

June 19, 2017

Rev A

# Continuous Improvement Workshop: Permanent Corrective Actions

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## Agenda

- **Purpose & Goals of this Workshop**
- **Guidelines on Initiation Criteria**
  - When and Why we issue a Corrective Action
- **Best Practices on How to Write a Corrective Action Response**
  - Section-by-section
- **TIPQA System**

## Moog Quality Team

- **Site Quality Manager – Dawn Salvatore**
- **Assembly Quality Engineers**
- **Quality Assurance Manager**
- **Purchasing Manager and Buyers**
- **Supplier Quality Engineers**
  - Steve Van Raay – Outside Processes
  - Bob Mietzner – Machined Parts
  - Daniel Hensel – Machined Parts & all other

# Purpose & Goals of this Workshop

## **Guidelines – Corrective Action Initiation Criteria**

- **After today you should know:**
  - Why a Corrective Action is issued by Moog
  - Who should respond to a Corrective Action
  - How to perform Containment
  - How to arrive at the Root Cause
  - The purpose of Objective Evidence
  - The difference between Corrective and Preventative Action
- **Our goal is to provide you with the toolbox to perform the best possible RCCA you can.**

# Guidelines: Why and When We Issue a Corrective Action Request

## Guidelines – Corrective Action Initiation Criteria

What is a “Corrective Action”?

A **Permanent Corrective Action** is immediate and preventative action taken on an assignable cause to **permanently** fix a systemic or process-related issue

## Guidelines – Corrective Action Initiation Criteria

What is a “Corrective Action”?

Ongoing Improvement initiatives

*Documents* the work done!

NOT a reprimand



## **Guidelines – Corrective Action Initiation Criteria**

**A Corrective Action is an  
Opportunity for *Improvement***

## Guidelines – Corrective Action Initiation Criteria

Moog Customer  
Request

Audit Finding

Significant  
Cost/Safety

Recurring Issue

## Guidelines – Corrective Action Initiation Criteria

- Corrective Actions that you submit are not only reviewed by internal parties, but are subject to review by others
  - Including customers, auditors, and regulating bodies from government agencies
- It is critical that all Corrective Actions submitted stand the scrutiny of the aforementioned parties
- It should be understood that what we do is necessary for the **safety and well-being of the public**

**What is our #1 priority?**



# **Best Practices:** **How To Perform a Good Root Cause Investigation and Permanent Corrective Action**

## Best Practices – Administrative Data

### Administrative Data

- Part Number, Part Quantity, Rejection #, Date CA issued, Date CA DUE
- BEST PRACTICES: Make sure the data is complete and correct. Be mindful of Bill Of Material (BOM) differences, revision, and affected quantity

It is Moog's responsibility to provide all the Administrative Data,  
But it is your responsibility to validate it is complete and correct.

## Best Practices – Description of the Finding

### Description of the Finding

- Describe the Non-Conformance. The action will only be as good as the finding!
- BEST PRACTICES: Be as objective and complete as possible. Make sure all references are noted, such as drawing feature sheet/zone, specification paragraph, PO provision, etc.
- Statement should describe “should be” condition and “is” condition.
- Again, the root cause will only be as good as the description of the finding.

It is Moog’s responsibility to provide as much information as clearly as possible,  
But it is your responsibility to make sure you **understand the problem!**



## Best Practices – Moog CA Worksheet

Moog		5 WHY PROBLEM SOLVING	
<b>Identification:</b>		<b>5 Why Analysis - Why did this happen?</b>	
Date: / /	Area/Location:	<b>Why?</b>	
Originator:	Part #:	1st Why	
Team:	WD/PO #:		
	Supplier #:		
	Customer:		
<b>Problem Category:</b>		<b>Why?</b>	
<input type="checkbox"/> Reject at Receiving Inspection	<input type="checkbox"/> Documentation	2nd Why	
<input type="checkbox"/> In-Process Reject	<input type="checkbox"/> Field Return		
	<input type="checkbox"/> Customer return		
<b>Problem Description:</b>			
<b>Containment Action</b>		Date: / /	<b>Why?</b>
		3rd Why	
<b>Immediate Corrective Action:</b>		Date: / /	<b>Why?</b>
		4th Why	
<b>Permanent Corrective Action:</b>		Date: / /	<b>Why?</b>
		5th Why	
<b>Preventative Corrective Action:</b>		Date: / /	<b>Root Cause:</b>

Input Problem Description in your own words here



**After Receiving a Corrective Action request the first thing you need to do is...what?**

# Best Practices: Containment

## Best Practices – Containment

- What is your organization going to put in place **IMMEDIATELY** to prevent Moog from receiving any further non-conforming product?
- **All** throughout the supply chain:
  - **YOUR** suppliers and sub-tier suppliers
  - Raw, WIP, and Finished Inventory
  - In-transit to Moog
  - Moog Incoming Stores
- What did **YOU** ship in case **WE** missed it?
- Identify **Inspection GAP** and implement **SHORT TERM detection method** to prevent non-conforming product from reaching Moog
- Ensure **ALL** necessary personnel in your organization/supply chain are **AWARE** of the non-conformity **AND** the Containment Actions

## Best Practices – Containment

- Has to happen in the 1<sup>st</sup> **24-48 Hours** after a Corrective Action request.
- Must ID **ALL** the product in the Supply Chain – from your suppliers; to your Raw, WIP and finished Inventories; to product in-transit to Moog; to Moog incoming stores
- **Awareness is key.** All necessary personnel must become immediately aware of the defect and the actions taken to contain it.



STOP THE  
BLEEDING.  
CONTAIN THE  
OIL.

Is this a  
Permanent  
Corrective  
Action?

## Best Practices – Containment

- **There are two parts to Containment:**

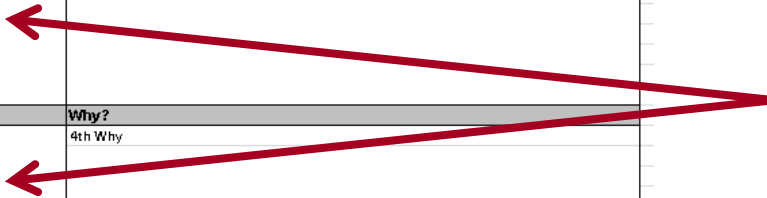
**Contain Defective Material**

**Identify Inspection Gap**

## Best Practices – Moog CA Worksheet

Moog		5 WHY PROBLEM SOLVING	
<b>Identification:</b>		<b>5 Why Analysis - Why did this happen?</b>	
Date: / /	Area/Location:	<b>Why?</b>	
Originator:	Part #:	1st Why	
Team:	WD/PO #:		
	Supplier #:		
	Customer:		
<b>Problem Category:</b>			
<input type="checkbox"/> Reject at Receiving Inspection	<input type="checkbox"/> Documentation		
<input type="checkbox"/> In-Process Reject	<input type="checkbox"/> Field Return		
<input type="checkbox"/> Tooling	<input type="checkbox"/> Customer return		
<b>Problem Description:</b>		<b>Why?</b>	
		2nd Why	
<b>Containment Action</b>		<b>Why?</b>	
Date: / /		3rd Why	
<b>Immediate Corrective Action:</b>		<b>Why?</b>	
Date: / /		4th Why	
<b>Permanent Corrective Action:</b>		<b>Why?</b>	
Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			

Input Containment Action within 24-48 hrs



# Break Out Session: Containment



# Best Practices: Root Cause Investigation

## FORM A TEAM

## Best Practices – Form a Team

Who should be on the team?



- Operator
- Manufacturing Engineer
- Design Engineer
- Quality Engineer
- Inspector
- Buyer
- Customer Service
- Management
- Others as applicable!

## Best Practices – Root Cause

- As a **TEAM**, determine the **Root Cause**
- At Moog, we look for evidence that critical thinking has occurred regarding Root Cause identification
  - Use of tools
    - 5 Why's
    - Cause and Effect (C&E)/Fishbone Diagrams
    - Process Maps
  - Include Objective Evidence
- A good root cause is not common
- **Corrective Action is only as good as the Root Cause identification!**

## Best Practices – Root Cause

**What is a Root Cause?**

## Best Practices – Root Cause

A Root Cause response must consider **2** facets:

### 1) Process Issue or Systemic Issue

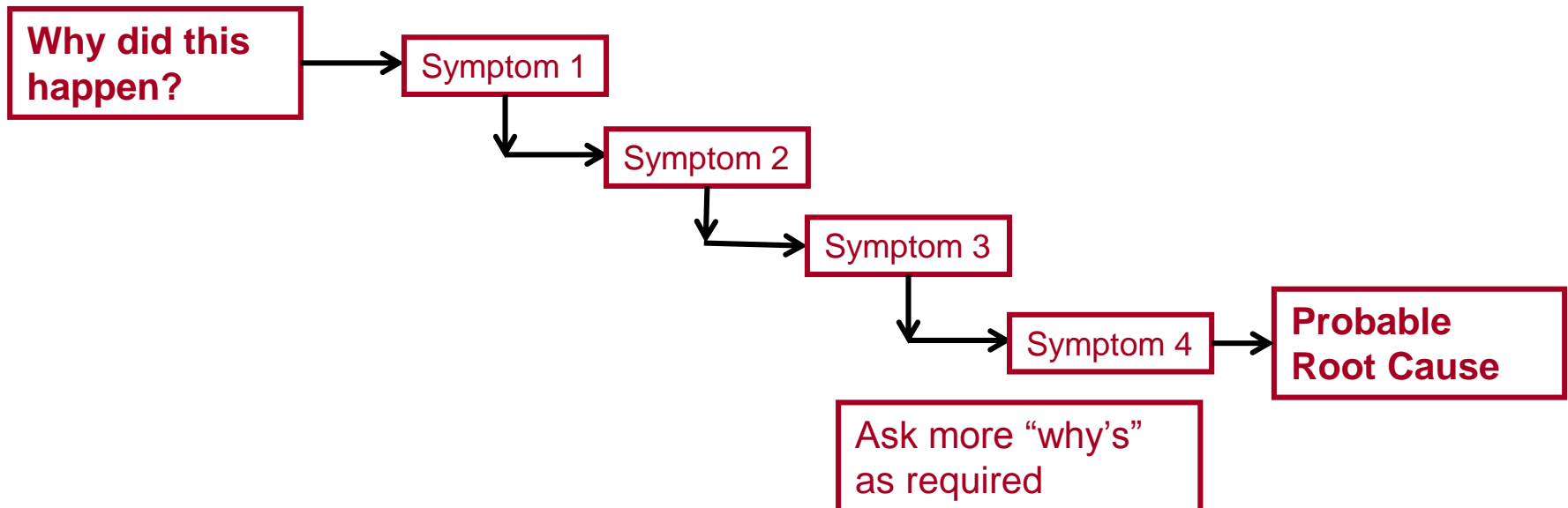
What caused the non-conformance in the first place? What allowed it to happen? What act or failure to act allowed the event to occur?

### 2) Failure to Detect

How did the part leave your building? Why was it certified as conforming?

## Best Practices – 5 Why

- Begin the questioning process with the “most likely” major cause
- Ask “Why does this defect occur or condition exist?”
  - Rule of Thumb is to ask “why” **5 times**
    - Early questions are usually superficial, obvious. As question continues, it becomes progressively more difficult and a more thought-provoking assignment
    - **Stop** when you reach an impacting yet achievable **action**. (don't fix the axis of Earth!).
    - May have to perform this exercise for different scenarios/paths.



## Best Practices – 5 Why Example

- **1. Why is the Jefferson Memorial in Washington D.C. deteriorating?**
  - Because harsh chemicals are frequently used to clean the monument
- **2. Why are harsh chemicals needed?**
  - To clean off the large number of bird droppings on the monument
- **3. Why are there a large number of bird droppings on the monument?**
  - Because the large population of spiders in and around the monument are a food source to the local birds
- **4. Why is there a large population of spiders in and around the monument?**
  - Because vast swarms of insects, on which the spiders feed, are drawn to the monument at dusk
- **5. Why are swarms of insects drawn to the monument at dusk?**
  - Because the lighting of the monument in the evening attracts the local insects



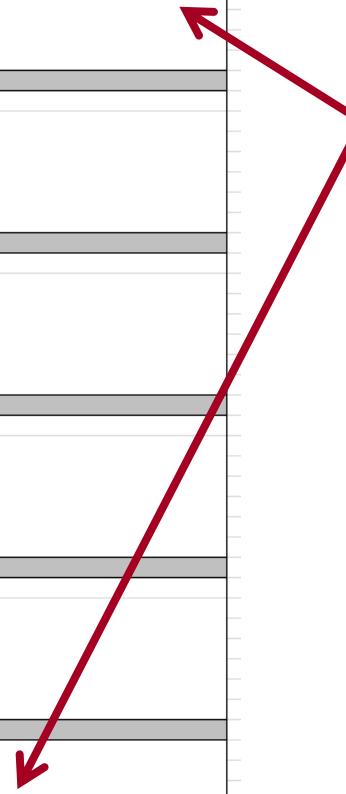
## Best Practices – 5 Why Example

- **1. Why** is there a high reject rate of widgets?
  - Because the plastic is stained.
- **2. Why** is the plastic stained?
  - Because there is excess oil in the cutting machine.
- **3. Why** is there excess oil in the cutting machine?
  - Because it is clogging as it has been months since it was cleaned.
- **4. Why** is it so long since it was cleaned?
  - Because we only service machines when they break down, not on a preventative basis.
- **5. Why** only service after breakdowns?
  - Because maintenance says it is cheaper
    - But what about the cost of rejects and rework?

## Best Practices – Moog CA Worksheet

Moog		5 WHY PROBLEM SOLVING	
<b>Identification:</b>		<b>5 Why Analysis - Why did this happen?</b>	
Date: / /	Area/Location:	<b>Why?</b>	
Originator:	Part #:	1st Why	
Team:	WD/PO #:		
	Supplier #:		
	Customer:		
<b>Problem Category:</b>		<b>Why?</b>	
<input type="checkbox"/> Reject at Receiving Inspection	<input type="checkbox"/> Documentation	2nd Why	
<input type="checkbox"/> In-Process Reject	<input type="checkbox"/> Field Return		
<input type="checkbox"/> Tooling	<input type="checkbox"/> Customer return		
<b>Problem Description:</b>			
<b>Containment Action</b>		<b>Why?</b>	
Date: / /		3rd Why	
<b>Immediate Corrective Action:</b>		<b>Why?</b>	
Date: / /		4th Why	
<b>Permanent Corrective Action:</b>		<b>Why?</b>	
Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			

Ask 5 Why's to arrive at Root Cause



## Best Practices – 5 Why

- Asking **Why** is a way of identifying the underlying root cause of a problem so that this can be tackled, rather than dealing only with superficial symptoms.
- It should be seen as a **simple** and **quick** alternative to Cause and Effect Analysis.
- The **5 Why** strategy is an easy and often effective tool for root cause identification, however for more complex issues C & E Analysis by be required.

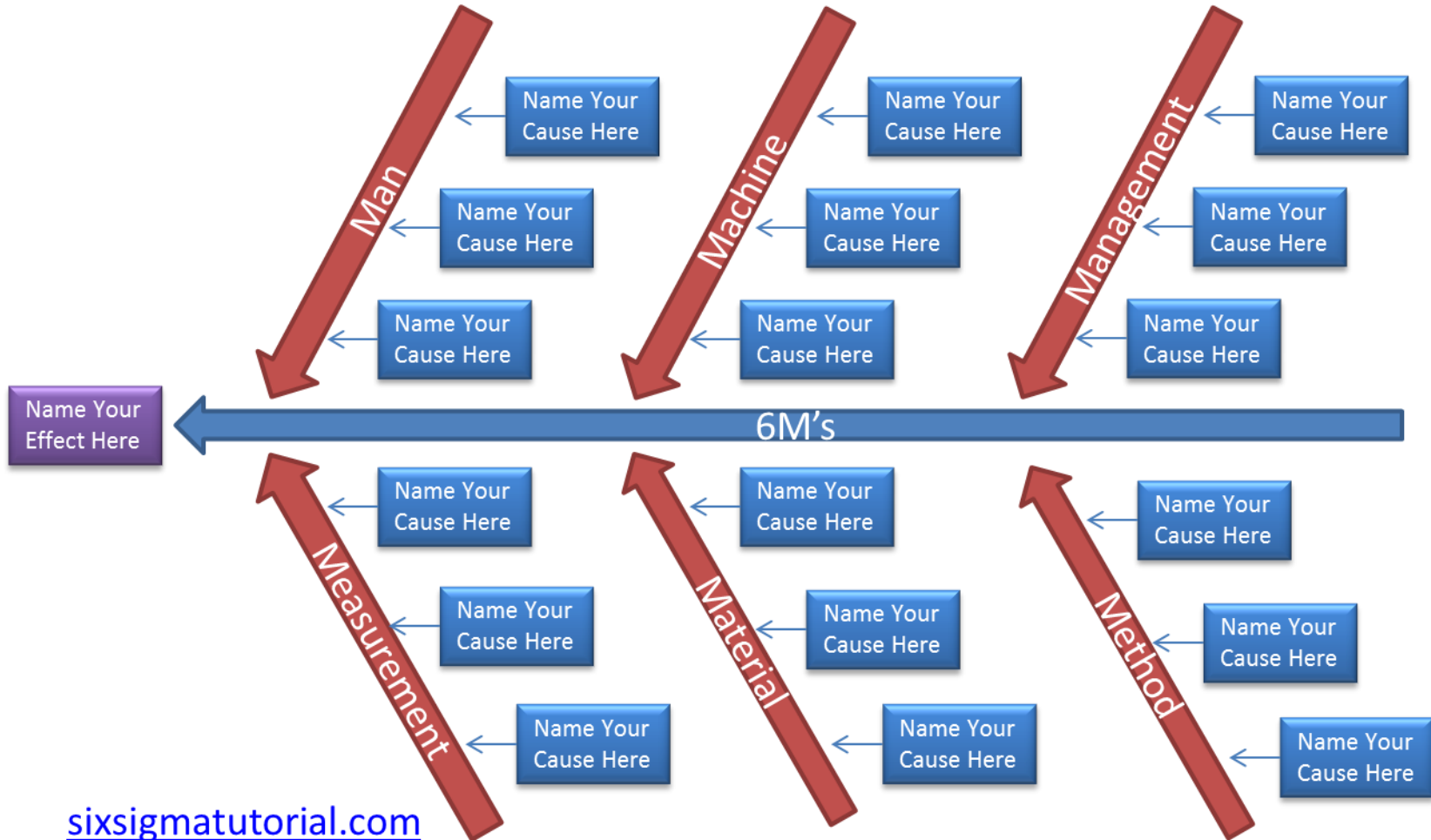
## Best Practices – Cause and Effect (C&E)/Fishbone Analysis

- **What is a Fishbone diagram or Fishbone Analysis?**
  - Fishbone diagram is an analysis tool to provide systematic way of understanding effects and the causes that create those effect. The design of the diagram looks like the skeleton of a fish hence, it is referred to as the fishbone diagram.
- **A fishbone diagram can be used when you:**
  - Want to study **all the possible reasons** why a process is having difficulties, problems, or breakdowns in the initial stages of the process.
  - Need to identify areas for data collection

# Best Practices – Cause and Effect (C&E) /Fishbone Analysis

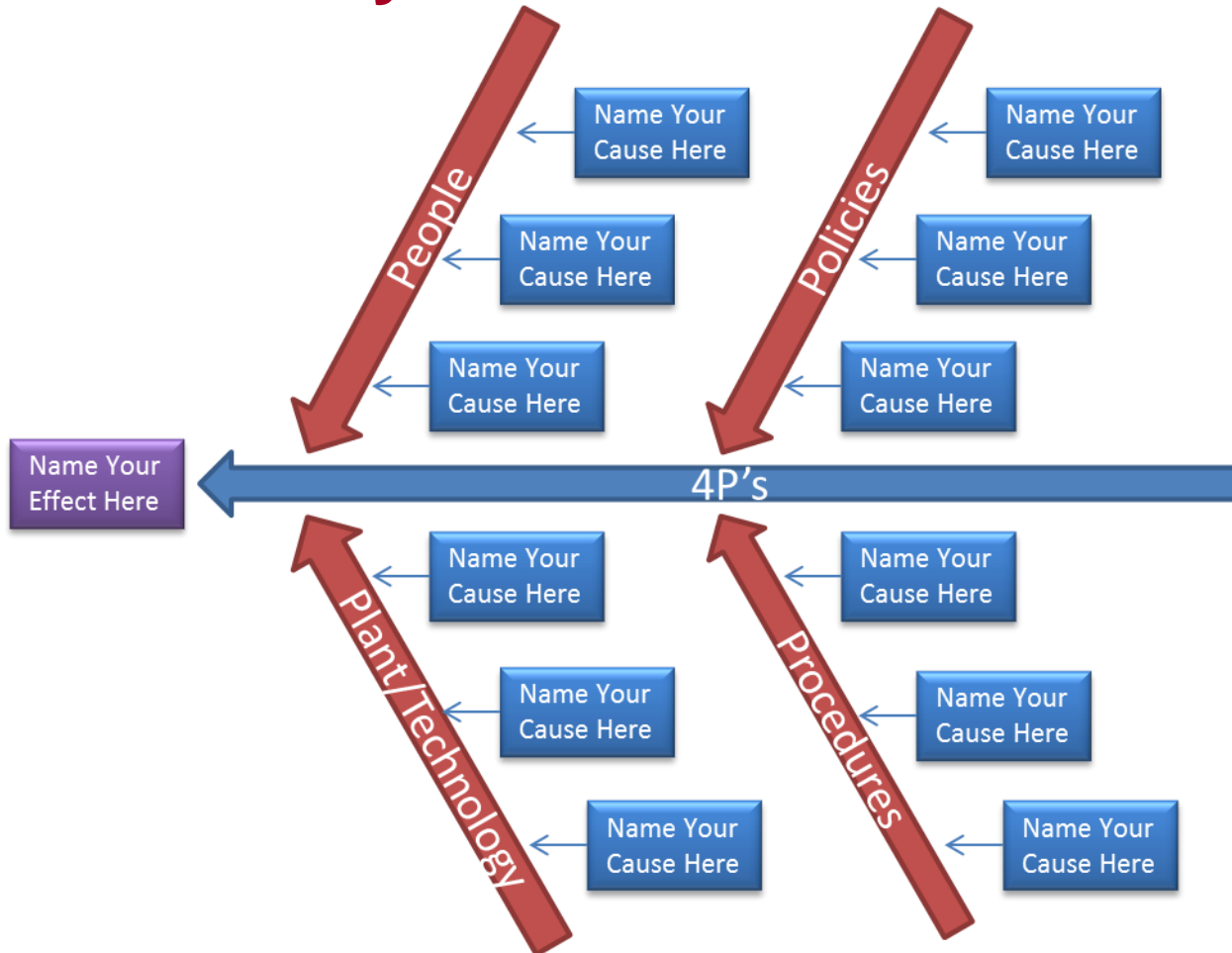
- **Creating a Fishbone Diagram:**
  1. List the problem/issue to be studied in the head of the fish
  2. Label each bone of the fish. The major categories typically used are:
    1. The 6 M's: Methods, Machines, Materials, Manpower, Measurement, Management
    2. The 4 P's: Place, Procedure, People, Policies
  3. Within the categories, brainstorm possible causes for the issue. List them on the fish.
  4. Analyze the results, identify the **most likely causes**.
  5. Evaluate the different most likely causes to identify the Root Cause.

## Best Practices – Cause and Effect (C&E) /Fishbone Analysis

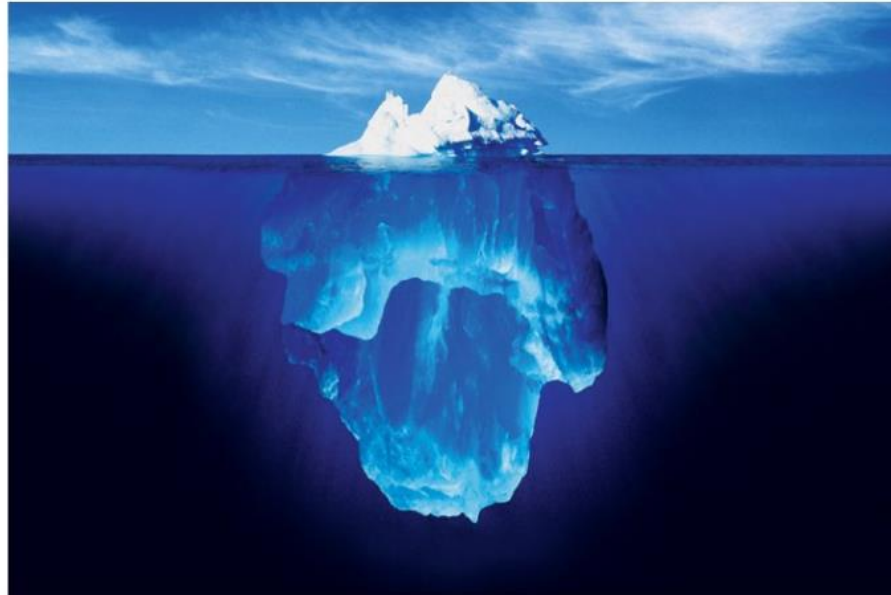


[sixsigmatutorial.com](http://sixsigmatutorial.com)

## Best Practices – Cause and Effect (C&E) /Fishbone Analysis



## Best Practices – Root Cause



Most problems are below the surface. Get to the **ROOT** cause.



## Best Practices – Root Cause

- Verify that you have identified both 1) the Process or Systemic Issue, and 2) the failure to detect.
- Once you have found the Root Cause, you must reassess the immediate Containment activity you took before, to ensure you have **FULL containment**.

# Break Out Session: Root Cause Investigation

# Best Practices: Permanent Corrective Action

## Guidelines – Permanent Corrective Action

What is a “Corrective Action”?

A **Permanent Corrective Action** is immediate and preventative action taken on an assignable cause to **permanently** fix a systemic or process-related issue

## Best Practices – Permanent Corrective Action

Actions must be credible – Objective Evidence (OE) must back up completed actions

Open Actions need Owners and Due Dates

Again, Permanent Corrective Actions will only be as good as the root causes defined!

(notice the repeated use of the word “permanent”)

## Guidelines – Permanent Corrective Action

Examples include:

- Updating Work Instructions/Procedures
- Drawing Changes
- Process Changes
- Tooling
- Visual Guides
- Poke-a-yoke assembly set ups
- Training *to the corrected action*

## Best Practices – Objective Evidence

Objective Evidence:

- Provides **tangible evidence of change**
- Provides **verification documentation**
- Can be audited and reviewed by **anyone** (such as independent **Corrective Action Board**, or **CAB**)
- Provides closure and feedback to **Moog** (and sometimes **Moog's Customer**)
- Stands on its own (what happens if you and your team win the **lottery** and leave work tomorrow? Will the Corrective Action still be in place?)
- **Always** provide Objective Evidence. We will ask for it!



# Let's look at some Examples



## Best Practices – Permanent Corrective Action

- **Root Cause:** “Inspection error (we missed it!).”
- **C/A:** “We will apply 100% Inspection from now on.”

Is this a good response?

## Best Practices – Permanent Corrective Action

- **Root Cause:** “Inspection error (we missed it!).”
- **C/A:** “We will apply 100% Inspection from now on.”
- **Quality cannot be inspected into a product.**
- **Did inspection create the feature?**
  - If not, it can NOT be the Root Cause!
- 100% Inspection by a person is *never* 100% effective.
- How does **Inspection** prevent the **problem** from happening again?
- “Why was it missed and what will prevent them from missing it again?” are just two of the questions to ask.
- Was the right tool (calibrated & capable) used to Inspect at the point of manufacture? (Gauge Reproducibility & Reliability, R&R)
- Inspection can be used as a **validation** of the Corrective Action.
- **Quality cannot be inspected into a product!**

## Best Practices – Permanent Corrective Action

- **Root Cause:** “Wrong gage, tool, machine or material used.”
- **C/A:** “We will use right gage, tool, machine or material from now on.”

Is this a good response?

## Best Practices – Permanent Corrective Action

- **Root Cause:** “Wrong gage, tool, machine or material used.”
- **C/A:** “We will use right gage, tool, machine or material from now on.”

**Remember to keep asking **Why!****

- Are the correct, gages/tools/machine/material called out and correctly identified on the instruction/router/job order?
- What will prevent this from happening again?
- Where is the **objective evidence**?

## Best Practices – Permanent Corrective Action

- **Root Cause:** “Operator screwed up. People make mistakes.”
- **C/A:** “We have reprimanded and trained the operator.”

Is this a good response?

## Best Practices – Permanent Corrective Action

- **Root Cause:** “Operator screwed up. People make mistakes.”
- **C/A:** “We have reprimanded and trained the operator.”

**Focus on the **Process** placed in the hands of the operator**

- What allowed the operator to do this?
- Has the process been mistake-proofed?
  - Is the program correct?
  - Are Work Instructions correct and do they have the right amount of detail?
- Never blame the operator
- Remember the Objective Evidence!

## Best Practices – Permanent Corrective Action

- **Root Cause:** “It wasn’t us, it was the Moog-directed sub-tier.”
- **C/A:** “We have requested an alternate sub-tier from Moog.”

Is this a good response?

## Best Practices – Permanent Corrective Action

- **Root Cause:** “It wasn’t us, it was the Moog-directed sub-tier.”
- **C/A:** “We have requested an alternate sub-tier from Moog.”
- Who is getting the Purchase Order contract from Moog?
- Who is signing the CofC?

**YOU ARE**

- You are responsible for the product you ship to Moog. If action must be taken against the sub-tier, you must take it.
- You may use sub-tier response in addition your own CA response. But be mindful of the due dates!



## Best Practices – Moog CA Worksheet

Moog		5 WHY PROBLEM SOLVING	
<b>Identification:</b>		<b>5 Why Analysis - Why did this happen?</b>	
Date: / /	Area/Location:	<b>Why?</b>	
Originator:	Part #:	1st Why	
Team:	WD/PO #:		
	Supplier #:		
	Customer:		
<b>Problem Category:</b>			
<input type="checkbox"/> Reject at Receiving Inspection	<input type="checkbox"/> Documentation		
<input type="checkbox"/> In-Process Reject	<input type="checkbox"/> Field Return		
<input type="checkbox"/> Tooling	<input type="checkbox"/> Customer return		
<b>Problem Description:</b>		<b>Why?</b>	
		2nd Why	
<b>Containment Action</b>		<b>Why?</b>	
Date: / /		3rd Why	
<b>Immediate Corrective Action:</b>		<b>Why?</b>	
Date: / /		4th Why	
<b>Permanent Corrective Action:</b>		<b>Why?</b>	
Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			

Input Permanent Corrective Action



# Break Out Session: Permanent Corrective Action

# Best Practices: Verification / Follow Up

## Best Practices – Verification / Follow Up

- **What is Verification?**

- You, someone on the Corrective Action **Team**, or an assigned Verifier, **verifies** – or Checks – that the Corrective Action was implemented **successfully and completely**, and that it is **working**

- The Verification **Plan** is submitted with the Permanent Corrective Action

- Actual Verification can be performed at a later date, usually about a month

## Best Practices – Verification / Follow Up

- What do you look for in Verification?

Two-step process:

- 1) Verify Corrective Action was implemented **successfully** and **completely**
- 2) Verify Corrective Action is **working**

## Best Practices – Verification / Follow Up

- What do you look for in Verification?

Verify that the applicable procedures were updated

Verify that open Actions were completed by Due Dates assigned

Verify that Corrective Action has eliminated non-conformity

Ask yourself:  
Could I reproduce the defect?

Step 1) Verification of Implementation

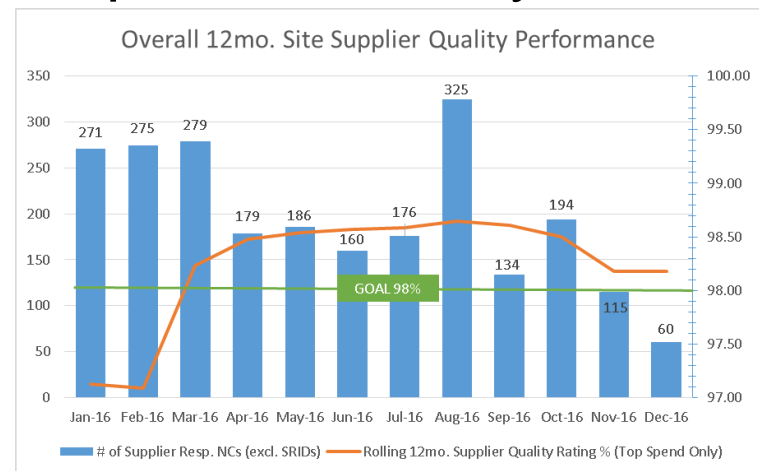
Step 2) Validation of Effectiveness

## Best Practices – Verification / Follow Up

- **Document** the Corrective Action results and provide informational **feedback**.
- Let everyone know what has been done, include it in the job folder, post it, make it visual!

## Best Practices – Verification / Follow Up

- Post measurement charts where everyone can see them
- Let everyone know what the charts mean to your business
  - By the way, this is a new AS9100 Rev D requirement, titled **Awareness**
- Keep the charts as simple as possible; always note which direction is good!





## Best Practices – Verification / Follow Up

- Train all affected employees to the Corrected Action
- Document the training on a training record
- Provide this record as **Objective Evidence** in your Corrective Action response

## Best Practices – Moog CA Worksheet

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<b>Containment Action</b>		<b>Why?</b>	
Date: / /		3rd Why	
<b>Immediate Corrective Action:</b>		<b>Why?</b>	
Date: / /		4th Why	
<b>Permanent Corrective Action:</b>		<b>Why?</b>	
Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			

Verification Plan should be included in Permanent Corrective Action Response



**Congratulations!**

## Congratulations!

You have completed and submitted a Root Cause and Corrective Action. You have also submitted Objective Evidence, and either Verified it is working, or provided a Verification plan.

Are you done?

**NO**

# Best Practices: Preventative Action

## Best Practices – Preventative Action

- Look across ALL Moog parts – could they see the same failure?
- Look across ALL your customers – having a “special process” for only one customer such as Moog is a **FAILURE MODE**.
- Why wait for a Corrective Action request? Take steps **now** to minimize risks later!



## Best Practices – Preventative Action

- If you produce a shaft of multiple lengths for Moog and you completed a Corrective Action on one part number, does it apply to all the other parts also?
  - Could they see the same failure mode?
  - Do those routers/processes need to be reviewed and updated as well?
- If you perform an Outside Process for Moog and you resolve a processing issue for one part number, does it apply to all other parts?
  - Be wary of any “special processes”!



## Best Practices – Moog CA Worksheet

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Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			

Input Preventative Corrective Action here



# Break Out Session: Preventative Action

# Best Practices: Using TipQA

## Best Practices – Using TipQA

CA Number SU00062766 Revision A Status WCA Assigned To s10821c Business Unit MNA

View Edit Subscribe Create CA Reassign Reverse Process

Identification Discrepancy Cause / Corrective Action Approvals Follow Up Closure Action History List Query

Pop Up Details Associated Part Numbers Revision Audit Trail Revision Revised By system Date Revised 12/09/2016 06:03:11AM

Summary  
A350 (scratches and paint overspray) Loc : X93R MRB Shop St

Associated Documents

Module	Document	Is Master
NC	RI00228591	<input checked="" type="checkbox"/>
NC	RI00228589	<input checked="" type="checkbox"/>
NC	RI00228641	<input type="checkbox"/>

Part Number :911-177B  
Part Revision : CB7731003 ASSY FLAP LEVER BR  
Commodity Code 31251507  
End Use Part / Model Number :911-177B  
Customer  
Part Criticality  
Quantity Lot 1  
Inspected 1  
Nonconforming 1

Supplier 10821 PD / Line Number 1608139301 1 Buyer 0029588  
Site Code  
Work Order Work Supervisor  
Supplier Part Number  
Manufacturer Part Number  
Drawing Number Date Response Due 12/23/2016  
Drawing Revision Date Response Received / /  
Location Code AG-WAF  
Factory Code X93R Date Extended / /  
Work Center Customer Due Date / /  
FRACAS Identifier Date Submitted to Customer / /  
Product Group Date Created 11/23/2016  
Product Sub Group Date Identified / /  
Generated From Module Code NC  
Sequence Number 228591  
Contract  
Project  
Priority Rating  
Planner 0000292  
Program Code  
End Item Serial Number  
Customer CA Number  
Sales Order Number  
Sales Order Line Item

Repeat Discrepancy

Use tabs to navigate. Left-to-right workflow

MNA - North America

## Best Practices – Using TipQA; Administrative Data

The screenshot displays the TipQA software interface with several key administrative data fields highlighted by red callouts:

- CA# & Status:** CA Number SU00062766, Revision A, Status WCA
- Brief Summary:** A350 (scratches and paint overspray) Loc : X93R MRB Jon St
- Part Number:** Part Number -911-177B
- Associated NCs:** A table listing associated Non-Conforming (NC) items with columns for Module, Document, and Is Master.
- Qty:** Lot 1, Inspected 1, Nonconforming 1
- PO # & Line:** PO / Line Number 1608139301 ... 1 ...
- Date Response Due:** Date Response Due 12/23/2016
- Date Created:** Date Created 11/23/2016

Module	Document	Is Master
NC	RI00228591	<input checked="" type="checkbox"/>
NC	RI00228589	<input checked="" type="checkbox"/>
NC	RI00228641	<input type="checkbox"/>

## Best Practices – Using TipQA; Description of the Finding

CA Number SU00062766 Revision A Status W/CA Assigned To s10821c Business Unit MNA

View Edit Subscribe Create CA Reassign Reverse Process

Identification Discrepancy Cause / Corrective Action Approvals Follow Up Closure Action History List Query

PopUp\_Details Standard\_Text

Revision Audit Trail Revision 2 Revised by 6024284 Date Revised 11/23/2016 12:35:28PM

Discrepancy Description  
The mounting holes appear to be scratched up from trying to remove the paint leakage. There is also a few fragments of masking left on a couple holes. Pictures are attached.

Letter A Highest Letter A

Category Code  
Process Code IQI7 Supply Chain

Defect Location

Detailed description of the finding

Each non-conformance for which the CA was issued will have its own "Letter" (e.g. NC line). Click to advance. Pay attention to – and respond to – each one!

Attachments, such as photos or test reports, will be attached here. If not, check the NC.

## Best Practices – Using TipQA; RCCA

Must be in Edit mode

Click blue triangle

Use the 5-Why spreadsheet to complete your Root Cause investigation

Re-state your Root Cause here

Remember to select the Cause Code and CA Code from the drop-down

Input Permanent Corrective Action here. Be as detailed as possible. Should include Verification plan!

Attach Objective Evidence

Input Preventative Action here. Be as detailed as possible.

Again, watch the letters

## Best Practices – Moog CA Worksheet

Moog		5 WHY PROBLEM SOLVING	
<b>Identification:</b>		<b>5 Why Analysis - Why did this happen?</b>	
Date: / /	Area/Location:	<b>Why?</b>	
Originator:	Part #:	1st Why	
Team:	WD/PO #:		
	Supplier #:		
	Customer:		
<b>Problem Category:</b>			
<input type="checkbox"/> Reject at Receiving Inspection	<input type="checkbox"/> Documentation		
<input type="checkbox"/> In-Process Reject	<input type="checkbox"/> Field Return		
<input type="checkbox"/> Tooling	<input type="checkbox"/> Customer return		
<b>Problem Description:</b>		<b>Why?</b>	
		2nd Why	
<b>Containment Action</b>		<b>Why?</b>	
Date: / /		3rd Why	
<b>Immediate Corrective Action:</b>		<b>Why?</b>	
Date: / /		4th Why	
<b>Permanent Corrective Action:</b>		<b>Why?</b>	
Date: / /		5th Why	
<b>Preventative Corrective Action:</b>		<b>Root Cause:</b>	
Date: / /			



## Best Practices – Using TipQA; RCCA

CA Number SU00062766 Revision A Status WCA Assigned To s10821c Business Unit MNA

View Edit Subscribe Create CA Reassign Revoke Process

Identification Discrepancy Cause / Corrective Action Approvals Follow Up Closure Action History List Query

Pop Up Details Task Planning Root Cause Text

Revision Audit Trail Revision 2 Revised By 0024284 Date Revised 11/22/2016 12:35:53PM

General  
Letter A Highest Letter A  
Cause Code  
Corrective Action Code  
Action Due // / /  
Cost of Quality  
Detail Task Planning Required for this line item

Classification  
Preventative Action  Corrective Action

Cause Description  
**Issuance of this report requires a formal Cause and Corrective Action response to Moog.** Failure to respond to removal from Moog's Approved Supplier List. Your response should be made to the Moog Commodity Supplier Qual made available on the Moog Supplier portal for your reference.

It is essential that you take positive and immediate corrective Action to prevent subsequent rejections at Moog.

Attached below for your use, is a 5-Why problem solving template that should be used to facilitate your C/CA activities. Please export the report, complete all sections, save it, and reattach it into your C/CA response to Moog.

Corrective Action Description

Preventative Action Description

When complete, you must select "Process". Status will change from WCA to WA

To confirm that you have submitted, observe the "Action History" tab

Once a Moog representative reviews and approves, it goes to the Corrective Action Board for review. After CAB acceptance, CA will enter Follow-Up (Verification) stage. You may observe the Approvals in the "Approvals" tab

If at any time the CA is not approved, it will be sent back to you for updates

# Summary

## Guidelines – Corrective Action Initiation Criteria

What is a “Corrective Action”?

A **Corrective Action** is immediate and preventative action taken on an assignable cause to permanently fix a systemic or process-related issue

## Summary

What is a “Corrective Action”?

Ongoing Improvement initiatives

*Documents* the work done!

NOT a reprimand

## Summary

- **FORM A TEAM**
- **CONTAIN** the non-conformance – Supply Chain, Detection, and Awareness
- A Corrective Action is only as good as it's **Root Cause**
- Get to the REAL **Root Cause** – there can be more than one!
- IMPLEMENT solid **Corrective Actions**
- SHOW **Objective Evidence**
- **VERIFY** the Corrective Action
  - Ask yourself if the defect could be reproduced by others
- **PREVENT** the defect from occurring on other parts

## Best Practices – Containment

- **There are two parts to Containment:**

**Contain Defective Material**

**Identify Inspection Gap**

## Best Practices – Root Cause

A Root Cause response must consider **2** facets:

### 1) Process Issue or Systemic Issue

What caused the non-conformance in the first place? What allowed it to happen? What act or failure to act allowed the event to occur?

### 2) Failure to Detect

How did the part leave your building? Why was it certified as conforming?

## Summary

Actions need Owners and Due Dates

Actions must be credible – Objective Evidence (OE) must back up completed actions

Again, Permanent Corrective Actions will only be as good as the root causes defined!

(notice the repeated use of the word “permanent”)



## Best Practices – Verification / Follow Up

- What do you look for in Verification?

Verify that the applicable procedures were updated

Verify that open Actions were completed by Due Dates assigned

Verify that Corrective Action has eliminated non-conformity

Ask yourself:  
Could I reproduce the defect?

Step 1) Verification of Implementation

Step 2) Validation of Effectiveness

## Best Practices – Preventative Action

- Look across ALL Moog parts – could they see the same failure?
- Look across ALL your customers – having a “special process” for only one customer such as Moog is a **FAILURE MODE**.
- Why wait for a Corrective Action request? Take steps **now** to minimize risks later!



## Summary

**A Corrective Action is an  
Opportunity for *Improvement***

**What is our #1 priority?**



**Congratulations!**

**Congratulations!**

You have completed a Continuous Improvement activity for the benefit of your organization, Moog, and all your other customers.

**Do you keep going?**

# YES



# Thank you!

**End.**

# Supplemental Material

## Summary

**Continuous Improvement is  
continuous**

# Best Practices – Cause and Effect (C&E)/Fishbone Analysis

- **Creating a Fishbone Diagram:**
  - List the problem/issue to be studied in the head of the fish
  - Label each bone of the fish. The major categories typically used are:
    - The 6 M's: Methods, Machines, Materials, Manpower, Measurement, Management
    - The 4 P's: Place, Procedure, People, Policies
  - Repeat this procedure with each factor under the category to produce sub-factors. Continue asking, “Why is this happening?” and put additional segments each factor and subsequently under each sub-factor.
  - Analyze the results, identify the ‘most likely causes’.
  - Evaluate the most likely causes to identify the true Root Cause.

## Best Practices – Permanent Corrective Action

“We talked to the operator, he/she will be trained”

Is training alone sufficient?

Never blame  
the operator

Focus on the  
process placed  
in the hands of  
the operator

What allowed  
the operator to  
do this?

Can the  
process be  
mistake-  
proofed?

# Best Practices: Good CA Example

## Best Practices – Good CA Example

# Revision Control



## Revision Control

Revision	Reason for Change	Release Date	Author
NC	Initial Release	2/1/2017	D. Hensel
A	Added Permanent CA examples, moved Containment ahead of Form a Team, added Containment slide		D. Hensel