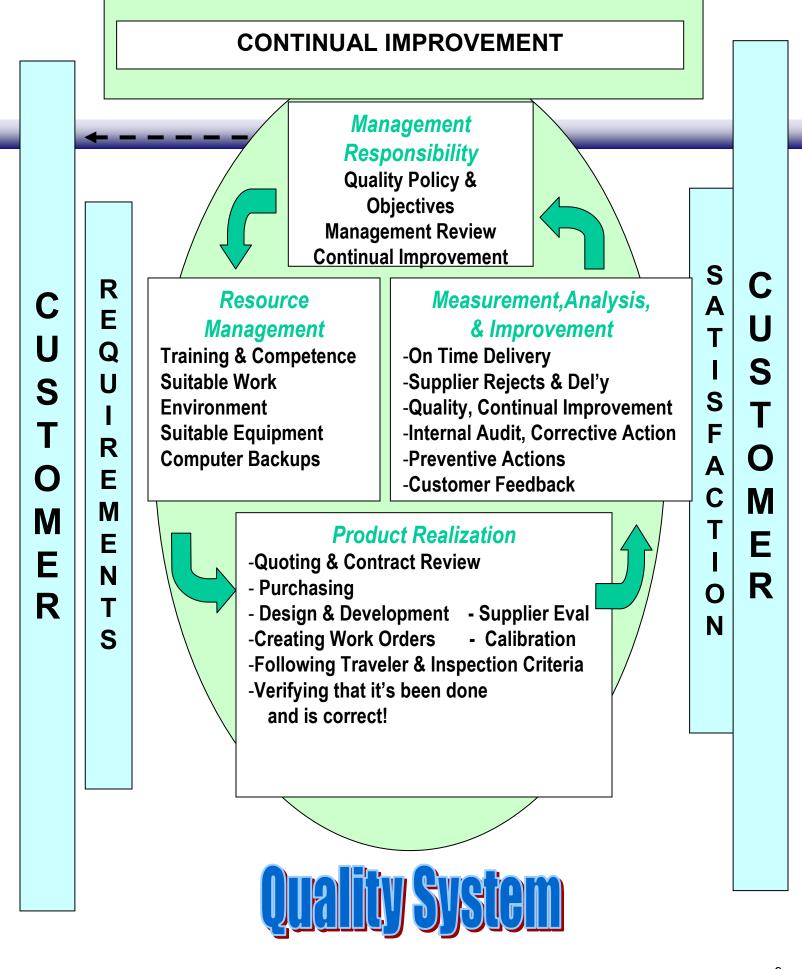
Continual Improvement

Taking Internal Auditing to Next Level

Process Audits

- Take a job that just shipped and check the entire system
- Use a repetitive problem, conduct root cause analysis, find the breakdowns in the process that allow this to happen
- Use Customer Identified Problem, Return, or something they feel is important
- Find recent screw up and audit that process

Sequence & Interaction of Processes



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