Lean Six Sigma Green Belt Training Developed for Starbucks Corporation

Written by Joan Ambrose, Tracy Camp, Bethany Quillinan and Will Mokszycki Lean Six Sigma Master Black Belts ETI Group

Presented by



Oregon: 503-484-5979 Washington: 360-681-2188 www.etigroupusa.com

Copyright © 2024 ETI Group all rights reserved

Lean Six Sigma Green Belt Course Table of Contents

Course Outline with Slide Numbers

Lean Six Sigma Overview Slide	
1.	Lean Overview
2.	Six Sigma Overview
3.	Why Combine Lean and Six Sigma?
De	fine Phase
4.	Project Scope and SIPOC
Me	easure Phase
5.	Observing the Current State
6.	Basic Process Mapping
7.	Value Stream Mapping
8.	X and Y Variables
9.	Data Collection and Sample Size Calculation
10.	Basic Statistics and Normal Distribution
11.	Measurement Variation
12.	Measurement System Analysis
13.	Categorical MSA 210
14.	Process Capability Indices
An	alyze Phase
15.	Root Cause Analysis
Im	prove Phase
16.	Developing and Prioritizing Solutions
17.	Lean Solutions
18.	Theory of Constraints
19.	Reviewing the Proposed Future State
Co	ntrol Phase
20.	Control Plan
21.	Statistical Monitoring
22.	Response Plan

1 Lean Overview

The goal	• Provide the greatest value for customers using the fewest resources
The methods	 Principles and practices based on the Toyota Production System (TPS)
The barrier	• Culture always defeats methodology
The path forward*	 Create a culture of continuous improvement (<i>kaizen</i>) Integrate improvement cycles into the daily work of all employees Improve all processes, every day

*See **Toyota Kata** (2010) by Mike Rother.

- *Value* is defined from the customer's point of view
 - \rightarrow Reduce or eliminate activities that do not add customer value

- *Value stream* all activities required to provide a specified family of products or services to the customer
 - \rightarrow Organize workflows by value stream, not by department

Customer defines value

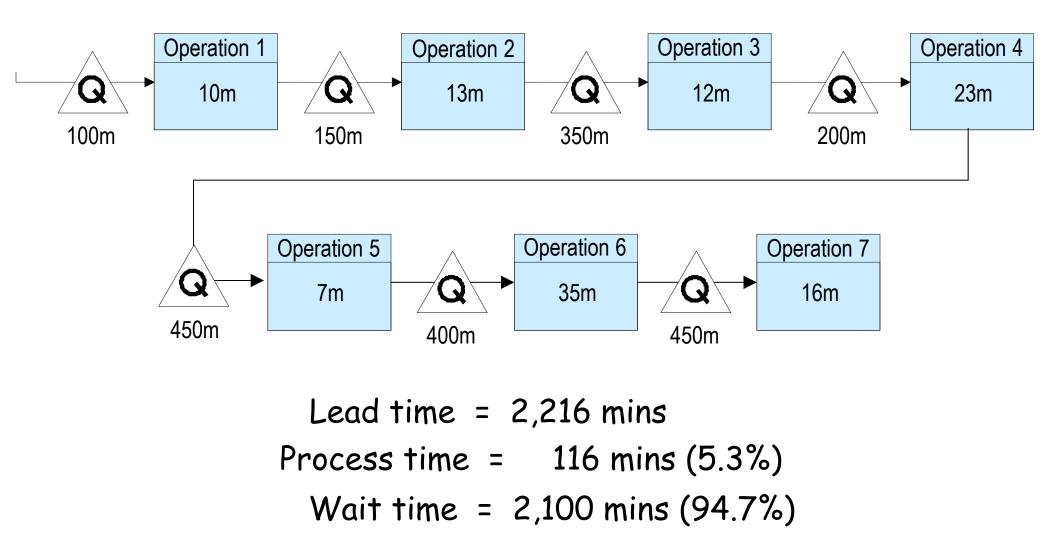
Customer value adding (CVA)

- Activities that are required, from the customer's point of view, to provide the desired products and services
- What the customer is willing to pay for
- Changes the form or function of the product
- Goal: Optimize and standardize these activities

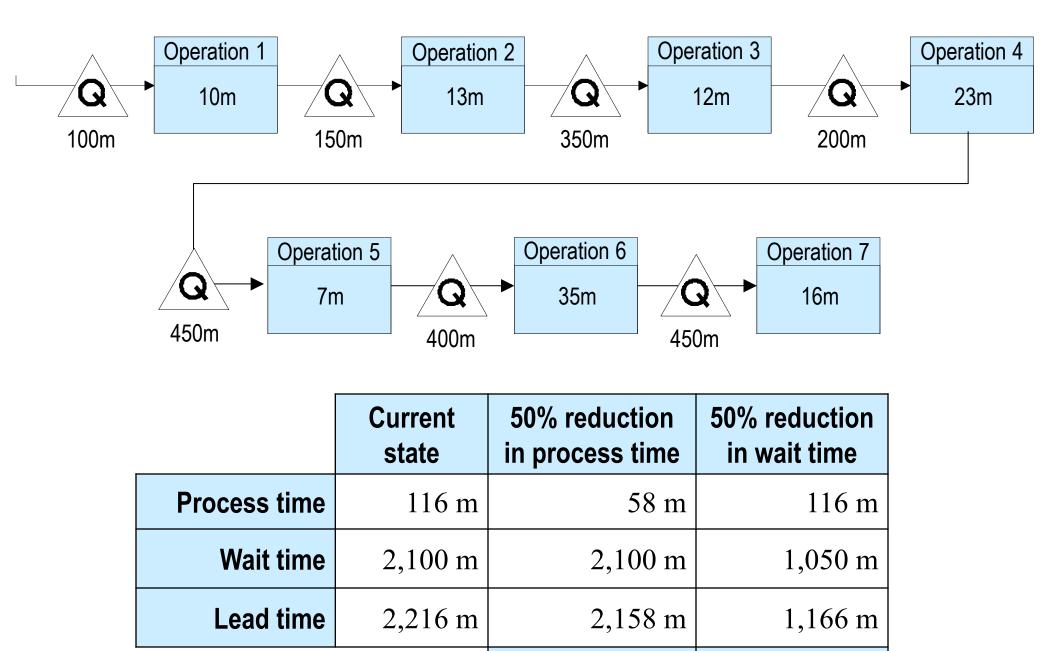
Non-value adding (NVA)

- There exists a feasible future state in which the desired products and services can be provided without these activities
- Goal: Eliminate or reduce
- Non-value adding but necessary
 - Activities that are not CVA, but cannot feasibly be eliminated under current constraints
 - Examples include audits, reporting, regulatory compliance, etc.
 - Goal: Question and reduce

Typical current state value stream



Queue (material or transactions waiting to be worked on) ightarrow 100% NVA

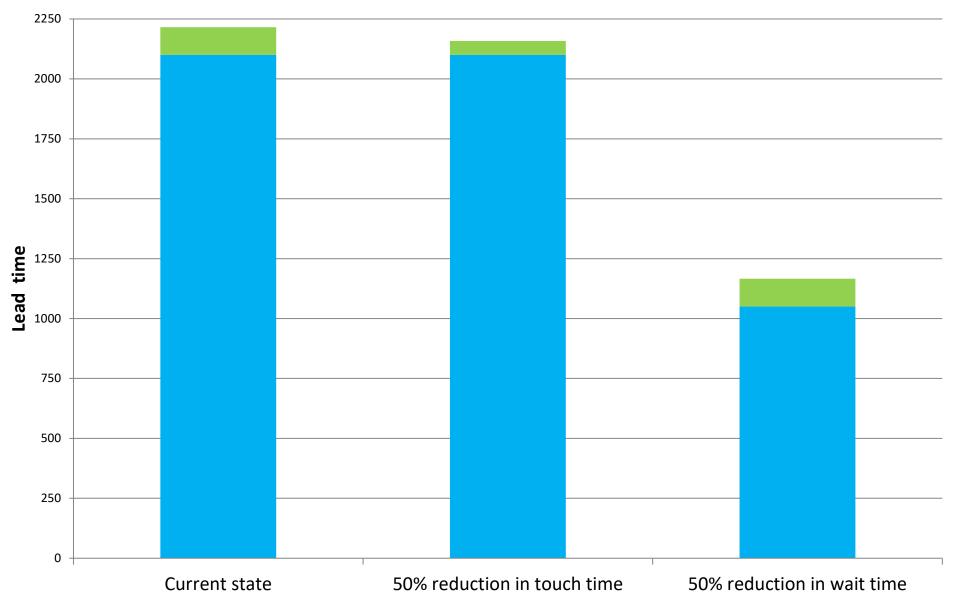


2.6%

47.4%

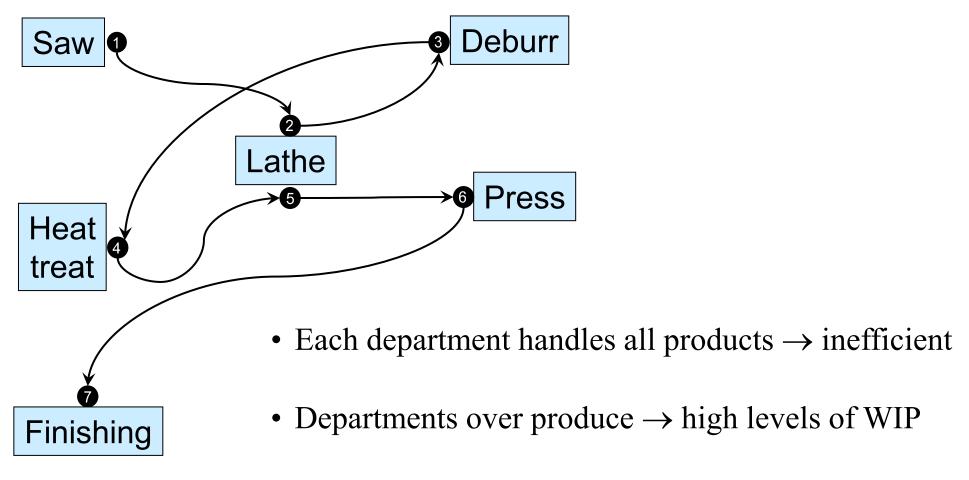
Reduction in lead time \rightarrow



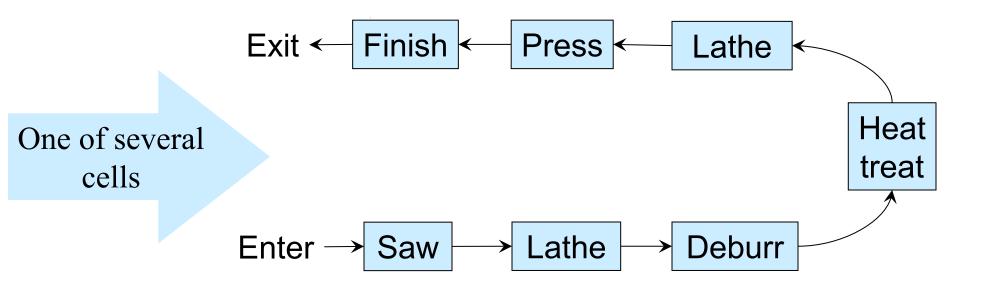


D	Defects: Failure to meet expected standards of quality or delivery	
0	Over production: Making or doing more than is needed at the time	
W	Waiting: People waiting to work, or things waiting to be worked on	
Ν	<i>Not utilizing creativity</i> : Failure to integrate improvement cycles into the daily work of all employees	
Т	<i>Transportation</i> : People or things being moved from one place to another	
I	Inventory: Supplies, WIP, or finished goods beyond what is needed	
Μ	<i>Motion</i> : Excessive motion in the completion of work activities	
Е	<i>Extra processing</i> : Producing or delivering to a higher standard than is required	

Example of organizing work by department



- WIP is valued as an asset in reality it's a cash sink
- WIP moves between departments in large batches → long lead times, long lags before defects are discovered
- Poor layout \rightarrow excessive transport

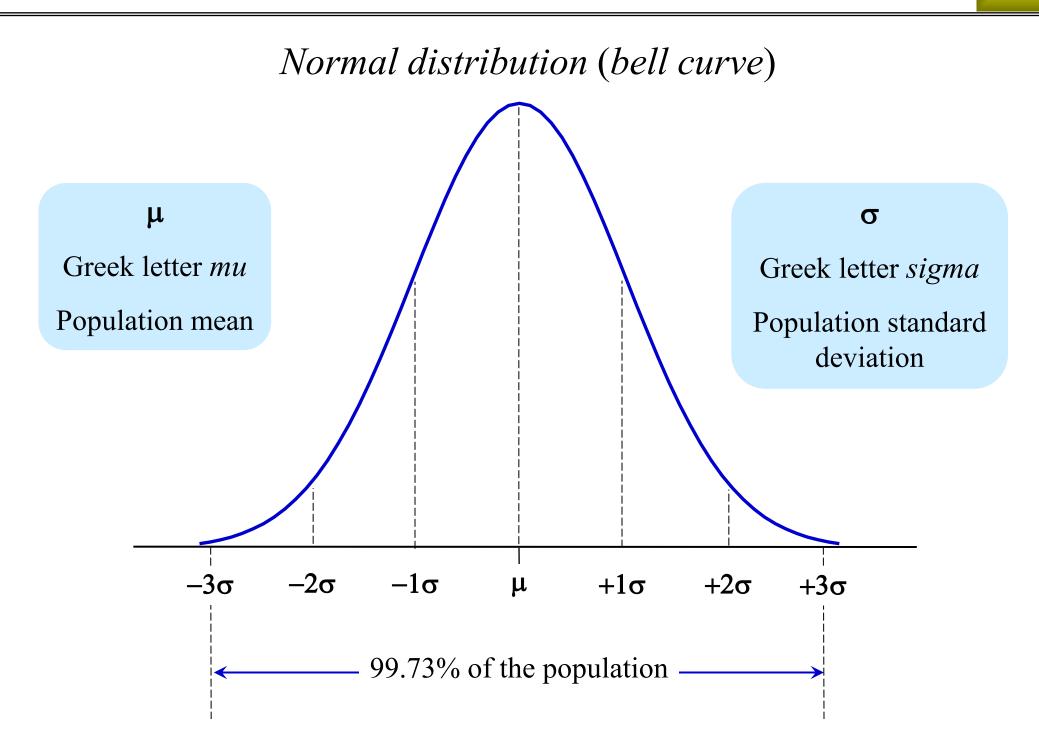


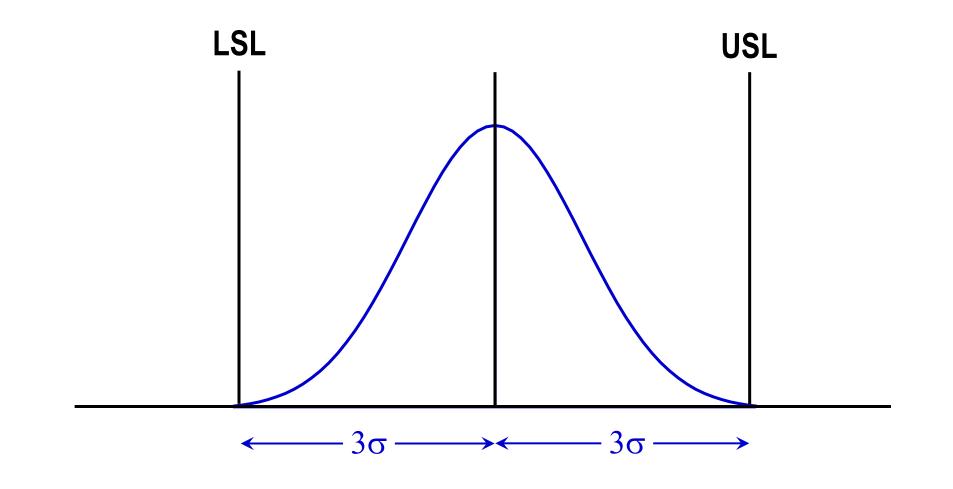
- Each cell handles particular, similar products \rightarrow efficient
- Cells produce only to current customer demand → low levels of WIP, less cash tied up
- WIP moves through each cell in small batches \rightarrow short lead times
- Proximity of operations → minimal transport, defects identified immediately

• Process spread

• Pursuit of perfect quality

• Pragmatic business initiative



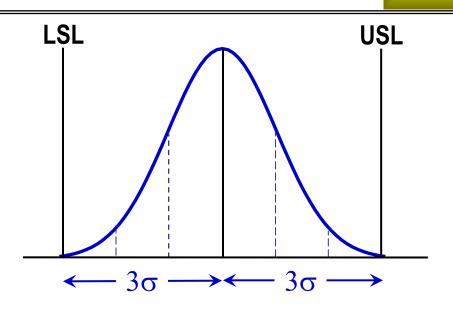


0.27% defective (first pass)

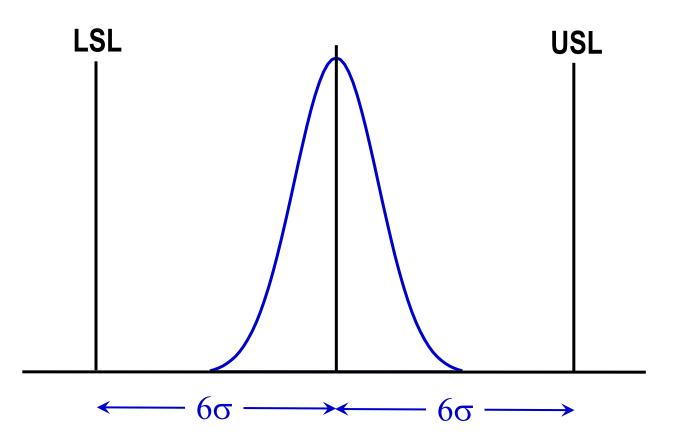
USL stands for *Upper Specification Limit*, LSL stands for *Lower Specification Limit*. Specification limits represent the Voice of the Customer with regard to measureable characteristics of products or services.

For the Normal distribution shown above, the mean (μ) is equal to the midpoint of the specification range, and the process spread (6 σ) is exactly equal to the width of the specification range (USL minus LSL). This means that 99.73% of product or service outcomes produced by this process satisfy the spec limits. Equivalently, 0.27% of outcomes lead to scrap, rework, do-overs, or other costly measures to prevent or respond to customer dissatisfaction.

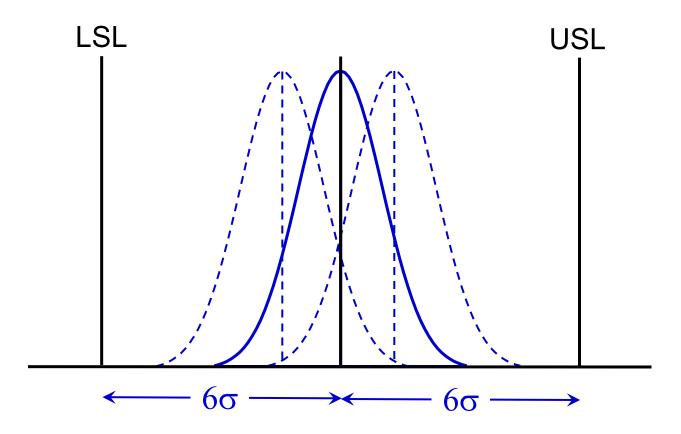
In the 1980s, Motorola questioned the adequacy of 0.27% defective as an improvement objective



2,700 defective parts per million 2,000 pieces of mail lost each hour 20,000 wrong prescriptions per year 15,000 newborn babies dropped per year No electricity or water 8.6 hours per month 500 incorrect surgical procedures each week Motorola proposed a more aggressive objective



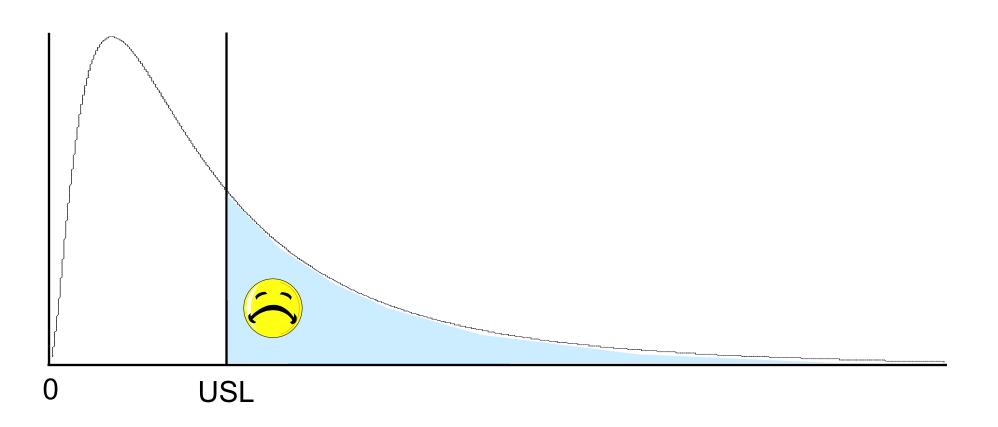
2 defective parts per billion



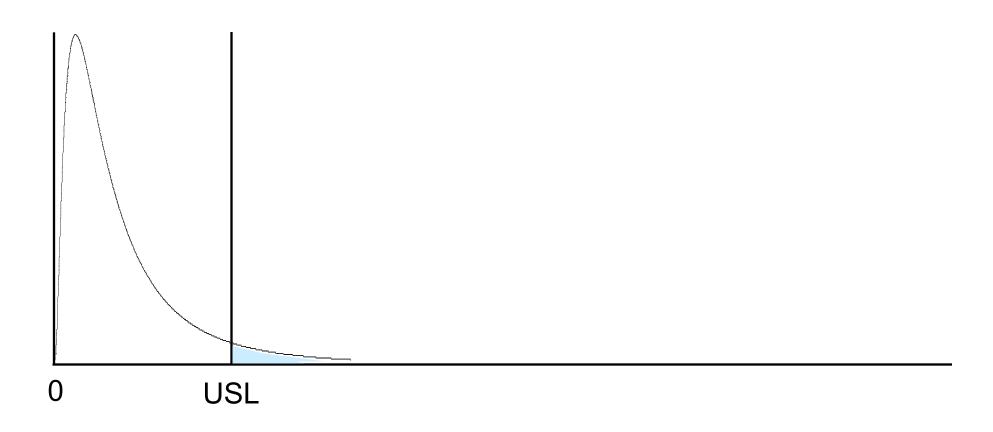
At most 3.4 defective parts per million (DPPM)

- Motorola backed away from 2 defective parts per billion as the stretch goal
- They allowed that the process mean might wander as much as 1.5σ away from the spec midpoint
- At these extremes, the process would produce 3.4 defective parts per million (DPPM)
- The $\pm 1.5\sigma$ offset was somewhat arbitrary, but 3.4 DPPM became the definition of "Six Sigma quality"

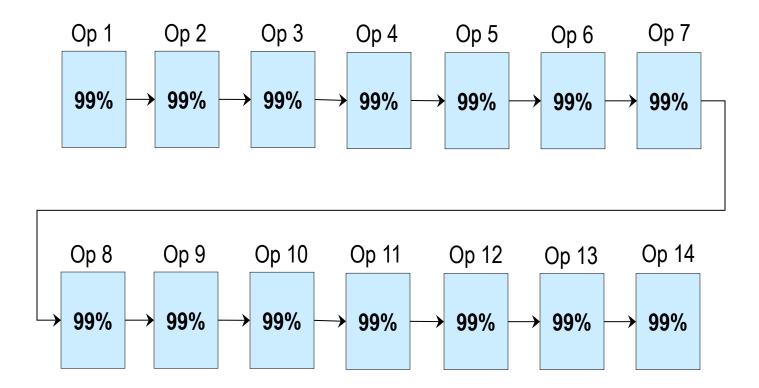
Before improvement project



After improvement project



Suppose we have 10,000 DPPM (99% yield) for each operation



Area manager: "Our overall yield is 99%"

Is this true?

Rolled Throughput Yield^{*} = Probability of no defect in 14 operations

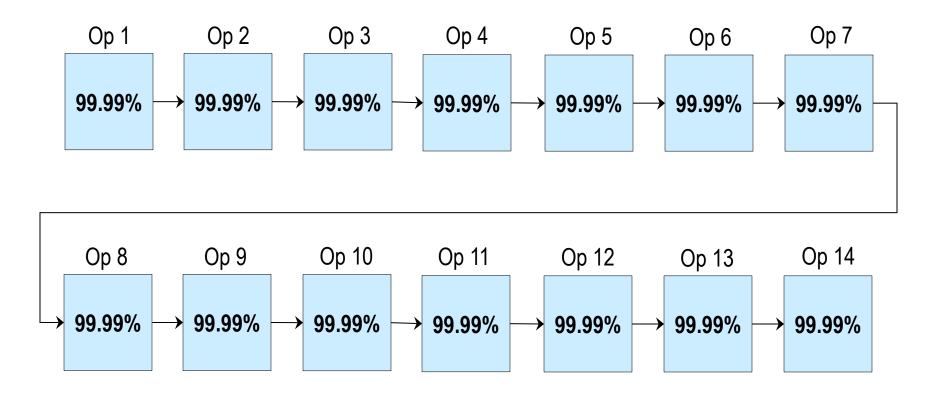
 $= 0.99 \times 0.99 \times \cdots \times 0.99$ (14 times)

 $= (0.99)^{14}$

 $= 0.868746 \rightarrow 86.9\%$

*Also known as overall yield, cumulative yield, and end-to-end yield

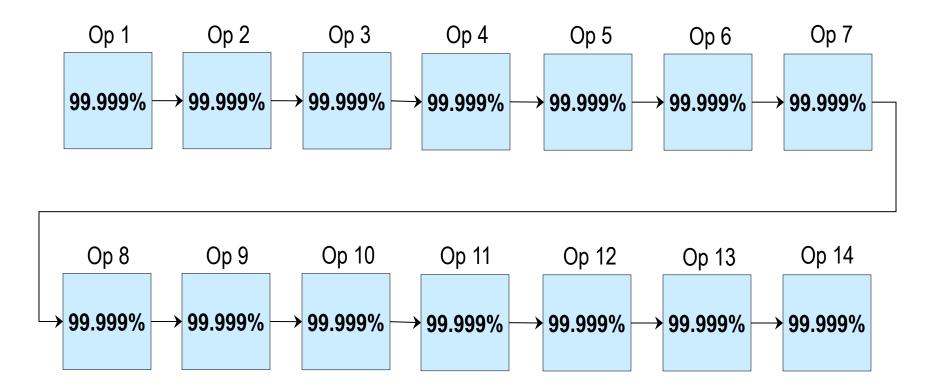
100 DPPM (99.99% yield) in each operation



Overall yield = $(0.9999)^{14} = 0.998601 \rightarrow 99.86\%$

1399 DPPM

10 DPPM (99.999% yield) in each operation

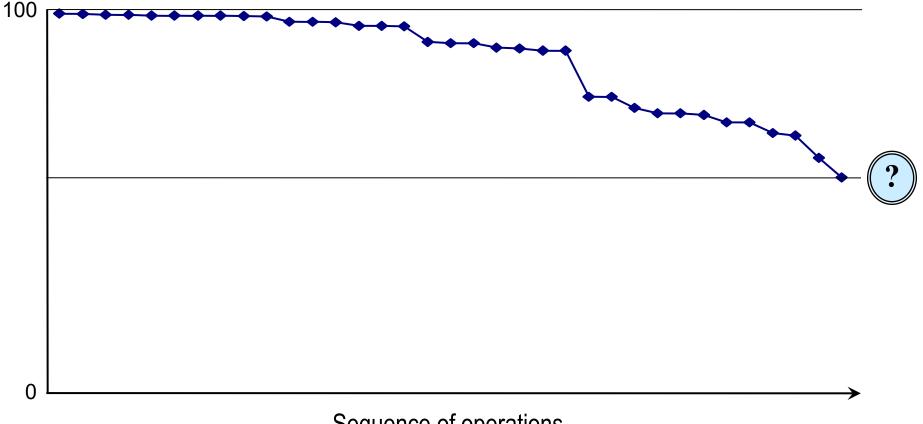


Overall yield = $(0.99999)^{14} = 0.999860 \rightarrow 99.986\%$

140 DPPM

Exercise 2.1

The average yield for 35 operations in an assembly process is 98.4%. Calculate the rolled throughput yield under the simplifying assumption that the yield for each operation is exactly equal to 98.4%. (The real answer would be the product of the actual operation yields.)



The area manager reported 98.4% as the overall yield of the operation. His reaction to the correct analysis followed the classic grief cycle:

Denial	"This can't be right. There must be a mistake in your calculation."	
Anger "This is ridiculous. You're wasting my time."		
Bargaining	ing "Isn't my method just as valid as your method?"	
Depression	'This is really bad. What am I going to tell everyone?"	
Acceptance	"I guess you can't solve a problem if you don't know you have it."	

We can count *defects* instead of *defective parts*

- Each potential defect on a part, or potential error in a transaction, is called an *opportunity*
- We can use DPMO (defects per million opportunities) instead of DPPM (defective parts per million)
- DPPM is more *customer* focused

The fact that **anything** is wrong is primary — the **number of things** wrong is secondary

• DPMO is more *process* focused

DPMO is a finer measure thanDPPM — it responds more rapidly to process changes

• Requirements for using DPMO

 \checkmark A finite number of identifiable opportunities per part or transaction

✓ Statistical independence of defect occurrence at different opportunities

In many cases, failure rates are quantified as percentages

2	7
/	

Definition of "opportunity"	Fraction defective	Expressed as a percentage	Focus
Each part	Defective parts All parts	% Defective	Customer
Each possible defect on a part	Defects (All parts) × (possible defects per part)	Defects per 100 opportunities (DPHO)	Process
Each transaction	Defective transactions All transactions	% Defective	Customer
Each possible error in a transaction	Errors (All transactions) × (possible errors per transaction)	Defects per 100 opportunities (DPHO)	Process

- In the 1990s, GE shifted the emphasis from the Six Sigma quality goal to *Six Sigma projects* the way to pursue the goal
- Leaders and Champions define *key performance indicators* (KPIs) — a "balanced scorecard" including but not limited to \$\$ measures
- KPIs drive a prioritization process
- Prioritization tells us which project(s) should be first in line
- "Black Belts" or "Green Belts" lead the project teams
- "Champions" provide resources and remove barriers for the teams

Examples of projects

Project	Annual \$\$ benefit
Reduce alpha case on large titanium castings	20,800,000
Reduce cost and lead time to develop extrusion tooling	2,000,000
Reduce wasted medication in hospital central pharmacy	1,100,000
Reduce roll stock inventory in box plant	768,000
Reduce cost of belt grinding in casting finishing	500,000
Improve the court collections process in city government	400,000
Reduce DOA replacement parts in field service	216,000
Reduce DPMO and amount of testing of circuit boards	192,000
Reduce electricity consumption in manufacture of airline storage bins	65,000
Reduce RFQ turnaround time (not counting increased PO hit rate)	34,000

3 Why Combine Lean and Six Sigma?

- They require the same culture
- They employ common strategies
- They focus on complementary problem areas
- They employ complementary methods
- They emphasize fact over opinion and use data to inform decisions
- One improvement infrastructure is better than two

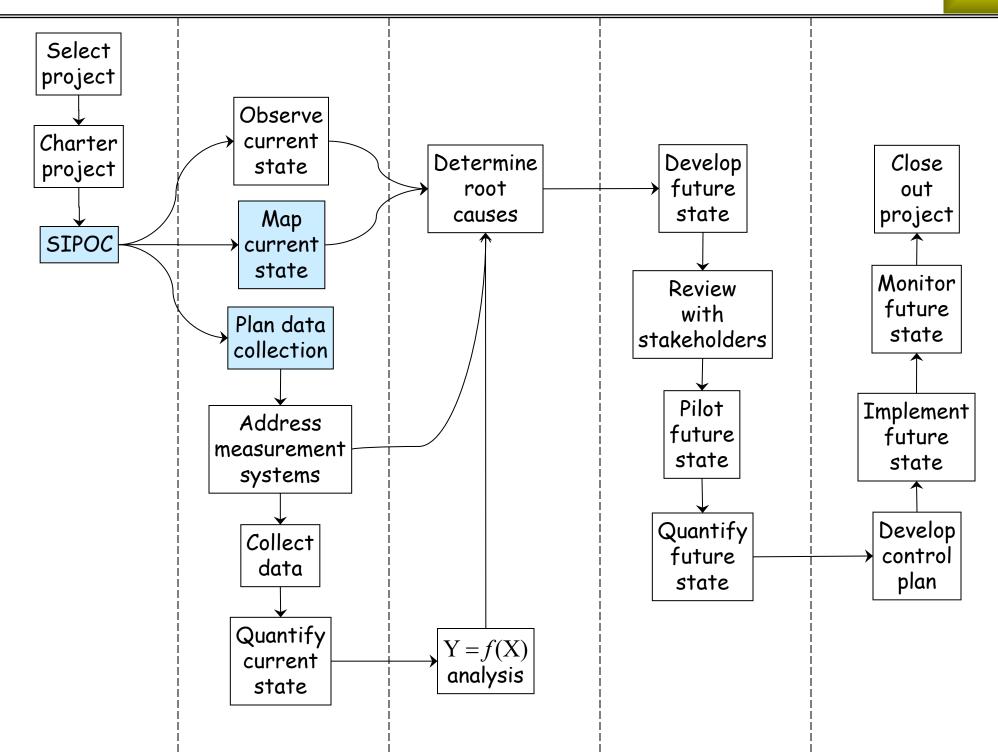
- Driven by Voice of the Customer
- Focus on eliminating waste
- Focus on processes and process improvement
- Improve processes via team projects
- Keep the improvement cycles going

Complementary problem focus and methods

Lean	Six Sigma
Lead time WIP Other visible waste	Defects "Invisible" waste
Defects caused by chaos and confusion	Defects caused by materials and equipment
Root causes easier to determine. (Processes directly observable.)	Root causes harder to determine. (Processes often not observable.)
Value stream mapping Geographic mapping	Basic process mapping Cross functional process mapping
Defines and standardizes the "Wisdom of the organization"	Data collection and analysis to discover a new solution
Common TPS solutions can be adapted to many circumstances	Project roadmap provides a method for finding solutions

Define Phase

4 Project Scope and SIPOC

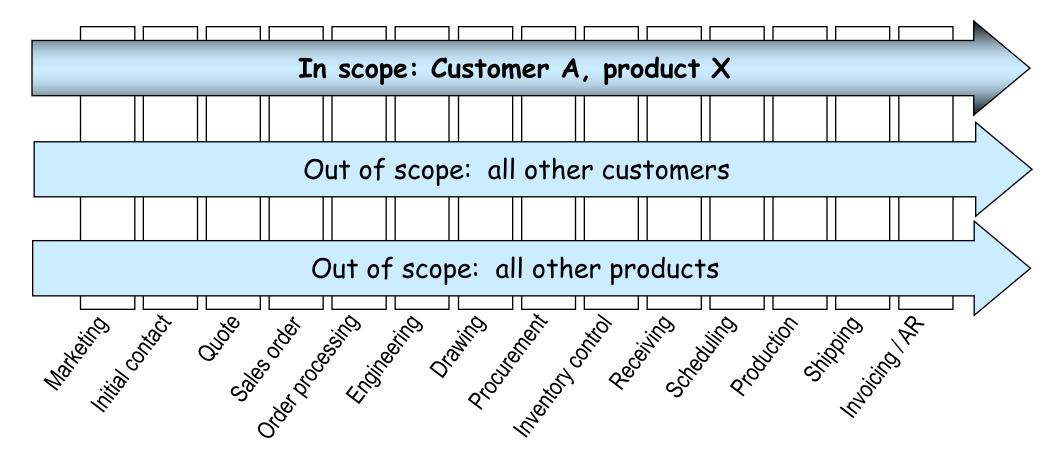


Defines the project scope in terms of . . .

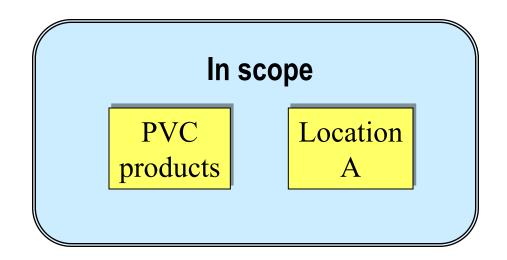
- ✓ Which customers?
- ✓ Which products?
- ✓ Which locations?
- ✓ Which materials?
- ✓ Which suppliers?

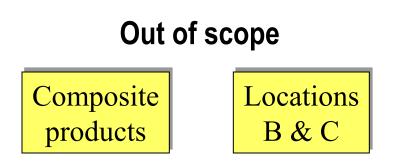
✓ ...

Value streamAll activities needed to provide a specified family of products or
services to customers



Project to reduce cost and lead time of extrusion tool development





Defines the project scope in terms of . . .

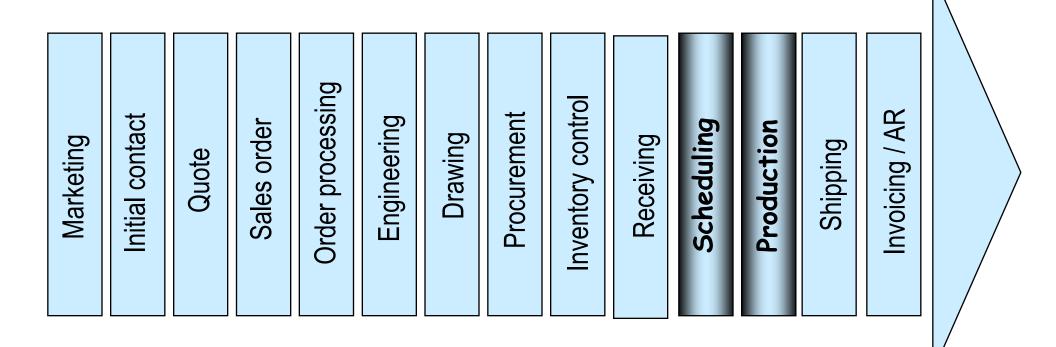
- ✓ Which activities?
- ✓ Which operations?
- ✓ Which processes?
- ✓ Which areas?

✓ ...

✓ Which departments?

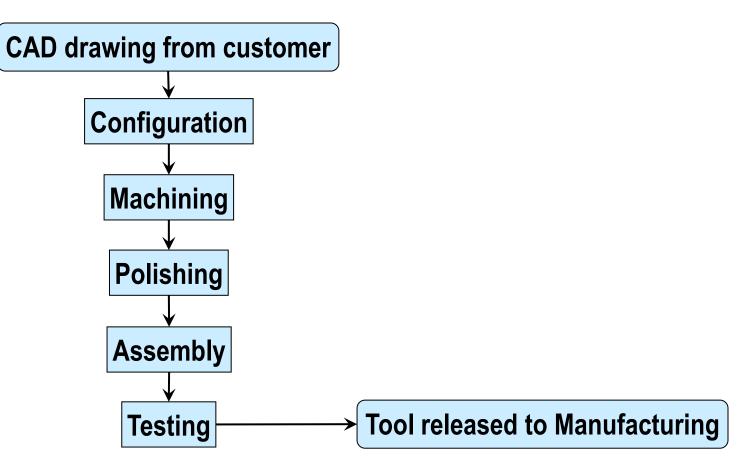


Which *activities* in the value stream are addressed by the project?

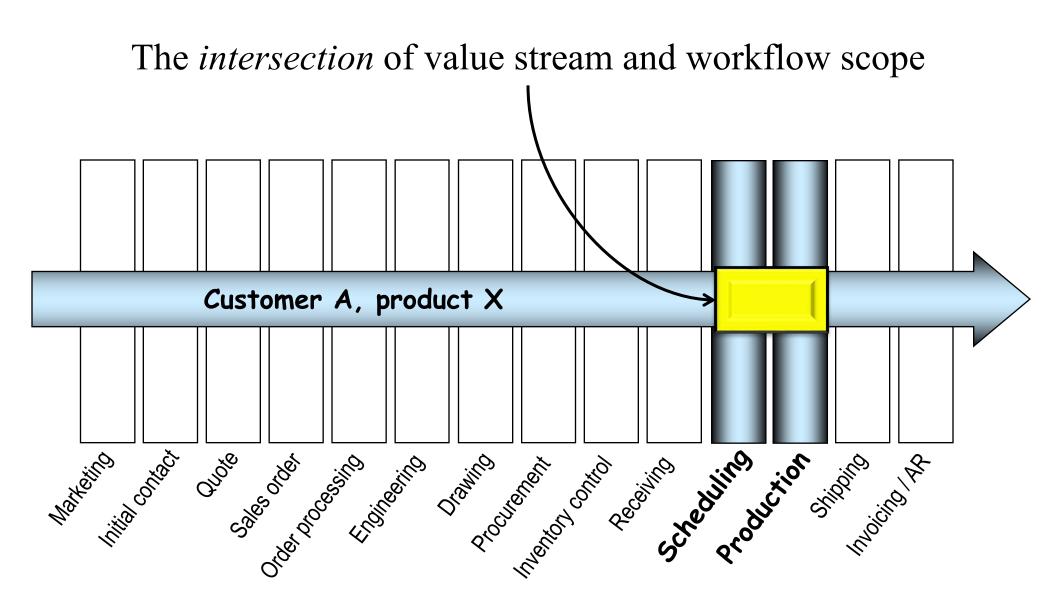


- Scheduling and Production are in scope
- Everything else is out of scope
- How will this affect the activities of the project team?

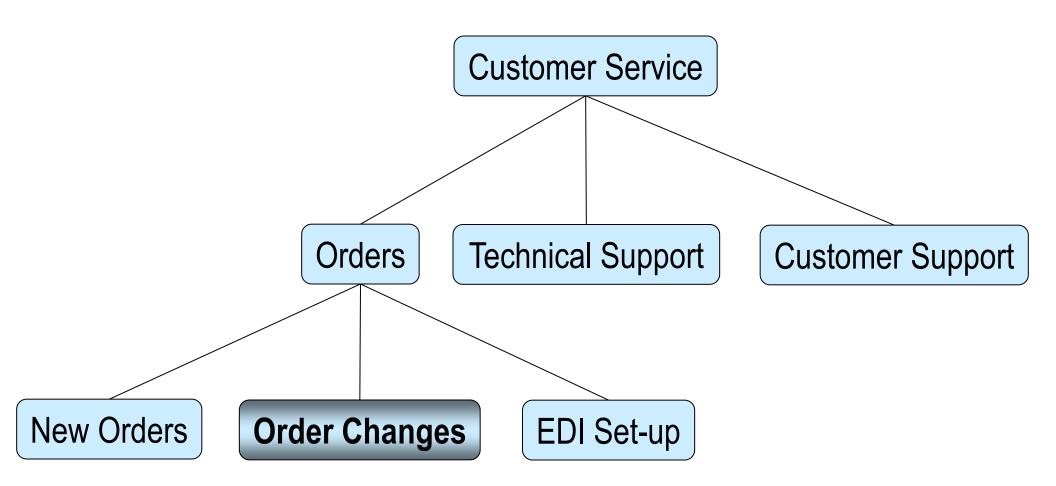
Project to reduce cost and lead time of extrusion tool development



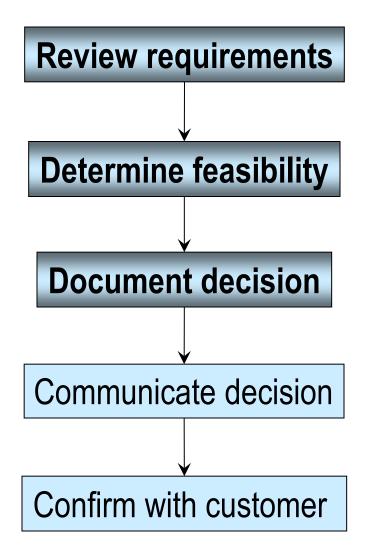
- Manufacturing is out of scope
- The project is not chartered to analyze and improve Manufacturing
- What is the relationship between Manufacturing and the workflow scope?



The project will address only order changes



The project will address only the *first three steps* of the order change process



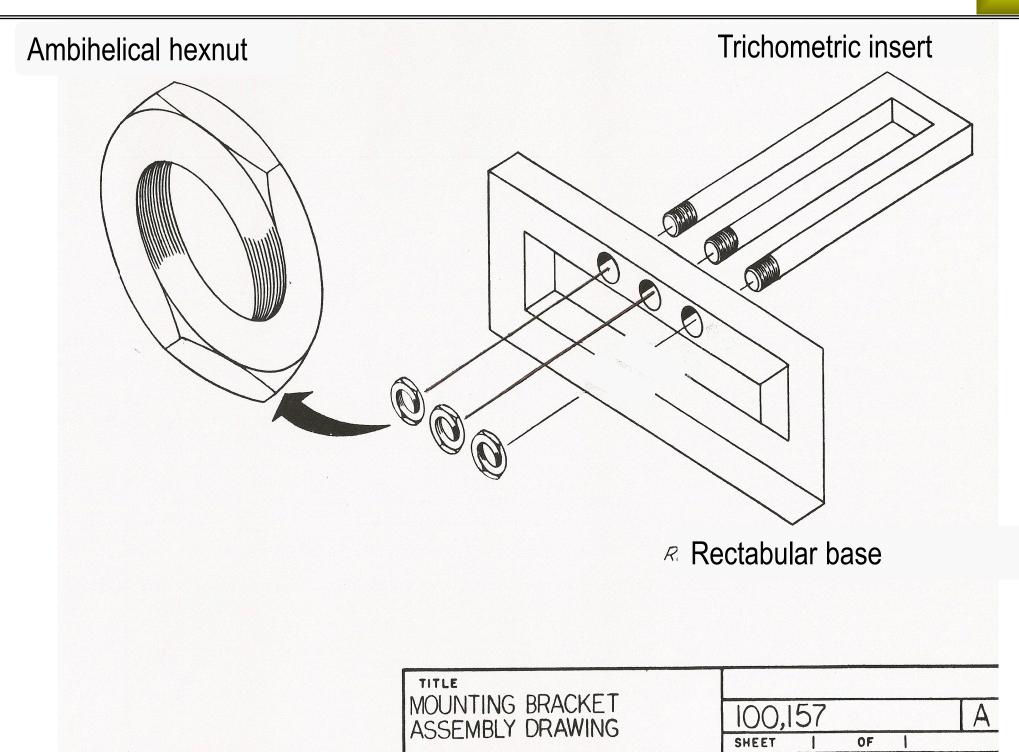
Exercise 4.1

Our company makes prototypes for various types of mounting brackets. These are classified as either standard or non-standard. A project has been launched to reduce the lead time for designing and building prototypes for non-standard brackets (see slide below for a typical example).

What is the value stream scope for this project?

What is the workflow scope for this project?

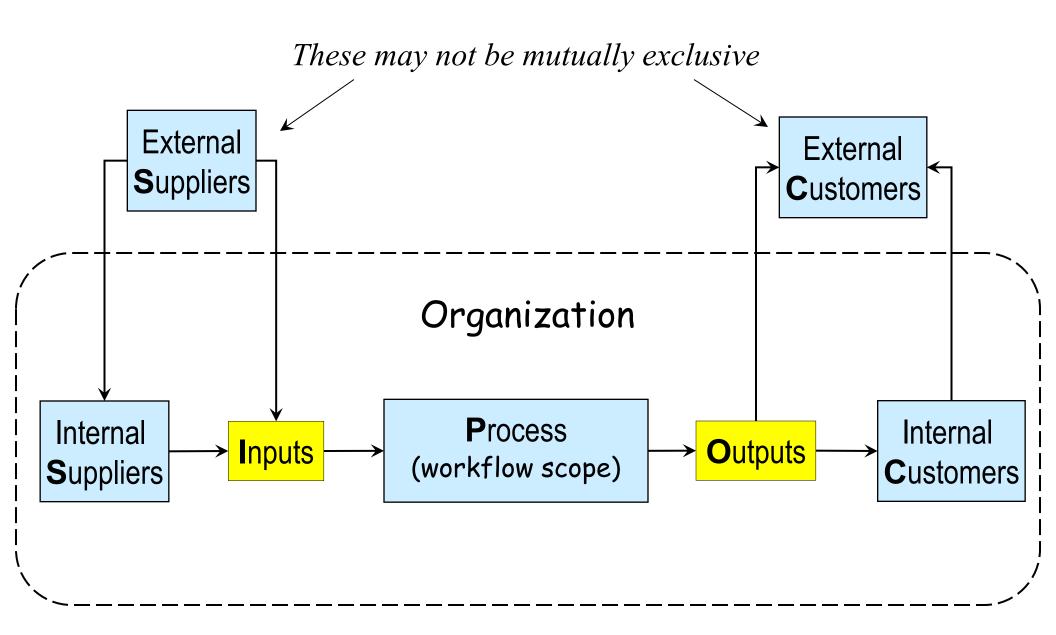
A non-standard mounting bracket



- SIPOC is a document that frames the project in the *process* space:
 Suppliers → Inputs → Process → Outputs → Customers
- SIPOC also documents the *data collection* needed for the project
- The five elements of SIPOC are defined on the slide below.
- The logical sequence for reading or creating a SIPOC:

 $\textbf{P} \rightarrow \textbf{O} \rightarrow \textbf{C} \rightarrow \textbf{I} \rightarrow \textbf{S}$

5) Suppliers	Entities who provide necessary <i>inputs</i> to the workflow scope. Suppliers may be internal or external to the organization.		
4) Inputs	Products, services, or information provided to the workflow scope by suppliers.		
1) Process	ProcessThe workflow scope: the activities to be analyzed and improved. A <i>high-level</i> description including first step, main intermediate steps, and last step.		
2) Outputs	Products, services, or information provided by the workflow scope to customers.		
3) Customers	Entities who receive <i>outputs</i> from the workflow scope. Customers may be internal or external to the organization.		



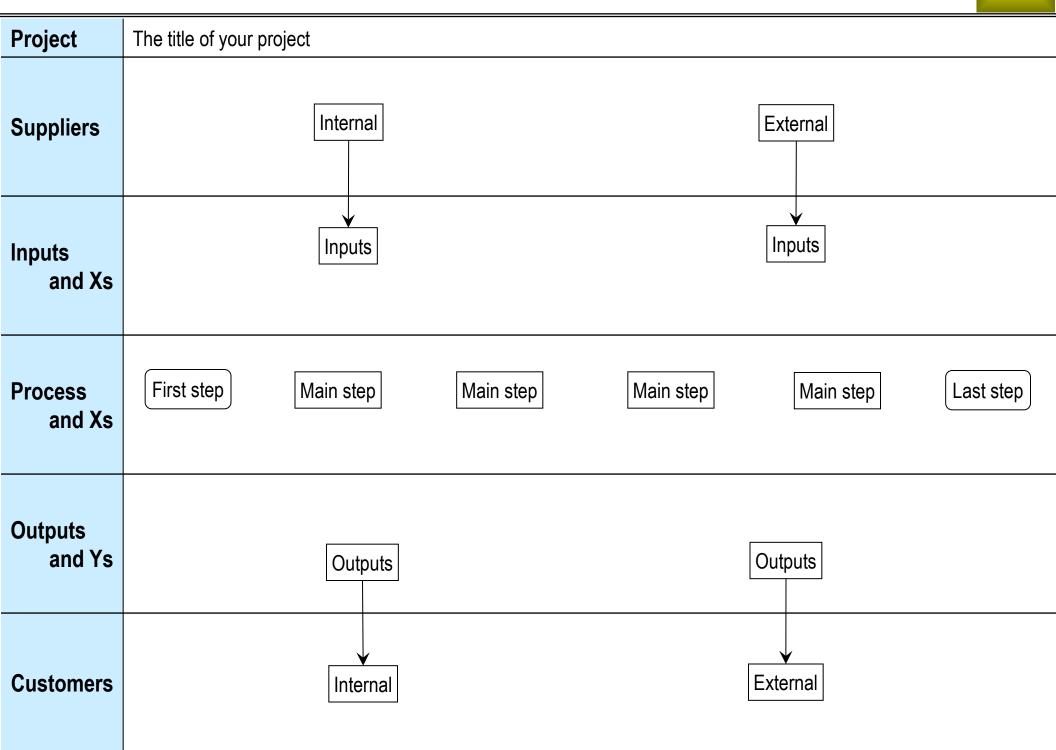
Y variables

- A *data variable* is measurable characteristic defined for individual parts or transactions (What does "variable" mean?)
- *Y variables* are measurable characteristics of *outputs* from the workflow scope
- They are the data variables from which the statistical **project metrics**, such as average or percent defective, are calculated
- Examples: lead time, pass or fail, quantitative measures of poor quality
- The Y variables are the reason we are doing the project (Why?)

- Data variables that are possible causes of variation in the Ys are called *X variables*
- Examples: Who, What, Where, When, Which, ...
- The greater the number of X variables identified, the greater the chance of solving the problem (Why?)
- The Fishbone Diagram will be used in the Measure Phase to identify and document the X variables

The SIPOC will contain only products, services, or information provided to the workflow scope by suppliers.

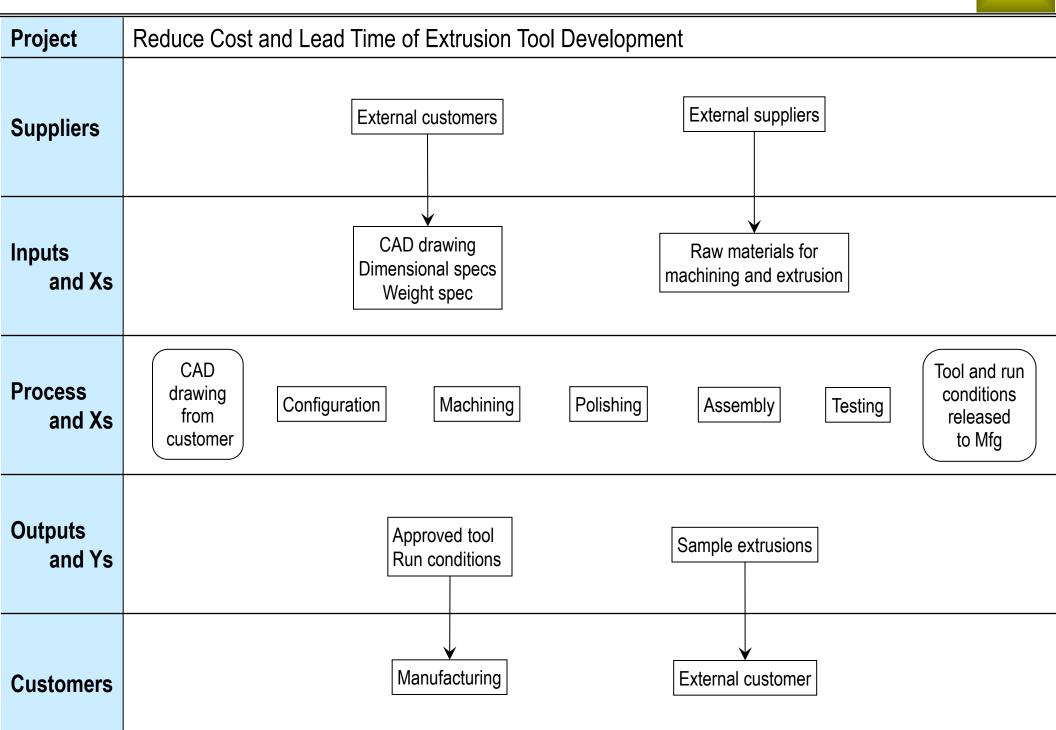
Blank SIPOC template



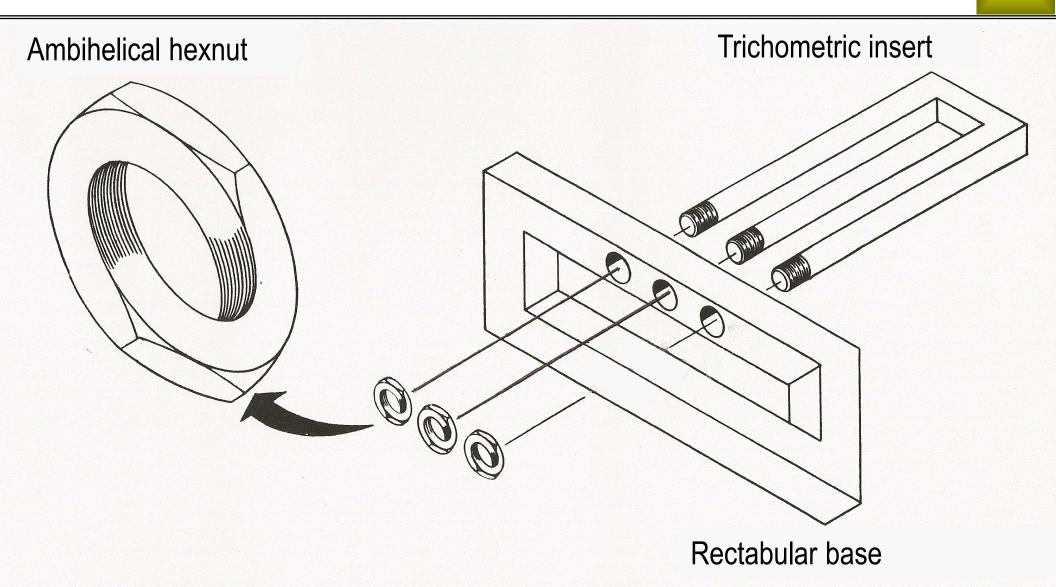
The slide shows a graphical SIPOC template. All you have to do is edit the various boxes and text. You can also add or delete boxes or text.

The following two slides show the graphical SIPOCs for two case studies.

SIPOC example 1



A non-standard mounting bracket



TITLE	
MOUNTING BRACKET ASSEMBLY DRAWING	100,157
AUDLI DIVININO	SHEET

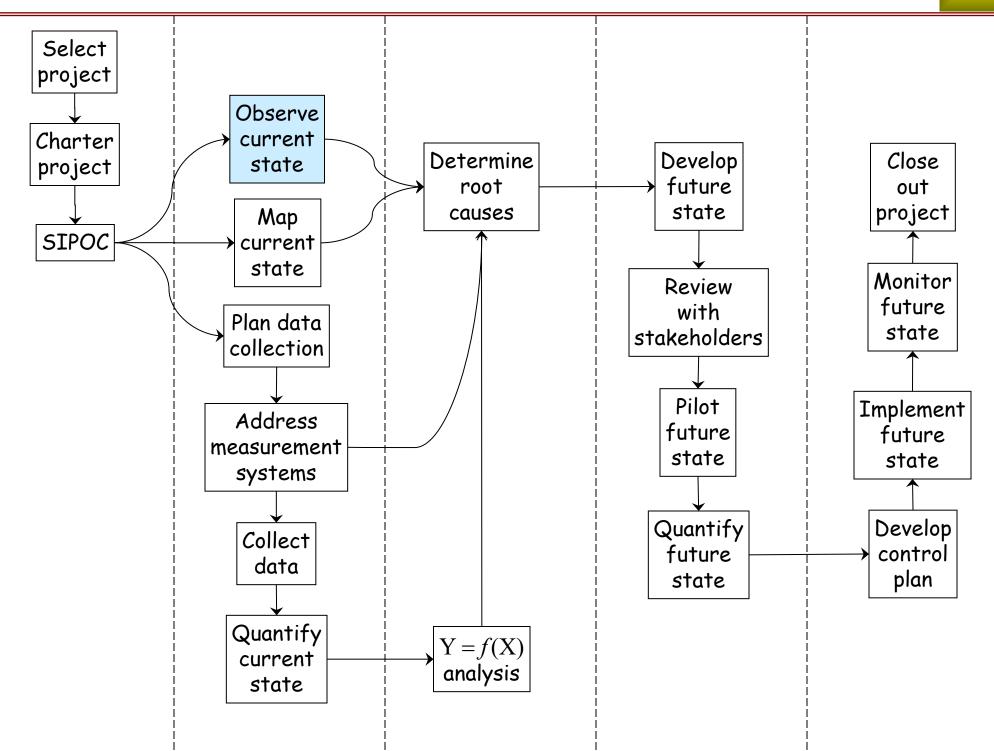
54

A

OF

Measure Phase

5 Observing the Current State



- The purpose is to improve the process, not to audit work performance
- Workflow observation periods should be scheduled in advance with appropriate supervisors and/or managers
- Workflow participants must be briefed on the project charter
- Participants must have adequate advance notice of observation periods
- Observations should be limited to the value stream and workflow scopes for the project

- Don't "gang up" on a few participants or process steps deploy team members effectively to get as many perspectives as possible
- Ask permission to take notes, photographs or videos this helps team members get the information they need without having to repeat questions later
- Observations should begin with introductions and guided tours, in some cases
- This should be done on all relevant shifts
- Subsequent "unguided" observations are often needed

- Interview workflow participants within the project scope
- Identify data variables and inspection points for inputs provided by internal suppliers
- Interview internal suppliers and customers of the workflow scope
- Identify data variables and inspection points for outputs provided to internal customers
- Identify NVA activities these may be opportunities for improvement within the project scope
- Confirm or revise process map(s)

Team roles & responsibilities

	Bob	Carol	Ted	Alice	Мое	Larry	Curly
Interview workflow participants	\checkmark			✓			
Observe and record changes to process map		✓			~		
Identify workflow data variables and inspection points			✓			~	
Identify data variables and inspection points for workflow inputs				✓			~
Interview internal customers	✓				~		
Identify data variables and inspection points for workflow outputs		✓				~	
Focus on measurement systems			✓				~

60

- The *way* you ask questions can affect the usefulness of the answers you get
- *Closed* questions can be answered with "yes" or "no" if the person is reluctant to talk to you, closed questions will not get you anywhere
- *Open* questions start with words like *what, why, when, where, who, which, how,* etc.
- Open questions are much better for eliciting information, ideas, opinions, etc.

Asking questions (cont'd)

Open questions	Closed questions			
"How do you do that?"	"Can you see from where you're sitting?"			
"Why is it done this way?	"Can you hear me in the back?"			
"How do you think that would help?"	"So, you agree with the schedule change?"			
"When you say , what do you mean?"	"Have we decided to meet on Fridays?"			
"What would be an example of that?"	"We covered that earlier, didn't			
"What are some possible causes of?"	we?"			
"Why do think that could be a cause?"	• Closed questions are useful for moving a conversation along			
"Why do you think that happens?"	• Try to phrase them so that the answer you want is "yes"			

Concentrate on what is being said.

Respond with eyes, voice, gestures, and posture to communicate empathy and understanding.

Reflect information by paraphrasing.

Elicit information by asking questions.

Control the urge to interrupt, judge, or change the subject.

- □ Are there opportunities for reducing batch size?
- □ Where is the greatest amount of work-in-process (WIP)?
- □ What are the most common do-overs?
- □ Is the physical layout causing excessive movement of people or material?
- □ Is there unnecessary complexity?
- □ Where are the most time-consuming changeovers?
- □ Are there opportunities for mistake proofing?

□ Are there serial activities that could be parallel?

- □ Are there separate steps that should be combined into a single step?
- □ Are there single steps that should be split into separate steps?
- □ Are work instructions missing, outdated, or not visible?
- □ Are there problems with availability of equipment or material?

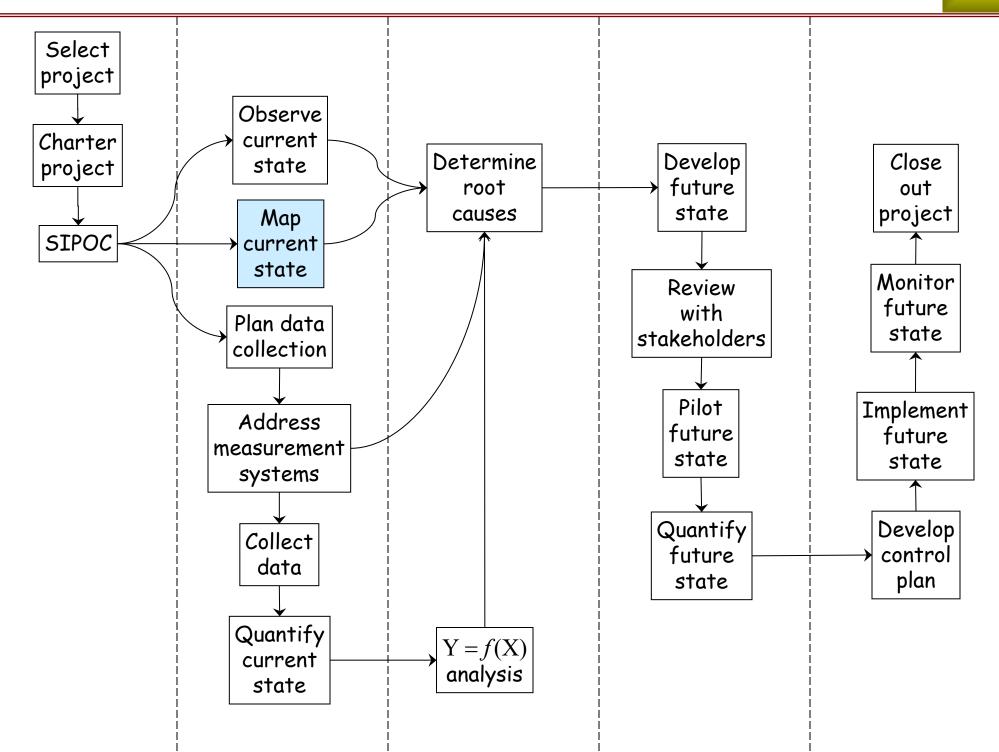
••••

- Team members may see possible causes of problems and solutions as soon as they start observing and mapping the current state
- These observations should *not* be publicized until the appropriate point in the project roadmap
- These observations *should* be logged as they arise, preferably in Excel (facilitates categorization and prioritization)
- The possible causes will be reviewed in the *Analyze* phase, along with data analysis results, to determine root causes
- The possible solutions will be reviewed in the *Improve* phase to develop the future state

Observation log (cont'd)

Team member	Date	Location	Possible cause	Possible solution

6 Basic Process Mapping

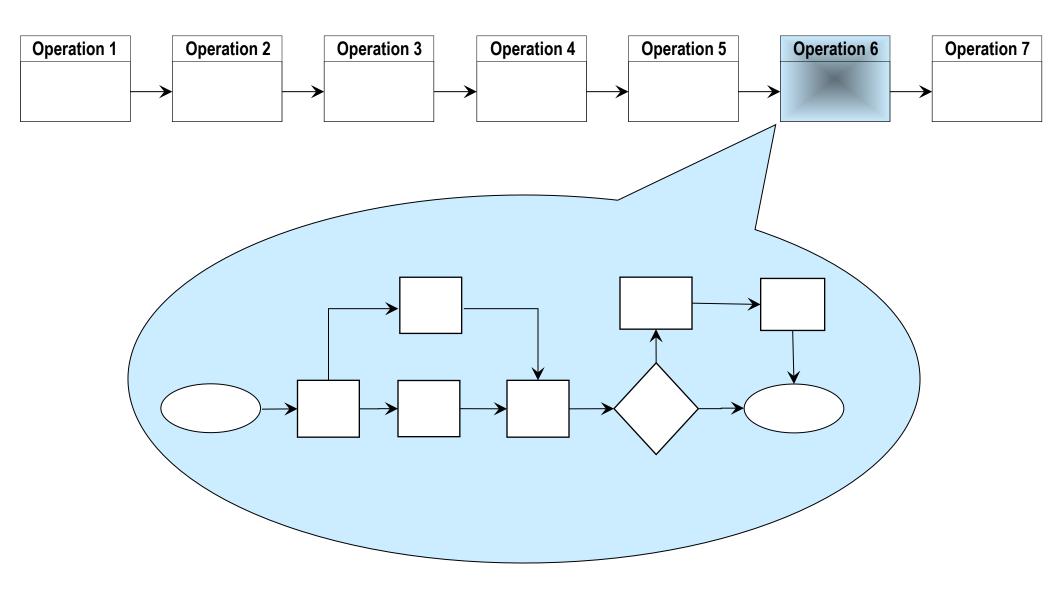


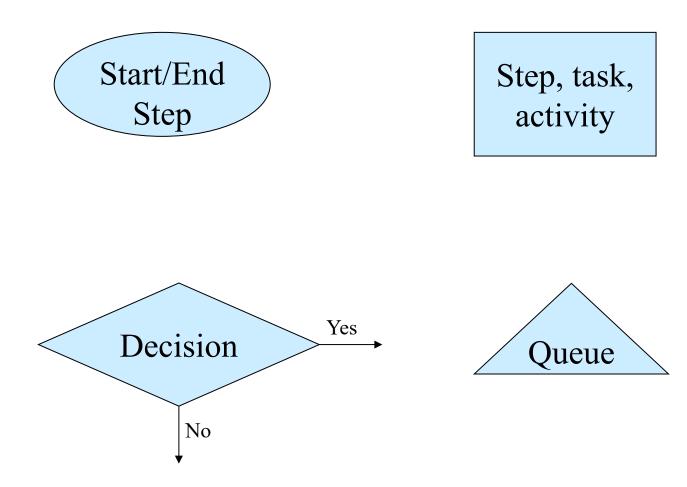
Process mapping is easy to learn and produces useful documentation of the current state. It is also a great team building activity.

The key to successful application of any mapping technique is to focus on the appropriate *level of activity* for your project. In SIPOC we identify the first, last, and main intermediate steps of the in-scope workflow. This gives you a high-level process map.

A high-level map is a good starting point for more detailed mapping. A basic process map, discussed in this section, shows individual tasks and decision points within the main steps. A cross functional or swimlane maps shows who is responsible for each task and decision. This and other common mapping formats are discussed in the next section.

A high-level map is also the usual starting point for value stream mapping (VSM). VSM combines visualization of what is happening with certain forms of data analysis. VSM will be discussed later in the program. Often, we want to create detailed maps for some or all of the main steps given in the SIPOC





Suspend your disbelief	Map the process the way it really is, not the way you think it should be.
Don't make assumptions	If you don't know what happens at a certain point, or can't agree on what happens, put a question mark there. Then, go ask someone who does know.
Solicit feedback	Ask participants of the in scope workflow, and their internal customers, to review the map for accuracy and clarity.
Document your work	Use mapping software to create an electronic version of the map.

Writing good narrative

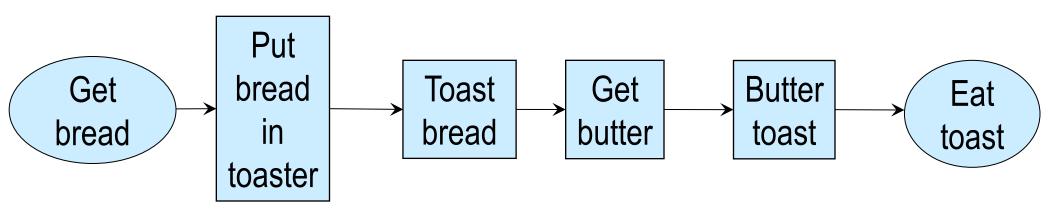
✓ Use active voice, not passive voice
 ☺ Order is entered
 ☺ Enter the order

✓ Use verb/object, not name of activity
 ☺ Order Entry
 ☺ Enter the order

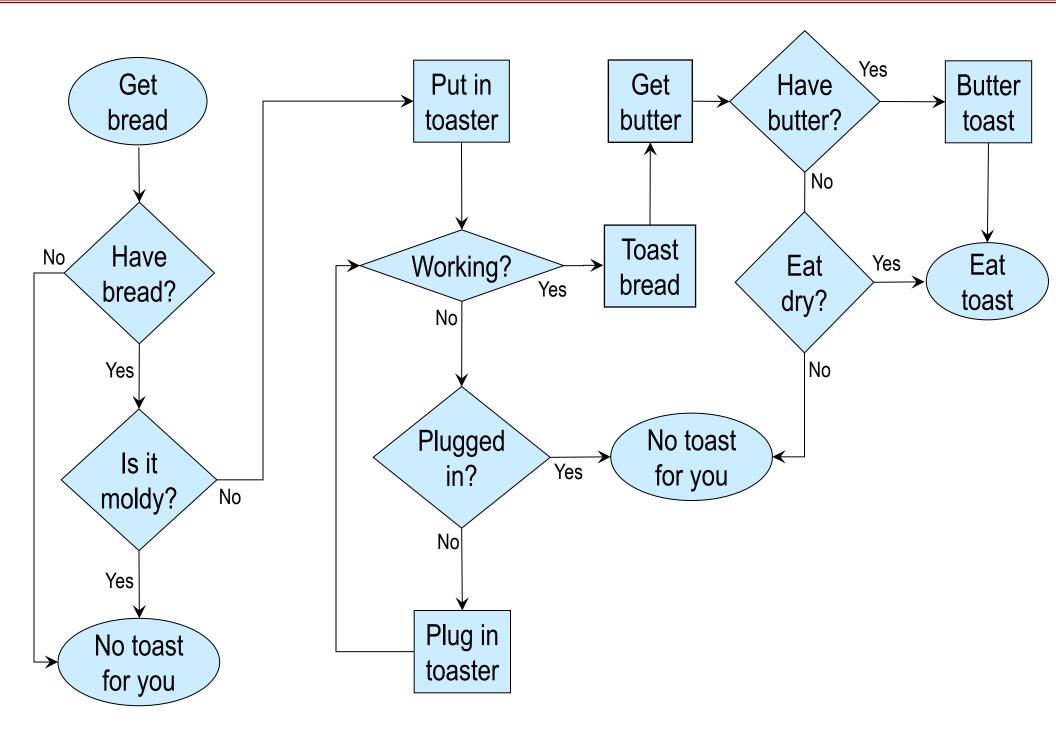
✓ Use short sentences with familiar words
 ☺ Twilight's last gleaming
 ☺ Dusk

✓ Use present tense

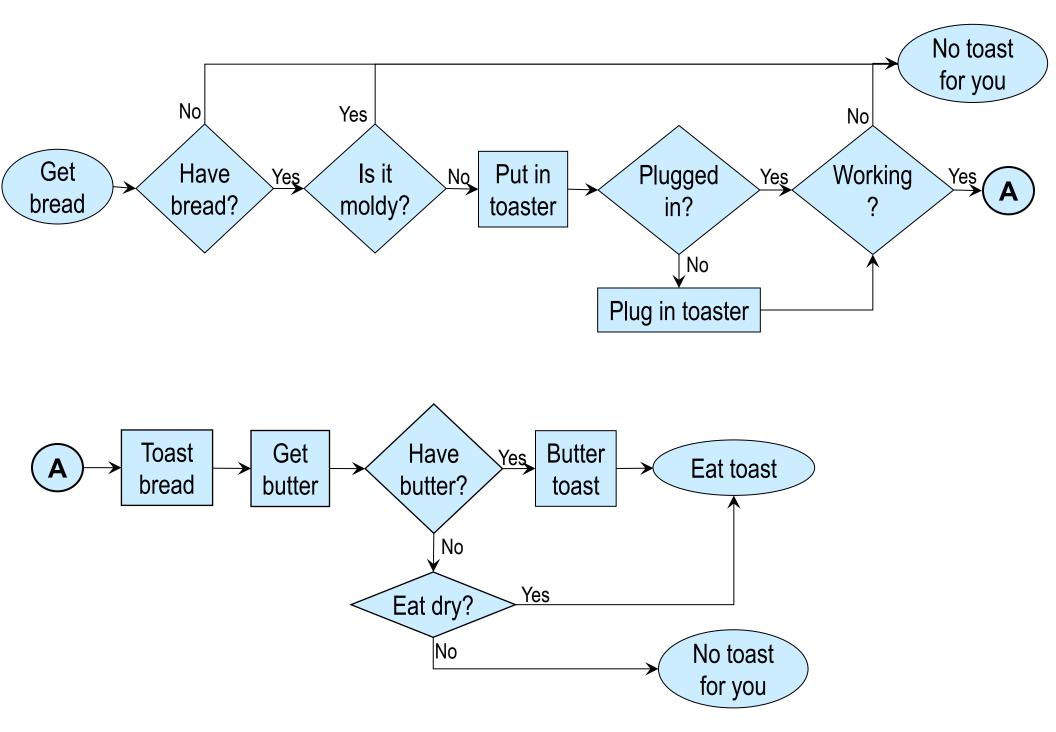
✓ Use logical, consistent layout



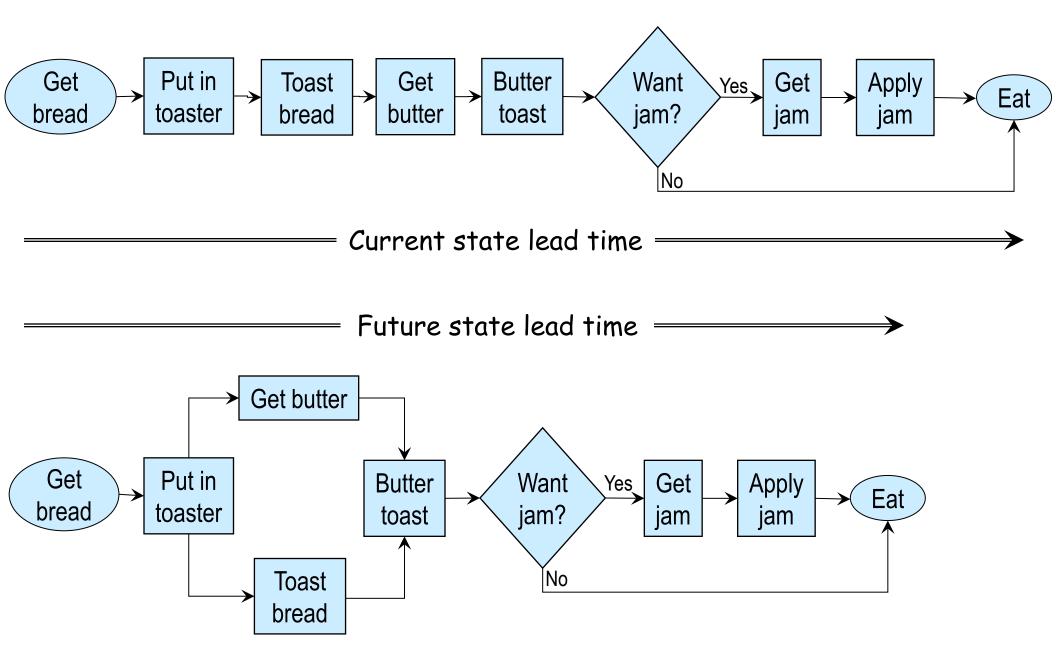
Decision steps show what really happens



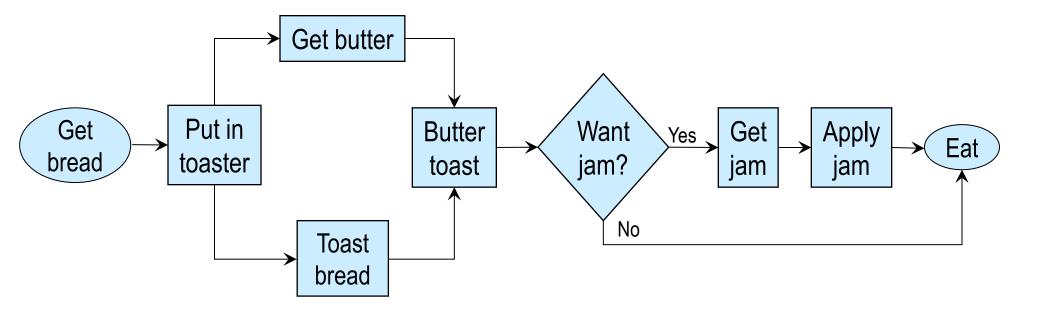
Best practice: follow a qualitative timeline



Common technique for reducing lead time: convert serial to parallel



How would you modify the toast-making process to further reduce the lead time?



You are to create a process map based on the information given on the slide below. It will be beneficial to work on this in small groups.

This is not *your* process. Someone else is describing *their* process to you. Do not make unwarranted assumptions!

Use a separate sheet of paper to draw your map. Use a qualitative timeline!

Exercise 6.2 (cont'd)

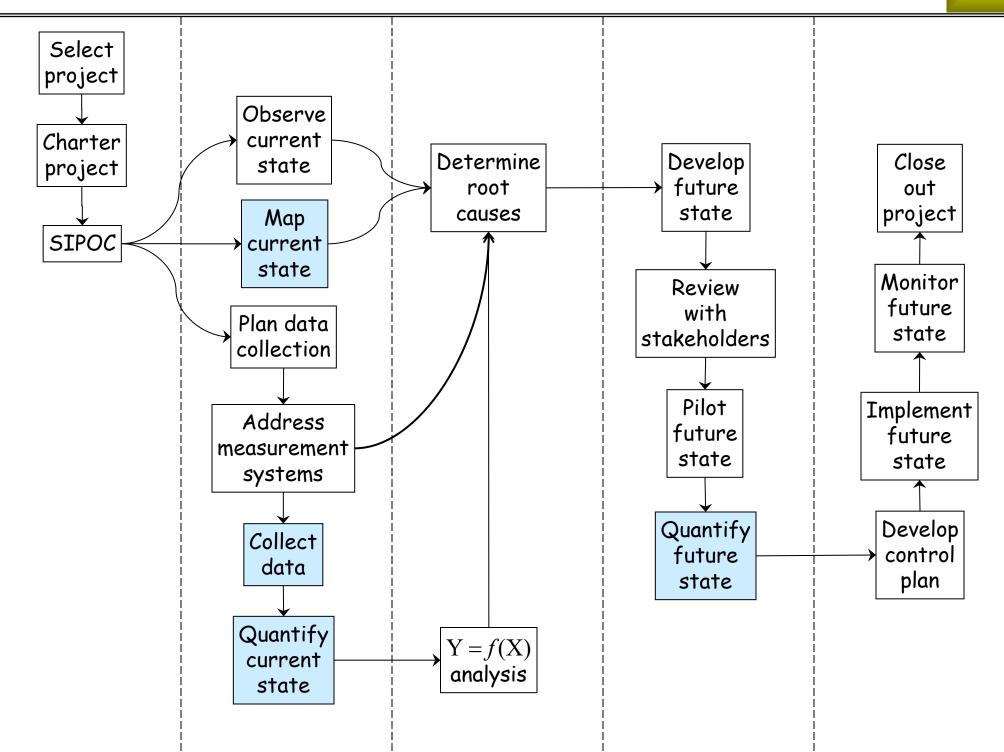
There are two types of material, A and B. The material must be processed before it can be used. There are two steps in this process. For Process 1, the A and B materials must be processed in separate Type 1 machines. If two Type 1 machines are available, load the A material into one machine, the B material into the other, and run the two machines at the same time. If there is only one Type 1 machine available, run the two loads sequentially in that machine.

When Process 1 is completed, unload the material, and move on to Process 2. Process 2 requires Type 2 machines. If two Type 2 machines are available, load the A material into one machine, the B material into another, and run the two machines at the same time.

Unlike the Type 1 machines, the A and B material can be processed together in the same Type 2 machine. If there is only one Type 2 machine available, load both the A and B material into that machine for processing. This will take longer than processing the A and B materials in separate machines, but not as long as running two loads sequentially.

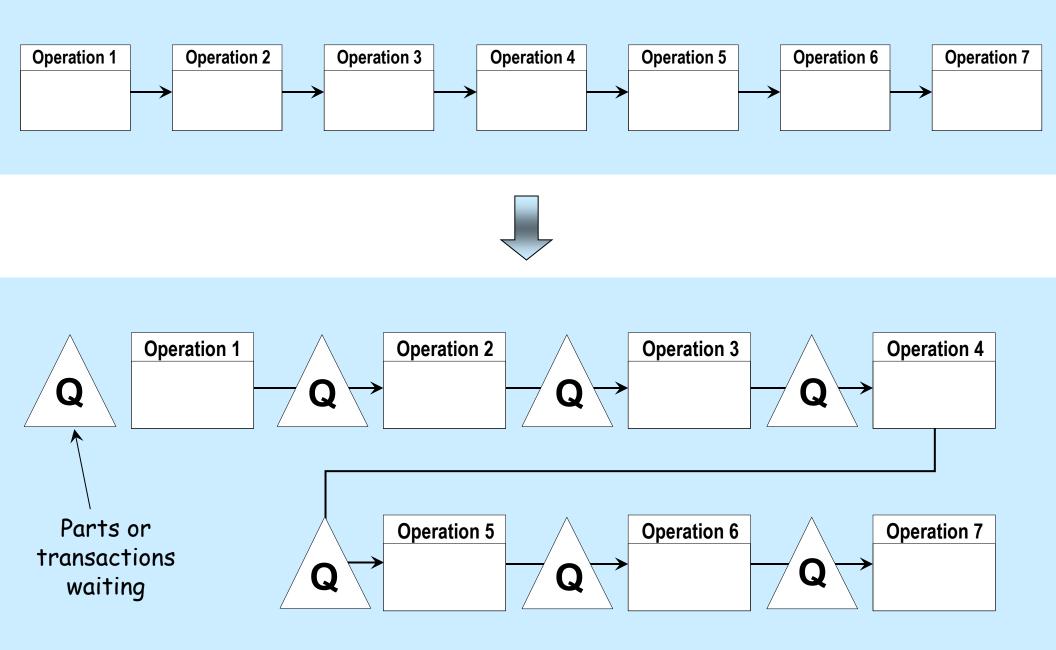
When Process 2 is completed, unload the material, separate the A and B materials if necessary, then store them for subsequent use.

7 Value Stream Mapping

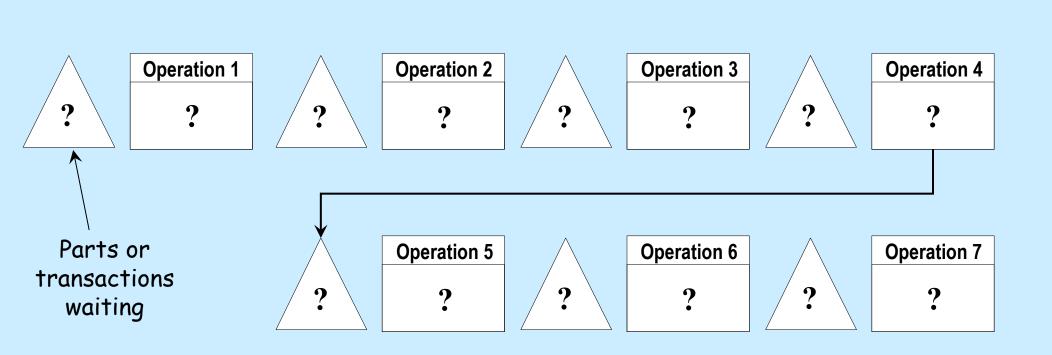


- Value stream mapping (VSM) combines several things:
 - \checkmark Visualization of the current state
 - \checkmark Documentation of the current state
 - \checkmark Certain types of data collection and analysis
- VSM is an effective way to identify improvement opportunities
 - Especially in projects involving WIP, capacity, and lead time reduction
 - \checkmark Also used to document the future state

High-level map from SIPOC



What is the average lead time? How much time is spent in each box or triangle? How do we get this information?



Definitions

Available Working Time (AWT)	 The time a process is available to conduct work AWT excludes time when work isn't occurring such as time for breaks, meetings, lunch, preventative maintenance, estimates of unplanned downtime, change overs, etc.
	• The average number of good parts or transactions completed over a period of time
Throughput (Tput)	 Typically measured as average over at least several days Throughput, lead time, and WIP are related through Little's
	Law

Definitions (cont'd)

Lead time (LT)	 The total elapsed time to produce one defect free product or transaction The time difference between when a part or transaction enters and leaves a process
Customer Demand Rate (CDR)	• The number of parts or transactions that the customer desires over a period of time (usually a day, week, or month)

Definitions (cont'd)

Takt time (TT)	 The pace at which an operation should complete products or transactions in order to meet customer demand during the Available Working Time. Available working time during a period divided by the number of products or transactions <i>required</i> during that same period
Cycle time (CT)	 The fastest repeatable time between part or transaction completions using the current processes and resources Shows how a process is capable of performing
	Combines with AWT to determine capacity

Process Cycle Efficiency (PCE)	• The percentage of time that WIP is being transformed by VA activities. In other words, the percentage of lead time that is value added.
Work In Progress (WIP)	• Includes items waiting to be worked on and items actively being worked on. WIP includes all of the inventory in the production system.

Available Working Time per day = 480 min - 90 min breaks, lunch, meetings

= 390 min

Avg. daily Customer Demand Rate = 32 units

Takt time = $\frac{390 \text{ minutes}}{32 \text{ units}}$ = 12.2 mins

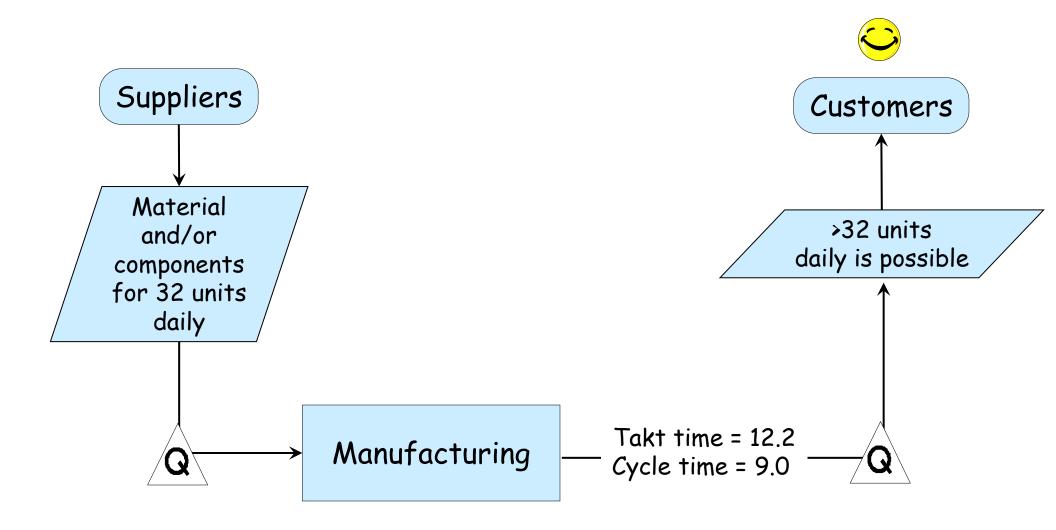
During a study of this process, parts were completed at the following times:

9:00, 9:09, 9:17, 9:28, 9:37, 9:46, 9:58, 10:07, 10:16, 10:24, 10:33, 10:42

Based on this, the elapsed time in minutes between completed units was:

9, 8, 11, 9, 9, 12, 9, 9, 8, 9, 9

Cycle Time = 9 minutes (the fastest repeatable value)



- Units of takt and cycle time: time divided by quantity
 - > Shorter cycle time \rightarrow more output
 - > Longer cycle time \rightarrow less output

- Cycle time *longer* than takt time
 - Cannot meet customer demand with current processes and resources

- Cycle time *shorter* than takt time
 - Can meet customer demand with current processes and resources, but may need to eliminate process variation

• Takt time longer than cycle time

• Downstream operations constrained to cycle time of upstream bottleneck

• Upstream operations pace themselves to cycle time of downstream bottleneck (pull system)

Using the information provided in Example 1, consider the scenario where the customer wants to increase their purchases from 32 to 42 units per day.

- a) What is the new takt time?
- b) What is the cycle time and is the new takt time faster or slower than the cycle time?
- c) Can you accommodate this demand increase?
- d) What problems might need to be solved?
- e) Why should cycle time measurements not typically be taken from process output data in an ERP system?

Method	Drawbacks	
Download accurate, time stamped	• The best scenario, if such data exists	
records from database	• Make sure WIP time is accounted for properly	
	• Tedious	
Shadow parts or transactions	 Logistically difficult 	
	• Time consuming for team members	
Tag documentation	 Anything identified as "special" is likely to be expedited 	
	• Data will not represent reality	
	• Tedious and time consuming	
Enter "file cabinet data" into Excel	 Likelihood of data entry errors 	
	• May not exist	
Little's Law	• Allows calculation of LT from WIP and T'put	

Lead Time = (WIP) / (Throughput)

- WIP is easy to count during process observation
- If WIP varies, count multiple times and use average or min/max to show range in lead time
- Throughput is the quantity completed during an observation period. Period should be at least several days.
- Lead time = amount of time that passes between when a piece enters and leaves a process or processes
- These values can be calculated for individual processes or for an entire production process chain

Average WIP for each box and triangle during an observation period



- Suppose in the system shown above, each operation has a throughput of 6 pieces per hour, so the entire production process is also making 6 pieces per hour
- We can use Little's Law to calculate the overall lead time for the process, for individual processes, or for subsets of processes

Applying Little's Law

	Avg. WIP
Queue 1	5.0
Operation 1	1.0
Queue 2	7.5
Operation 2	1.5
Queue 3	15.0
Operation 3	3.0
Queue 4	10.0
Operation 4	2.0
Queue 5	22.5
Operation 5	4.5
Queue 6	22.5
Operation 6	4.5
Queue 7	5.0
Operation 7	1.0
Total	105.0

The previously described process was studied and the average WIP counts are shown here. They are measured as follows:

- Queue WIP is the average pieces waiting to be processed. For example, Queue 1 WIP is the typical amount of work waiting to be processed by Operation 1.
- Operation WIP is the average pieces actively being processed. For example, Operation 1 is typically processing one piece.
- The Total WIP in the process is the sum of all of the Queue and Operation WIPs

Applying Little's Law

	Avg. WIP
Queue 1	5.0
Operation 1	1.0
Queue 2	7.5
Operation 2	1.5
Queue 3	15.0
Operation 3	3.0
Queue 4	10.0
Operation 4	2.0
Queue 5	22.5
Operation 5	4.5
Queue 6	22.5
Operation 6	4.5
Queue 7	5.0
Operation 7	1.0
Total	105.0

We can apply Little's Law to the entire process, an individual process, or a subset of processes. Remember:

Lead Time = (WIP) / (Throughput)

Since each operation, and therefore the entire process sequence, averages 6 pieces per hour, Little's Law lets us calculate lead times as follows:

• For the entire process:

Lead Time = 105 pieces / 6 pieces per hour = 17.5 hours or 1050 minutes

• For Queue 1 and Operation 1:

Lead Time = 6 pieces / 6 pieces per hour = 1 hour or 60 minutes

VSM with waiting and process times



Lead time = 1050 minutes or 17.5 hours

Waiting time = Sum of time in queue = 50 + 75 + 150 + 100 + 225 + 225 + 50 + 10 = 875 minutes

Process time = Sum of time the pieces are being worked on = 10 + 15 + 30 + 20 + 45 + 45 + 10 = 175 minutes

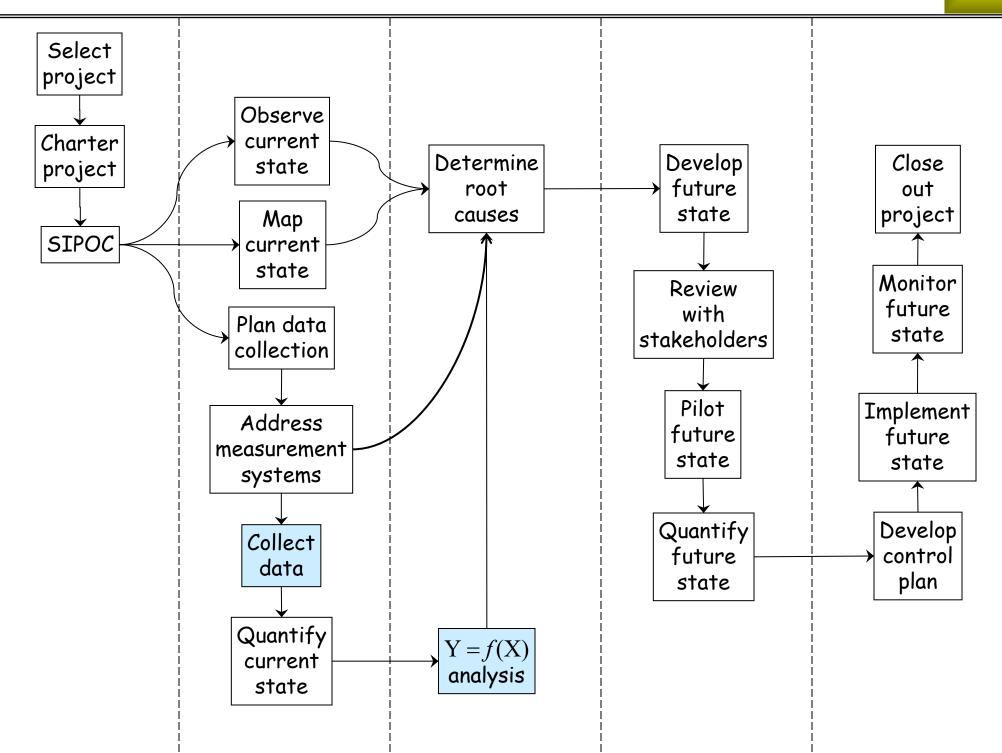
Process Cycle Efficiency = The percent of lead time that a part is being worked on = (175 / 1050)*100 = 16.7% a) A manufacturing process completes an average of 45 defect-free parts each day. The average WIP is 15 parts. Calculate the average lead time in hours.

b) Supposing in the example above, the company works one 8-hour shift per day. Under what conditions would this impact the lead time calculations and when would it not?

c) A manufacturing operation runs 365 days a year. They produce about 416 defectfree units of a particular product per year. The average WIP for this product is 40. Calculate the average lead time in days.

d) Should raw materials be counted as WIP?

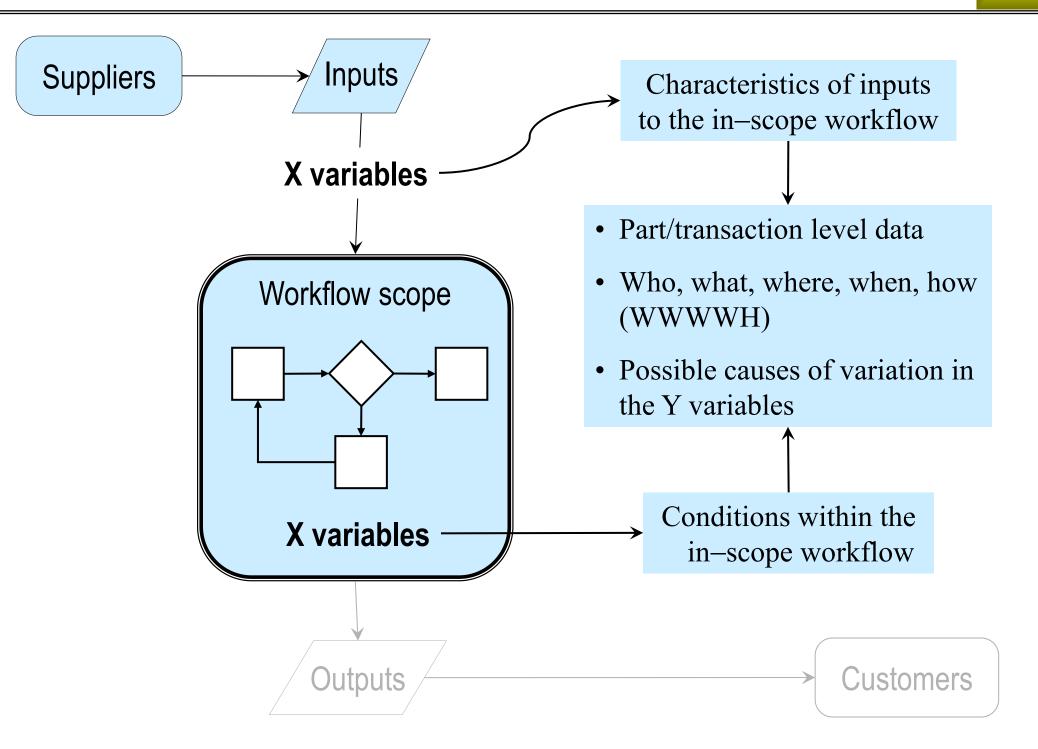
8 X and Y Variables



Topics

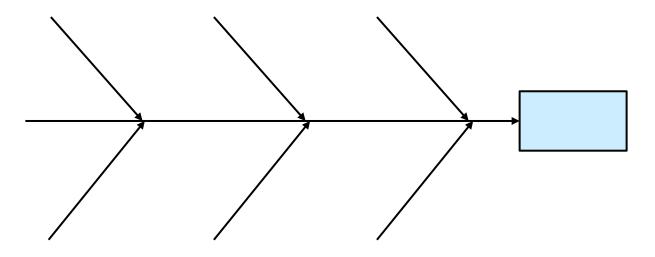
- X variables
- Fishbone Diagram
- Prioritizing X variables
- Y variables
- Operational definitions for data variables
- "Big Y" and "little y"

X variables



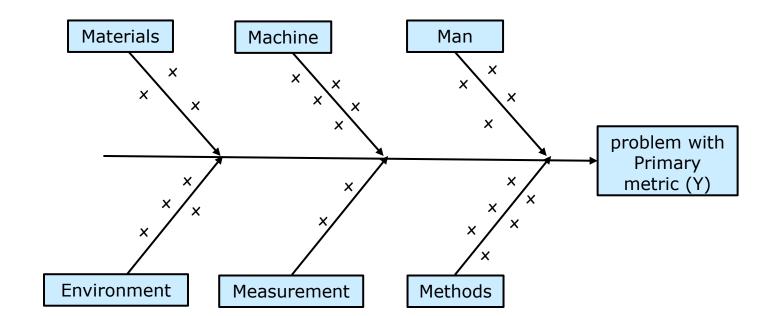
The Fishbone Diagram is:

- used to identify all potential causes (X's or inputs) of the effect (output or problem of interest), usually the primary metric.
- part of identifying process inputs during the Measure Phase
- most often associated with root cause analysis
- also known as Cause-and-Effect Diagram and Ishikawa Diagram



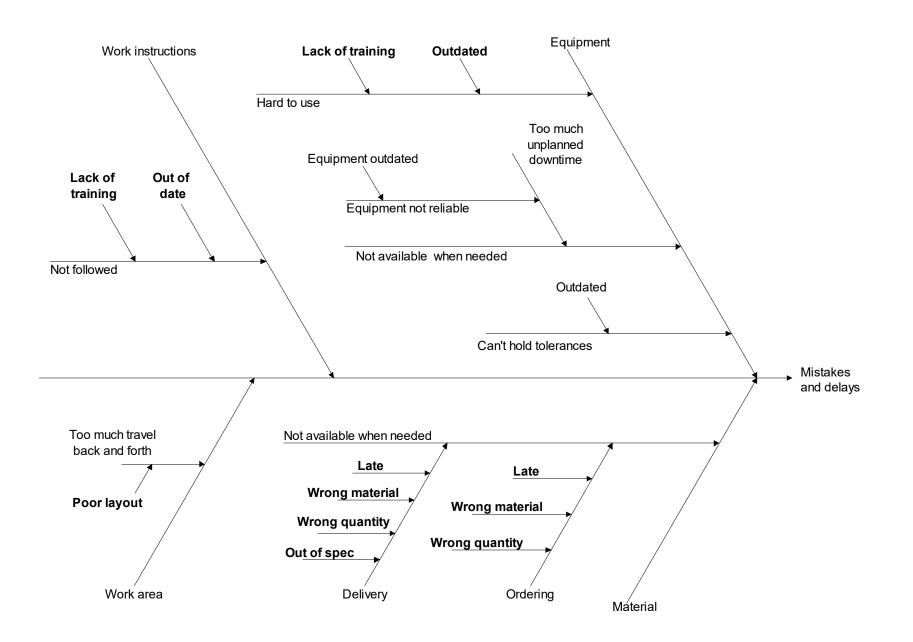
The Fishbone Diagram is created with the project team.

- It focuses the team on the particular effect, shown in the "head of the fish"
- All ideas for potential causes (critical x's) are collected using brainstorming
- Categories on the main "bones" help trigger ideas
 - Standard categories are Man, Machine, Materials, Methods, Measurement and Environment ("5 M's and an E")
 - The team can choose to use different categories
 - Standard categories (with minor modifications) are recommended for your first uses



The Fishbone Diagram must be visible to the entire team during the brainstorming (creation) session.

- 1. Put output of interest (usually primary metric) in the "head of the fish."
- 2. Choose categories for "bones"
 - Standard Categories: Man, Machine, Materials, Methods, Measurement, Environment
 - The team can choose to use other categories
- 3. Brainstorm all possible inputs (x's) that could cause the problem seen in the output (primary metric—Y)
 - Rules for Brainstorming: Accept all stated ideas and add to diagram; No ideas are evaluated or rejected during the brainstorming session
- 4. Break broad categorical x's into more useful, more measurable features
 - Measurable features can be verified as causes of performance issues in the primary metric during the Analyze Phase
 - We can act upon them to improve the process
 - They need to be identified early in the project
 - Example: Work instructions not followed—out of date; lack of training
- 5. Highlight those x's deemed most important by the team



A project has been launched to improve the mounting bracket development process (MBDP) in a company that makes mounting brackets. Background on the project and process may be found in the following files in the *Student Files* folder:

MBDP charter

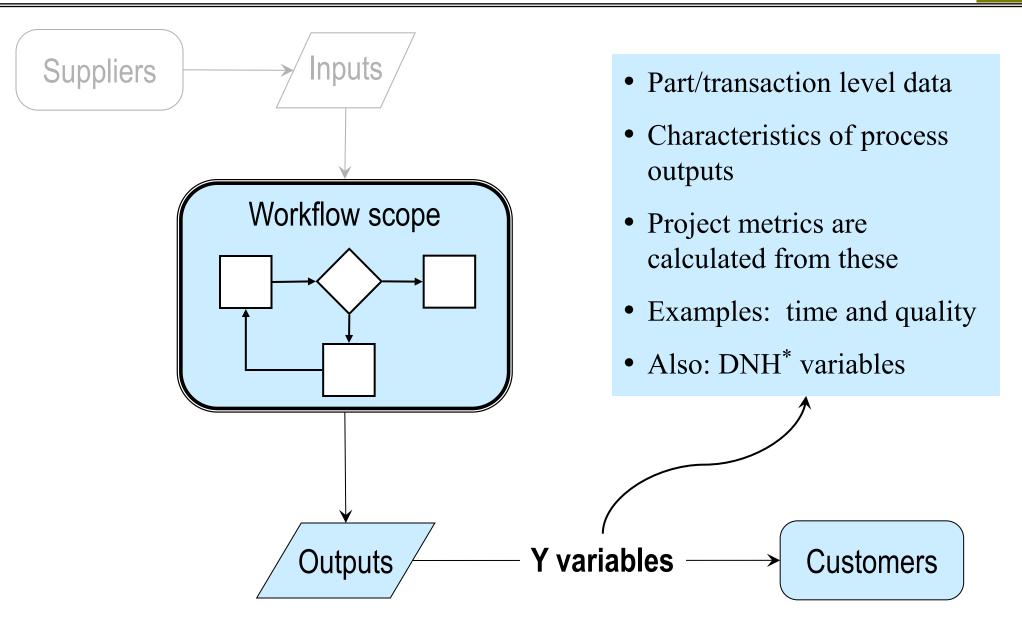
MBDP description for process map

Based on the information in these documents and the process map you created earlier, create a Fishbone Diagram for this project.

Another method for prioritizing X's for data collection is to use multi-voting:

- 1. Count the number of X's
- 2. Divide the total number of X's by 3. Each team member gets that many "votes"
- 3. Each team member decides how they will apply their votes, giving one vote to each X they think is a most likely main contributor to the problem
 - Give a marker to each team member and have them write their votes on the fishbone diagram or list
 - Use a *secret ballot* if there are concerns of undo influence among team members
- 4. Focus data collection on those X's that rise to the top

Y variables

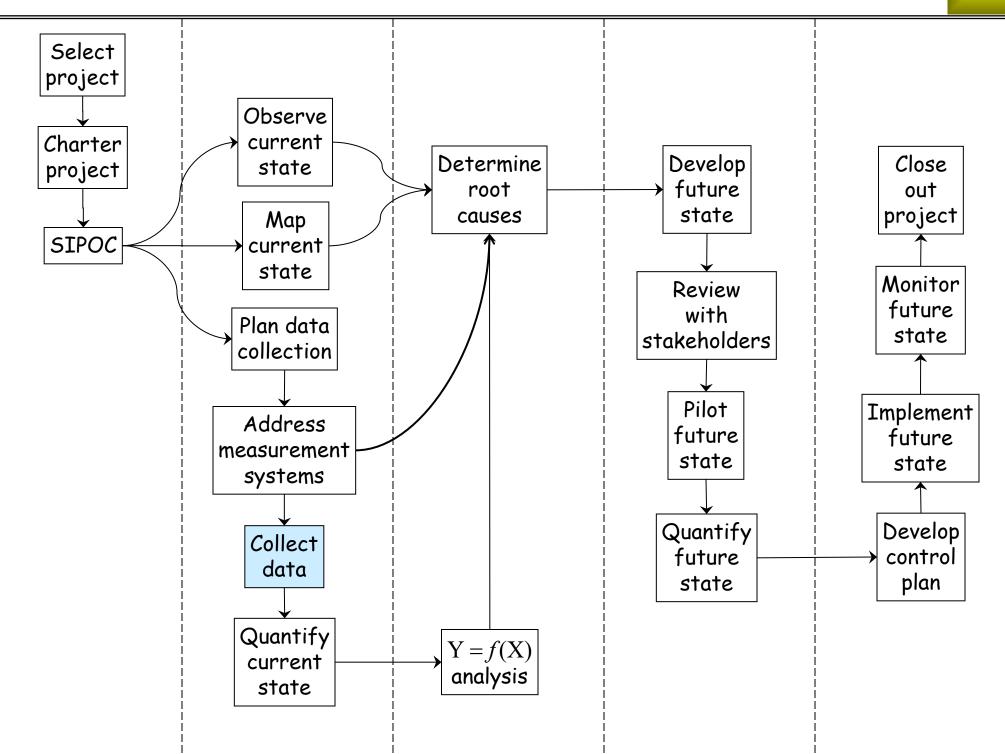


^{*}Do No Harm — not trying to improve them, just don't want to make them worse. These are your secondary project metrics.

Examples of questions to be answered

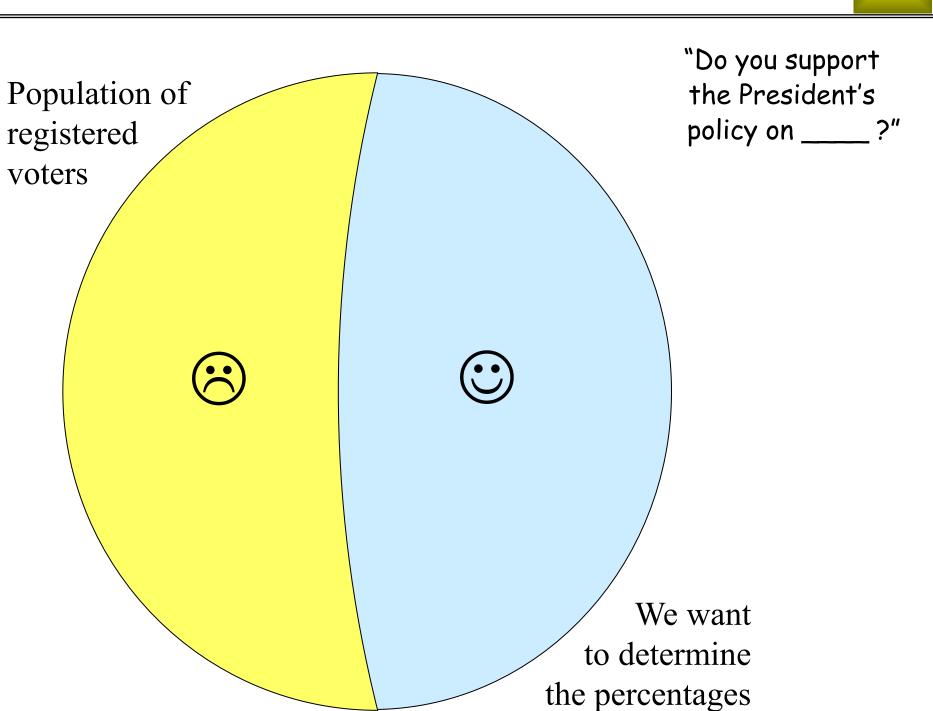
- How, and from what basic quantities, will Y be calculated?
- What measurement system will be used?
- If Y is a lead time, what are the starting and stopping points?
- If Y is pass/fail, what are the possible defects?
- If you are going to count defects per opportunity, how are the opportunities defined?
- If Y is unplanned downtime, how will you record your data: hourly/daily/weekly summaries or event log?
- If there is existing data, can you use it with minor modifications to your operational definition(s)? (Data readily available will jump start your project. Use it whenever possible, even if minor adjustments to the project scope are needed.)

9 Data Collection and Sample Size Calculation

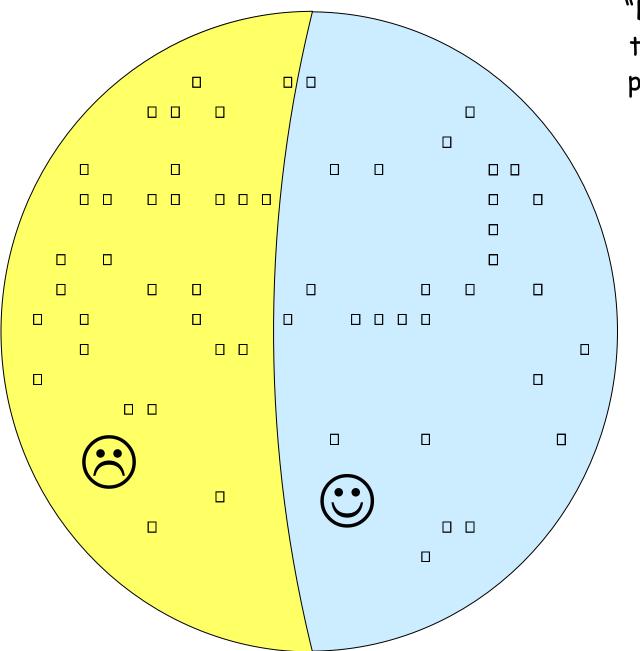


- Calculate project metrics for the current state
- Pareto analysis of defect types, error types, failure reasons, etc.
- Comparisons within the current state (stratification analysis)
- Correlation of X and Y variables
- Use analysis results to help identify root causes

Population	• A specified collection of people or things
Sample	 A subset of a population Usually relatively small Intended to represent the population



The sample must be representative



"Do you support the President's policy on ____?"

- Examples of obvious biases: sample includes only
 - ✓ Democrats
 - ✓ Republicans
 - ✓ Men
 - ✓ Women
 - \checkmark Residents of Wyoming
 - $\checkmark\,$ Convicted white collar criminals
 - $\checkmark\,$ Relatives of elected government officials
- Standard survey sampling technique
 - ✓ All counties are categorized into something like 30 groups ("strata") according to population density
 - Each stratum (group of counties with similar population density) is randomly sampled in proportion to its population
- This is an example of *stratified random sampling*

Decide whether the proposed sample in each case below will be representative of the population. If not, note obvious or possible biases on the slide below.

Population	Purpose	Proposed sample				
(a) Former Enron employees	Opinion on culpability of top Enron executives	Those with the largest retirement accounts, comprising 85% of lost value				
(b) A year, make, and model of car	Surreptitiously determine % with a given defect	Offer a free until 100 cars have been inspected at each US dealership				
(c) ER patients at a hospital last year	Customer satisfaction survey	Those whose last names begin with the letter M				
(d) Lambs born in New Zealand last year	Determine % with "mad lamb" disease	Random sample of each ranch in NZ, proportional to # of lambs				
(e) Registered voters	Opinion on presidential candidate	Generate telephone numbers at random, call those people				

(a)

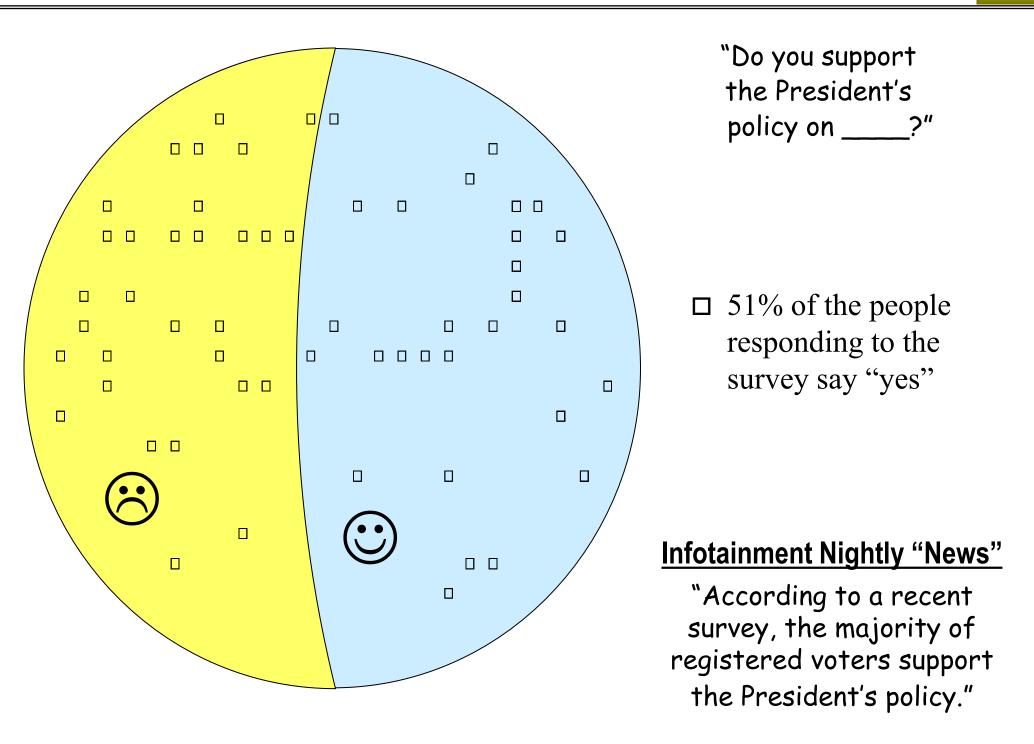
(b)

(c)

(d)

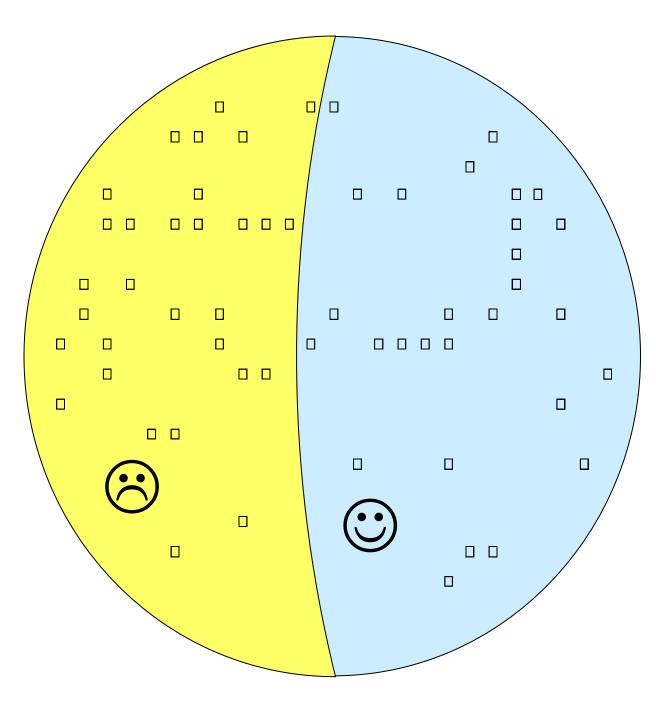
(e)

Incorrect interpretation of survey data



- Suppose the sampling plan was perfectly representative of the population
- Still, we cannot say that what is true in the sample is true in the population
- The sample data does *not* prove that 51% of registered voters agree with the President's policy

Must quantify the uncertainty



"Do you support the President's policy on ____?"

Rational Public Radio

"51% of sampled voters supported the President's policy. The margin of error is ±3 percentage points, so the survey is inconclusive."

- "Margin of error" (MOE) is how we quantify our uncertainty about the population in light of the sample data
- The most we can say: "The percentage of registered voters agreeing with the President's policy is between 48% and 54%"
- The data fails to demonstrate a majority on *either* side of the question

Process

A predetermined sequence of actions and decisions intended to produce a desired outcome. (A way of doing something.)

Manufacturing process

✓ Service process

✓ Business process

 \checkmark Transactional process

✓ Decision process

✓ Design process

For any process, there is an associated *population*

Process sampling (cont'd)



All parts or transactions — past, present, or future— within the project scope

How do we know the extrapolation is valid?

Process

Materials, methods, equipment, operators, environments

Sample

Outcomes the process has produced or is now producing • 100% sampling for a period of time, is the most common method

• What are some situations where 100% sampling is not possible?

• The sample must cover a representative time period

• The sample must capture all *typical sources of variation* (see slide below)

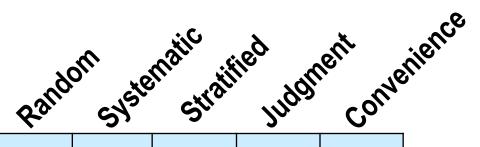
Process participants "Identical" pieces of equipment Time of day, week or month Batches or lots of raw material **Different** suppliers Production lots, work orders, ... Different locations Changing environmental conditions Multiple measurement systems

Random	Items are selected by a random number generator
Systematic	Items are selected at regular intervals
Stratified random*	Items are sampled from homogeneous subpopulations, in proportion to subpopulation size
Judgment	Items are selected using knowledge of the process
Convenience	Items are selected based on cost or ease of access

*Usually considered to be the most representative sampling method.

Exercise 9.2

Check the sampling methods that apply in each case based on the given information.



	7	s,	2.	U
Pulled 10 parts off the high-volume production line at the top of each hour				
Reviewed Enron electricity trades during periods of highest demand				
Used random numbers to select 10% of patient charts for the past year				
Monitored every 1000 th customer service call				
Downloaded invoices with numbers ending in 0 or 5				
Inspected the first 3 parts from each production lot				
Took a sample from the top of each barrel on the top layer of the stack				

• Amount of data: more is better than less

• Time period: longer is better than shorter^{*}

• Capturing all typical sources of variation usually gives an adequate sample size

• You should do a sample size calculation just to make sure

Sample size calculation: opinion poll example

1	The fraction (proportion) of people in the population who would say yes to the survey question if asked.						
φ	We don't know, and will never know, the exact value of ϕ . However, we can get an accurate estimate of ϕ if we collect enough data.						
Sample	The people who respond to the survey. Usually, this is a very small subset of the population.						
	The fraction (proportion) of the respondents who say yes to the survey question. This is our estimate of ϕ .						
Ф sample	We don't know this now, but we will after we get the data.						
MOE	Margin of error: the amount by which ϕ_{sample} could differ from ϕ , based on an established statistical standard of evidence.						
MOL	The most common standard of evidence is called "95% confidence."						
Ν	The number of people who respond to the survey — the <i>sample size</i> .						
Ĩ	The required sample size depends on ϕ_{sample} and the desired MOE.						

In most opinion polls, ϕ_{sample} is assumed to be close to 0.5 when determining sample size. This gives the largest sample size needed to achieve the desired margin of error (MOE). If ϕ_{sample} is not 0.5, the MOE will be smaller, which is desirable. The approximate formula for the MOE (with 95% confidence) is:

MOE = 1.96
$$\sqrt{\frac{\phi_{\text{sample}}(1 - \phi_{\text{sample}})}{N}} = 1.96 \sqrt{\frac{0.5(0.5)}{N}} = \frac{0.98}{\sqrt{N}}$$

We can solve this equation for N:

 $N = (0.98 / MOE)^2$

MOE	Ν
0.05	384
0.04	600
0.03	1067
0.02	2401
0.01	9604

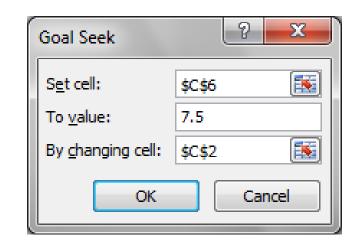
- In process applications, ϕ represents the fraction defective
- In this case, the margin of error on the high side is of greatest interest:

 $\phi_{sample} + MOE_{upper} = Upper bound on \phi$ (with 95% confidence)

- To do a sample size calculation, we must provide two inputs:
 - a) A guess for ϕ_{sample}
 - b) An acceptable upper bound on ϕ (giving the desired MOE, which is the difference between this upper bound and ϕ_{sample})
- Open Student Files \rightarrow calculator sample size \rightarrow % Defective

7

- We think ϕ_{sample} will be close to 0.05 (5% defective)
- If this turns out to be true, we want to be able to say (with 95% confidence) that ϕ is no larger than 0.075 (7.5% defective)
- Enter 1 in cell C2, 5 in C3, and 7.5 in C5
- We want to set cell C6 to 7.5 by changing cell C2
- Select $Data \rightarrow What If Analysis \rightarrow Goal Seek \rightarrow set$ up as shown to the right \rightarrow click OK



	Α	В	С	D	E	F	G	Н	1
1									
2		Sample size (N)	319)←					
3		Guess for sample % defective	5						
4		Defectives in the sample	16						
5		Desired upper bound on population % defective	7.5						
6		Actual upper bound on population % defective	7.50	95	% Confidence level				

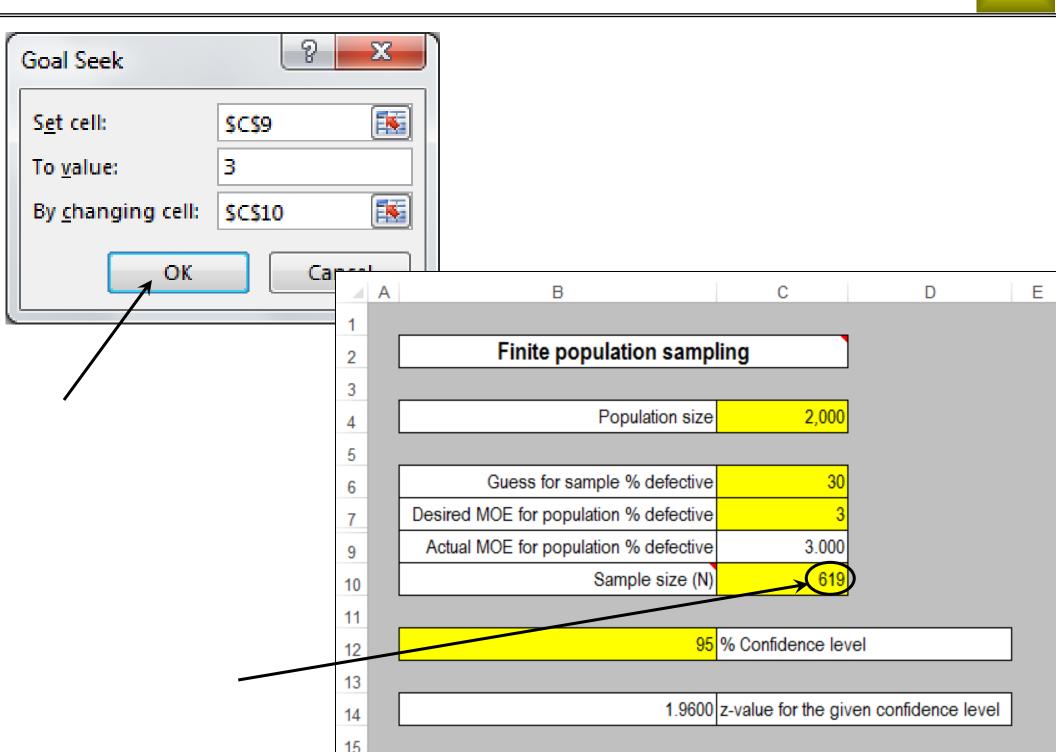
We want to get an accurate estimate of the population % defective. Find the required sample size in the following scenarios.

	Guess for sample % defective	Desired upper bound on population % defective	Sample size
(a)	10	20	
(b)	10	15	
(c)	10	13	
(d)	1	4	
(e)	1	3	
(f)	1	2	

Open Student Files \rightarrow calculator - sample size \rightarrow Finite population sampling

- We want to determine the % defective in a finite population of size 2000
- Enter the values shown below in cells C4, C6, and C7
- We want to set cell C9 to 3 by changing C10 ۲ 🔺 A В С D Data 1 Finite population sampling 2 What If Analysis 3 Population size 2,000 4 Goal Seek 5 Guess for sample % defective 30 6 Desired MOE for population % defective Set up as shown on 7 Actual MOE for population % defective 89.817 the next slide 9 Sample size (N) 10 11 95 % Confidence level 12 13 1.9600 z-value for the given confidence level 14

Finite population sampling (cont'd)

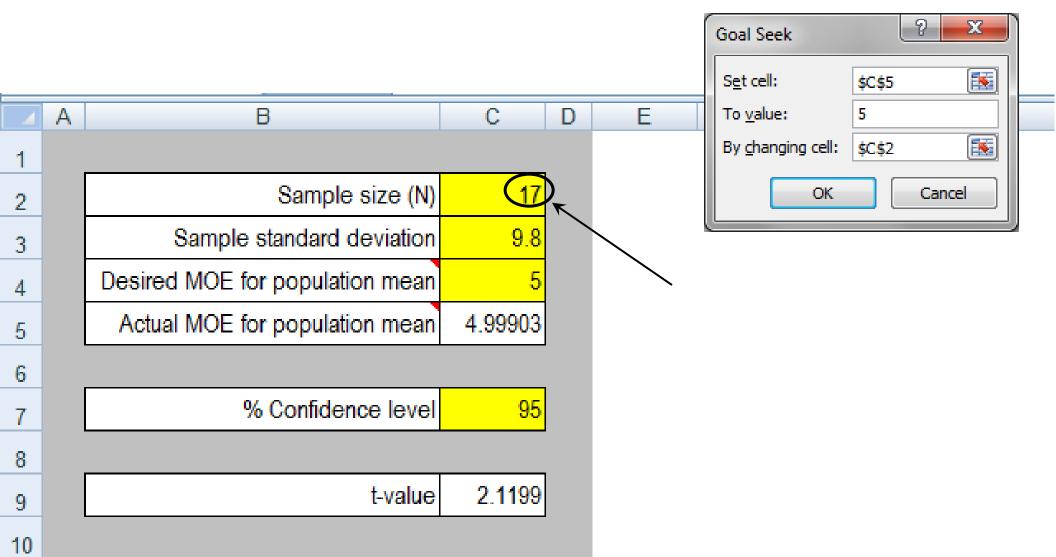


- This requires an estimate of the standard deviation
- Common practice:

✓ Collect a small amount of data, calculate the standard deviation
✓ Do a sample size calculation to see how much more you need
✓ You can also get a rough estimate of the mean from this data

- Suppose our rough estimates are $\mu = 50.4$ and $\sigma = 9.8$
- We want our MOE to be 10% of the mean \rightarrow MOE = .1 * 50.4 = 5
- Go to the sheet *Pop. mean for quant. Y*→ enter the value 2 in cell C2, 9.8 in C3, and 5 in C4
- Select $Data \rightarrow What If Analysis \rightarrow Goal Seek$

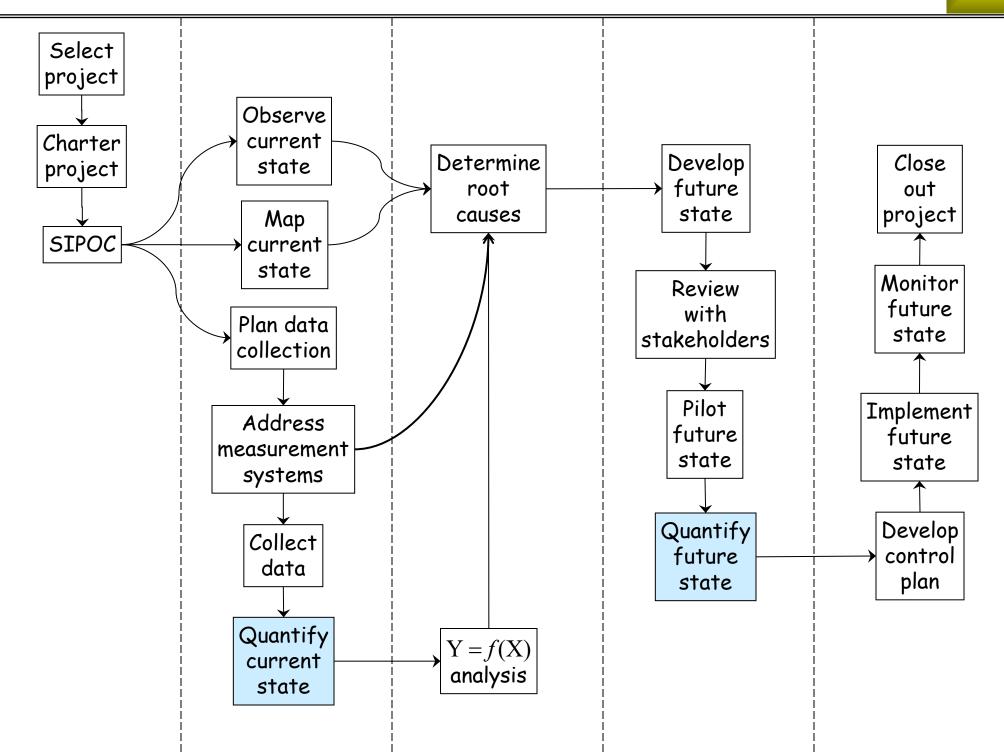
- We want to set cell C5 to 5 by changing cell C2
- Set *Goal Seek* up as shown here, click OK



a) For the previous example, calculate the sample size assuming we want our MOE to be 5% of the mean instead of 10%.

b) Calculate the sample size assuming we want MOE to be 1% of the mean.

10 Basic Statistics and Normal Distribution



Average = (Sum of N numbers)/N

Sample mean = Average of a sample from a population

A set of numbers: 76, 80, 80, 81, 82, 82, 88, 92

$$N = 8$$

Average = (76 + 80 + 80 + 81 + 82 + 82 + 88 + 92)/8= 661/8= 82.6

Minimum = 76

Maximum = 92

Sample standard deviation =

$$(76-82.6)^{2} + (80-82.6)^{2} + (80-82.6)^{2} + (81-82.6)^{2} + (82-82.6)^{2} + (82-82.6)^{2} + (82-82.6)^{2} + (88-82.6)^{2} + (92-82.6)^{2}$$

7

$$= 5.04$$

Average and standard deviation in Excel

	C2		•	(•	<i>f</i> _∞ =AVE	RAGE(A2:	A9)		
	А	В		С	D	E	F		
1	Data			Average	Std. Dev.				
2	76			82.6	5.0)			
3	80								
4	80								
5	81								
6	82			D2	- 		fx =STDE	V.S(A2:A9)	
7	82			Α	в	C	D	E	F
8	88		1	Data	0	Average	Std. Dev.	L	
9	92		2	76		82.6			
			3	80					
			4	80					
			5	81					
			6	82					
			7	82					
			8	88					
			9	92					

Open Student Files \rightarrow anatomy of STDEV

	А	B C	D	E	F	G	Н		J	
1										
2		Data		Average		Difference				_
3		76		82.6		-6.6				
4		80		82.6		-2.6				
5		80		82.6		-2.6				
6		81		82.6		-1.6		C	0 000000	
7		82	_	82.6	=	-0.6		Sum =	0.000000	
8		82		82.6		-0.6				
9		88		82.6		5.4				
10		92		82.6		9.4				
11	Sums of Squares (SS)	54793.0	-	54615.1	=	177.9				
12	Degrees of Freedom (DF)	8	_	1	=	7				
13	Mean Square (MS) [*]	(SS ÷ DF)			25.41				
14	Standard Deviation	(Square r	oot of MS	S)		5.04				
15										

*Also known as Variance

This sheet lays out the calculation of the sample standard deviation (the STDEV.S function in Excel).

The *Data* column contains 8 independent measurements (no constraints among them). We describe this by saying this column has 8 *degrees of freedom* (DFs).

The *Average* column contains a single value, repeated 8 times. We describe this by saying this column has 1 DF.

The *Difference* column is mathematically constrained to sum to 0, so it contains only 7 mathematically independent values. From any 7 values in this column, we can calculate the remaining value. (What is the formula?) We describe this by saying this column has 7 DFs.

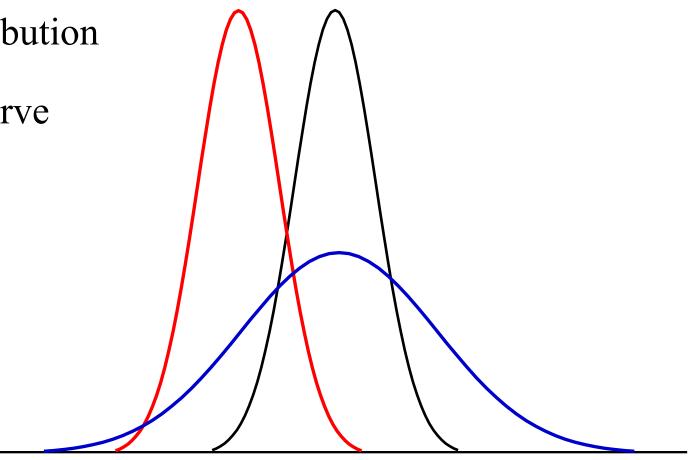
This is why the sum of the squared differences is divided by 7 rather than 8. Dividing by 8 would bias it downwards.

a) Open *Data Sets* \rightarrow *solution properties*. Calculate the average and standard deviation for *Spec grav*. Save your work.

b) Open *Data Sets* \rightarrow *ED patient visits*. Calculate the average and standard deviation of *Visits*. Save your work.

Also known as

- Gaussian distribution
- Bell-shaped curve



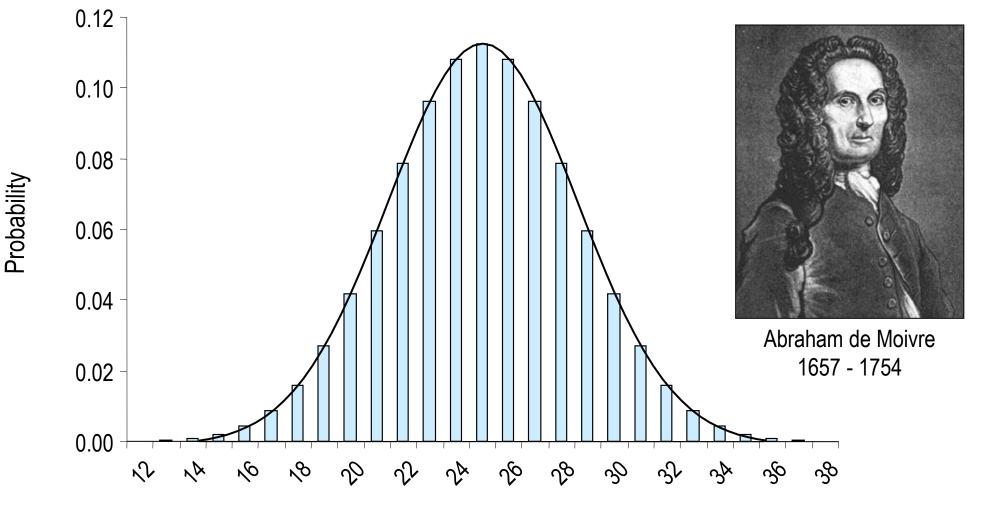
Everyone believes in the Normal curve: experimenters think it is a mathematical theorem, mathematicians think it is an experimental fact. —G. Lippman

The Normal distribution is an abstraction, an idealization, a mathematical construct. At the same time, it has been a device of great practical value in Statistics.

It's called the Gaussian distribution because the German mathematician Carl Friedrich Gauss made important early applications to astronomy in the 1820s. As we will see, it was actually discovered a century earlier by the French mathematician Abraham de Moivre.

I guess life really isn't fair.

As the number of tosses of a fair coin increases, the probability distribution of the number of heads approaches a bell shaped curve.



Number of heads in 50 tosses of a fair coin

Origin of Normal distribution (cont'd)

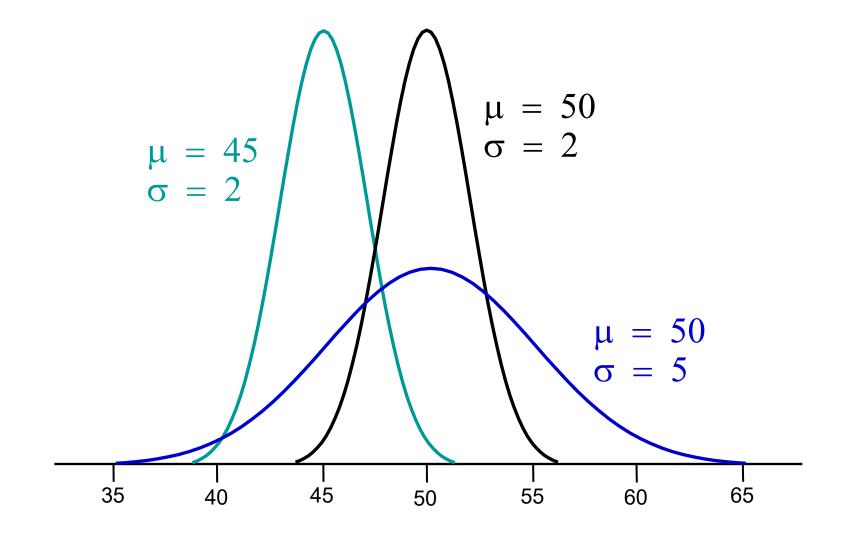
The statistical model for the number of heads in N tosses of a coin is called the Binomial distribution. In 1730, the French mathematician Abraham de Moivre discovered the bell-shaped curve as the limiting form approached by the Binomial distribution as the sample size N increases without bound. He never made any money on his discovery of the Normal distribution, and in fact died a pauper. To add insult to injury, it was eventually named after someone else (Gauss).

Over the next 200 years, de Moivre's discovery was extended far beyond coin tossing. Today, we know that many quantitative measurements are sums of large numbers of small, independent, possibly unobservable contributing factors. Measurements of this type in a stable population will follow the Normal distribution, at least as a good approximation. Statisticians call this phenomenon the Central Limit Theorem.

The Normal distribution is the default population model for quantitative measurements.

 μ = Greek letter mu \rightarrow Population mean

 σ = Greek letter *sigma* \rightarrow Population standard deviation

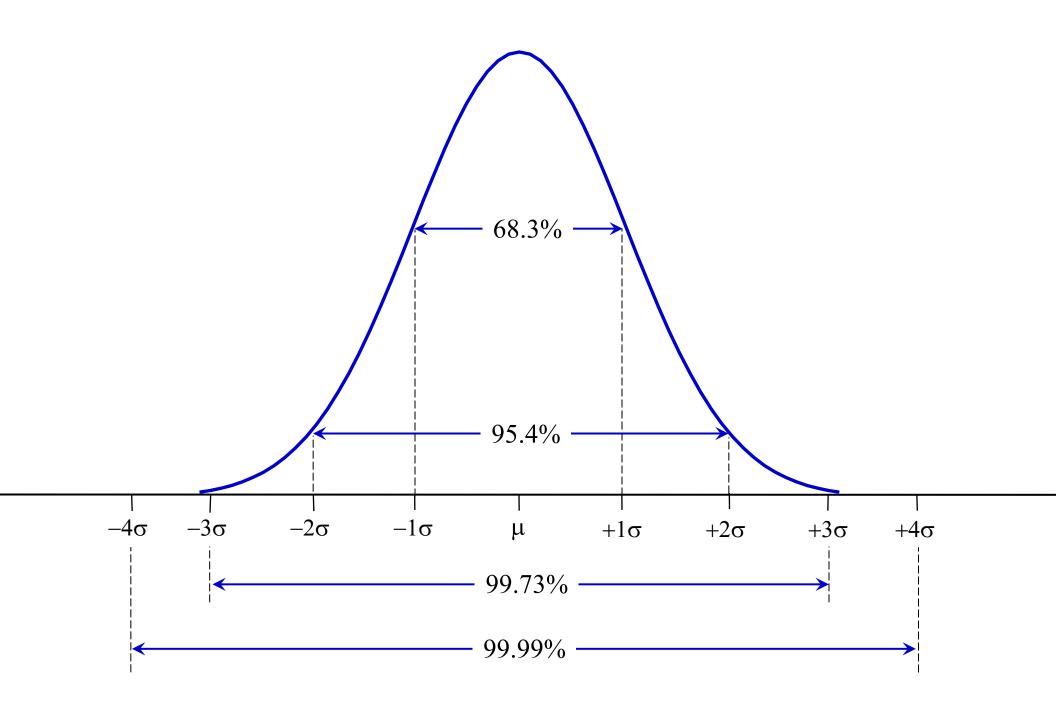


A population model is an equation that can be used to make predictions about a population. When we represent the mean and standard deviation by Greek letters, as above, we are thinking of the mean and standard deviation of the entire population, not just the numbers in our data set. It means we are thinking of the Normal distribution as a population model.

The formula for the bell shaped curve is given below. In this equation, f(y) is the height of the curve above the value y on the horizontal axis.

$$f(y) = \frac{1}{\sqrt{2\pi}} \frac{1}{\sigma} e^{-\frac{1}{2} \left(\frac{y-\mu}{\sigma}\right)^2}$$

You may have been graded "on the curve" at some point in your academic career. Well, this is the curve.

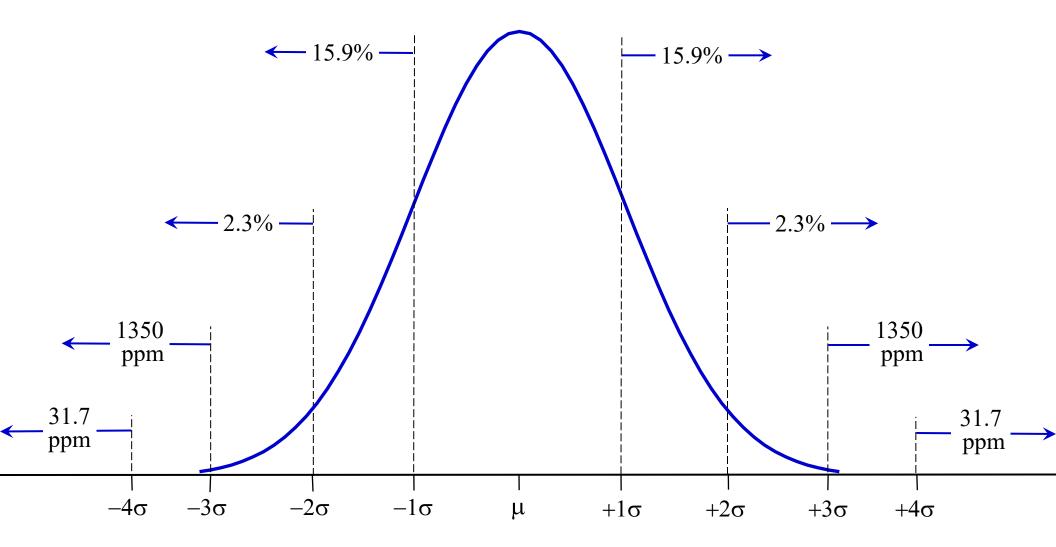


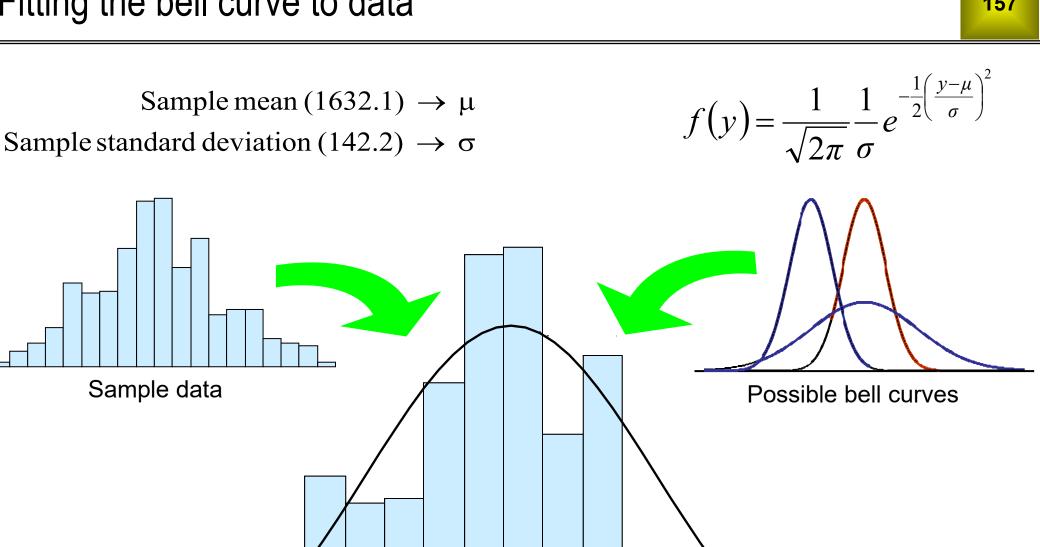
For a Normal population:

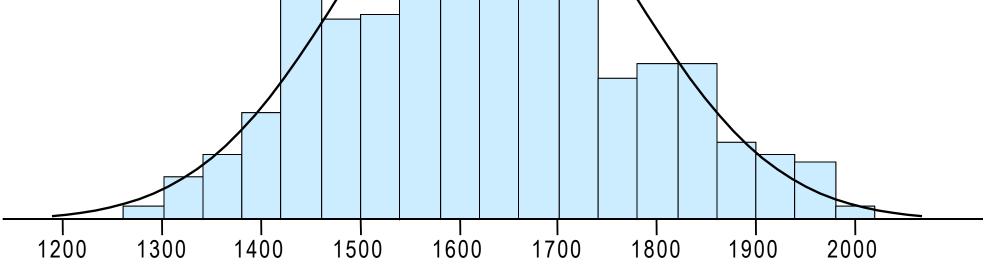
- The 1.960σ limits contain 95% of the population.
- The 2σ limits contain 95.45% of the population.

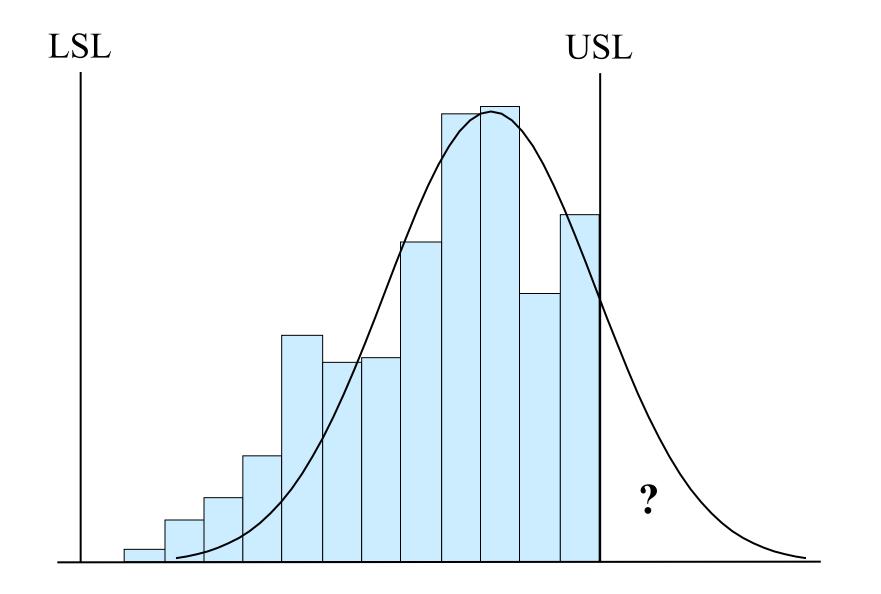
- The 2.576σ limits contain 99% of a Normal population
- The 3σ limits contain 99.73% of the population.

Usually we care mostly about % beyond certain points





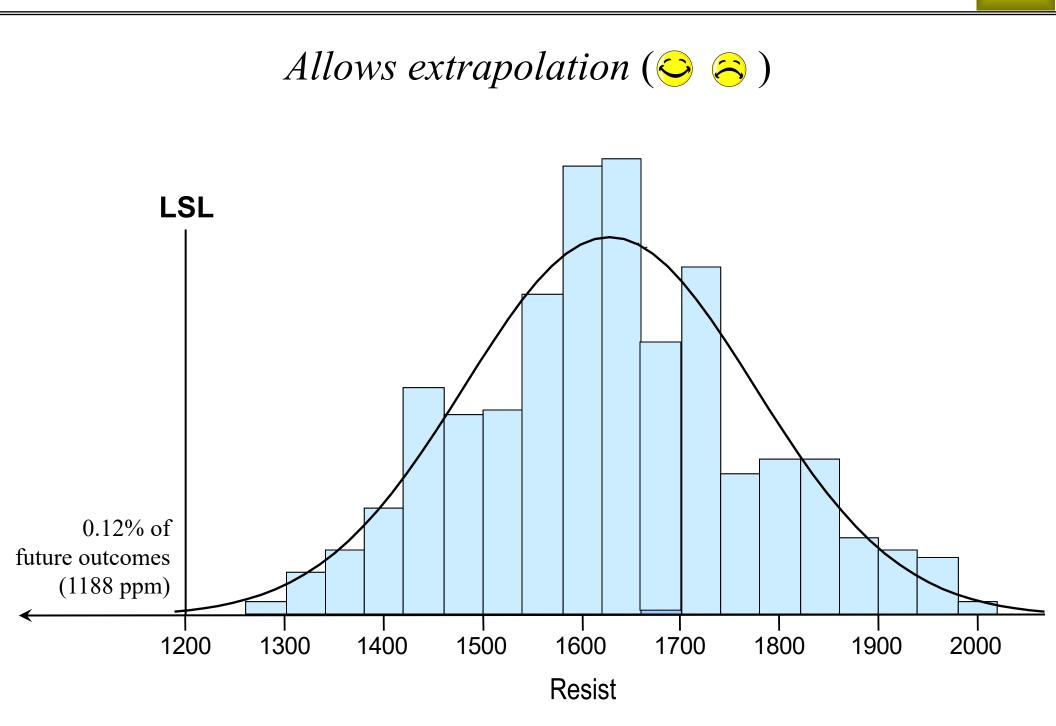




The practice of calculating % defective or DPPM by means of fitted distributions instead of raw data came about historically as a crude but effective way for customers in the aerospace and automotive supply chains to expose the "hidden factories" of their suppliers.

Suppliers would present final inspection data to customers to document their process capability. In the example shown above, the supplier claims 100% yield. When plotted as a histogram, the data mysteriously disappears right at the upper spec limit. This is because parts exceeding the upper limit are either scrapped or reworked to the limit. Often the rework is done by the inspector and not recorded as rework. In many cases, the first pass data is not recorded.

A distribution curve pays no attention to spec limits and will always produce a positive value for % defective or DPPM. This gives an estimate of the supplier's first pass yield. In the example shown above, it is obvious that the first pass yield is far below 100%.



% below 1200

Student Files \rightarrow calculator - Normal distribution

	Α	В		С	D	E	F	G	Н	
1		1. Enter the quantities in the YELLOW cells.								
2		2. The other values a	tre c	alculated	for y	'ou.				
3				\frown	`	_				
4		LSL		/ 1200			LSL	USL	Total	
5		USL				Population % out of spec	0.119	0.000	0.119	
6		Mean		1632.1		Population PPM out of spec	1188.1	0.0	1188.1	
7		Standard deviation		142.2	/					
8				\bigtriangledown						
9	_									
		These calculations of	an 1	be sensitiv	ve to	round-off error. Don't round of	ff the mean			

These calculations can be sensitive to round-off error. Don't round off the mean and standard deviation when you enter them into the calculator. The best thing to do is copy them from a basic statistical summary, then use *Paste Special* \rightarrow *Values*. 161

10

Student Files \rightarrow calculator - Normal distribution

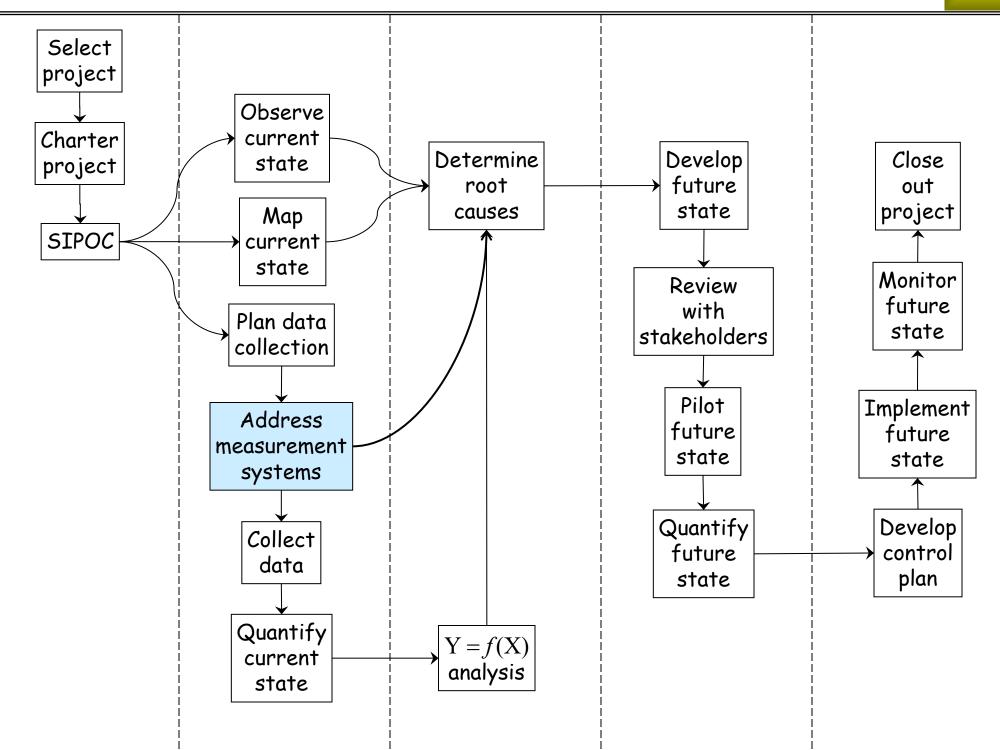
4	А	В	С	D	E	F	G	Н		
1	1. Enter the quantities in the YELLOW cells.									
2	2. The other values are calculated for you.									
3					_					
1		LSL	1200			LSL	USL	Total		
5		USL	2000		Population % out of spec	0.119	0.484	0.603		
6		Mean	1632.1		Population PPM out of spec	1188.1	4838.0	6026.0		
7		Standard deviation	142.2							
8										

These calculations can be sensitive to round-off error. Don't round off the mean and standard deviation when you enter them into the calculator. The best thing to do is copy them from a basic statistical summary, then use *Paste Special* \rightarrow *Values*. **162**

10

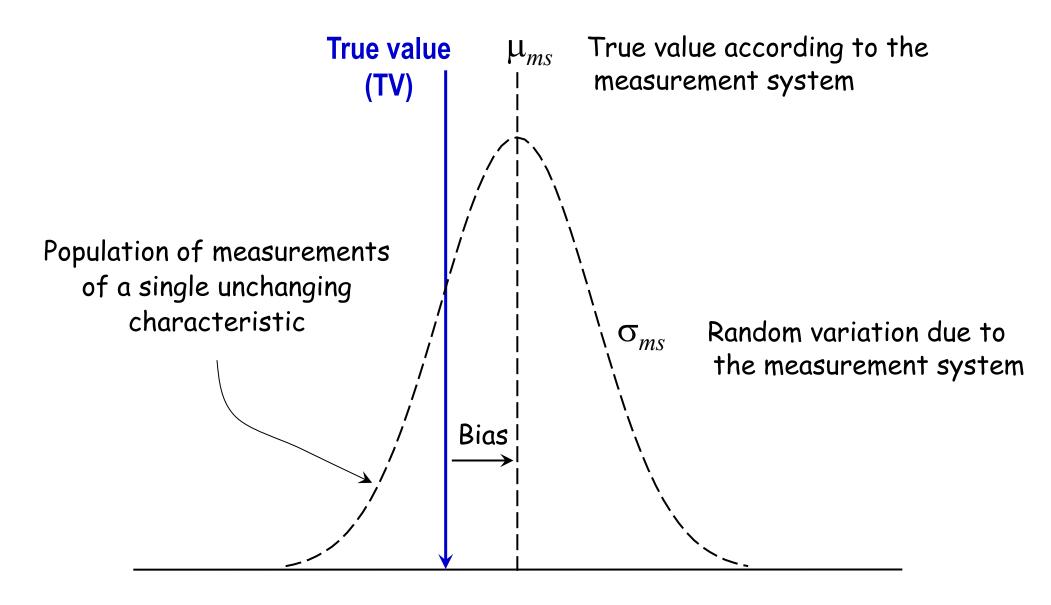
9

11 Measurement Variation



- Population model for measurement variation
- How components of variation add up
- Calculating measurement variation*
- Degrees of freedom

*In the situation where there is only one appraiser.

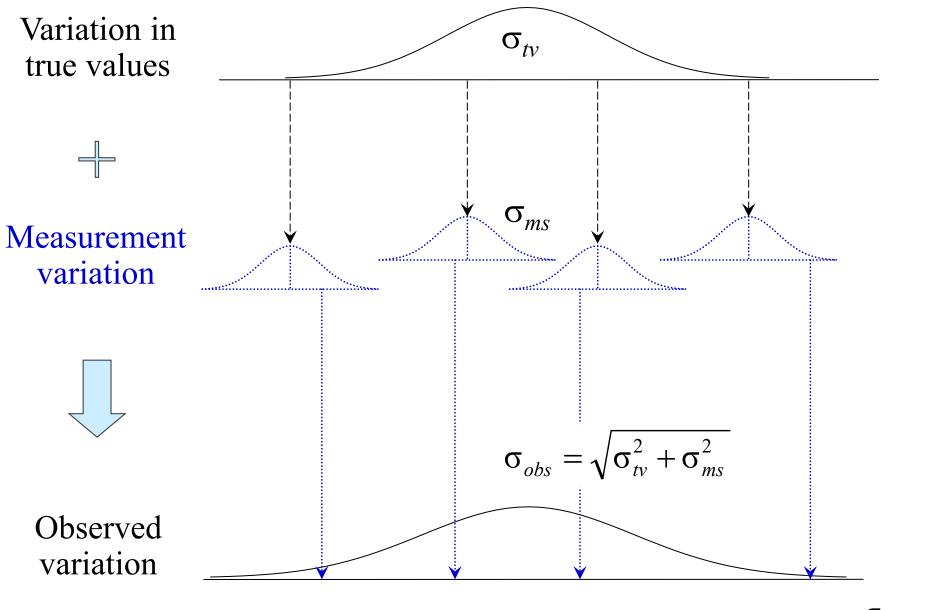


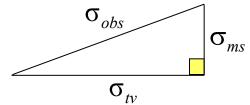
Measurement error = Systematic error (bias) + Random error

- The purpose of calibration is to eliminate gage bias
- Calibration requires standards (measurable items whose true values are known) or a calibrated second gage of higher accuracy
- The primary objective of quantitative measurement system analysis (MSA) is to determine the variation contributed by the measurement system, σ_{ms} , which is *more than gage bias*

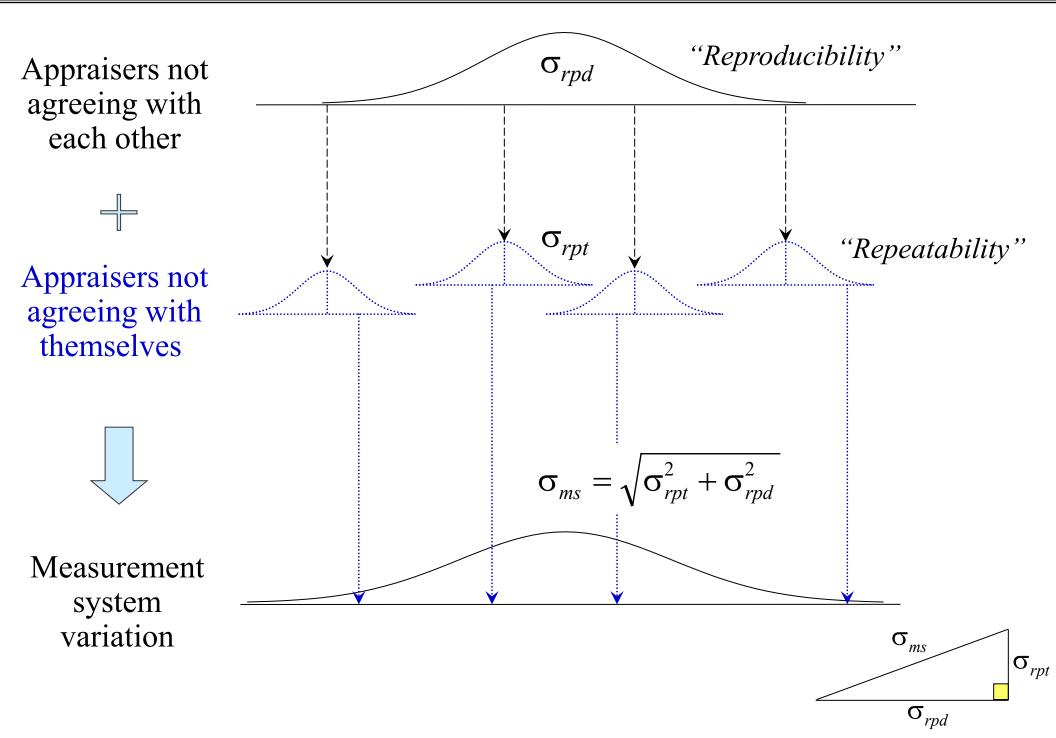
To be clear, calibration is not enough!

- Quantitative MSA does not require standards
- If gage bias is constant during the MSA, the resulting σ_{ms} will be accurate
- If gage bias changes during the MSA, the resulting σ_{ms} will be biased upwards





Components of measurement system variation



STDEV revisited

	А	В	С	D	E	F	G	Н		J	K	L
1			Data		Average		Difference	e				
2			9.61		9.691		-0.081					
3			9.71		9.691		0.019					
4			9.54		9.691		-0.151					
5			9.67		9.691		-0.021					
6			9.75		9.691		0.059					
7			9.49		9.691		-0.201					
8			9.55		9.691		-0.141					
9			9.42	=	9.691	+	-0.271		Sum =	0.00000000		
10			9.58		9.691		-0.111					
11			9.61		9.691		-0.081					
12			9.87		9.691		0.179					
13			9.93		9.691		0.239					
14			9.81		9.691		0.119					
15			9.89		9.691		0.199					
16			9.94		9.691		0.249					
17	Degrees of freedom (DF)		15	=	1	+	14					
18	Sum of squares (SS)		1409.220	=	1408.829	+	0.391					
19	Mean square (MS)		(SS / DF)				0.028					
20	Square root of MS						0.167					
21						. .	↑					
22 23						Sample	standard d (STDEV)	levi	ation			
23							(GIDEV)					

STDEV (cont'd)

The slide above is a screen shot of the worksheet *Observed variation* in *Student Files* \rightarrow *MSA - one appraiser*. This sheet reviews the calculation of the sample standard deviation. In MSA, this is called the "observed variation." In other types of data analysis, it is called the "total variation."

Recap of degrees of freedom (DFs)

- The *Data* column has 15 DFs because it consists of 15 independent measurements.
- The *Average* column has 1 DF because it consists of a single value repeated 15 times.
- The *Difference* column is constrained to sum to 0, so it contains only 14 independent values, so it has 14 DFs.
- DFs have to add up. For example, 15 = 1 + 14.

	A	В	С	D	E	F	G	Н	I J	K	L
1						Part	M	leasuremer	nt		
2	<u> </u>	<u>Part</u>	D	ata		averages		variation			
3		1		9.61		9.656		-0.046			
4		1		9.71		9.656		0.054			
5		1		9.54		9.656		-0.116	Sum =	0.00000000	
6		1		9.67		9.656		0.014			
7		1		9.75		9.656		0.094			
8		2		9.49		9.530		-0.040			
9		2		9.55		9.530		0.020			
10		2		9.42	=	9.530	+	-0.110	Sum =	0.00000000	
11		2		9.58		9.530		0.050			
12		2		9.61		9.530		0.080			
13		3		9.87		9.888		-0.018			
14		3		9.93		9.888		0.042			
15		3		9.81		9.888		-0.078	Sum =	0.00000000	
16		3		9.89		9.888		0.002			
17		3		9.94		9.888		0.052			
18	Degrees of freedom	(DF)		15	=	3	+	12			
19	Sum of squares ((SS)	140	9.220	=	1409.159	+	0.061			
20	Mean square ((MS)	(SS	/ DF)				0.005			
21	Square root of	f MS						0.072			
22 23								↑			
						(σ of me	asurement	variation		
24											

Μ

The slide above is a screen shot of the sheet *Measurement variation*. It lays out the calculation of σ_{ms} when each of 3 parts is measured 5 times by one appraiser.

The *Part averages* column has 3 DFs because it consists of 3 independent values (the part averages).

In the *Measurement variation* column, the values for each part are constrained to sum to 0, so any 4 of them determine the remaining value. There are 3 parts, so there are only $3 \times 4 = 12$ independent values in this column, so it has 12 DFs.

Because the calculation of σ_{ms} involves only 12 independent values, we sometimes refer to σ_{ms} itself as having 12 DFs. The greater the DFs for σ_{ms} , the more accurate it is.

As before, DFs have to add up: 15 = 3 + 12.

	А	В	С	D	E	F	G
1	Part 1	Part 2	Part 3				_
2	9.61	9.49	9.87		Excel data format for MSA with one ap	nraiser	
3	9.71	9.55	9.93		Excel dura formar for more with one ap	pi disci	
4	9.54	9.42	9.81				
5	9.67	9.58	-				
6	9.75	9.61	9.94		Data > Data Analysis > ANOVA Single Factor		
7	L					_	
8				ſ	Anova: Single Factor		
9					Input		
10					Input Range: \$A\$1:\$C\$6 K		
11	Tn	atouatio	ong for		Cancel		
12		structio	•		Grouped By: O Columns		
13	doi	ng the c	analysis		© <u>R</u> ows <u>H</u> elp		
14					Labels in first row		
15 16					Alpha: 0.05		
17							
18					Output options		
19					Output Range:		
20					New Worksheet Ply:		
21					○ New Workbook		
22							
23							
24							
25 26		Sc	creen s	hot o	f the sheet Data format & analysis –		
27					· · · · ·		
28			rile:	Stuc	lent Files \MSA-one appraiser		
20							

173

	А	В	С	D	E	F	G	Н	
1	Anova: Single Factor								
2									
3	SUMMARY								
4	Groups	Count	Sum	Average	Variance				
5	Part 1	5	48.28	9.656	0.00688				
6	Part 2	5	47.65	9.53	0.00575				
7	Part 3	5	49.44	9.888	0.00272				
8									
9									
10	ANOVA								
11	Source of Variation	SS	df	MS	F	P-value	F crit		
12	Between Groups	0.329773	2	0.164887	32.22541	1.5E-05	3.885294		
13	Within Groups	0.0614	12	0.005117					
14									
15	Total	0.391173	14						
16									
17									
18									
19									
20									
21									
22			1 .		1				
23		Scree	n shot	ot the s	sheet D	etault	output		
24							_		
25									
26									
27									

	А	В	С	D	E	F	G	Н	
1	ANOVA: Single Factor								
2									
3	SUMMARY								
4	Groups	Count	Average						
5	Part 1	5	9.656						
6	Part 2	5	9.530						
7	Part 3	5	9.888						
8									
9									
10	ANOVA								
11	Source of Variation	SS	df	MS					
12	Between Groups	0.330	2	0.165					
13	Within Groups	0.061	12	0.005	(o ms) ²				
14				0.072	σ_{ms}	=SQRT(D13)		
15				0.215	$3\sigma_{ms}$	=3*D14			
16									
17									
18									
19		Screen	shot of	the che	et Edi	ted out	nut		
20		JUICEN	5101 01	THE SHE			Pui		
21									
22									
23									

Open file *Student Files* \ *MSA-one appraiser*

Perform the analysis shown in the last three slides.

The value $3\sigma_{ms}$ is the *measurement error* — the amount by which a single measurement could vary (+ or -) from the true value.

• Let: N = sample size of an MSA (total number of measurements)

I = number of items in the MSA (parts, transactions, samples, . . .)



- In the previous example: N = 15, I = 3
- DF for $\sigma_{ms} = N I = 15 3 = 12$

For each scenario below, give the total number of measurements and the degrees of freedom for σ_{ms} .

	Ν	DF
(a) 1 item is measured 15 times		
(b) Each of 15 items is measured 1 time		
(c) Each of 3 items is measured 5 times		
(d) Each of 3 items is measured 10 times		
(e) Each of 15 items is measured 2 times		
(f) Each of 4 items is measured 10 times		
(g) Each of 20 items is measured 2 times		
(h) Each of 8 items is measured 8 times		
(i) Each of 36 items is measured 2 times		

• Let: N = sample size of an MSA (total number of measurements)

I = number of items in the MSA (parts, transactions, whatever)

A = number of appraisers

S = number of *sessions* (measurements per item per appraiser)

• In general: DF for σ_{ms} N – I DF for σ_{rpt} (repeatability) IA(S – 1) DF for σ_{rpd} (reproducibility) I(A – 1)

• Note that the DFs for σ_{rpt} and σ_{rpd} add up to the DF for σ_{ms} (because N = IAS)

Example

• 5 items, 7 appraisers, 2 sessions

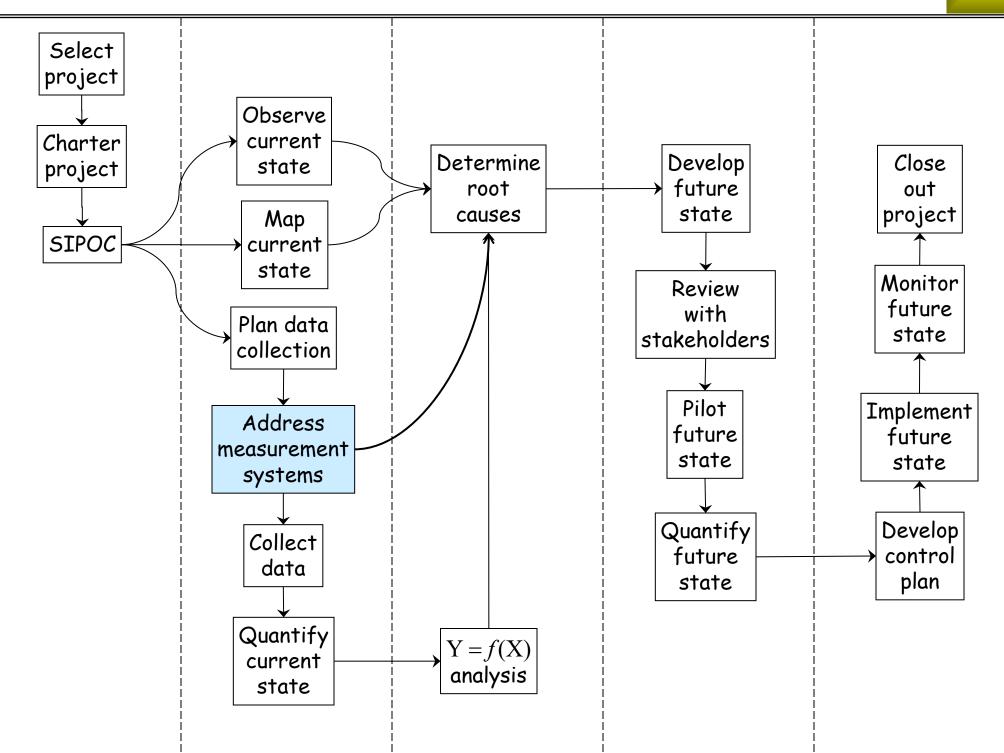
• N =
$$(5)(7)(2) = 70$$

- DF for $\sigma_{ms} = N I = 70 5 = 65$
- DF for σ_{rpt} (repeatability) = IA(S 1) = 5(7)(1) = 35
- DF for σ_{rpd} (reproducibility) = I(A 1) = 5(6) = 30

Exercise 20.3

Repeat these calculations for 10 items, 3 appraisers, and 3 sessions.

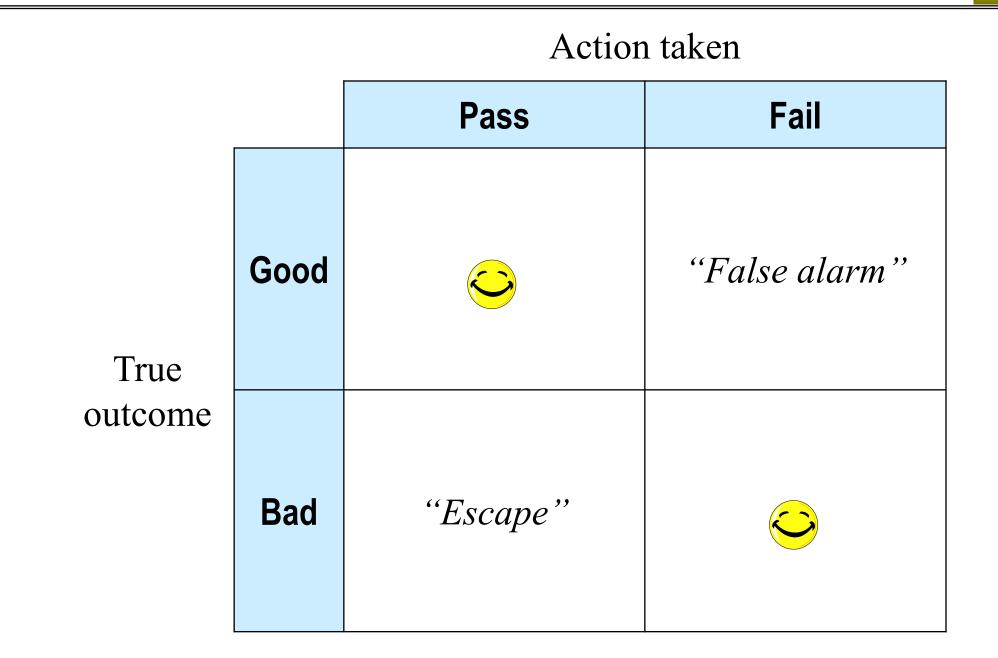
12 Measurement System Analysis



- Gages
- Measurement systems
- Statistical model for measurement variation
- Impact of measurement variation
- Measurement system analysis (MSA)
- Basic assumption for MSA
- MSA for quantitative measurements

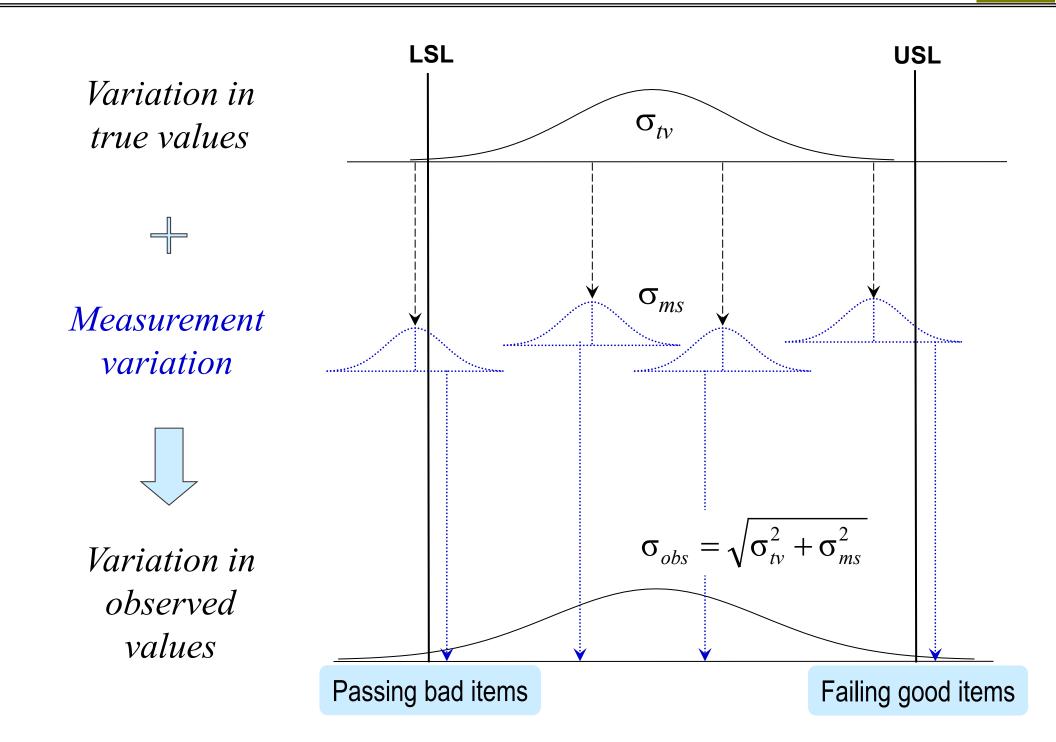
- A gage is a measurement device
- Gages can produce quantitative measurements or categorical classifications
- The people who use the gages are usually called *appraisers*, *inspectors*, or *operators*
- For visual inspections, the appraisers are themselves the gages, but they are not called that
- For automated measurement systems, the appraisers may not play a significant role in producing the results

- A set of gages used to measure defined characteristics of a defined class of objects or events
- The gages produce the same type of data
- For quantitative measurements, the gages provide the same data resolution (x.x, x.xx, x.xx, xx.x, ...)
- The appraisers are part of the system
- The methods and documentation are part of the system
- If there are standards, they are part of the system



Which type of error is more costly? For which is the cost easier to quantify?

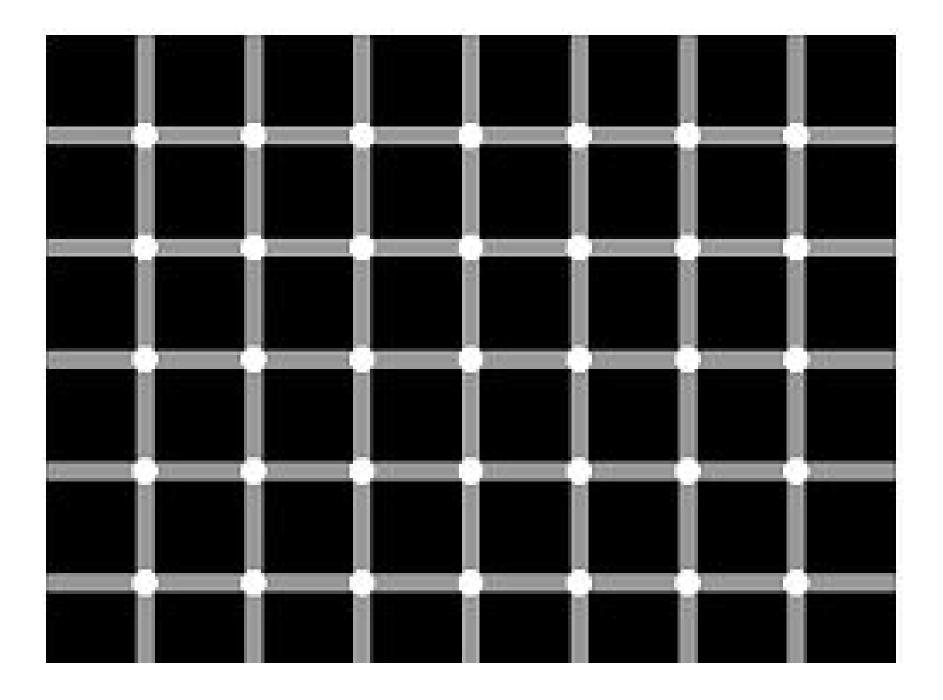
Impact of measurement variation (cont'd)



- Companies should make decisions based on data
- Bad data \rightarrow bad decisions
- One large company estimated the annual cost impact of excessive measurement variation as \$33M
- MSA quantifies and classifies measurement variation
- MSA → corrective action → reduced measurement variation
 → reduced cost

- Improving procedures and fixtures
- Improving gages
- Training appraisers
- Acquiring better gages

Exercise: count the black dots



- MSA requires multiple measurements of "unchanging objects"
- This is not always possible
 - Measurement process may destroy measured items
 - Measurement process may change measured characteristics
 - Measured characteristics may change over time
- In such cases, ad hoc workarounds are used
 - Treat contiguous material samples as the same sample
 - Treat items categorized as "very similar" as the same item
- Workarounds bias σ_{ms} upwards

Measurement system looks worse than it really is

% Tolerance	$100 \times \frac{3\sigma_{ms}}{(\text{USL}-\text{LSL})/2}$	 Most common metric Must have both LSL and USL (usually product or process specs)
% Tolerance LSL only	$100 \times \frac{3\sigma_{ms}}{\mu - LSL}$	 Use when there is only LSL Process mean (μ) should be based on historical data, not the MSA data
% Tolerance USL only	$100 \times \frac{3\sigma_{ms}}{USL - \mu}$	 Use when there is only USL Process mean (μ) should be based on historical data, not the MSA data
% Process	$100 \times \frac{\sigma_{ms}}{\sigma_{obs}}$	 Doesn't require spec limits Process standard deviation (σ_{obs}) should be based on historical data, not the MSA data
Measurement error	$3\sigma_{ms}$	 Has units of the measured characteristic Intrinsic capability, not relative to product or process requirements

10% or less	Excellent
10-20%	Good
20-30%	Acceptable
Greater than 30%	Unacceptable

Open *Student Files* \rightarrow *calculator* - *sample size* \rightarrow *MSA* sheet

Number of items	10	
Number of appraisers	3	
Number of sessions	3	
# Opportunities for appraiser self-agreement	<mark>60</mark>	These should be at least 30 for
# Opportunities for appraiser cross-agreement	20	quantitative, at least 60 for categorical.
Total sample size	90	

- The standard automotive gage study ("10 3 3")
- Not enough opportunities for appraiser cross agreement
- Unnecessarily many opportunities for appraiser self agreement

A better plan

Number of items	15	
Number of appraisers	3	
Number of sessions	2	
# Opportunities for appraiser self-agreement	45	These should be at least 30 for
# Opportunities for appraiser cross-agreement	30	quantitative, at least 60 for categorical.
Total sample size	90	

- Better balance of opportunities for self and cross agreement
- Same total sample size

Best plan, assuming there are actually 7 appraisers

Number of items	5	
Number of appraisers	7	
Number of sessions	2	
# Opportunities for appraiser self-agreement	35	These should be at least 30 for
# Opportunities for appraiser cross-agreement	30	quantitative, at least 60 for categorical.
Total sample size	70	

- Adequate opportunities for self and cross agreement
- Smaller total sample size

1. Perform this sequence for each session:

First appraiser measures all items once Second appraiser measures all items once

Last appraiser measures all items once.

2. The order in which the items are measured should be reversed each time the appraiser changes. Or, better yet, randomize the order each time.

Analyzing a quantitative MSA

- Open Data Sets \rightarrow msa velocity gage
- Measurements are of Drop Velocity
- This is the data format required for continuous MSA in Excel
- The standard analysis requires that every appraiser measures every part the same number of times
- I = 8, A = 3, S = 2
- Was this a well designed MSA?

	Α	В	С	D	E	
1	Session	Part	Oper A	Oper B	Oper C	
2	1	1	9.61	9.54	9.67	
3	1	2	9.49	9.44	9.58	
4	1	3	9.87	9.77	9.89	
5	1	4	9.78	9.66	9.74	
6	1	5	9.89	9.91	9.89	
7	1	6	10.15	10.12	10.16	
8	1	7	9.96	9.87	9.97	
9	1	8	9.80	9.72	9.72	
10	2	1	9.71	9.61	9.75	
11	2	2	9.55	9.42	9.61	
12	2	3	9.93	9.81	9.94	
13	2	4	9.75	9.63	9.72	
14	2	5	10.03	9.84	9.93	
15	2	6	10.31	10.08	10.18	
16	2	7	10.05	9.96	9.97	
17	2	8	9.87	9.74	9.78	
18						

What do the numbers in cell range C2:C9 represent: part variation, measurement variation, or observed variation?

What do the numbers in cell range C2:E2 represent: part variation, measurement variation, or observed variation?

Worked example

- 1. Sort the da the right (th this).
- 2. Data \rightarrow Data **Two-Factor**
- 3. Set up as sh

	A	В	С	D	E
1	Session	Part	Oper A	Oper B	Oper C
2	1	1	9.61	9.54	9.67
3	2	1	9.71	9.61	9.75
4	-	2	9.49	9.44	9.58
5	2	2	9.55	9.42	9.61
6	1		9.87	9.77	9.89
			9.93	9.81	9.94
	-	-			9.74
	2				9.72
	1				9.89
11	2	5			9.93
					10.16
					10.18
					9.97
					9.97 9.72
$\overline{}$	Ca	ncel			9.72
			9.01	9.14	9.10
		elp			
				co curco	r hara
					,
			riig		ns runge
			\searrow		
			of	sessions	s here
	2 3 4 5	1 Session 2 1 3 2 4 1 5 2 6 1 7 2 8 1 9 2 10 1 11 2 Call	1 Session Part 2 1 1 3 2 1 4 1 2 5 2 2 6 1 3 7 2 3 8 1 4 9 2 4 10 1 5 11 2 5 ØK 7 ØK 7	1 Session Part Oper A 2 1 1 9.61 3 2 1 9.71 4 1 2 9.49 5 2 2 9.55 6 1 3 9.87 7 2 3 9.93 8 1 4 9.78 9 2 4 9.75 10 1 5 9.89 11 2 5 10.03 OK 7 9.96 OK 7 9.96 7 0K 7 9.80 9.87 Help Help Pla hig Help Help Pla hig	1 Session Part Oper A Oper B 2 1 1 9.61 9.54 3 2 1 9.71 9.61 4 1 2 9.49 9.44 5 2 2 9.55 9.42 6 1 3 9.87 9.77 7 2 3 9.93 9.81 8 1 4 9.78 9.66 9 2 4 9.75 9.63 10 1 5 9.89 9.91 11 2 5 10.03 9.84 3 10.15 10.12 5 10.31 10.08 7 9.96 9.87 7 10.05 9.96 3 9.80 9.72 6 0K 7 9.80 9.72 3 9.87 9.74

4. Scroll down to the ANOVA table as shown here.

	A	В	С	D	E	F	G
58							
59	ANOVA						
60	Source of Variation	SS	df	MS	F	P-value	Fcrit
61	Sample	1.729748	7	0.247107	103.23	2.37E-16	2.422629
62	Columns	0.096329	2	0.048165	20.12097	7.39E-06	3.402826
63	Interaction	0.028371	14	0.002026	0.846575	0.618209	2.129797
64	Within	0.05745	24	0.002394			
65							
66	Total	1.911898	47				
67							
68							

5. Open Student Files \rightarrow calculator – Gage R&R.

Example (cont'd)

6. Copy the shaded area.

4	A	В	С	D	E	F	G	Н
1	ANOVA							
2	Source of Variation	SS	df	MS				
3	Sample	22.4742	7	3.2106				
4	Columns	84.5409	2	42.2704				
5	Interaction	73.5770	14	5.2555				
6	Within	233.2751	24	9.7198				
7								
8	Total	413.8672	47					
9								
10		σ^2		3σ				
11	Reproducibility	2.3134	19.2%	4.5630				
12	Repeatability	9.7198	80.8%	9.3530				
13	Measurement System	12.0332	100.0%	10.4067		Conv	this area.	
14					Γ P		ANOVA to	ıble.
15	N	48						
16	Items	8						
17	Appraisers	3						
18	Sessions	2						
19								

Example (cont'd)

7. Paste the shaded area below your ANOVA table as shown.

 $3\sigma_{ms} = 0.2179$

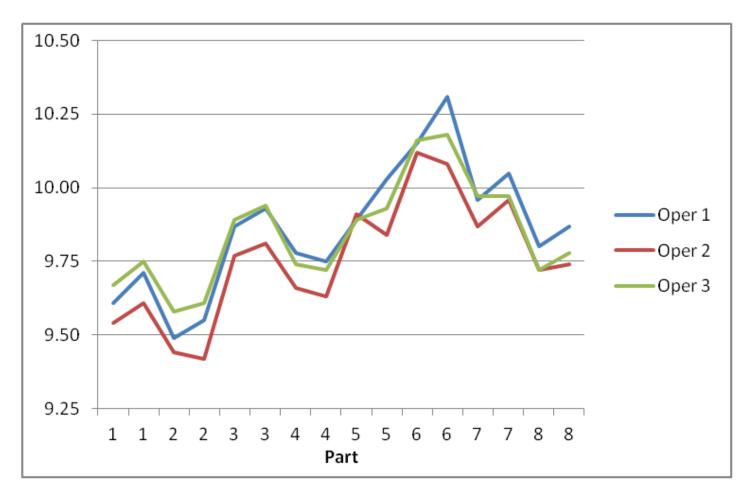
Reproducibility is the dominant component, but not by much.

	4	A	В	С	D	E
5	8					
5	9	ANOVA				
6	0	Source of Variation	55	ď	MS	F
6	1	Sample	1.729748	7	0.247107	103.23
6	2	Columns	0.096329	2	0.048165	20.12097
6	3	Interaction	0.028371	14	0.002026	0.846575
6	4	Within	0.05745	24	0.002394	
6	5					
6	6	Total	1.911898	47		
6	7					
6	8		σ^2		3σ	
6	9	Reproducibility	0.0029	54.6%	0.1611	
7	0	Repeatability	0.0024	45.4%	0.1468	
7	'1	Measurement System	0.0053	100.0%	0.2179	
	2	····				
	3	Ν	48			
-	-					
-	'4	Items	8			
- 7	'5	Appraisers	3			
- 7	6	Sessions	2			
7	7					

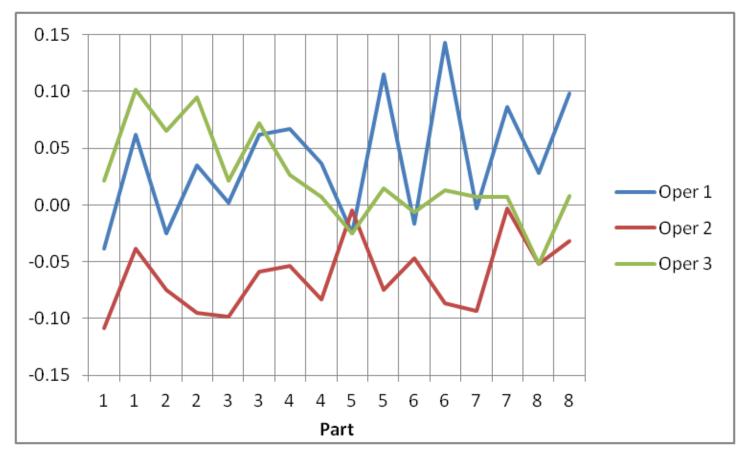
8. For this measurement "Drop Velocity," (USL-LSL)/2 = 1.65. Use Excel to calculate the % *Tolerance* metric.

$$\%$$
Tol = 100 × $\frac{3\sigma_{\rm ms}}{1.65}$ = 13.2%

- 9. Create a line chart of the operator columns by part (Highlight columns > Insert Line Chart)
- 10. This is what a good one looks like. The operator curves are close together and roughly parallel, showing they are getting similar measurements for each part.

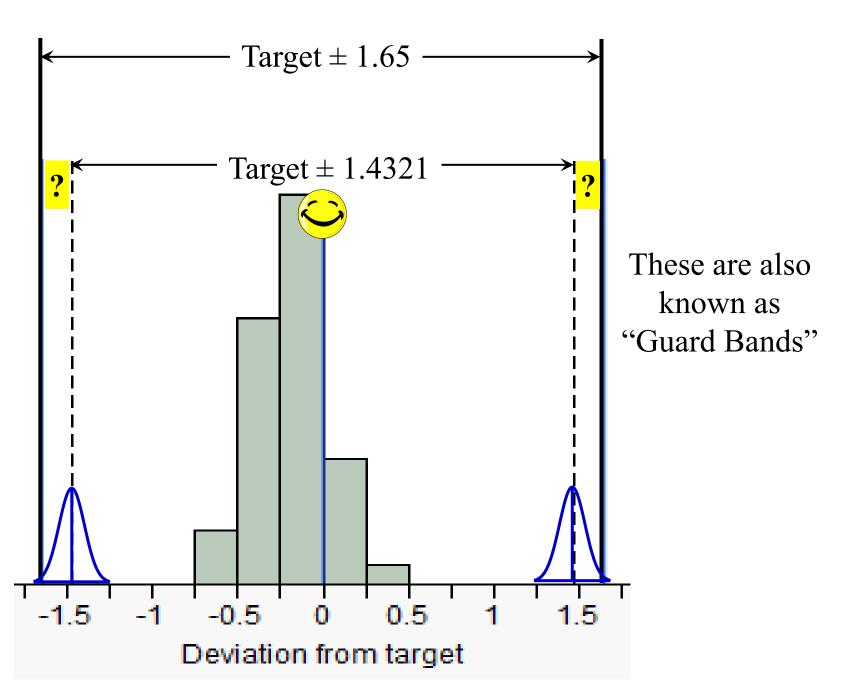


- If part variation is large enough relative to measurement variation, the lines on the previous chart will appear to be superimposed on each other
- The file *Data Sets* \rightarrow *msa velocity gage with charts* gives the calculations for the chart below, which shows the data with the part averages subtracted out.
- This helps you see what's going on with the measurements by each operator, when part variation in the study is large compared to measurement variation.



- In this example, $3\sigma_{ms} = 0.2179$
- For a given measurement *m*, the true value lies in the interval $m \pm 0.2179$ with 99.7% confidence

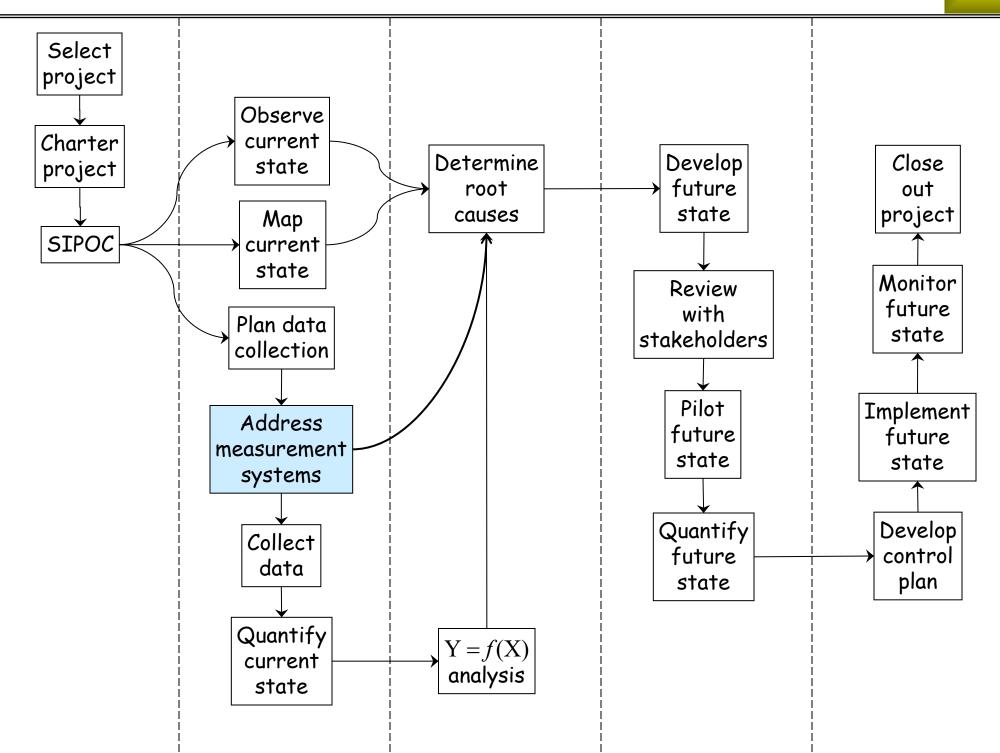
- The tolerance for drop velocity is ± 1.65 (Given on previous slide)
- 1.65 0.2179 = 1.4321
- To be confident that a drop velocity is in spec, it must be within 1.4321 of the target value (see next slide)



Open *Data Sets* \rightarrow *msa calipers*. These are dimensional inspections of PVC extrusions made with a hand-held digital caliper.

(a) The tolerance for this dimension is Target $\pm 0.020''$. Find $3\sigma_{ms}$ and calculate the % *Tolerance* metric. Classify the measurement system as excellent, good, acceptable, or unacceptable.

13 Categorical MSA



- Also known as *Attribute Gage Study*
- Applied most often to pass/fail inspections
- The terms *repeatability* and *reproducibility* are not used in this context
- In this section we assume that the study is based on *standards* (items for which we know the true value)
- Primary objective in this case:

Determine the % agreement with standard (Also known as % correct)

- 1. Choose <u>at least 10</u> items (parts, samples, documents...) to be inspected. There should be roughly equal numbers of items that are clearly passing, borderline passing, borderline failing and clearly failing. Choose an expert appraiser to function as the reference standard.
- 2. If the measurement system has only a few appraisers, include them all in the study. If there are many appraisers, include as large a representative sample as possible.
- 3. Let I = the number of items, A = the number of appraisers, and S = the number of measurements per item per appraiser.
 - The quantity IA(S 1) is the number of independent opportunities for appraisers to agree *with themselves*. It should be at least 60.
 - The quantity I(A 1) is the number of independent opportunities for appraisers to agree *with each other*. It should be at least 60.

It is best to satisfy these requirements by increasing A with I = 10 and S = 2. If this is not possible, increase I.

- 4. If the measurements are taken by devices, and operators have no influence on the results, the devices are the appraisers.
- 5. If devices are used to aid human inspection, combinations of devices and human inspectors should be treated as the appraisers. The ideal is to use all possible combinations of human inspectors and devices. If this is not possible, a DOE matrix with an acceptable number of combinations should be created.

Open *Student Files* \rightarrow *calculator* - *sample size* \rightarrow *MSA* sheet

Number of items	30	
Number of appraisers	3	
Number of sessions	2	
# Opportunities for appraiser self-agreement	90	These should be at least 30 for quantitative
# Opportunities for appraiser cross-agreement	60	data, at least 60 for categorical data.
Total sample size	180	

Best plan if there are only 3 appraisers

Number of items	10	
Number of appraisers	7	
Number of sessions	2	
# Opportunities for appraiser self-agreement	70	These should be at least 30 for quantitative
# Opportunities for appraiser cross-agreement	60	data, at least 60 for categorical data.
Total sample size	140	

Best plan if there are 7 appraisers

1. Perform this sequence for each session:

First appraiser measures all items once Second appraiser measures all items once

Last appraiser measures all items once.

2. The order in which the items are measured should be reversed each time the appraiser changes.

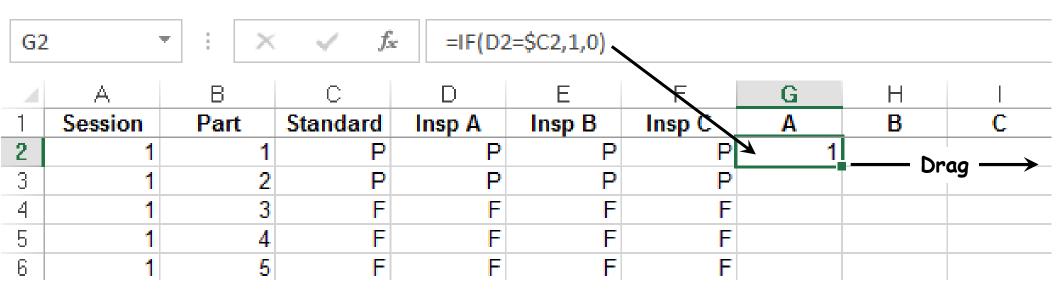
*Same as for quantitative MSA

Analyzing a categorical MSA

- Open Data Sets \ msa passfail
- I = 50, A = 3, S = 3
- Did they follow the best plan for 3 appraisers?
- P = pass, F = fail
- *Standard* gives the correct answer for each part inspected
- The analysis is based on % agreement with the standard

	Α	В	С	D	E	F
1	Session	Part	Standard	Insp A	Insp B	Insp C
2	1	1	Р	P	P	P
3	1	2	Р	Р	Р	P
4	1	3	F	F	F	F
5	1	4	F	F	F	F
6	1	5	F	F	F	F
7	1	6	P	P	Ρ	Р
8	1	7	Р	P	Р	Р
9	1	8	Р	P	P	P
10	1	9	F	F	F	F
11	1	10	P	P	P	P
12	1	11	P	P	P	P
13	1	12	F	F	F	F
14	1	13	P	P	P	P
15	1	14	P	P	P	P
16	1	15	P	P	P	P
17	1	16	P	P	P	P
18	1	17	P	P	P	P
19	1	18	P	P	P	P
20	1	19	P	P	P	P
21	1	20	P	P	P	P
22	1	21	P	P	P	F
23	1	22	F	F	F	P
24	1	23	P	P	P	P
25	1	24	P	P	P	P
26	1	25	F	F	F	F
27	1	26	F	F	F	F
28	1	27	P	P	P	P
29	1	28	P	P	P	P
30	1	29	Р	P	P	P

The first step is to define new columns indicating whether A, B, and C agree or disagree with *Standard* in each case (1 = agree, 0 = disagree)

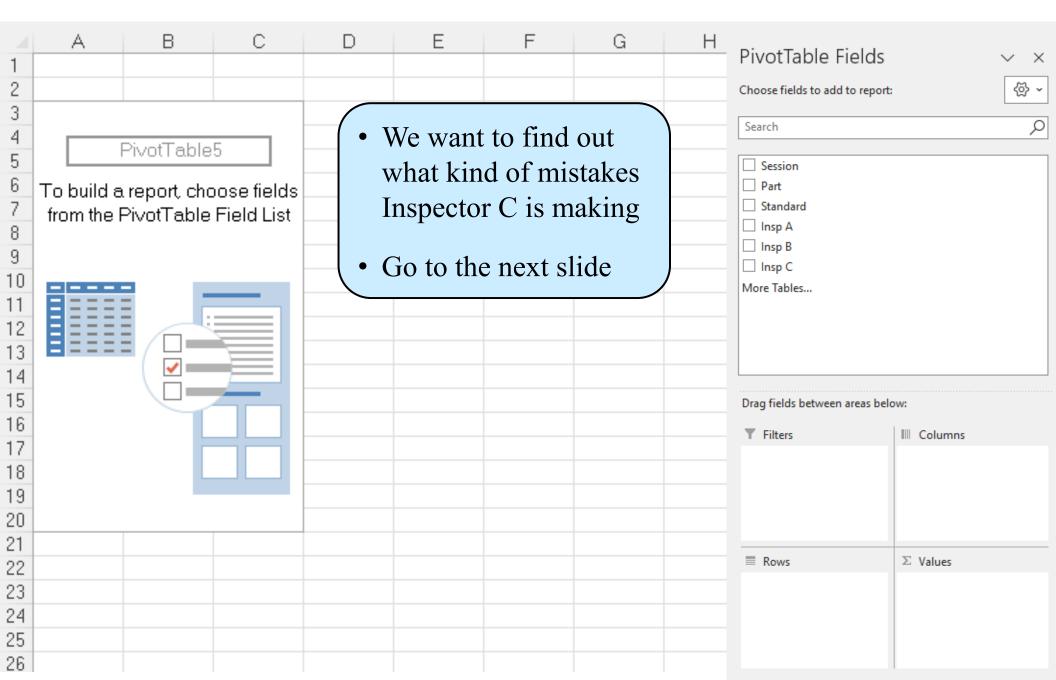


12		÷÷×	√ f _×	=IF(F2	=\$C2,1,0)				
	A	В	С	D	E	F	G	Н	
1	Session	Part	Standard	Insp A	Insp B	Insp C	Α	B	С
2	1	1	P	Р	P	P	1	1	≯ 1
3	1	2	P	Р	Р	P			/
4	1	3	F	F	F	F			
5	1	4	F	F	F	F			
6	1	5	F	F	F	F			
							Double cl	ick /	

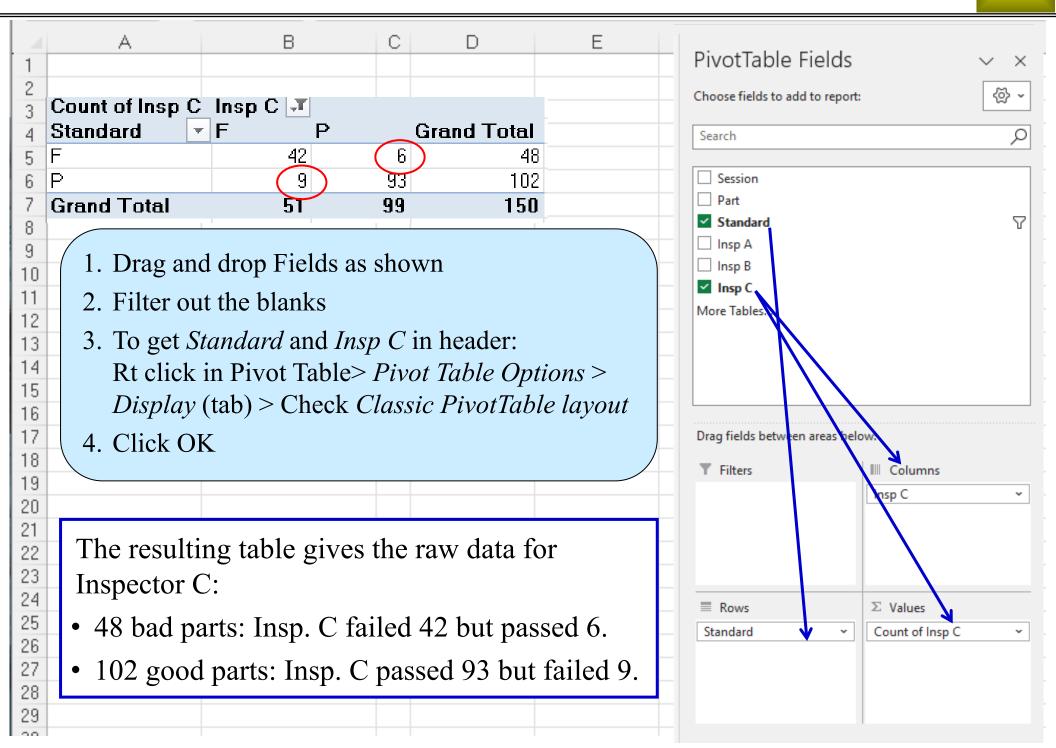
- Use the *Average* function on the *AutoSum* button to get the % agreement with standard for each inspector (cells G152 through I152)
- Use it again to get the overall % agreement with standard (cell J152)
- If improvement is needed, Inspector C is the greatest opportunity

J15	52 🔻	- ÷	× 🗸 fs	=AVEF	RAGE(G152:	152)				
	A	В	С	D	E	F 🔪	G	Н	I [J
1	Session	Part	Standard	Insp A	Insp B	Insp C	A	В	С	
143	3	4	2 F	F	F	F	1	1	1	
144	3	4	3 P	Р	Р	F	X	1	0	
145	3	4	4 P	Р	Р	Р	1	1	1	
146	3	4	5 F	F	F	F	1	1	1	
147	3	4	6 P	Р	Р	Р	1	1	1	
148	3	4	7 P	Р	Р	Р	1	\uparrow	1	
149	3	4	8 F	F	F	F	1	1	1	
150	3	4	9 P	Р	Р	Р	1	1	1	
151	3	5	0 F	F	F	F	1	1	X	
152							94.7%	96.7%	90.0%	× 93.8%
153										

Highlight columns A-F \rightarrow select the *Insert* ribbon \rightarrow select *PivotTable* \rightarrow OK



Example (cont'd)



Example (cont'd)

Count of Insp C Column La Row Labels 💌 F F P Grand Total	P Grand Total 87.50% 12.50% 100.00% 8.82% 91.18% 100.00% 34.00% 66.00% 100.00%	PivotTable Fields ∨ × Choose fields to add to report: & Search ♀
 Inspector C passed 12.5% of the bad parts and failed 8.8% of the good parts 	Click dropdown on Count of Insp C > Value Field Settings > Show Values As > % of row total	 Session Part Standard Insp A Insp B ✓ Insp C ✓ More Tables
• Inspector C needs further training to reduce both types of errors	Value Field Settings Source Name: Insp C Custom Name: Count of Insp C Summarize Values By Show Values As Show values as % of Row Total No Calculation % of Grand Total % of Grand Total % of Row Total % of Parent Row Total Tisp D Insp C	Drag fields between areas below: ▼ Filters IIII Columns Nnsp C IIII Columns Nesp C Values Standard

Open *Data Sets* \rightarrow *msa print samples 1*. These are visual inspections of print samples by 3 inspectors. The standards were determined by a committee of experienced print quality evaluators.

(a) Calculate the % agreement with standard by inspector and overall.

(b) Which inspector offers the greatest opportunity for improvement? Make a pivot table to determine whether the main problem is passing bad samples, failing good ones, or both.

(c) Save your work.

Open *Data Sets* \rightarrow *msa print samples 2*. These are visual inspections of new print samples by the same 3 inspectors after additional training.

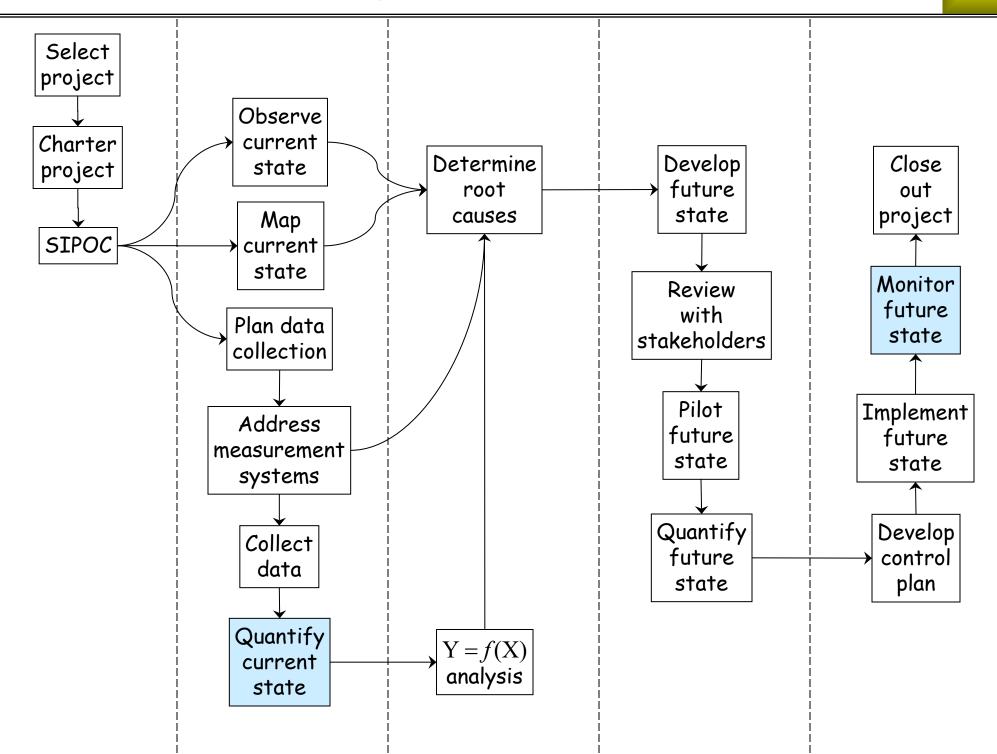
(a) Calculate the % agreement with standard by inspector and overall. Have we improved?

(b) There is something interesting about the data for sample 18 (not row 18). What are the possible explanations? (Sorting by sample number will help.)

(c) It turns out the standard for sample 18 was wrong. Reclassify the standard for sample 18 as passing. What is the % agreement now?

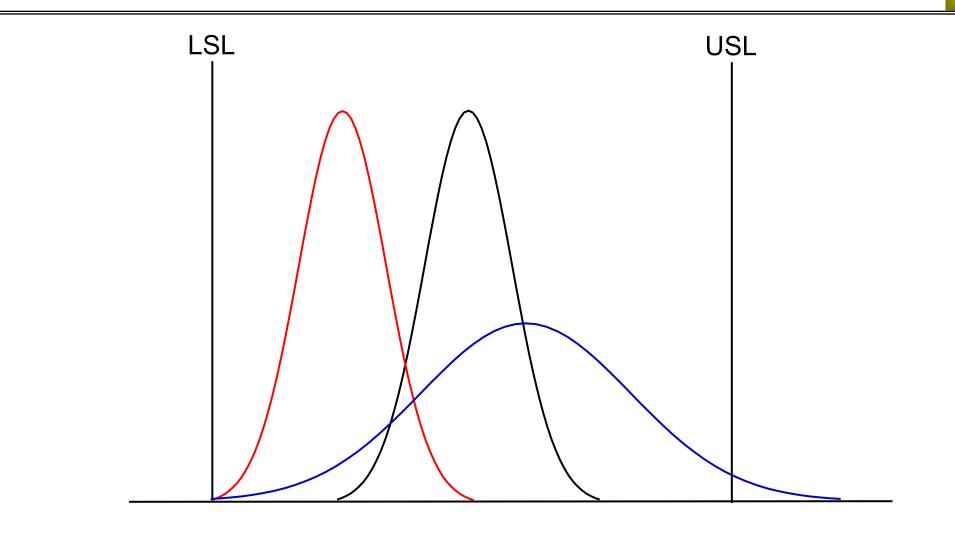
(d) Save your work.

14 Process Capability Indices



- Purpose of Process Capability Indices
- Commonly used indices
- Important assumptions for validity

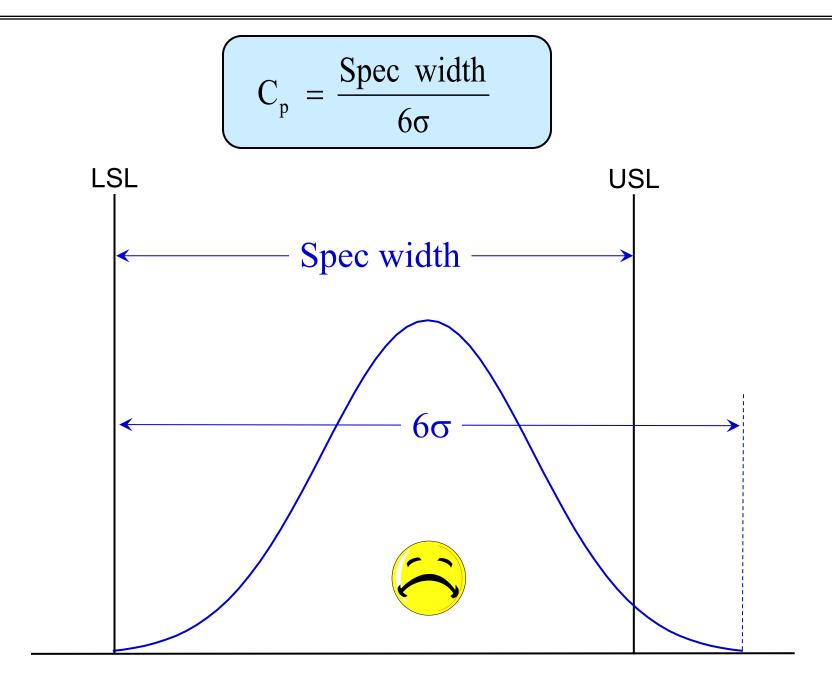
Process capability indices



- Some industries focus on "capability indices" instead of DPPM or DPMO
- These are calculated from the specification limit(s) and a fitted distribution
- Back in the day, the distribution was always assumed to be Normal

- Do your organization's external customers ask for process capability reporting?
- Are there internal requirements or needs for process capability reporting?

 C_p index



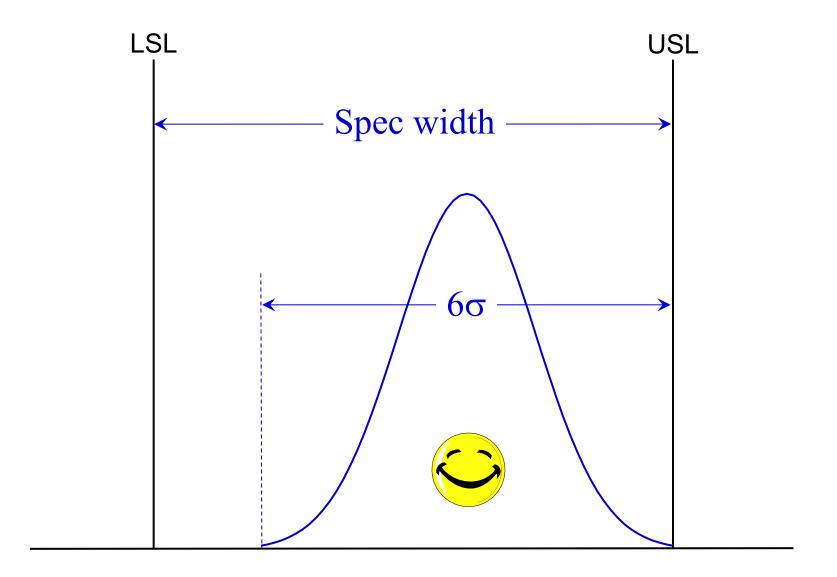
In this case, C_p is less than 1

C_p (cont'd)

The C_p index was historically the first to be used. It is defined as the specification width (USL – LSL) divided by the process spread (6 σ). It set the precedent for capability indices to be defined so that "higher is better."

In the example above, the process spread is greater than the spec width, so C_p is less than 1. It is common for customers to push suppliers to achieve index values of 1.33 or higher for key Y variables.

 C_p (cont'd)



In this case, C_p is greater than 1, but what's a potential problem here?

In the example above, the process spread is less than the spec width, so C_p is greater than 1.

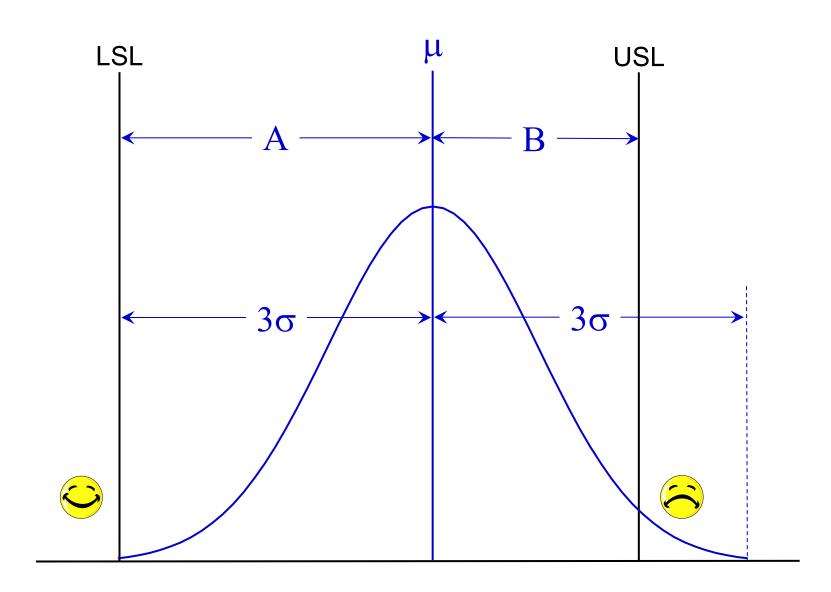
The limitation of C_p is that it doesn't depend on the process mean. If the process mean is equal to the midpoint of the specification range, then C_p is directly related to first pass yield.

If the process mean does not equal the midpoint of the specification range, C_p represents the capability that could be attained by moving the process mean to the midpoint.

 C_{pl} and C_{pu} indices

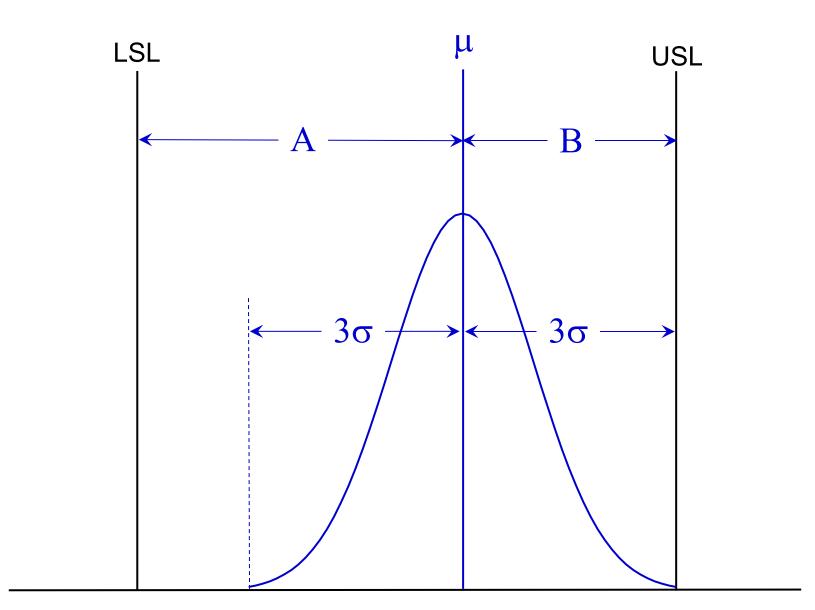
$$\left(C_{pl} = "C_{p} \text{ lower"} = \frac{A}{3\sigma} \right) \qquad \left(C_{pl} = C_{pl} \right)$$

$$\left(C_{pu} = "C_{p} upper" = \frac{B}{3\sigma} \right)$$



The indices C_{pl} and C_{pu} , pronounced " C_p lower" and " C_p upper", were introduced to overcome the deficiency of C_p . They depend on both the mean and standard deviation of the process. If we know both C_{pl} and C_{pu} we can determine the first pass yield of the process.

Like the C_p index, C_{pl} and C_{pu} are defined so that "higher is better." In the example shown above, the main problem is on the high side, with C_{pk} less than 1.



 $C_{\rm pl}$ is greater than 1, $C_{\rm pu}$ is equal to 1

 C_{pk} is equal to 1 in the example above. If improvement is needed, the opportunity is on the high side.

Many people have asked what the k in C_{pk} stands for. To everyone's great disappointment, the k seems to have been chosen arbitrarily and may not stand for anything.

There is, however, a bit of historical trivia that may give us a clue:

- > C_{pk} was first popularized by a man named Victor Kane.
- > Is it possible Victor simply used the first letter of his last name?

- Use C_{pl} if you have only a lower spec limit
- Use C_{pu} if you have only an upper spec limit
- Use C_{pk} (smaller of C_{pl} and C_{pu}) if you have both lower and upper spec limits
- As noted previously, C_p indicates what C_{pk} would be if the process mean were equal to the midpoint of the spec range.
 - > If this is not the case, C_p represents a potential capability.
 - Centering a process at this midpoint may not always be desirable.

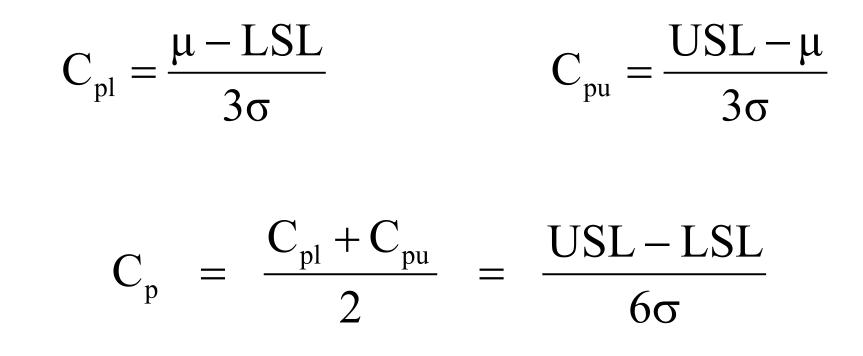
For Process Capability indices to be valid, the following must be true:

- The process is in statistical control (we will cover this during the Control phase)
- The measurement data is normally distributed*
- The sampling method used is representative of day-to-day process operation

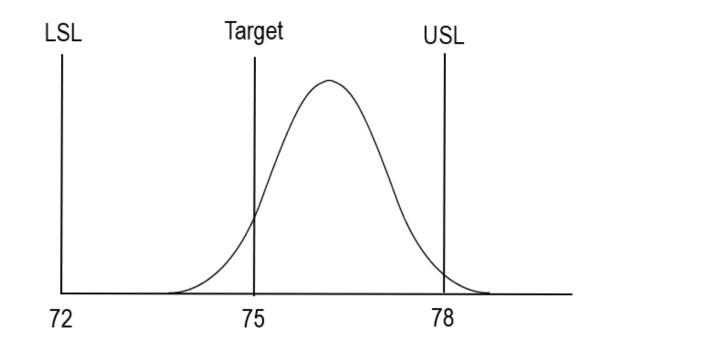
There are times when we want to calculate process capability before the process is under control, for example to set an initial baseline or make a rough prediction.

• The purpose of a process capability study should always be communicated along with the numbers.

*Handling situations when the data is not normally distributed is beyond the scope of this course. Some statistical software packages offer options for calculating Process Capability for non-Normal distributions, along with indices for other special cases.



$$C_{pk} = \min(C_{pl}, C_{pu})$$



For this distribution, the mean = 76 and the standard deviation = 1.

$$C_{p} = \frac{\text{USL}-\text{LSL}}{6\sigma} = \frac{78-72}{6*1} = \frac{6}{6} = 1.0$$

$$C_{\text{pl}} = \frac{\mu - \text{LSL}}{3\sigma} = \frac{76 - 72}{3*1} = \frac{4}{3} = 1.33$$
$$C_{\text{pu}} = \frac{\text{USL} - \mu}{3\sigma} = \frac{78 - 76}{3*1} = \frac{2}{3} = 0.67$$

$$C_{pk} = \min(C_{pu}, C_{pl}) = 0.67$$

(a) Calculate C_p and C_{pk} for a process with mean = 55, standard deviation = 1, USL = 60 and LSL = 50. Sketch the distribution.

(b) Calculate C_p and C_{pk} for a process with mean = 100.20, standard deviation = 0.20, USL = 101.00 and LSL = 100.00. Sketch the distribution.

<u>Capability</u>	How good is this?	<u>Sigma Level</u>
$C_{p} = 1.0$	Marginally capable	3 sigma
$C_{p} = 1.33$	Good	4 sigma
$C_{p} = 2.0$	World-class	6 sigma

The indices C_p and C_{pk} are assumed to be measures of the long-term capability of the process. Therefore,

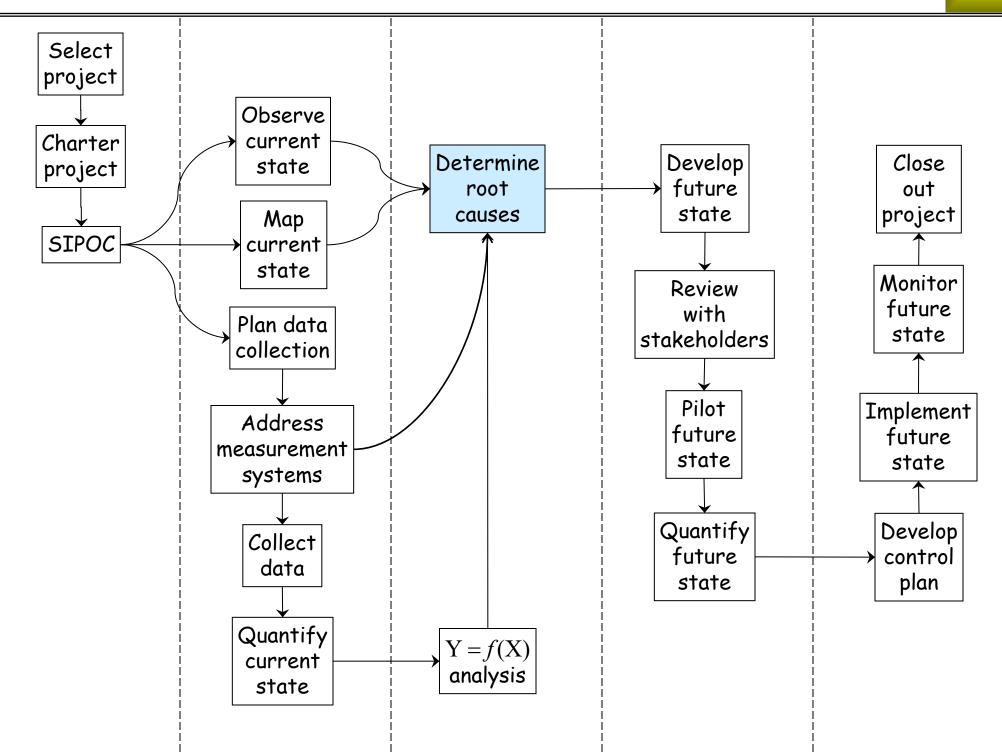
- the data needs to be gathered over a long enough period of time to capture all regular contributors to process variation,
- *and* a sample size of at least 70 is needed, with 100 preferred.

C _p , C _{pk} Value	C _p Fallout (centered)	C _{pk} Fallout (not centered)		
.5	133,620 PPM	66,810 PPM		
.6	71,860	35,930		
.7	35,720	17,860		
.8	16,400	8,200		
.9	6,940	3,470		
1.0	2,700	1,350		
1.1	966	483		
1.2	318	159		
1.3	96	48		
1.33	66	33		
1.4	26	13		
1.5	7	3		
1.6	2	800 PPB		
1.7	340 PPB	170		
1.8	60	30		
1.9	12	6		
2.0	2	1		

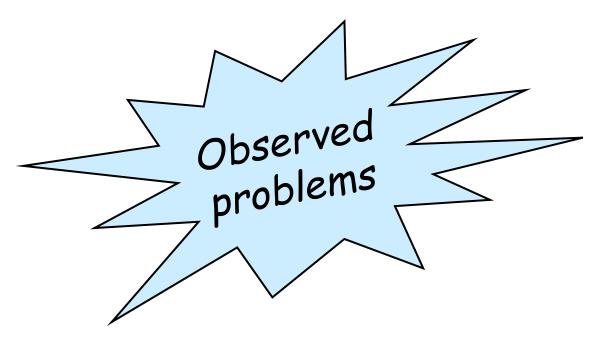
PPM = Parts Per Million PPB = Parts Per Billion Note: 1%=10,000 PPM

Analyze Phase

15 Root Cause Analysis

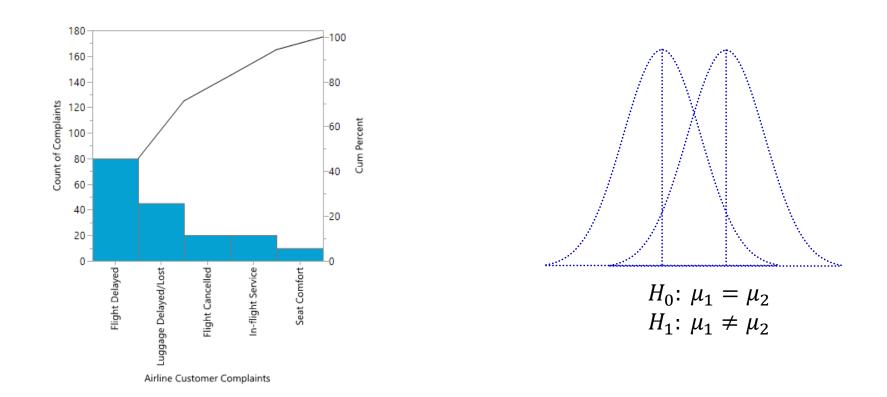


We usually identify problems while mapping and observing the current state during the *Measure* phase



(*a.k.a.* opportunities for improvement)

Analyses such as Pareto Charts point us in the direction of the root causes or critical x's



But, we usually need to dig deeper . . .

Additional tools and techniques to identify root causes:

- Failure Modes and Effects Analysis (FMEA)
- Multi-level Pareto Analysis
 - Five whys
 - Five whys based on Y = f(X)

FMEA can be used in the Analyze Phase to prioritize x's

- It is used at the *beginning* of the Analyze Phase:
 - to identify the inputs that are likely to have a significant impact on the primary metric Y, and to remove from consideration those that are deemed trivial
 - data collection and analysis are required for verification of those failure modes with high RPNs, to validate their significant impact on Y, as FMEA is an opinionbased tool
- Actions for remedying failure modes with high RPNs are *not* discussed or taken in Analyze
- We will learn about FMEA in the Improve Phase, when it is used to evaluate risk and prevent problems before they occur in the proposed process, its original application.

- Principles and steps are the same
 - Purpose and objectives differ
- Rating scale definitions will differ

Design

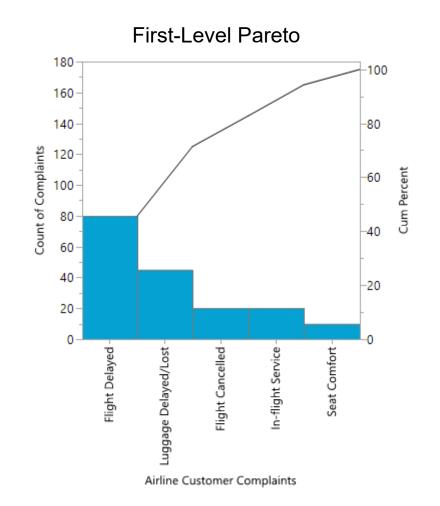
Discover potential problems with the design of the product that will result in safety concerns, malfunctions, or shortened life

Process

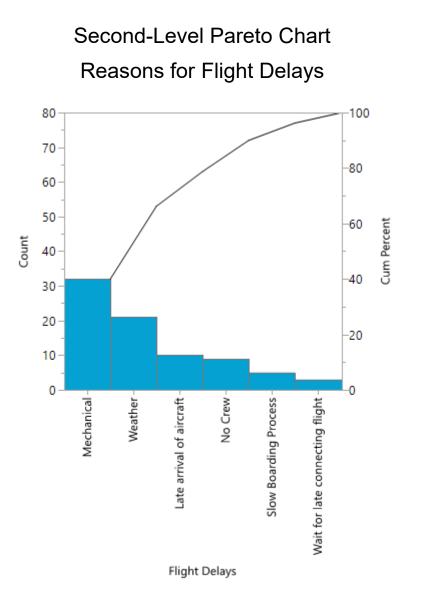
Discover potential problems related to the manufacture of the product that will affect the product, safety, or processing efficiency

We can drill down to root causes using a series of Pareto Charts

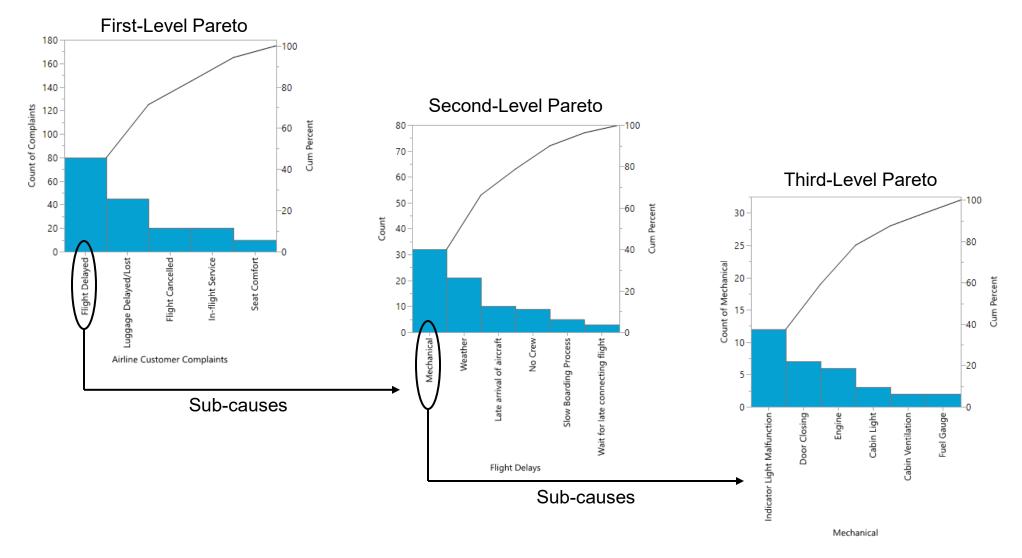
• From a first-level Pareto Chart, we can see which categories are contributing the most to our problem



The highest bar(s) from the first-level Pareto can be broken down further into a second-level Pareto Chart:

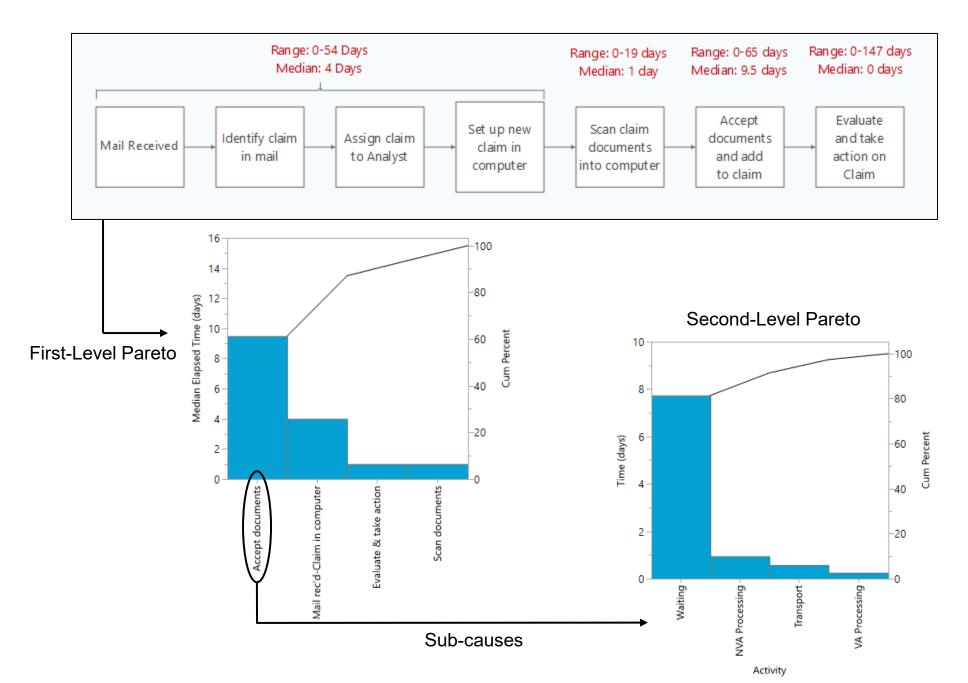


By continuing to drill down, we can determine root causes of most frequently occurring defects.

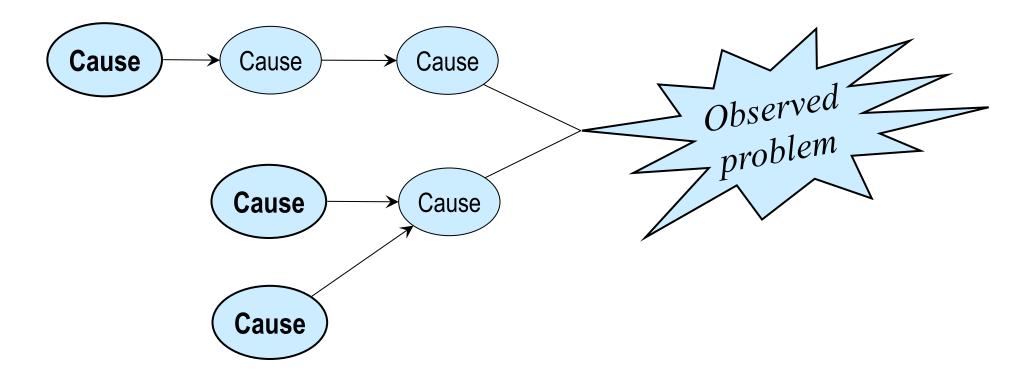


When data is not available for multi-level Pareto analysis, use the firstlevel Pareto Chart with 5 Whys to determine root causes.

Lead time by high-level process step is measured:



• We work our way back to root causes by asking "why" questions



- This process is called "5 whys" because it usually takes no more than 5 questions
- The goal of 5 Whys is to get to a deep, actionable cause.

"The number of accidents in the plant was way up last month"

Do you know what caused the increase?	Workers are slipping and falling in Aisle 7 next to the molding machine.
Why are workers slipping and falling?	There's a puddle of water on the floor.
Where did the water on the floor come from?	It's dripping from the ceiling.
What caused it to start dripping from the ceiling?	A pane of glass is broken in the skylight.
How did the glass get broken?	A tree branch broke the glass during a storm.
How did the tree branch manage to hit the skylight?	The tree it came from was close to the building.

"There's too much scrap in the Coiling Department"

What kinds of defects are causing the scrap?	The vast majority is due to bad welds.
Why do we have so many bad welds?	The welders aren't very good.
Why aren't they very good?	Well, they're hired off the street, and they don't get much training.
You don't hire certified welders?	Are you kidding? We would have to pay them too much.
In that case, why aren't your welders given more training?	I don't know. I guess there isn't enough time. This is the way we've always done it.
Don't they get better as they become more experienced?	Well yeah, but they don't stay in this department long enough for that to help.

Why do they leave this department so soon?	There's another department where welders are used. As soon as there's an opening over there, everybody here applies for it.
Why are they so eager to work in the other department?	For one thing, the working conditions over there are much better. We have the highest accident rate in the company.
Is there another reason?	Over there they pay a dollar an hour more than here.

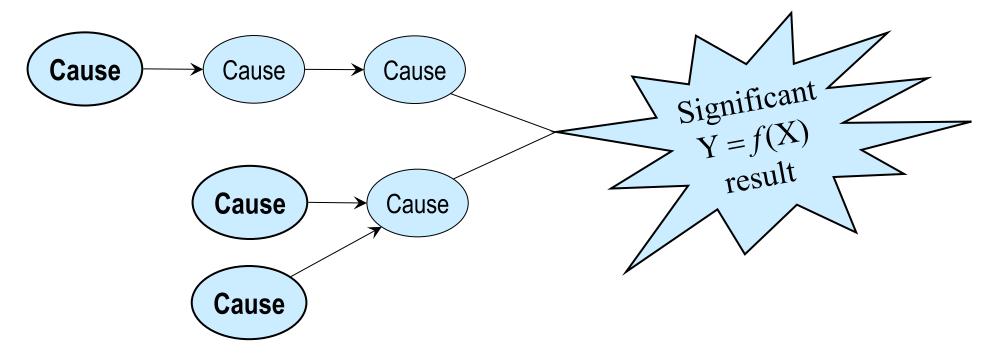
"I was late for work today."

Why were you late for work today?	I overslept.
Why did you oversleep?	My alarm didn't go off.
Why didn't your alarm go off?	The power went out last night.
Why did the power go out last night?	There was a thunderstorm.

What is wrong with this 5 Whys path?

If you get to a non-actionable root cause, back up and try to find a different path to an answer.

- Data analysis provides the basis for penetrating questions
- After we have completed our Y = f(X) analyses, we should interview process participants again to determine the causes of significant comparisons or correlations.
- 5 Whys and Cause and Effect (Fishbone) Diagrams are helpful "interviewing" tools.



- **Q** "There is a significant correlation between dwell time and DPPM. What causes the variation in dwell time?"
- A "The dwell time stretches out when operators are called away to do other things while they're getting ready to mold parts."
- Q "Isn't there an upper spec on the dwell time?"
- A "Yes. The operators are supposed to purge the tank if the dwell time gets too long, but they don't always do that."

Q ...

Whenever we can collect data to verify the root cause found through 5 Whys, that should be done.

- Q "The turnaround time is significantly longer for some account managers than for others. What do you think causes that?"
- A "They don't all use the same quotation preparation process."
- Q "Why not?"
- A "There is no standard process. They have all developed their own way of doing it."

Q ...

Whenever observation can verify the root cause found through 5 Whys, that should be done

- Q "The turnaround time is significantly longer for some business units than for others. What do you think causes that?"
- A "Some of the business units aren't using the automated configuration tool."

Q "Why not?"

Α...

Whenever observation or data collection can verify the root cause found through 5 Whys, that should be done.

- **Q** "The tool development process often results in slow line speeds and overweight material. What causes that?"
- A "The testers slow the line down and increase the weight to get the dimensions on target."
- **Q** "Why do they use weight and line speed instead of other variables?"
- A "They're usually in a hurry. They've discovered that manipulating weight and line speed is the fastest way."

Q ...

Whenever observation or data collection can verify the root cause found through 5 Whys, that should be done.

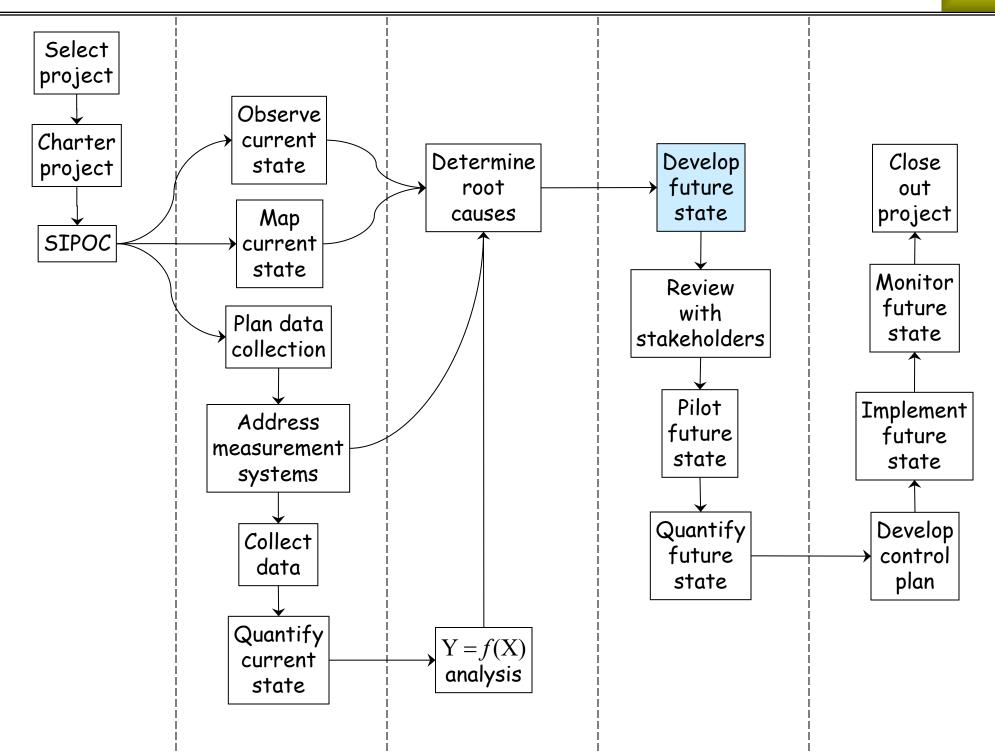
At the conclusion of the Analyze Phase, the team must list those specific root causes or critical x's to be acted upon during the Improve Phase

- Review the analyses completed to:
 - determine those critical x's and root causes that have been validated as significant contributors to unsatisfactory performance in the primary metric
 - ✓ list those that are no longer under consideration

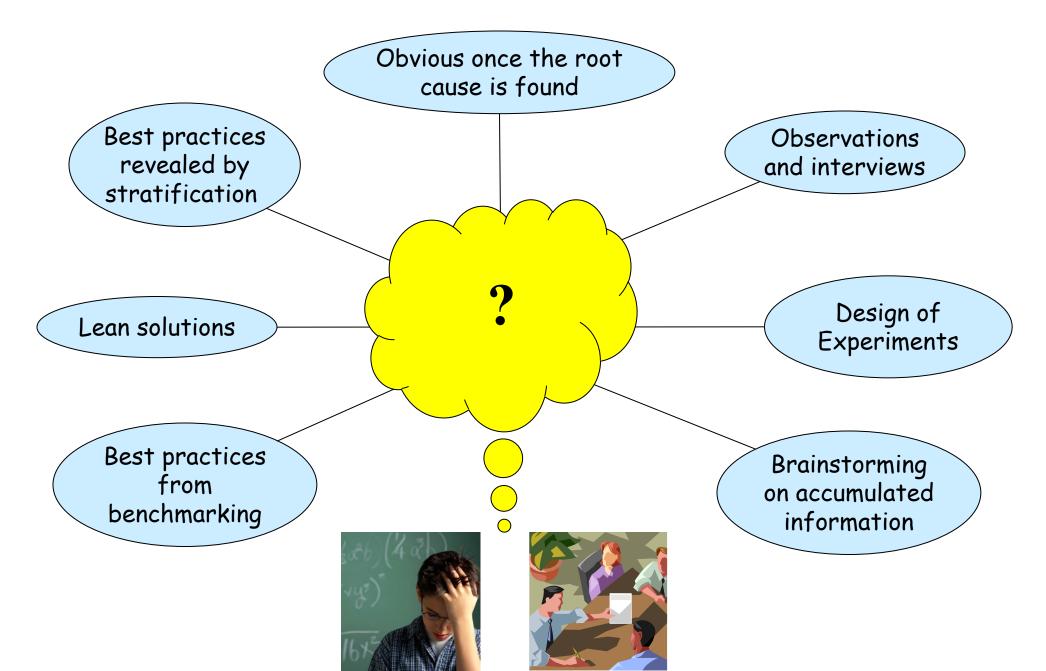
• The team should show the analyses that support their decision on which opportunities to address in the Improve Phase

Improve Phase

16 Developing and Prioritizing Solutions



Solutions come from many sources



Improvement ideas can come from many sources. Some ideas will contribute more to the success of the future state than others. The greater the number of ideas, the greater the probability of discovering successful solutions. The team should generate as many improvement ideas as possible.

The nature of this process is that the initial list gets shorter. Some ideas are discarded along the way, others are retained intact, still others are modified or combined. This process leads to a future state that is likely to be best available within the constraints of the project.

- Technology upgrades
- Lean solutions (we'll learn more about these in the next section of the course)
- Standardization
- Modification of procedures
- "Just do it" solutions that haven't yet been implemented

LSS projects address problems for which solutions are not known. Nevertheless, there are commonly occurring categories.

A common example of technology upgrade would be switching to a better measurement system.

We don't need a LSS project to tell us that Lean is good. But what if the organization lacks consensus on the benefits of these methods? A high priority LSS project that makes significant improvements by applying Lean solutions could help the organization recognize the value of Lean across the board.

The same applies for "just do it" solutions. Everyone knows what needs to be done, but it isn't getting done. A LSS project identifying and quantifying the need for the "just do it" solution might get some high-level attention, cut through the lethargy, and stimulate action on the issue.

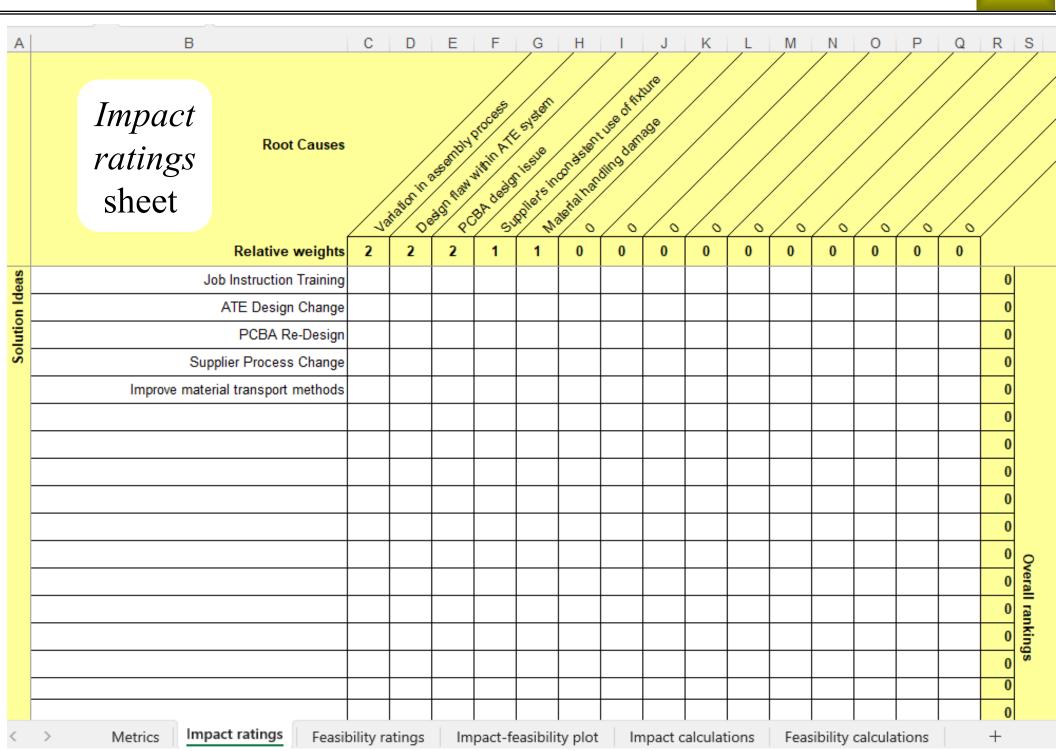
- Uses the impact/feasibility method same as prioritizing projects
- Defines "impact" as addressing the root causes identified by the project team
- Gives the organization a basis for making sound decisions in light of project findings
 - Opportunity to expedite implementation of solutions with high impact or high feasibility
 - ✓ Opportunity to postpone implementation of solutions with low impact and low feasibility

- 1. Open Student Files \rightarrow blank C&E matrix impact & feasibility.
- 2. In the *Metrics* sheet, change *Impact metrics* to *Root causes*.
- 3. List your prioritized root causes and relative weights (overall rankings).
- 4. List your feasibility metrics and relative weights.
- 5. Go to the Impact ratings sheet, change Items to be ranked to Solutions.
- 6. List the solutions you wish to rank.
- 7. Rate each solution for impact on each root cause.
- 8. Go to the *Feasibility ratings* sheet, rate each solution for each feasibility metric.
- 9. Go to the sheet *Impact feasibility plot* to evaluate the results.

Root causes of Long Lead Time

B	С	DE		F (G H I J
Root Causes	Relative weights	Feasibility n	netrics Relati	ive weights	Metrics
Variation in assembly process	2	Inexpens	sive	2	
Design flaw within ATE system	2	Fast		2	sheet
PCBA design issue	2	Easy	K	1	
Supplier's inconsistent use of fixture	1				These are common
Material handling damage	1				feasibility metrics,
					but you can define the metrics and weights to suit your own situation.
ATE = Automated Test Equipment	Feasibility ratings	Impact-feasibility plot	Impact calculations	Feasibility c	Root causes determined in Analyze Phase. Weights indicate relative impact of cause on project metric

Prioritizing solutions (cont'd)



272

Prioritizing solutions (cont'd)

A	В		С	D	E	F	G	Н		J	K	L	M	N	0	P	Q	R	S
	<i>Feasibility</i> <i>ratings</i> sheet	Feasibility metrics	m	afteria ve	* 5	Set o	////	////	///	///	///	///	/	0	/	///	0		
		Relative weights		2	1	0	0	0	0	0	0	0	0	0	0	0	0		
on Id		ATE Design Change																0	
Solution Id		PCBA Re-Design																0	
So	Sup	plier Process Change																0	
	Improve mater	ial transport methods																0	
		0																0	
		0																0	
_		0																0	
-		0																0	
-		0																0	
-		0																0	Ove
-		0																0	Overall rankings
-		0																0	rank
-		0																0	
-		0																0	
-		0																0	
-		0 0																0 0	
_					6								1						
<	> Metrics I	mpact ratings Fea	asidilit	y ratin	gs	Impac	t-feasi	bility p	lot	Impa	ct calci	ulation	s F	easibil	ity calo	ulatio	ns	-	⊢

Prioritizing solutions (cont'd)

A	в	С	D		E			F		(G		Н			I
Solution Ideas	Tag	Impact	Feasibility													
Job Instruction Training	А	0	0		1											
ATE Design Change	В	0	0		1 -											
PCBA Re-Design	С	0	0		1 -											
Supplier Process Change	D	0	0		1 -											
Improve material transport methods	Е	0	0													
0	F	0	0	Feasibiilty	, 1-											
0	G	0	0	Fea	1 -											
0	Н	0	0		0 -											
0	1	0	0		0 -											
0	J	0	0													
0	к	0	0		0 -											
0	L	0	0		0 -											
0	М	0	0		0 1	×	0	0	0	0	1	1	1		1	
0	Ν	0	0		-											
0	0	0	0		Impact											
0	Ρ	0	0													
0	Q	0	0													
0	R	0	0			Impact forsibility plat										
0	S	0	0	Impact-feasibility plot												

Impact calculations

Feasibility calculations

+

E (🖷

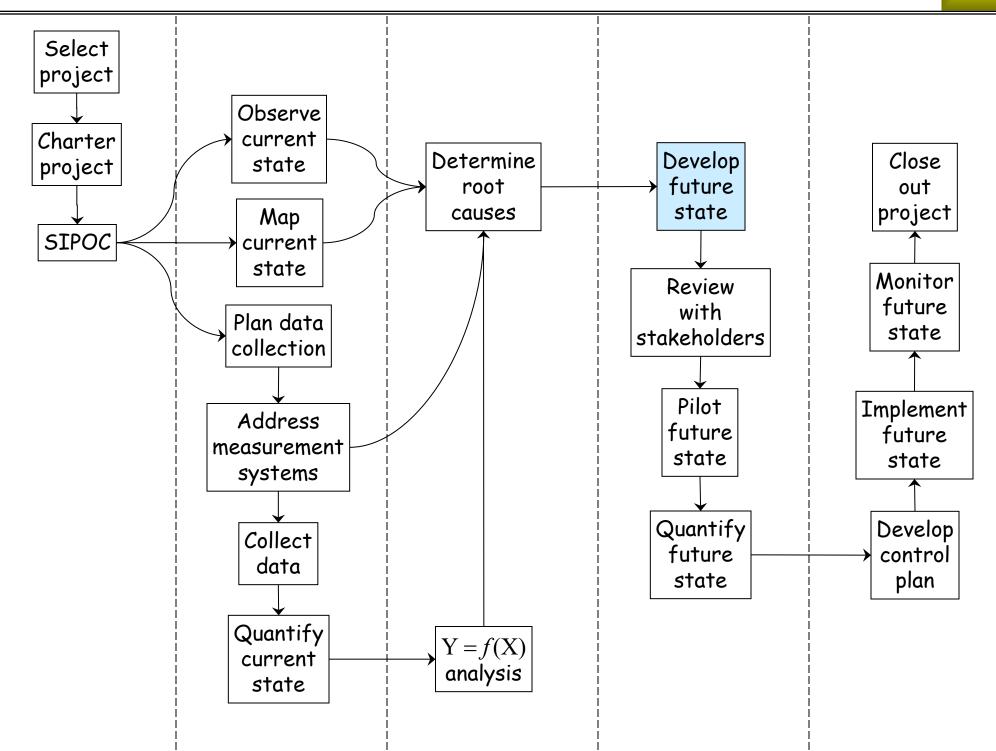
Open Student Files \rightarrow prioritizing solutions - exercise.

Use the root causes and solution ideas as provided. Note that the first row of each sheet is frozen for ease of use during ranking.

Use your knowledge and experience to complete the following tasks:

- a) Change the relative weights for the feasibility metrics as you see fit.
- b) Fill out the Impact ratings sheet using H, M, L or blank.
- c) Fill out the *Feasibility ratings* sheet using H, M, or L.
- d) Use your impact-feasibility plot to decide which solution ideas should be implemented sooner, which should be implemented later, and perhaps, which should not be implemented.

17 Lean Solutions



Continuous Flow Production – 8 Steps

- Parts Quantity (PQ) Analysis
- Establish Standard Routing
- Standardize Work
- Takt Time, Cycle Time, Lead Time, Processing Time
- Work Balancing (leveling)
- Pull System

Standard Work

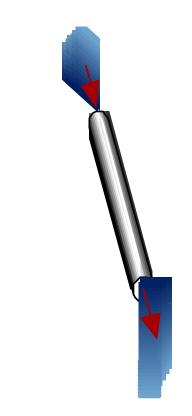
5S

Mistake proofing (Poke Yoke)

Reducing Batch Sizes

Traditional: Batch Production—meandering stream with many stagnant pools and waterfalls Lean Production: Pipeline with fast-flowing water





Definition

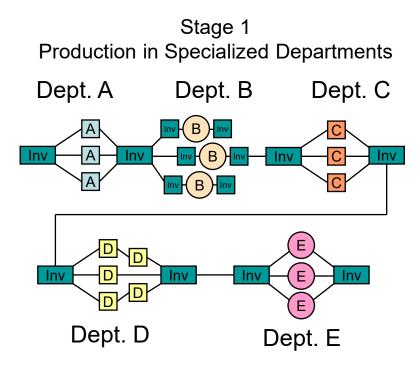
Flow of products in a *level* manner through the production operations—the *ideal* situation is *one-piece* flow at and between processes

Benefits

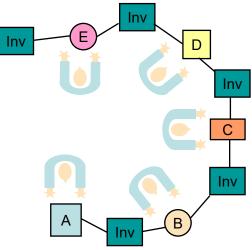
CFP increases the *velocity*, *predictability* and *flexibility* of the production cycle

- 1. Collect data and analyze workflow
- 2. Design process sequence and cell
- 3. Standardize the work
- 4. Produce and move one piece at a time
- 5. Produce at the rate of customer's consumption
- 6. Separate people from machines
- 7. Train people to operate multiple processes
- 8. Balance operations in the cell

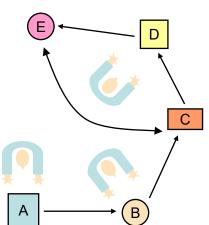
Continuous Flow Journey



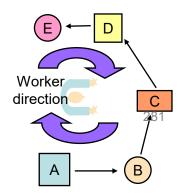
Stage 2 Production in Product Cell



Stage 3 Production in Compact Cell with One-Piece Flow



Stage 4 Production in Compact Cell with One-Piece Flow and Separation Man/Machine



Conduct a Parts Quantity (PQ) Analysis

Prepare a Process Flow Analysis

Group products with similar routings

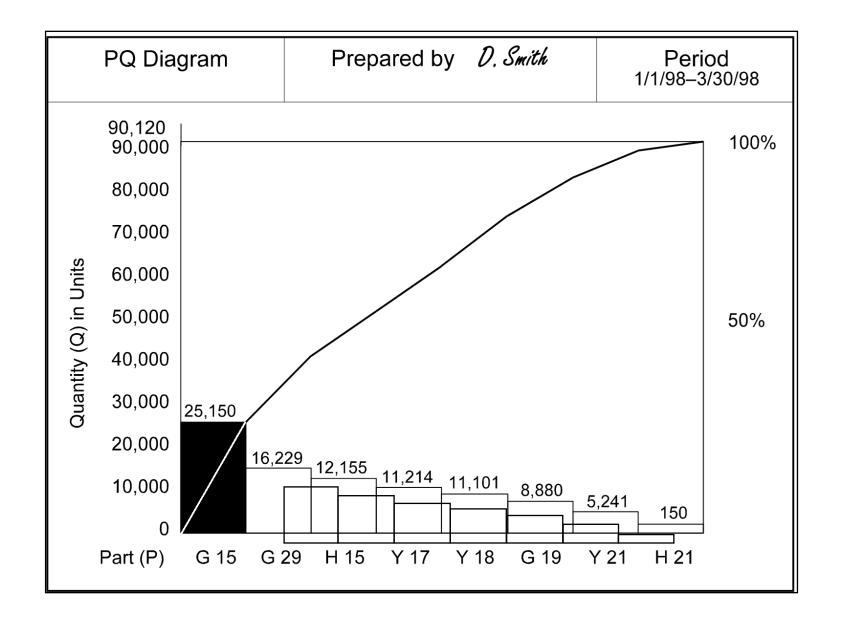
Determine possible cells for product groupings

Evaluate capacity bottlenecks and equipment that can be shared or dedicated

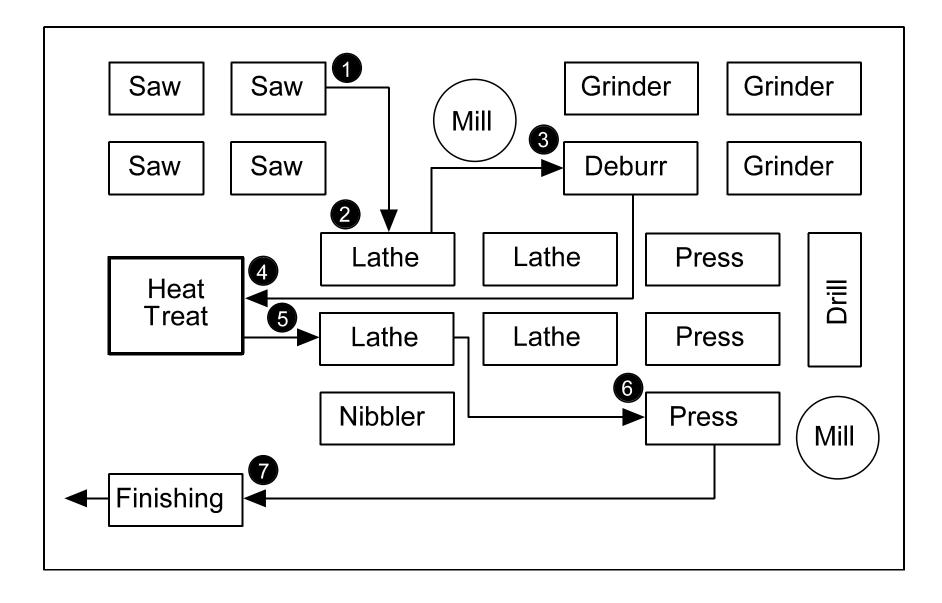
EXAMPLE: PQ Analysis List

Prepared by D.S	Smith		Period 1/1/98-3/30/98							
Part Number	Qty	Cumulative	%	Cum %						
G 15	25,150	25,150	27.9	27.9						
G 29	16,229	41,379	18.0	45.9						
H 15	12,155	53,534	13.5	59.4						
Y 17	11,214	64,748	12.4	71.8						
Y 18	11,101	75,849	12.3	84.1						
G 19	8,880	84,729	9.9	94.0						
Y 21	5,241	89,970	5.7	99.7						
H 21	150	90,120	0.3	100.0						

List all part numbers processed in order of decreasing quantity



Visually display the part number distribution in a Pareto Chart



May require changes to existing "routers"

Part. No.	Saw	Drill	Mill	Lathe	H.T.	Grind	Assy	Cell 1	Cell 2	Cell 3
G15	1	2		3 5	4		6		x	
G29	1	2		3	4		5	Х		
H15	1	2	3	4	5	6	7	Х		
Y17	1		2		4	3	5			x
Y18	1		2		4	3	5			x
G19	1	2		3	_4		5	Х		

Record the process sequence for each part number displaying all "loop-backs" and out-of-area processes

Based on common routings, products can be grouped into cells. The PQ Analysis, process cycle times, and equipment quantities determine how many cells can be made and what equipment must be shared

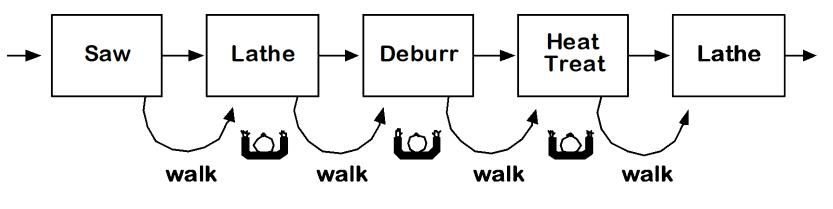
The objective is to identify the minimum number of product groups/cells

Establish standard routing

Locate equipment in proper sequence

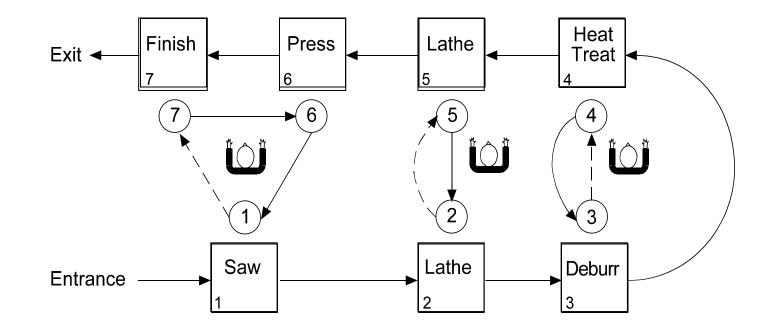
Reduce waste

- Distance: equipment close together
- Inventory: implement pull
- Handling: eliminate wherever possible
- Walking: parts, tools, supplies at point of use



Straight line arrangement disadvantages

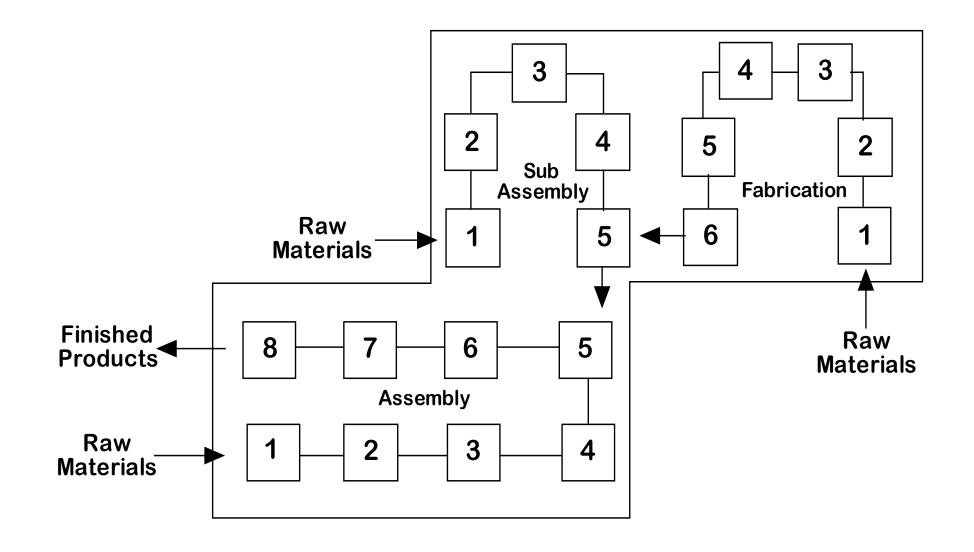
- People usually become dedicated to one machine
- Less flexibility between people
- Normal way to increase output is to work faster
- Work is usually not well balanced between machines
- Quality and output problems noticed at the end of the line are difficult to communicate to the front



Some advantages:

- Shortest distance for people to walk
- Work enters and leaves near the same location
- Improved communication
- Easier to balance the workload between people

289



To sustain and continue improvement, standard work is needed.

<u>Definition:</u> The documentation of how work is performed by both humans and machines to produce products. Standard work has three prerequisites:

- People are performing work
- Processes or tasks must be repetitive
- Work procedures must be documented

Standard work has three components:

- Takt time—The pace at which an operation should complete products or transactions to meet customer demand during the Available Working Time.
- Work sequence—order of operations
- Standard quantities—lot sizes, raw materials, work-in-process

LMT- Administration	M T	
Owner All SBCL MY employeen Stel Management	Managing Improvement Activities	
Four improvement activities (20) on evoluter Feedback	State and the train the state	and the second second
Simble coupleyers in IN @c	the Store in word In and I a	2 manual
Ruptigers organing	Parente TRO AND TO A	· Designed
Manifer, seatain, + Communicate results	A man and the top and the	A manufacture in
way.	The winter as the tame	
6. GROUN		
A		
	man and and and and and and and and and a	
b		
	Ctal	
Ţ	R	
_ _		
	the second secon	
	2 出口出生"世"	
	The second se	
明日間を聞き	前前に思っ 読げ	
The		
at a		

Move one piece at a time

Just try it!

Reduce lot sizes to one

Eliminate constraints to small lots

Make cell compact

- Minimize distance between equipment
- Install casters on equipment

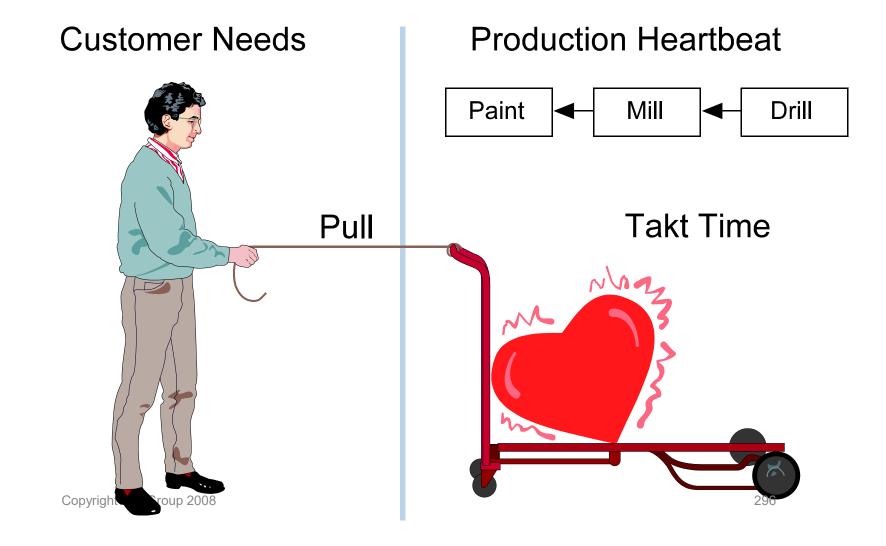
Deal with monuments

Move operations as close together as possible:

- Eliminates WIP buildup
- Eliminates travel distance
- Encourages work sharing... line balance

Put equipment on casters where possible:

- Easy to reconfigure line if demand changes
- Move equipment out of cell when not being used
- Encourages continuous improvement



Takt time

• The pace of production required by the *customer*, calculated as:

Net working time per day

Daily quantity required

Cycle time

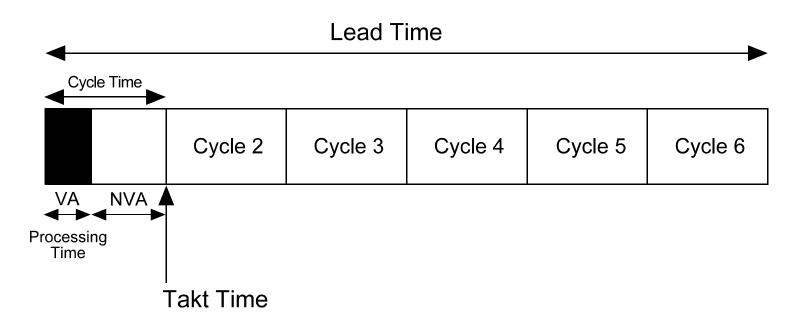
 The current pace of production, measured as the time from completion of one good unit to completion of a second good unit

Lead time

 Total elapsed time from start of a process to the end; the boundaries of the process may be very broad, or very narrow

Processing time

The actual value-added component of cycle time



Strive to make cycle time equal takt time

Takt time > *Cycle time*

- Slow down production
- Redeploy people

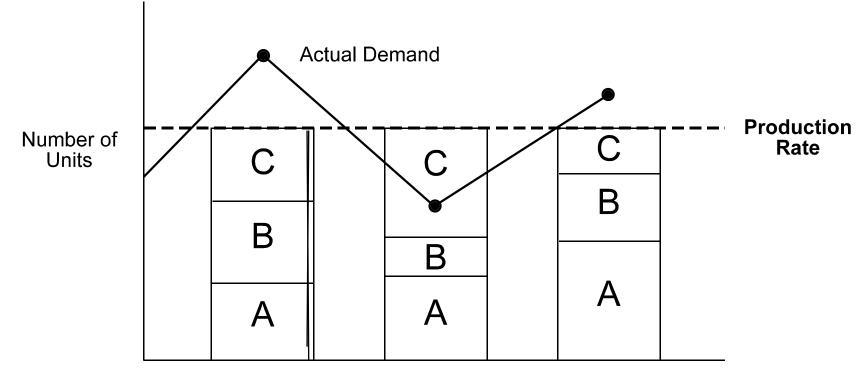
Takt Time =Net Working Time per Day
Daily Quantity Required

"Heartbeat/Drumbeat/Cadence" of Production

Example:

Net working time per day = 7 hours = 420 minutes Daily quantity required = 56 pieces

Takt Time =
$$\frac{420 \text{ minutes}}{56 \text{ pieces}}$$
 = 7.5 minutes



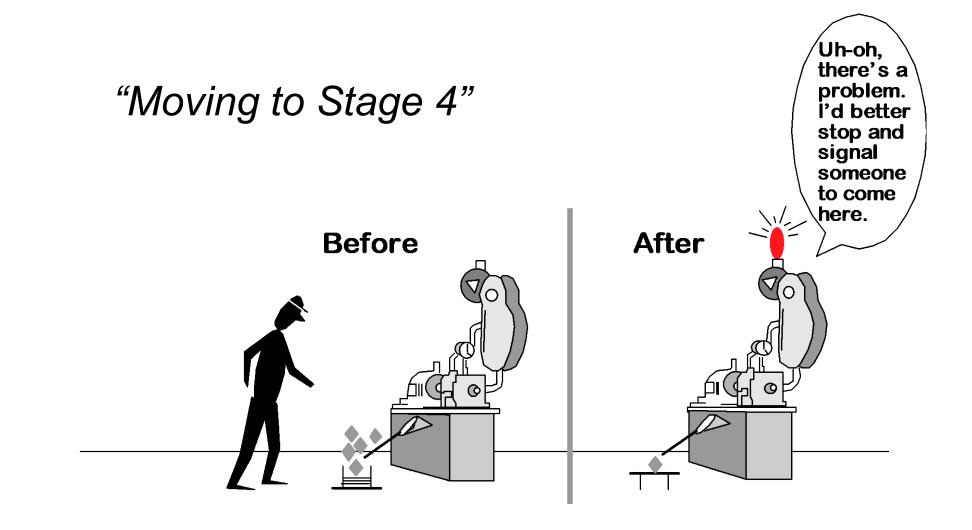
Daily Production Schedule

Process Data

- Total working time per day = 960 minutes
- Time per day for breaks and other manufacturing activities = 160 minutes
- Monthly Customer Demand = 1600 pieces
- Working days per month = 20

Process	Machine Cycle Time
A	10 min
В	10 min
С	3 min
D	2 min
Е	5 min

What is takt time? Do we have capacity? How many operators?



Apply autonomation (automation with a human touch)

Install devices to stop machine when finished or problem occurs (Jidoka or autonomation)

Install mistake-proofing devices (poka-yoke)

Eliminate watching machines and waiting

- Enable people to leave machine and move to next process
- Perform another value-adding task while machine is working

Autonomation (Jidoka)

 Machines and assembly operations can stop production when errors are detected or at the end of a production cycle

Mistake-proofing (Poka-Yoke)

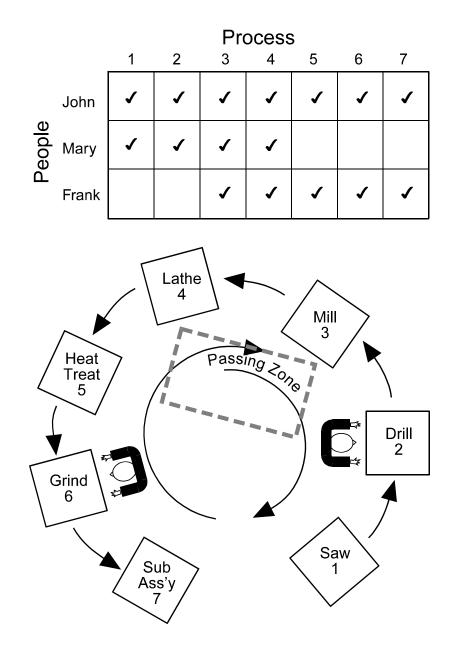
Methods for error detection and (especially) prevention

What skills are needed to operate the cell?

Who performs what tasks?

How will we work together in the cell?

Design for mutual assistance and use defined "passing zones"



Make cycle time = takt time

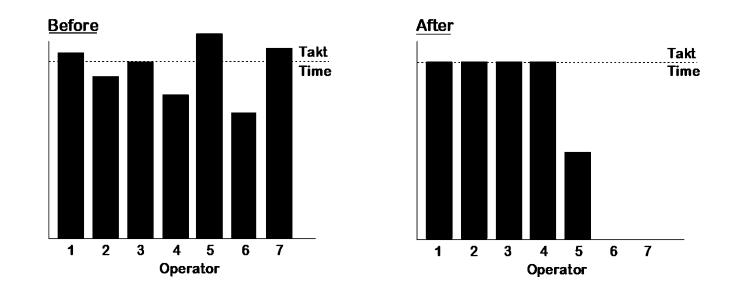
- Tasks
- Machines
- People

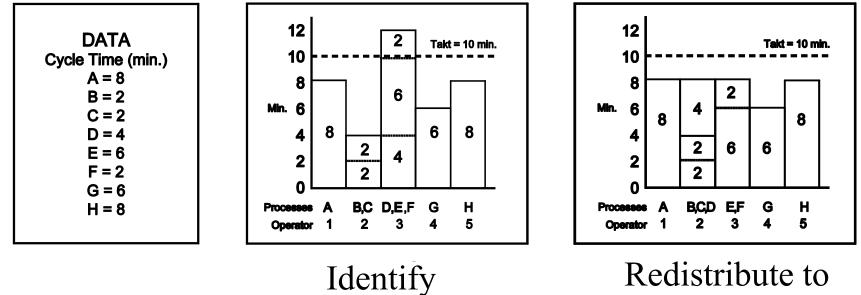
Smooth production

- Level schedule
- Mixed lot production



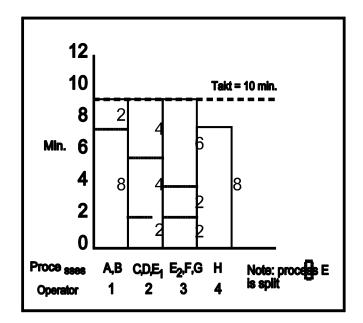
Example: Sum of Cycle Times = 50.40 seconds Takt Time = 11.35 seconds Number of People Required = $\frac{50.40 \text{ seconds}}{11.35 \text{ seconds}}$ = 4.4



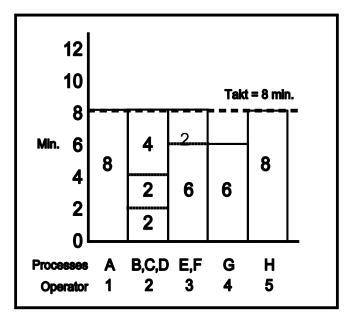


Bottlenecks

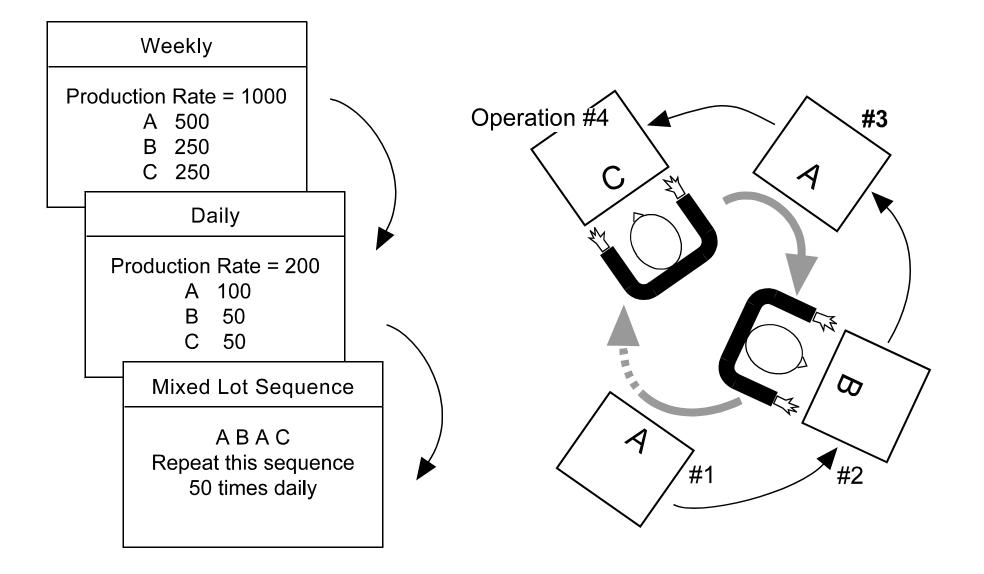
Redistribute to eliminate bottlenecks



Identify opportunities to increase productivity reduce from 5 to 4 people



Adjust production line to decrease in takt time increase the number of people or improve processes to meet the 8-min. takt time



Standard Work

Defines and standardizes the best current work practices

Provides consistency in the operation, and low variation in the output

Provides a basis for improvement

Takt time

Work sequence

Standard quantity (WIP)

Takt time

- The pace at which an operation should complete products or transactions to meet customer demand during the Available Working Time.
- Available working time during a period divided by the number of products or transactions *required* during that same period

Cycle time

- The fastest repeatable time between part or transaction completions using the current processes and resources
- Shows how a process is capable of performing
- Combines with AWT to determine capacity

Lead time

- The total elapsed time to produce one defect free product or transaction
- The time difference between when a part or transaction enters and leaves a process

Processing time

• The actual value-added component of lead time

Takt Time = $\frac{\text{Net Working Time per Day}}{\text{Daily Quantity Required}}$

"Heartbeat" or "Drumbeat" of Production

*Example:*Net Working Time per Day = 7 hoursDaily Quantity Required = 56 pieces per day

Takt Time =
$$\frac{(7 \text{ hours})(60 \text{ min/hour})}{56 \text{ pieces}} = 7.5 \text{ minutes}$$

- Some equipment may need to be slowed down to produce at takt time!!.
- Takt time is expressed as **time**!
- Takt time is always **calculated** from rate of customer demand

The sequence of **operations** to produce a product (e.g., #1 cut, #2 bend, #3 drill, #4 assemble)

The sequence of *activities* necessary to accomplish a single *operation* (e.g., sequence of tasks to bend a part)

Time Observation Form

Standard Work Analysis Sheet

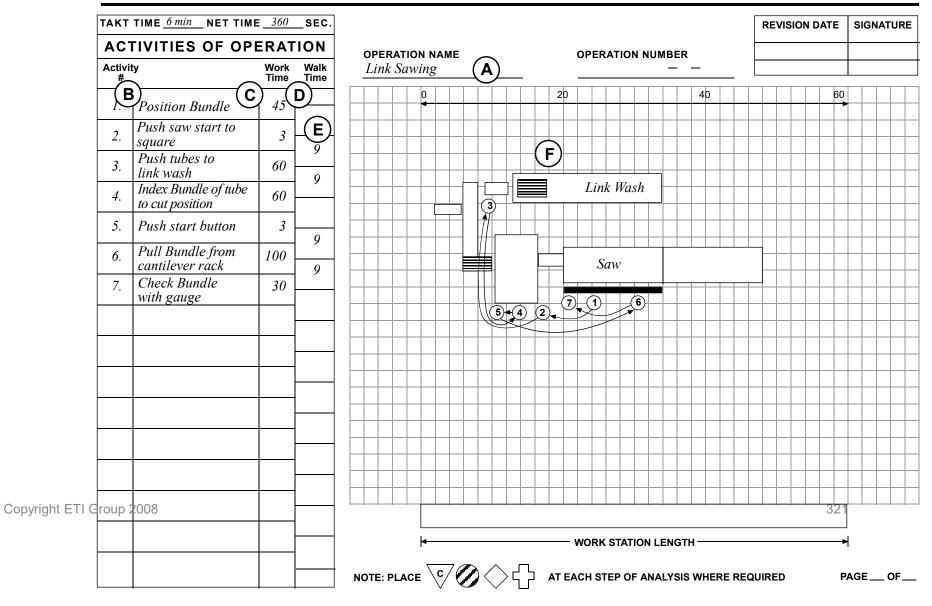
Standard Work Combination Chart

Line Balance Analysis Chart

EXAMPLE: Time Observation

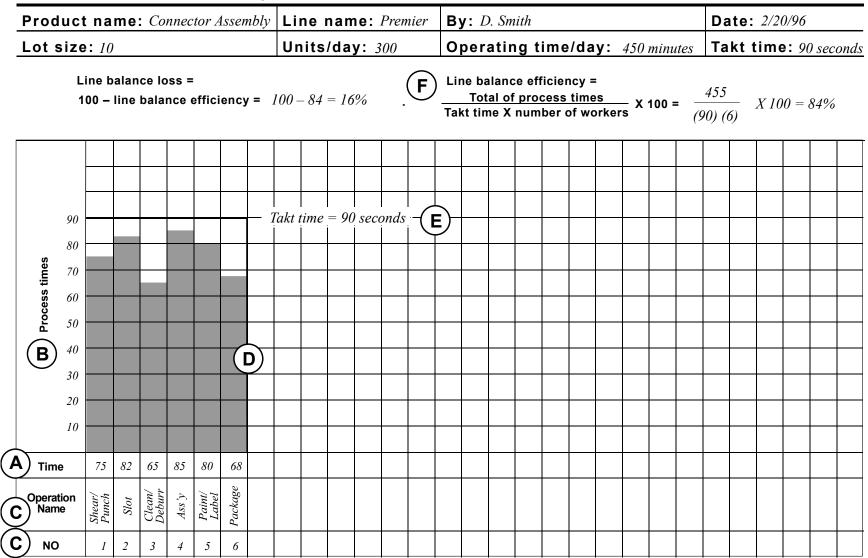
Proc	Process Plate Fab (A) Observer DS Date 2/20/96 (B)													e 2/20/96 B		
		(I	b)_													
No.	Operation/Activity	1	2	3	4	5	6	7	8	9	10	11	12	Average	Range	Comments
1	Cut C	15.0	14.8	14.8	15.4	15.1	15.2	14.7	15.0	15.0	15.1	15.2	14.8	15.0 (F	.7 (G
2	Bend	14.8	14.5	15.5	15.2	15.1	15.5	14.8	14.7	15.1	15.0	14.9	14.7	15.0	1.0	
3	Drill	20.5	20.0	19.8	19.7	19.2	20.5	20.0	21.0	20.2	19.5	20.1	20.2	20.1	1.8	
4	Тар	15.2	15.0	15.0	15.1	15.1	14.9	14.9	14.8	14.8	14.8	15.0	15.0	15.0	.4	
5	Clean/Deburr	15.5	16.0	16.0	14.0	14.0	15.0	15.5	16.0	15.0	14.0	14.2	15.7	15.1	2.0	Variation in burr removal time
	\frown															
Time	for Cycle	81.0	80.3	81.1	79.4	78.5	81.1	79.9	81.5	80.1	78.4	79.4	80.4	80.1	3.1	

EXAMPLE: Standard Work Analysis Sheet



Standard Work Co Model No. A 42-11						ate) bare	d	(C			Quota per E 1500/day												•	Manual					-			_
Оре	eration B Lii	nk sa	awin	g		e p'			(D			Та	akt tim			F) 6	ó m	nin	36		50 Sec					utomatic /alking		~~~~		~~~	 ~	
#	Activity	Manual	Time Auto	5	-	Time (in seconds)																												
G	H		\sim	K)																													Τ
1	Position bundle	45	 	 																														T
2	Push saw start to square			9			ž		••••															T										T
3	Push tubes to link wash	60	! 	9			Ę			-74																								T
4	Index bundle of tube to cut position	60	 	 						ž			7									(Ĺ)										T
5	Push start button		90	 									Ļ				• • • •																	T
6	Pull bundle from cantilever rack	100	 	9									Ž					-72																T
7	Check bundle with gauge	30	 	9															2															T
			! 	 																														T
			 	i I I																														T
		M	(\mathbf{N})	\bigcirc																														T
	Totals	\sim	Wait 180	\sim																			3,	50'	Tak	t ti	me						<u></u>	<u> </u>
	Total Operating Time			I	L																	I			101									

EXAMPLE: Line Balance Analysis Chart



A Workplace that is:

- Clean, organized, orderly
- Safe
- Efficient and pleasant
- The foundation for all other improvement activities

Resulting In:

- Fewer accidents
- Improved efficiency
- Improved quality
- Workplace control

And therefore:

- Reduced waste
- Reduced cost

5S

Sort – Sort through and Sort out

- Keep what is needed Eliminate what is not
- Reduce quantity of items to what is needed

Set in Order – A place for everything and everything in its place

- Identify best location and relocate out-of-place items
- Make locations visually identified easy to see missing items
- Set height, quantity, and size limits
- Organize for safety

Shine – Shine and Inspect through cleaning

- Filthy work environments lead to poor morale
- Spills and debris are safety hazards
- Its easier to identify a maintenance need on clean equipment

Standardize

- Build the framework for maintaining Sort, Set in Order, and Shine
- Clarity about what is and is not normal with simple action plans

Sustain

• Incorporate 5S into the daily work cycle

• Material usage should be first-in-first-out (FIFO)

• Supply orders are triggered by *kanbans* (cards, empty bins, or other signals)

• The objective is to minimize stock-outs without keeping excessive supply quantities on hand

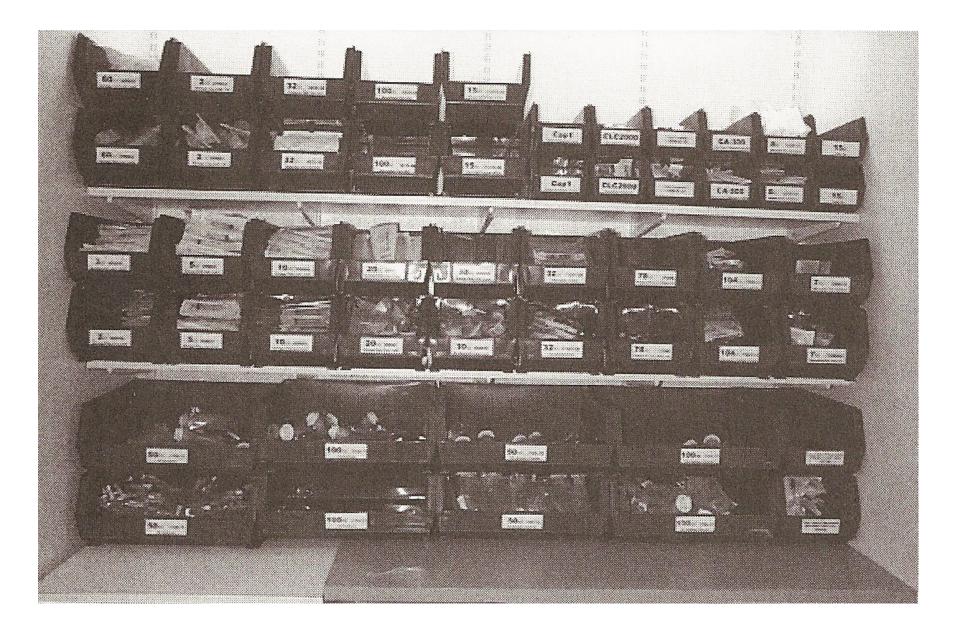
- An order is triggered when the minimum quantity is reached^{*}
- A kanban card goes with the order, returns with the delivery
- The minimum quantity should represent what is needed to span the delivery cycle time

Item Name	
Max. Quantity	
Min. Quantity	
Re-order Qty.	(Max – Min)
Vendor	
Catalog Pg. No	

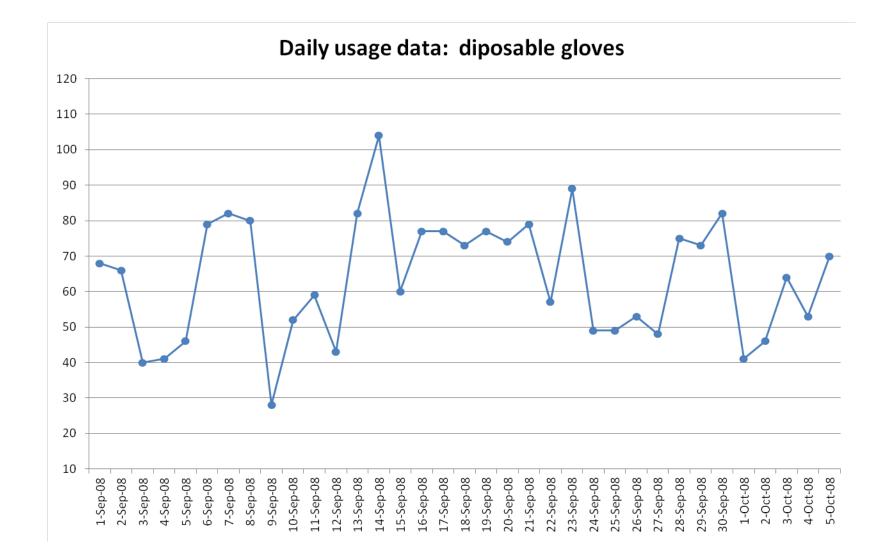
• The maximum quantity should represent a desired upper bound for supply quantity on hand

What can cause this system to fail?

- Two bins for each item (see next slide)
- Amount in each bin = min. quantity = order quantity
- Order when top bin is empty, move bottom bin to top
- Visual system, easy to use
- The max and min quantities can be determined by trial and error
- If usage data is available, there is a better way



- Required inputs
 - \checkmark Time basis for usage data (hourly, each shift, daily, weekly, . . .)
 - ✓ Average usage per time period
 - \checkmark Standard deviation of usage per time period
 - ✓ Minimum order quantity
 - \checkmark Min. value (number of orders)
 - ✓ Max. value (number of orders)
- Values calculated in the simulation
 - \checkmark Starting quantity for each period
 - \checkmark Quantity received during each period
 - \checkmark Quantity used during each period
 - \checkmark Ending quantity for each period
 - \checkmark Quantity ordered during each period



Average = 63.9 Std. dev. = 17.2

	A	В	С	D	E	F	G	Н		J	K	L	М	N	0	Р	Q	R	S
1	Avg. usage each period																ss and	average	e them.)
2	Std. dev. of usage each period	17.2	(lf usa	ge data	canno	ot be ob	tained,	get be	st-gue	ss high	and lo	w value	s and u	use (hig	h-low)/	6.)			
3	Minimum order quantity	100																	
4	Min. value (# orders)	1																	
5	Max. value (# orders)	2																	
6																			
7	Period (hours, shifts, days, weeks)	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
8	Starting quantity	200	136	86	140	74	97	134	83	128	57	42	61	130	77	108	63	79	137
9	Quantity received	0	0	100	0	100	100	0	100	0	100	100	100	0	100	0	100	100	0
10	Quantity used	64	50	46	66	77	63	51	55	71	115	81	31	53	69	45	84	42	42
11	Ending quantity	136	86	140	74	97	134	83	128	57	42	61	130	77	108	63	79	137	95
12	Quantity ordered	0	100	0	100	100	0	100	0	100	100	100	0	100	0	100	100	0	100
13																			
14																			
15			180 -																
16																			
17			160 -					• 1						•					
18			140 -				•		•					-1-	•				
19		≥	120 -	Λ	• •	Λ.	Λ -	T		•	• •	•	1		Λ 🕈				
20		Ending quantity	120 -	M M		. M	ΝT	TWT				• • /			///	- 1	P 🕈 🏌		
21			100 -	VIÓV	1 11			↓∐↓	• / / / /		h h	1 / / 	- 	+ 1	* 7	7 ? 	11 +		
22			, 9 80 -	• y 4	11		- 1171		<u>\ 🕈 V V</u>	1/1 4	$\mu \mu \mu$	1014	↓ /↓	V V	- 14	AA =	W		
23				•		V 14	•V V		1 +1	VIT.	L+T ~~	V VE	[•	4 × .	11	• V \ /•	VV		
24			60 -		•	•			1.	•11			•		•	•//	•1		
25			40 -		¥		•		•	•	- <u>\</u>	••	_			+	•		
26			20 -												•	•			
27										-									
28			0 -		111111														
29			,	~ 6	~ ~	\$ \$	20	১ ৯০	Å	\$ 5	~ ~So	ଚ୍ଚ୍ଚ୍ଚ୍ଚ୍	$^{\prime}$	1° 8	°~ ⟨	5 6	õ		
30										Per	iod								
31										Per	lou								
32																			
33																			
34				Averag	e endir	ng qty.	90.7		# Or	ders	61		# Stoc	k-outs	0				
35																			

	A	В	С	D	E	F	G	Н		J	K	L	Μ	Ν	0	Р	Q	R	S	Т
1	Avg. usage each period	63.9	(lf usa	ge data	a canno	t be ob	tained,	get in	depend	ent bes	st gues	ses fror	m peop	le close	e to the	proces	ss and	average	e them)
2	Std. dev. of usage each period	17.2	(If usa	ge data	a canno	t be ob	tained,	get be	st-gues	ss high	and lo	w value	s and u	ise (hig	h-low)/	6.)				
3	Minimum order quantity	100																		
4	Min. value (# orders)	1																		
5	Max. value (# orders)	2																		
6																				
7	Day of week		Tue	Wed	Thu	Fri	Sat	Sun	Mon	Tue	Wed	Thu	Fri	Sat	Sun	Mon	Tue	Wed	Thu	Fri
8	Beginning quantity	200	120	59	103	63	93	32	-34	12	58	89	126	61	8	-38	-17	31	70	103
9	Quantity received	0	0	100	0	100	0	0	100	100	100	100	0	0	0	100	100	100	100	0
10	Quantity used	80	61	56	40	70	61	66	54	54	69	63	65	53	46	79	52		67	61
11	Ending quantity	120	59	103	63	93	32	-34	12	58	89	126	61	8	-38	-17	31	70	103	42
12	Quantity ordered	0	100	0	100	100	0	0	100	100	100	0	100	0	0	100	100	100	0	100
13																				
14																				
15			200																	
16																				
17			150											1						
18			100	1	<u> </u>				1					Δ	•		•			
19				VV.	<u>^</u>	4	*	Δ	<i>T</i>	Λ			*	r \	/		<u>/</u>			
20		Ending quantity	50	*•	**	1	$-\Lambda$	71	*	11	•		Λ	• • •	+					
21		n n n n n n n n n n n n n n n n n n n	- 0	Ţ		r	<u>• 1</u>	<u>/ </u>		4			<u>/ \ </u>			* L	T			
22		g	, č	at o	l⊇V		¢=¥	e d	at a	∠ = •	s∱¥€	*		¥≣	2 9	•\⊆/`	: =			
23		- i	-50	ž "		ட்ரு	<u>• • · · ·</u>	 	<u>6 0 </u>	r i l i	-/ 6		<u> </u>	i	<u>- • /</u>	-6	<u> </u>			
24		_ <u> </u>	-100			<u> </u>	•	<u> </u>		k	• \/				<u> </u>	<u> </u>	<u> </u>			
25											↓ •	V V			. * .	•				
26			-150														-			
27			-200																	
28			200																	
29																				
30										D										
31										Day of	week									
32																				
33																				
34				Averag	e endin	ig qty.	4.5		# Or	ders	62		# Stoc	k-outs	49					
35																				

- Designing connecting cables and ports so that a cable cannot be plugged into the wrong port
- Programming software so that the user cannot proceed unless necessary information is filled in
- Auto fill of previously entered information on electronic forms
- Pull down menus in computer programs especially for data entry
- Using feedback control systems and alarms on equipment
- Fixturing to prevent incorrect placement and hold things in place

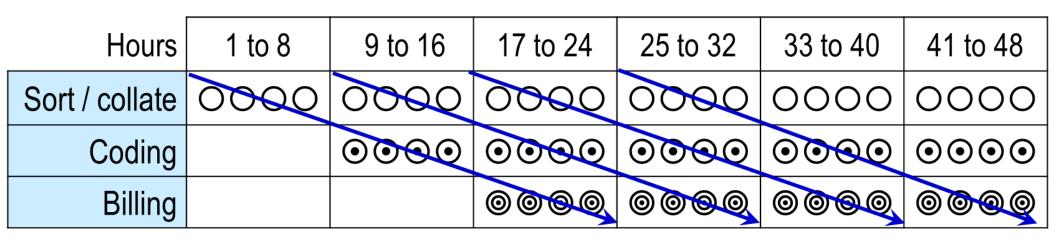
Don't do things in batches. The ideal is to do one thing at a time. Come as close to this as you can.

- Wait a minute batching is supposed to be "efficient"
- Maybe, but here are some problems with batching:
 - ✓ A customer who wants just one item has to wait for a whole batch to be completed
 - ✓ Reduces flexibility in building different products.
 - ✓ Items accumulate until the batch quantity is reached wastes space, creates opportunities for defects

Of course, there can be a legitimate problem with reducing batch sizes: it increases the number of changeovers.

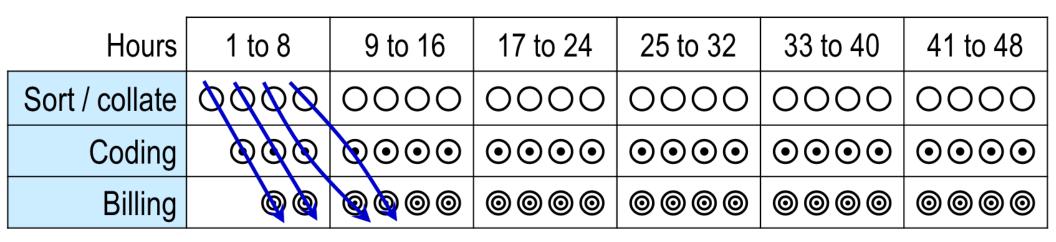
Fortunately, this is a problem for which Lean has excellent solutions. Lean projects have reduced changeover times by 80% or more.

3 operations 2 hours per transaction per operation



Lead time = 24 hours (3 days)

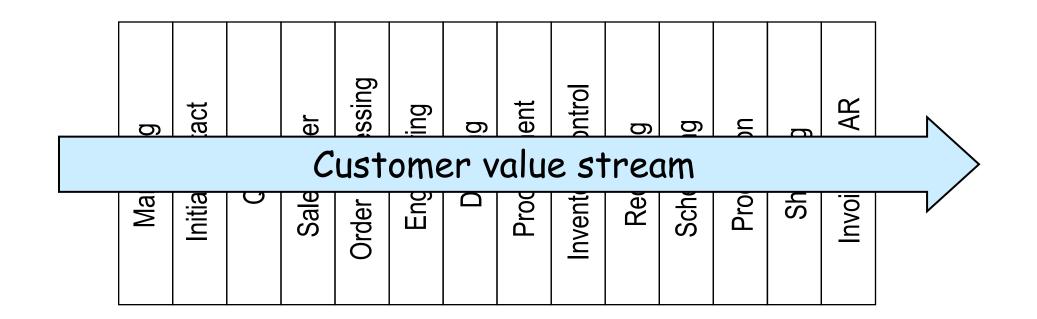
3 operations 2 hours per transaction per operation



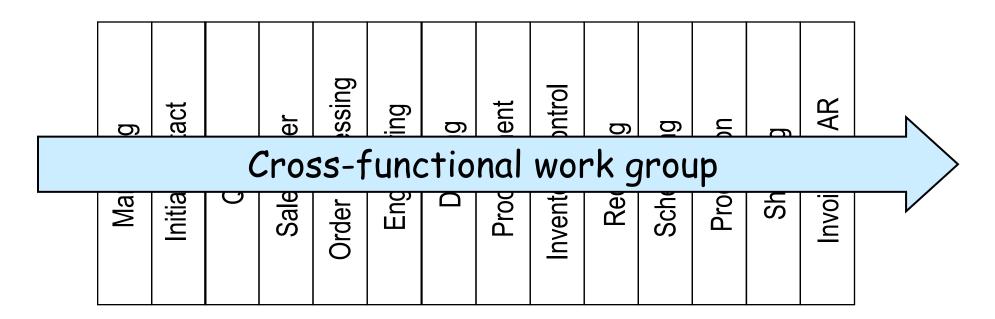
Lead time = 6 hours (less than one day)



- Departmental boundaries create "silos"
- Vestige of industrial revolution need for specialization
- Silos are "islands" of responsibility
- Hand offs between silos are opportunities for poor communication and lack of coordination

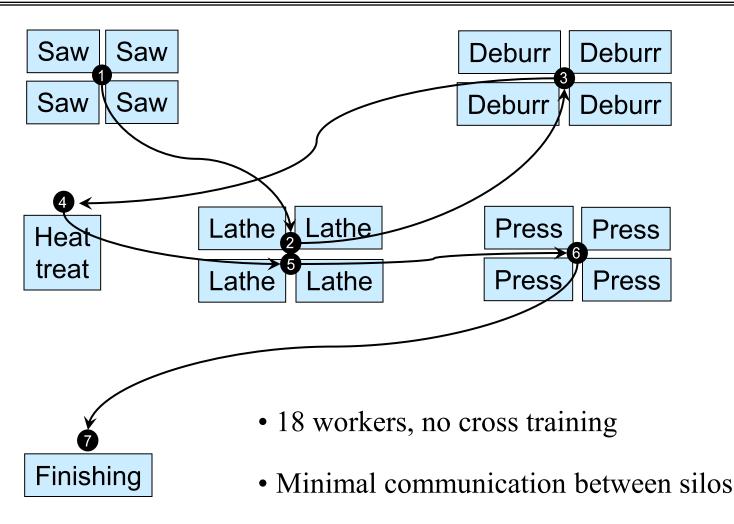


- Customer value stream spans all silos
- Often, no single entity has overall responsibility for customer satisfaction



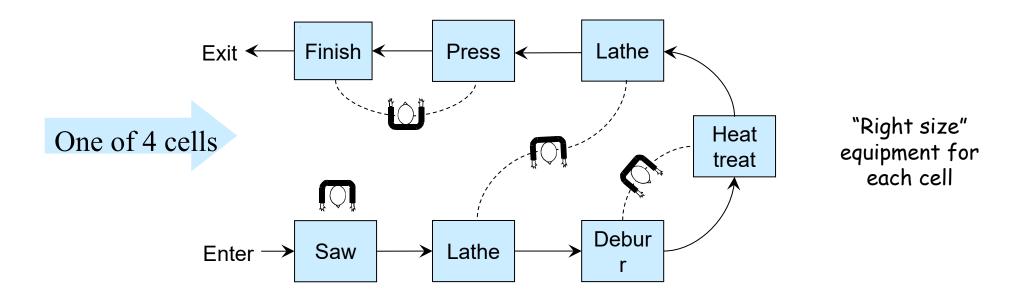
- Responsible for entire value stream for a product/service "family"
- Physical co–location is ideal
- Alternative: "value stream team"
- Stand-up meetings: every day, shift, or other frequent interval
- Alternative: virtual meetings

Manufacturing operation in silos



- Each silo handles all products
- Silos produce as much as possible, all the time (push system)
- WIP moves between silos in large batches \rightarrow long lead time

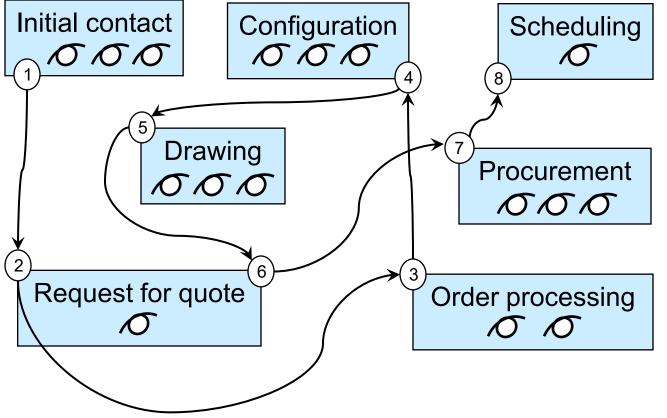
Manufacturing operation in U-shaped work cells*



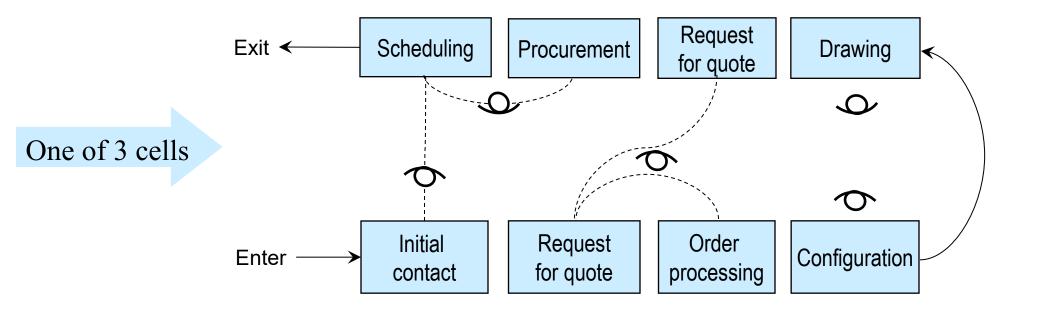
- Each cell handles all operations for one product family, and produces just what is needed to meet current demand (pull system)
- Continuous flow \rightarrow minimal WIP \rightarrow short lead time
- Rapid response to workflow or quality problems
- 16 workers instead of 18 what happened to the other 2?

*Physical co-location is not always possible in process industries, where equipment determines capacity and is difficult or impossible to relocate. See **Lean for the Process Industries** by Peter King for ideas on how to apply Lean in this situation.

Transactional process in silos

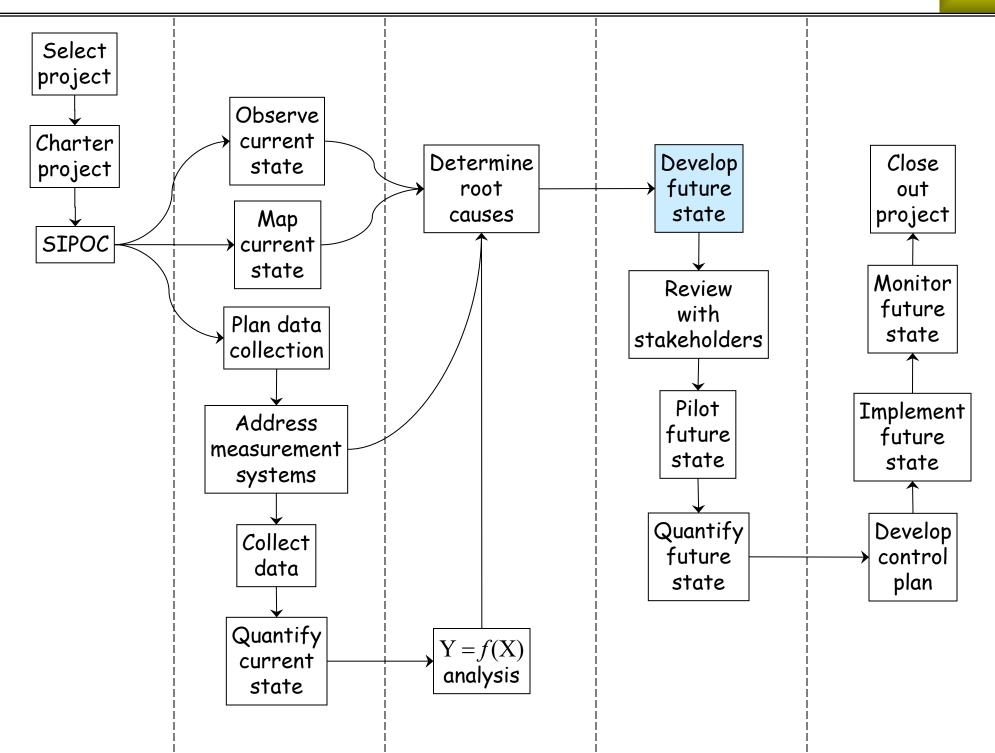


- 16 workers (σ), no cross training
- Each silo handles all transactions
- Minimal communication between silos
- Lots of do overs (not shown in diagram)
- Lots of WIP \rightarrow long turnaround time



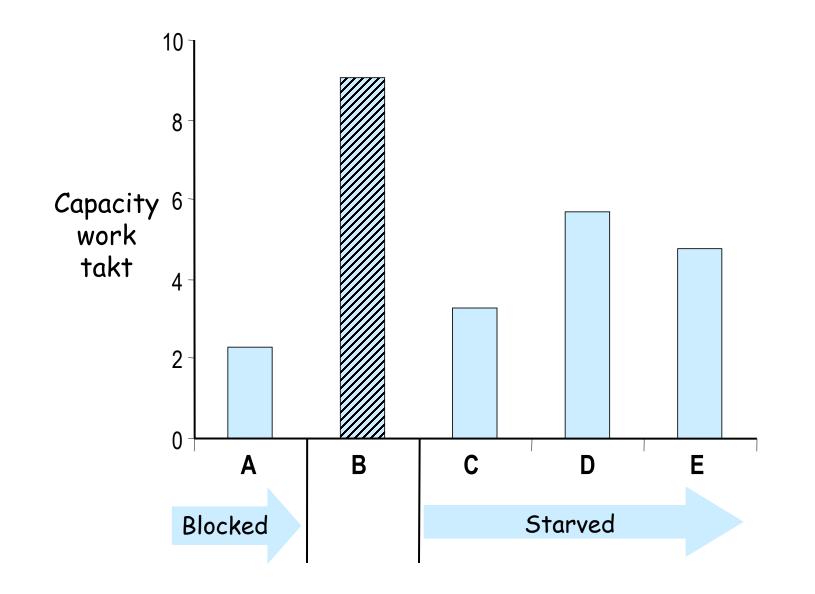
- Each cell handles all steps for one transaction family
- Continuous flow \rightarrow minimal WIP \rightarrow short turnaround time
- Rapid response to errors or workflow problems
- 15 workers instead of 16 what happened to the other one?

18 Theory of Constraints (TOC)



TOC (cont'd)

What if cross training is not feasible? What if the bottleneck is machine capacity?



TOC improvement cycle	Lean terminology
1. <i>Identify</i> the system constraint (the "drum")	Find the bottleneck ("pacemaker")
2. <i>Exploit</i> the identified constraint (includes establishing the "buffer")	 Move resources to the bottleneck Minimize NVA at the bottleneck Maintain needed level of "safety" WIP
3. <i>Subordinate</i> everything else to the constraint (establish the "rope")	Pull system synchronized with the takt time of the bottleneck
4. <i>Elevate</i> the constraint	Add enough resources to eliminate the bottleneck
5. Return to step #1	Find the new bottleneck, repeat same steps

- Greatest WIP
- Longest cycle time
- Longest process time
- Highest % utilization

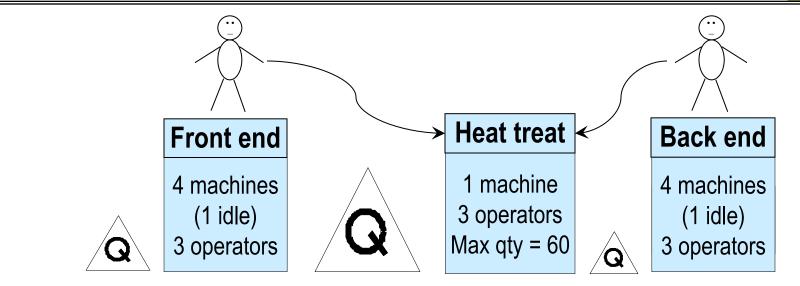
	\bigwedge	Heat treat	
Front end			Back end
		1 machine	
4 machines		1 operator	4 machines
4 operators		Max qty = 60	4 operators

Lead time				120 mins		
"process" time	0	4 min	0	90 mins (load change = 30 mins)	0	4 min
Cycle time		4 mins/4 pcs = 1 min		120/60 = 2 mins		4 mins/4 pcs = 1 min
Constrained cycle time		1 min		2 mins		2 mins

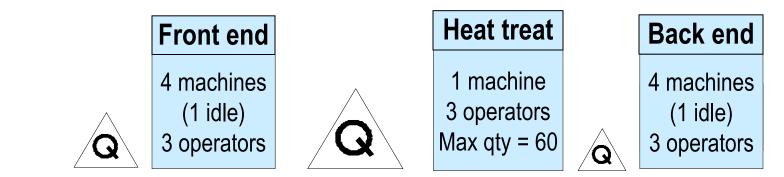




Future state #1: reallocate resources



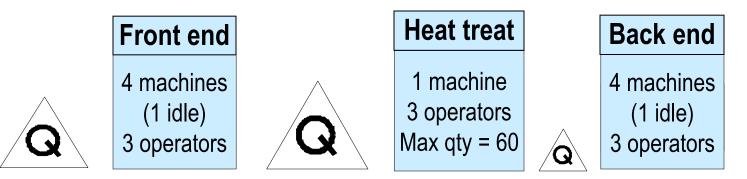
Lead time				100 mins		
"process" time	0	4 min	0	90 mins (load change = 10 mins)	0	4 min
Cycle time		4 mins/3 pcs = 1.33 mins		100/60 = 1.67 mins		4 mins/3 pcs = 1.33 mins
Constrained cycle time		1.33 mins		1.67 mins		1.67 mins
		Less over- production	Less WIP	Faster load change		



Lead time				95 mins		
"process" time	0	4 min	0	90 mins (load change = 5 mins)	0	4 min
Cycle time		4 mins/3 pcs = 1.33 mins		95/60 = 1.58 mins		4 mins/3 pcs = 1.33 mins
Constrained cycle time		1.33 mins		1.58 mins		1.58 mins

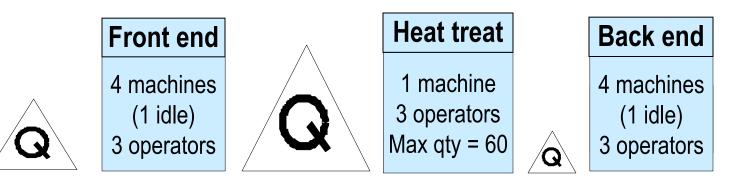
Even faster load change

0



Lead time				95 mins		
"process" time	0	4 min	0	90 mins (load change = 5 mins)	0	4 min
Cycle time		4 mins/3 pcs = 1.33 mins		95/60 = 1.58 mins		4 mins/3 pcs = 1.33 mins
Constrained cycle time		1.58 mins		1.58 mins		1.58 mins
	60 WIP at heat treat					

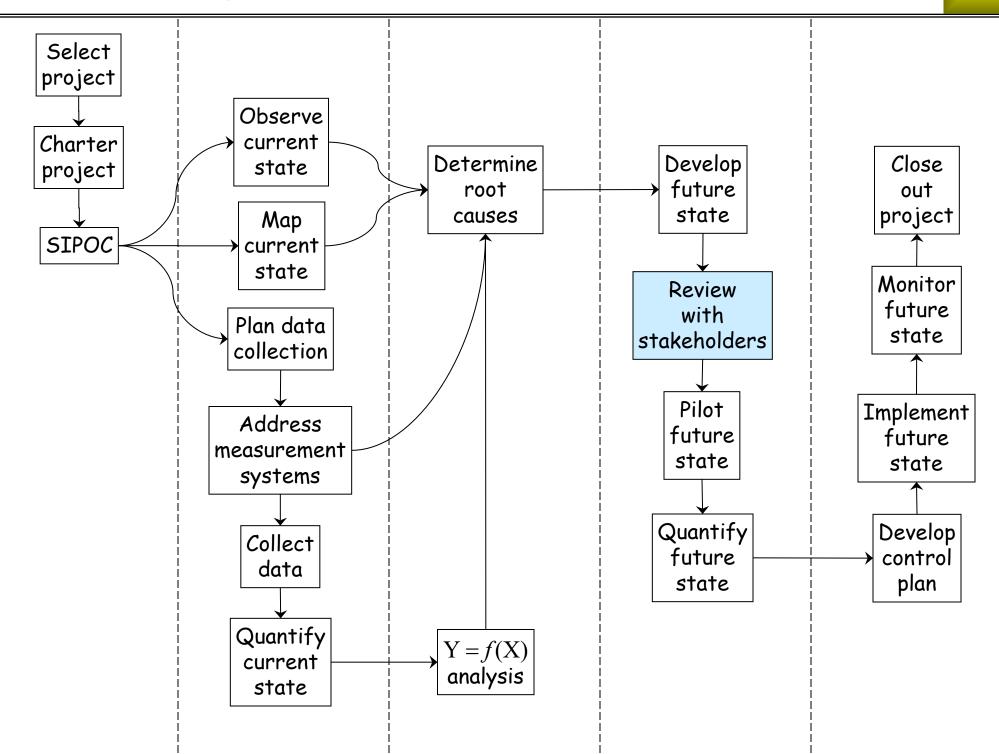
 \leftarrow 95 mins \rightarrow



Lead time				95 mins						
"process" time	0	4 min	0	90 mins (load change = 5 mins)	0	4 min				
Cycle time		4 mins/3 pcs = 1.33 mins		95/60 = 1.58 mins		4 mins/3 pcs = 1.33 mins				
Constrained cycle time		1.58 mins		1.58 mins		1.58 mins				
WIP at heat treat 60										

 \leftarrow 95 mins \rightarrow

19 Reviewing the Proposed Future State



- Use *Failure Modes and Effects Analysis* to identify problems (failure modes) that could occur in your new process and their impact (effects)
- Put things in place in the new process, to prevent or mitigate these failure modes, before they happen
- After you develop your proposed future state, the next step is to review it with stakeholders
 - Give them an opportunity to voice concerns or suggest enhancements prior to piloting
 - This can be an informal process of presentation and discussion

- Principles and steps are the same
 - Purpose and objectives differ
- Rating scale definitions will differ

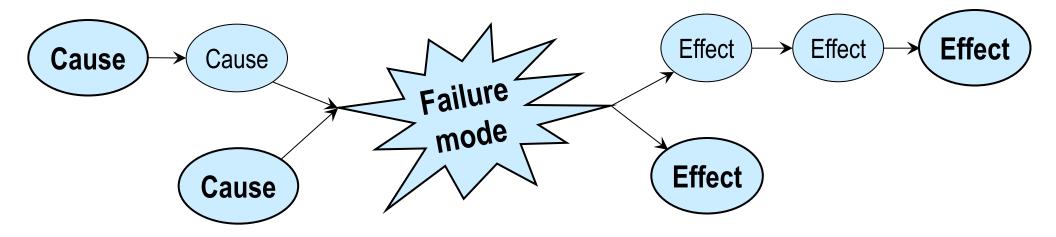
Design

✓ Discover potential problems with the design of the product that will result in safety concerns, malfunctions, or shortened life

Process

✓ Discover potential problems related to the manufacture of the product that will affect the product, safety, or processing efficiency

1. Identify potential failure modes before deploying a new product, service, or process



3. Identify and prioritize root causes of potential failure modes 2. Identify and evaluate ultimate effects of potential failure modes

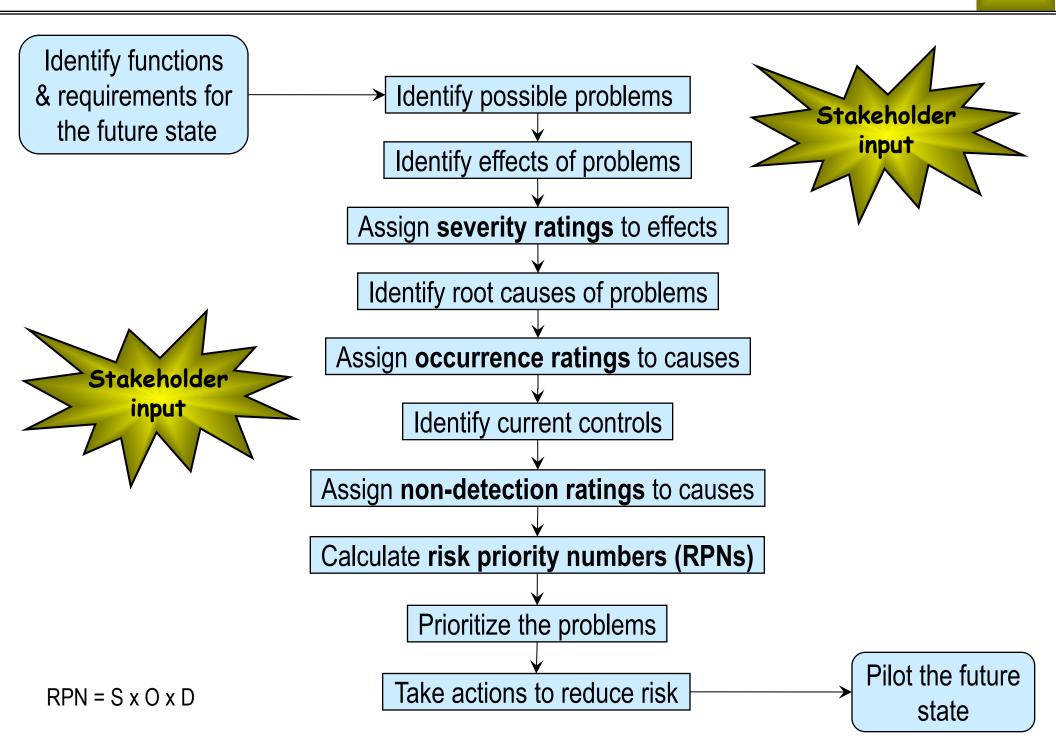
4. Identify and take corrective actions to eliminate or reduce the occurrence of root causes

• Identify and prioritize stakeholder concerns with the proposed future state

• Take appropriate corrective action prior to piloting the future state

• Use results to strengthen the control plan for the future state

Detailed FMEA process



Level		Description
10	Hazardous, no warning	May endanger machine or assembly operator. Failure causes unsafe product operation or noncompliance with government regulation. Failure will occur without warning.
9	Hazardous, warning	May endanger machine or assembly operator. Failure causes unsafe product operation or noncompliance with government regulation. Failure will occur with warning.
8	Very high	Major disruption to production line. 100% of product may have to be scrapped. Product is inoperable with loss of Primary Function.
7	High	Minor disruption to production line. Product may have to be sorted and a portion scrapped. Product is operable but at a reduced level of performance.
6	Moderate	Minor disruption to production line. A portion of the product may have to be scrapped (no sorting). Product is operable but comfort or convenience item(s) are inoperable.
5	Low	Minor disruption to production line. 100% of the product may have to be reworked. Product is operable but comfort or convenience item(s) operate at a reduced level of performance.
4	Very low	Minor disruption to production line. Product may have to be sorted and a portion reworked. Fit/finish or squeak/rattle item does not conform. Most customers notice defect.
3	Minor	Minor disruption to production line. Some product may require rework on-line but out-of- station. Fit/finish or squeak/rattle item does not conform. Average customers notice defect.
2	Very minor	Minor disruption to production line. Some product may require rework on-line but in-station. Fit/finish or squeak/rattle item does not conform. Discriminating customers notice defect.
1	None	No effect.

Example of an Occurrence rating

Lei	vel	Description	Failure Rate		
10	Vory high	Failure is almost inevitable.	≥□ 1 in 2		
9	Very high	Fallure is almost mevitable.	1 in 3		
8	High	Generally associated with processes similar to	1 in 8		
7	High	previous processes that have often failed.	1 in 20		
6		Generally associated with processes similar to	1 in 80		
5	Moderate	previous processes which have experienced	1 in 400		
4		occasional failures, but not in major proportions.	1 in 2000		
3	Low	Isolated failures associated with similar processes.	1 in 15,000		
2	Very low	Only isolated failures associated with almost identical processes.	1 in 150,000		
1	Remote	Failure is unlikely. No failures ever associated with almost identical processes.	≤ 1 in 1,500,000		

	Level	Description
10	Almost impossible	No known controls available to detect failure mode or cause.
9	Very remote	Very remote likelihood current controls will detect failure mode or cause.
8	Remote	Remote likelihood current controls will detect failure mode or cause.
7	Very low	Very low likelihood current controls will detect failure mode or cause.
6	Low	Low likelihood current controls will detect failure mode or cause.
5	Moderate	Moderate likelihood current controls will detect failure mode or cause.
4	Moderately high	Moderately high likelihood current controls will detect failure mode or cause.
3	High	High likelihood current controls will detect failure mode or cause.
2	Very high	Very high likelihood current controls will detect failure mode or cause.
1	Almost certain	Current controls almost certain to detect failure mode or cause. Reliable detection controls are known with similar processes.

- The previous three slides give examples of traditional 1–10 ratings for severity, occurrence, and non–detection
- Note the detailed quantitative operational definitions
- Customers or regulatory agencies may require this level of detail
- For the application to LSS projects, qualitative 1–5 ratings are often sufficient:
 - 1. Very low
 - 2. Low
 - 3. Moderate
 - 4. High
 - 5. Very high

	Doguiromonto	Failure Modes	Effects	SEV	Courses	000	CN	Current	DET	ווחס	Actions Planned	Deeneneihle	Due Date	Actions Taken
Process Functions Reagent lot creation	Requirements New lot information distributed to OPS team	Printer malfunction	Delay in distribution to the OPS team	1	Causes Electrical	1	1	Controls One printer	1 1	RPN 1		Responsible		Такеп
Reagent creation	New reagent created based on processing demand	-	Processing delay, wasted sub-reagents, time lost, labor money	5	Did not use trained witness	1	5	SOP requires trained witness for procedure	1	5				
Reagent storage	Storage of new reagent at point of use (laboratory)	Insufficent storage space in freezer or fridge	Reagent stock-out	4	Freezer space not reconciled	5	20	No control.	5	100				
	and reagents in	Insufficient shelf space for materials.	Material stock-out	3	Too many items on shelving	3	9	Shelving units with four shelves	5	45				
Material storage	designated location within the functional laboratory	Staff is unclear where material items should be stored	Materials not stocked in designated location within the functional area	2	Insufficent labeling system to designate material and reagent locations	3	6	Labels on shelving only	3	18				
Material Distribution	Distribution of materials based on MIN/MAX forecasting	MIN/MAX values not accurate	Material shortage	2	Forecasting not accurate	3	6	Master Science Forecasting	5	30				

Problem statement

Operations staff within the Gene Expression Lab (GEL) are experiencing frequent material stock outs while performing procedures. They have to stop processing samples until the missing material is delivered. This increases process cycle time and reduces the quality of the data deliverables. Other labs directly affected by this problem are:

- ✓ Tissue Homogenization
- ✓ Experiment Processing
- ✓ Sample Processing

Goal statement

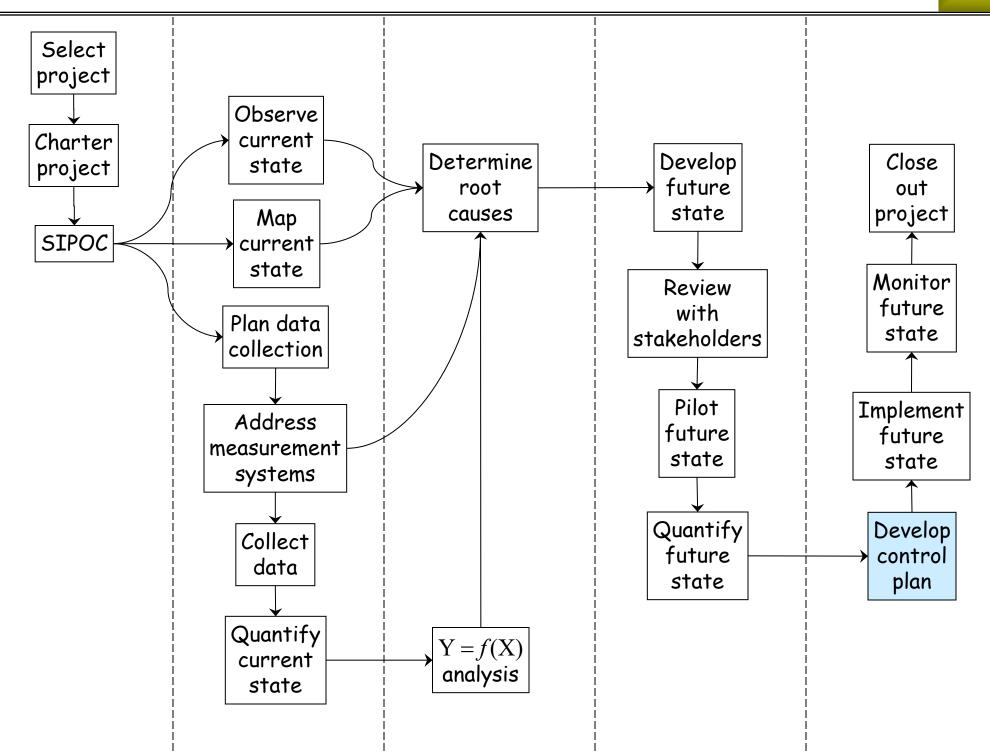
- Reduce frequency of stock outs by 50%.
- Reduce time lost due to stock outs by 50%.

Constraint

No increase in labor cost.

Control Phase

20 Control Plan



368

- A summary of the plan to sustain the gains from a LSS project
- The project team helps the in–scope process owner and participants develop the plan
- Project team advises the in-scope process owner and participants on statistical monitoring issues
- Most common control methods: training, auditing, control chart
- Most common control chart quantities: *individual measurements*, *averages*, and *percentages*

Process name:	
Process owner:	
Revision date:	

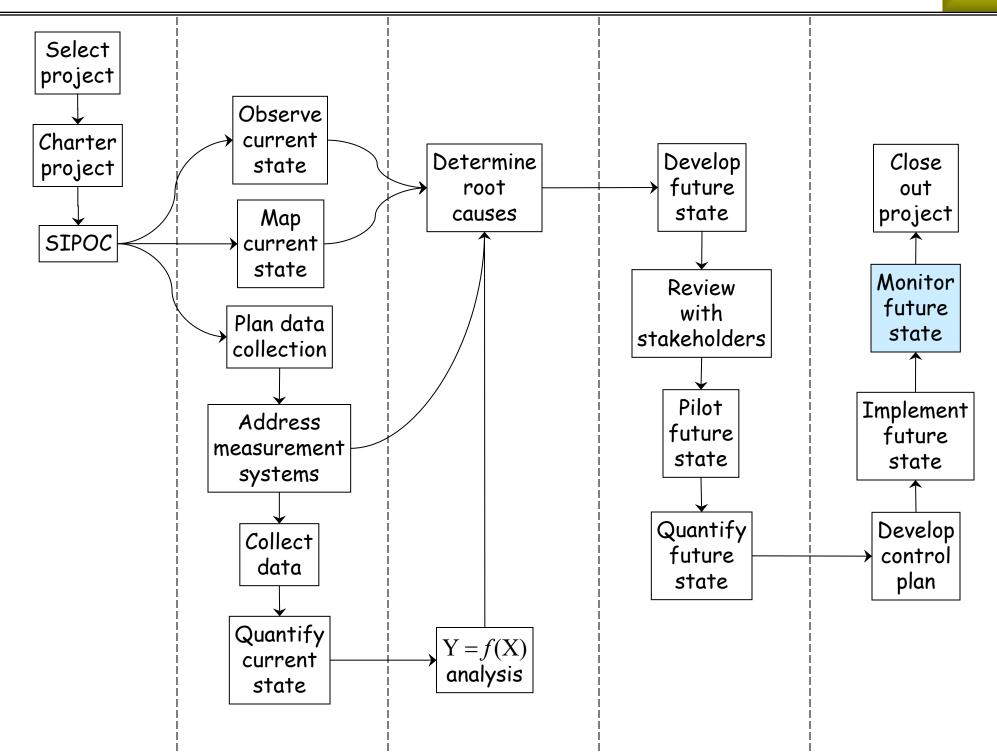
Process step	Control	Fraguancy	Data	Meas.	Metric to	Contro	l limits	Response plan	Response plan	
riocess siep	method			Lower	Upper	owner	location			
	•	•	•	•	•	•	•			

Student Files \rightarrow tool development control plan

Process name:	Tool Testing Process
Process owner:	Testing Area Manager
Revision date:	

Durana atau	Control method	F	Data variable	Meas.	Metric to monitor	Control limits		Response plan	Response
Process step	Control method	Frequency	Data variable	system	Metric to monitor	Lower	Lower Upper		plan location
Determine run conditions	Audit compliance with new procedure requiring special approval to change weight or line speed	Monthly, then Quarterly	Run conditions						
Determine run conditions	Disable weight and line speed controls on test line								
Release to manufacturing	Control chart	Weekly	Number of days in testing	Database	Average		TBD	Testing area manager	TBD
Release to manufacturing	Control chart	Weekly	Number of rework cycles	Database	Average		TBD	Testing area manager	TBD
Dimensional inspection	Install DVT gage and train testers to use it								
Dimensional inspection	Periodic gage R&R	TBD	Spec dimensions	DVT	% of Tolerance		TBD	Testing Engineer	TBD

21 Statistical Monitoring



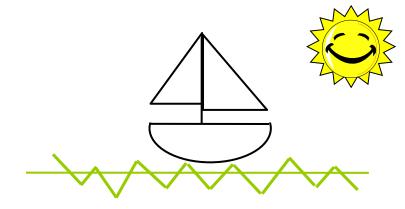
- Two kinds of variation
- Quantifying common cause variation
- Establishing control limits
- Commonly used control charts
- Interpreting control charts
- Response plans
- Relationship to Process Capability

*The more commonly used term is Statistical Process Control (SPC), even though it has nothing to do with "control" in the usual sense.

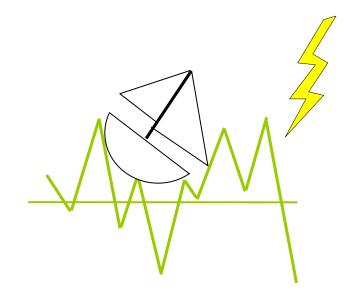
a) Sign your name five times in the space provided below.

b) Put your pencil or pen into the other hand. Sign your name once in the space provided below.

Variation due to common causes



Variation due to assignable causes



- Random variation
- Inherent in the process as currently defined
- Many small fluctuations
- Outcomes are statistically predictable
- Causes for individual fluctuations cannot be determined

- Systematic variation
- Mistakes, malfunctions, miscommunications, external factors . . .
- Relatively few large fluctuations
- Outcomes are not predictable
- Causes of individual fluctuations *can* be determined

• Common cause variation is usually represented by upper and lower *control limits*

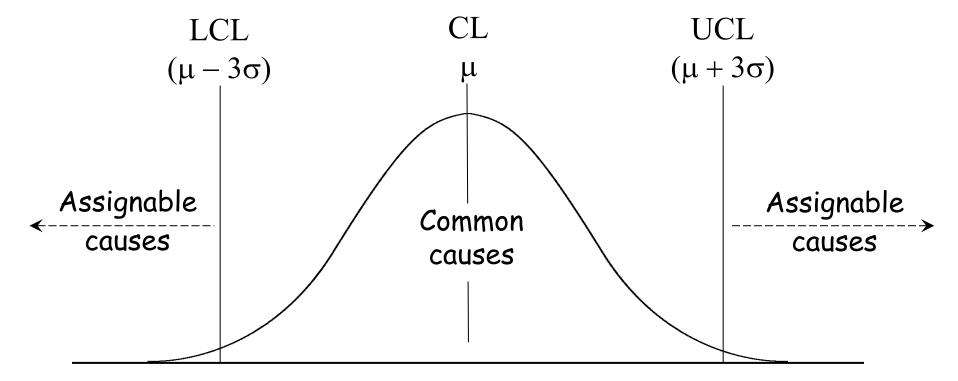
• Upper control limit (UCL) = $\mu + 3\sigma$

• Lower control limit (LCL) = $\mu - 3\sigma$

• These are also called *three-sigma limits*

• Center Line (CL) = μ

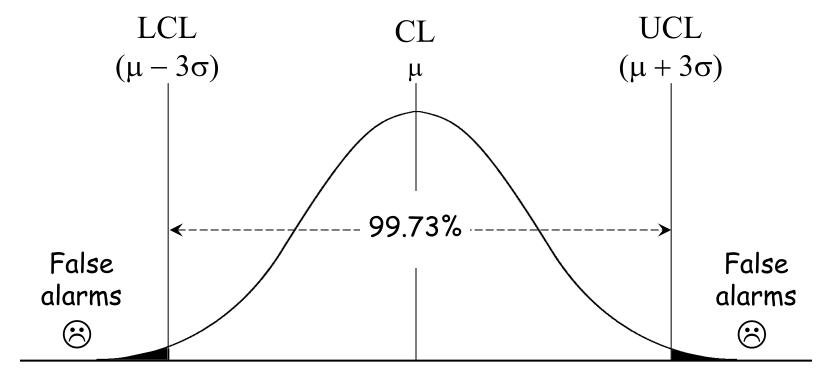
Control limits provide an *operational definition* of common cause variation



Baseline variation in the quantity to be monitored

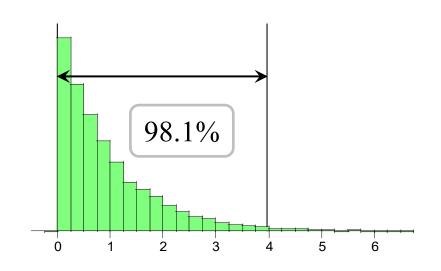
 μ = average σ = standard deviation

If the quantity to be monitored follows a Normal distribution, the chance of a *false alarm* is 0.27%

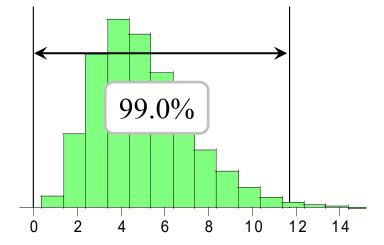


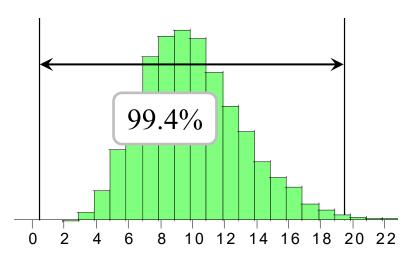
Baseline variation in the quantity to be monitored

Common cause variation (cont'd)



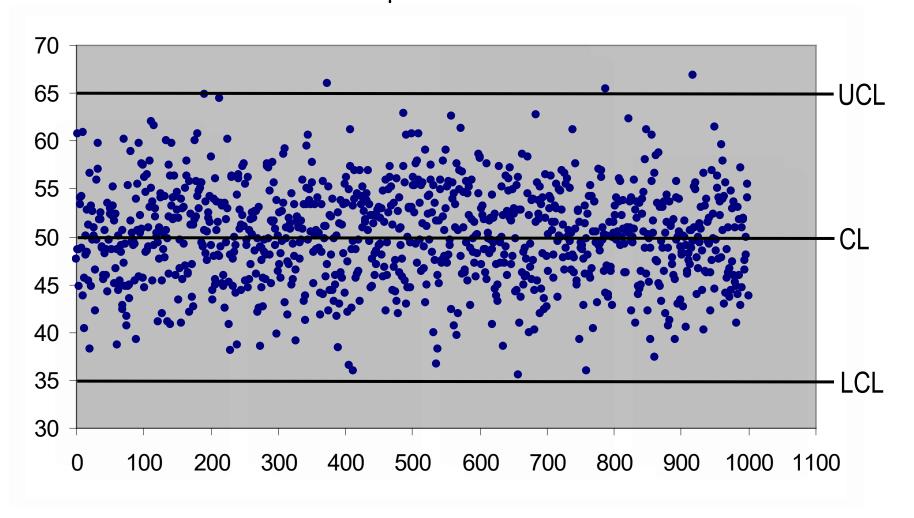
- 3σ limits are shown for three non–Normal distributions
- Data doesn't need to be Normally distributed for most charts
- The *Central Limit Theorem* also greatly reduces the effect of non-Normality when samples are used
- 3σ limits are an economic compromise between *false alarms* and *missed signals*





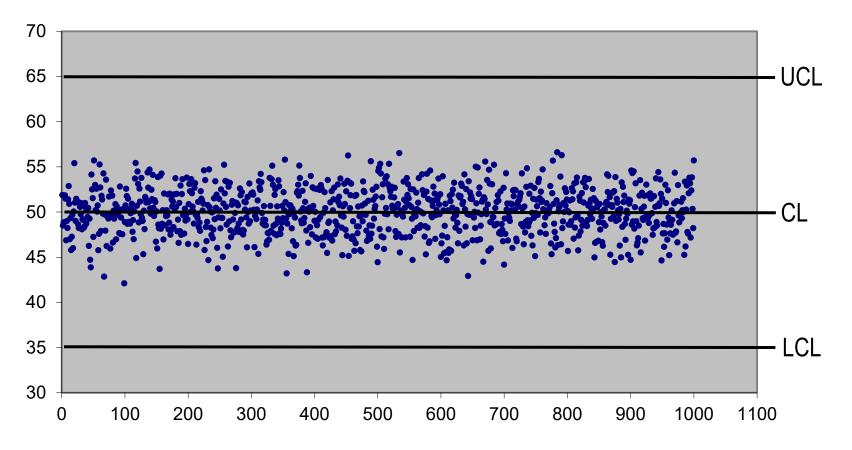
Behavior of Averages: the "Central Limit" effect

Individual data values sampled from a population with $\mu = 50$ and $\sigma = 5$



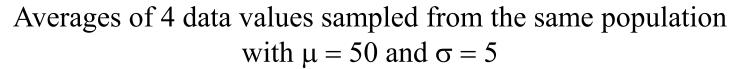
Behavior of Averages: the "Central Limit" effect (cont'd)

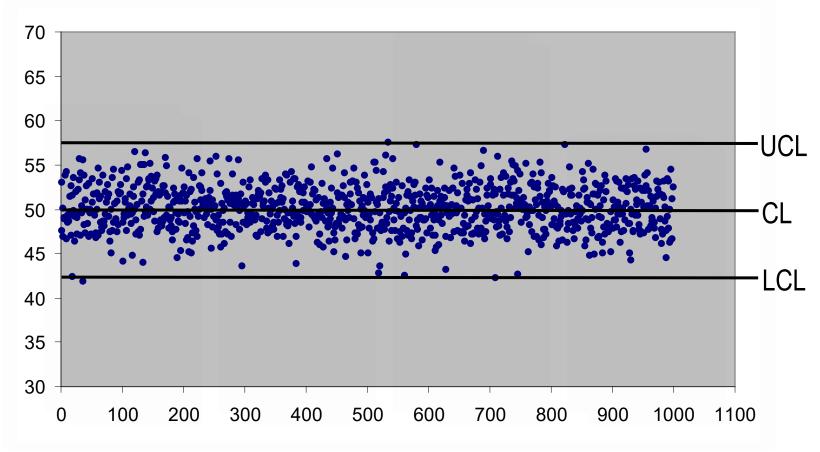
Averages of 4 data values sampled from the same population with $\mu = 50$ and $\sigma = 5$



- Why would the limits shown above be ineffective for statistical monitoring of the averages?
- Control limits for a distribution of averages must be calculated a different way.

Behavior of Averages: the "Central Limit" effect (cont'd)

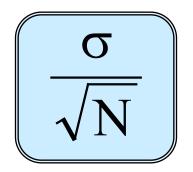




- These are the true control limits for the averages.
- In addition to the obvious narrowing of the distribution, the Central Limit Theorem (stated simply), concludes that subgroup averages converge to a Normal distribution, even if the underlying distribution is non-Normal.

If we repeatedly sample sets of N individual data values from a population with mean μ and standard deviation σ , and calculate the average in each case, the *standard deviation of*

the averages is:



If we repeatedly sample sets of N individual data values from a population with mean μ and standard deviation σ , and calculate the average in each case, the *three-sigma limits for*

the averages are:

$$UCL = \mu + 3\frac{\sigma}{\sqrt{N}}$$
$$LCL = \mu - 3\frac{\sigma}{\sqrt{N}}$$

- Control Limits are calculated using data *representative* of day-to-day process operation
- The exact calculation for three sigma limits depends on the type of control chart being used
- The type of control chart used depends on the type of data and the sampling method
- At least 20 25 sample subgroups should be used to set control limits
- Data from a pilot run can be used to set control limits for the "future state" process, if the pilot is representative of the process that will be implemented.
 - If not, run the "future state" process long enough to gather a sufficient sample.

Control limits are *not* the same as specification limits!

To detect process shifts, we need to take a *reasonable* sample of the process.

- Samples should estimate, or try to represent, the population.
- Samples need to be taken in the order of production and as soon as possible in an operation to get an early warning of defects.
- The chance of variation from assignable causes should be *minimized within* an individual sample set (pull parts for a sample close together in time).
- The chance of variation from assignable causes should be *maximized between* samples (time separation between samples).
- Pulling subgroups of parts at a predetermined interval works best.
 - Do not pre-identify which parts will form the SPC sample before they are manufactured (avoid bias).
 - > Do not adjust the process during sampling.

Quantitative measurement:

- \overline{X} & s (sample average and standard deviation)
- \overline{X} & R (sample average and range)
- IX and MR (individual values and moving range

Categorical classification:

• p (fraction defective)

With quantitative control charts, we pull samples from the process and use them to estimate how the process as a whole is performing.

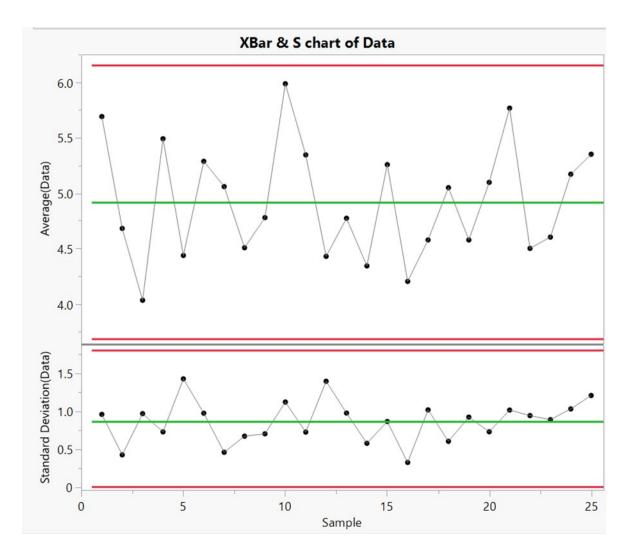
We can then answer two important questions using two graphs:

- 1. Is the process staying centered?
- 2. Is the process staying consistent?

SCENARIO A		SCENARIO B	
\bigwedge	The centering is		The centering is
	The consistency is		The consistency is

Control Chart	Statistics Plotted	Sample Size	Description				
X-bar & R Average & 2–5 Range		2–5	The X-bar and R chart was the first and most common quantitative control chart used in SPC, only because in the days before calculators and statistical software, Range was easier to calculate than Standard Deviation.				
			The X-bar and R chart can be useful for monitoring product, process or environmental characteristics when the sample size is fairly small (say 5 or less).				
			But given the prevalence of software tools available, it should really be replaced by the X-bar and s chart unless there is a particular need for spotting "outlier" range values.				
X-bar & s	X-bar & s Average & 5–15 Standard Deviation		The X-bar and s chart is useful for monitoring product, process or environmental characteristics, especially when the sample size is larger (say, more than 5).				
			Again, the standard deviation chart will be more robust than range because all data are used, not just the highest and lowest numbers.				
IX & MR	Individual & Moving	1	The IX and MR chart is used when the sample size is one. A single sample may need to be taken because:				
	Range		• It is expensive to take samples.				
			• The measurement method is destructive.				
			• It is the only sample size that makes sense for that process.				
			Because an average cannot be calculated for a sample size of one, the individual data points are used.				
			When there is only one number, standard deviation and range cannot be calculated. Instead, we use what is called the <i>Moving Range</i> .				

For each sample, the average is plotted on the \overline{X} chart (centering) and the standard deviation (consistency) is plotted below on the s chart.



JMP Output of \overline{X} s Chart of *control chart diameter*

Monitoring frequency	Metric to monitor	Statistic(s) Needed	Control limits
Hourly Daily	\overline{X} chart: Average	Average (µ)	UCL = $\mu + 3\frac{\sigma}{\sqrt{N}}$
Weekly	Twerage	Standard deviation (σ)	$CL = \mu$ $LCL = \mu - 3\frac{\sigma}{\sqrt{N}}$
Monthly Quarterly etc.	<i>s chart:</i> Standard Deviation	Standard deviation (σ)	UCL = $\overline{\sigma} + 3 \frac{\sigma}{\sqrt{2(N-1)}}$ CL = $\overline{\sigma}$ LCL = $\overline{\sigma} - 3 \frac{\sigma}{\sqrt{2(N-1)}}$

We want to use \overline{X} and s control charts to monitor a critical dimension, diameter, of the parts we are producing. Open *Data Sets* \rightarrow *control chart diameter*. Does the baseline data appear to be adequate to represent process variation? Use Excel formulas for the following:

- a) Calculate the average (\bar{x}) and standard deviation (s) for each subgroup of five parts.
- b) Calculate the overall average, which will be the center line (CL) of the \overline{x} chart. There are two ways to do so: take the average of all the data points or take the average of the subgroup averages. The name given to the statistic from the second method is $\overline{\overline{X}}$ (X-double bar) or the Grand Average.
- c) Calculate the average of the subgroup standard deviations, (\overline{s}) , which will be the Center Line (CL) for the standard deviation chart.

Exercise 21.2 (cont'd)

- d) The estimates of the standard deviation of the distribution of averages and the distribution of standard deviations have been calculated for you. They are used in the "3-sigma" quantities that are added to and subtracted from the Center Lines.
- e) Use the numbers found above to calculate the upper and lower control limits for each chart.

$$UCL_{x} = UCL_{s} = UCL_{s} = CL_{x} = CL_{s} = LCL_{x} = LCL_{s} = LCL_{s$$

$\overline{\mathbf{X}}$ Chart Control Limits:

 $UCL = \overline{\overline{x}} + A_2 \overline{R}$ $CL = \overline{\overline{x}}$ $LCL = \overline{\overline{x}} - A_2 \overline{R}$

R Chart Control Limits:

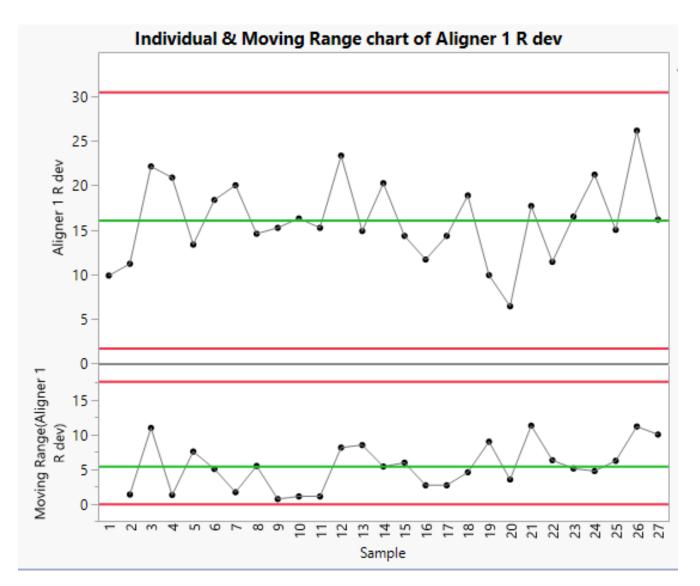
$UCL = \overline{R}D_4$	
$CL = \overline{R}$	
$LCL = \overline{R}D_3$	

Constants for sample size n

n	A ₂	D ₃	D ₄	d ₂
2	1.880	0.000	3.267	1.128
3	1.023	0.000	2.574	1.693
4	0.729	0.000	2.282	2.059
5	0.577	0.000	2.114	2.326
6	0.483	0.000	2.004	2.534
7	0.419	0.076	1.924	2.704
8	0.373	0.136	1.864	2.847
9	0.377	0.184	1.816	2.97
10	0.308	0.223	1.777	3.078

From Introduction to Statistical Quality Control by Douglas C. Montgomery

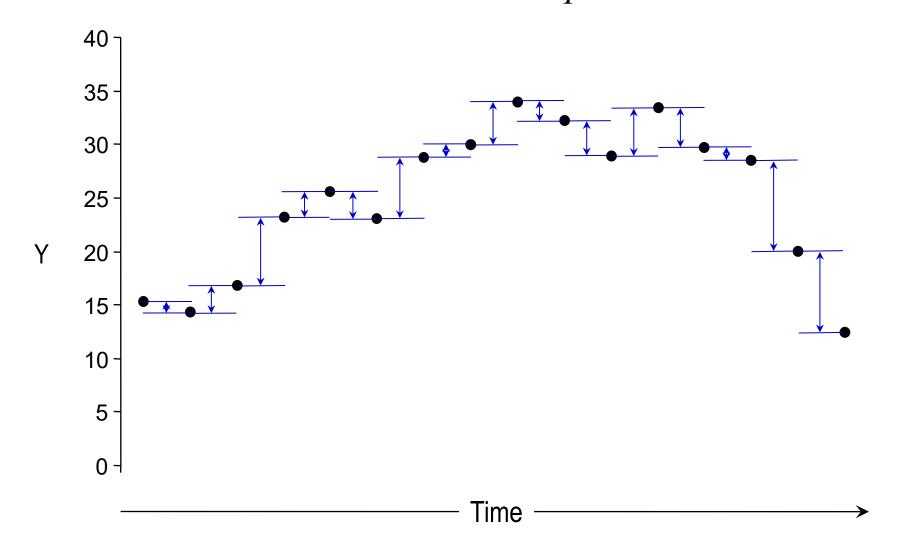
For each unit, the measurement is plotted on the Individual chart and the Moving Range is plotted below.



Why is the first point missing on the MR chart?

Estimating standard deviation using the moving range method

Each moving range is the absolute value of the difference between consecutive data points.



Individual Chart Control Limits:

UCL =
$$\overline{x} + 3 \frac{\overline{MR}}{d_2}$$

CL = \overline{x} MR = $|x_i - x_{i-1}|$
LCL = $\overline{x} - 3 \frac{\overline{MR}}{d_2}$

The value of d₂ is 1.128 since the range is between two consecutive points.

Moving Range Chart Control Limits:

 $UCL = D_4 \overline{MR} = 3.267 \overline{MR}$

$$CL = \overline{MR}$$

 $LCL = D_3 \overline{MR} = 0$

To make it easier to calculate the moving range, open Student Files \rightarrow calculator – individual moving range chart

B4	`		$f_x \sim = ABS(h$	A4-A3)					
	А	В	¢	D	E	F	G	Н	I.
1		Formula	s for n=2	Individual	Measureme	ents Chart	Movin	g Range C	hart
2	Data	Moving Ranges	Average Moving Range	LCL	CL	UCL	LCL	CL	UCL
3			0.0000	#DIV/0!	#DIV/0!	#DIV/0!	0.0000	0.0000	0.0000
4		0.0000							
5									
6									
7									
8									
9									
10		• •	'aste you	ir data ii	nto cell	A3			
11									
12		 Copy cell B4 down to the end of your data 							
13									
14									
15									

Excerpted data from *Data Sets*→ *solution properties*

A	В	С	D	Е	F	G	Н	I.
	Formulas for n=2		Individual Measurements Chart		Moving Range Chart			
Data	Moving Ranges	Average Moving Range	LCL	CL	UCL	LCL	CL	UCL
0.9239		0.0006	0.9214	0.9230	0.9246	0.0000	0.0006	0.0019
0.9233	0.0006							
0.9236	0.0003							
0.9224	0.0012							
0.9231	0.0007							
0.9224	0.0007							
0.9231	0.0007		 If Y ≥ 0 and LCL < 0, ignore LCL With MR calculations, the number of 					
0.9236	0.0005							
0.9230	0.0006							
0.9233	0.0003		decimal places shown may need to be					
0.9229	0.0004		increased					
0.9232	0.0003							
0.9225	0.0007							
0.9218	0.0007							

We want to use \overline{IX} and MR control charts to monitor radial deviation. This measurement requires special equipment and is very time-consuming, hence the sample size of one.

Open Data Sets \rightarrow control chart aligner

Open Student Files \rightarrow calculator - individual moving range chart

a) Copy the R dev data into the calculator (Paste Values).

b) Copy the calculation in cell B4 down Column B, in order to calculate the moving range for R dev. What is the average moving range?

 $\overline{MR} =$

Exercise 21.3 (cont'd)

c) What are the control limits for the Individual Chart?

UCL =

CL =

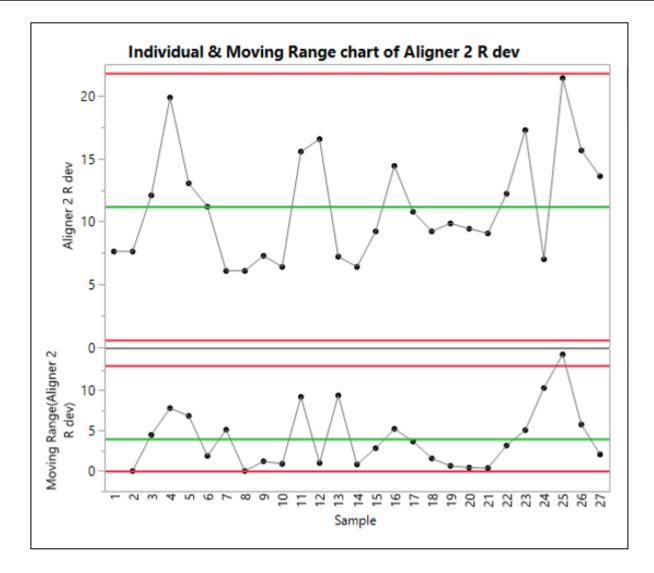
LCL =

d) What are the control limits for the Moving Range Chart?

UCL =

CL =

LCL =



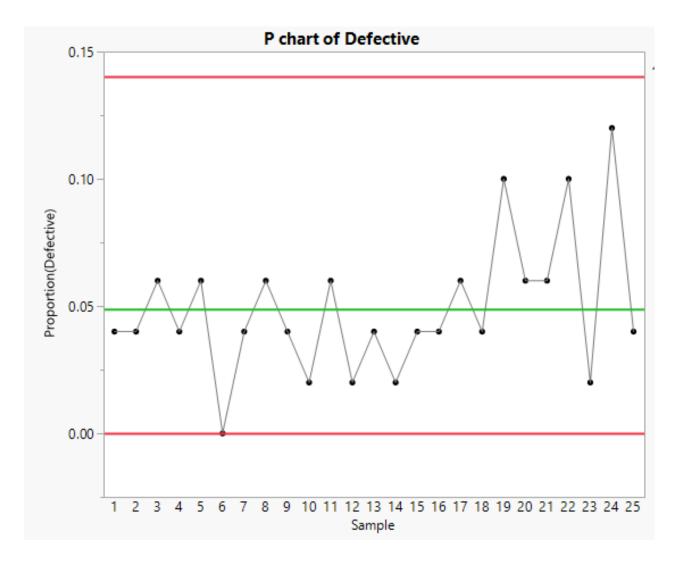
JMP Output of Individuals & MR Chart of Aligner 2 R dev Data Sets \rightarrow control chart aligner

The p Chart is used when:

- Samples are periodically taken and it's determined whether each unit in the sample is good or bad
- The data plotted is fraction or percent defective
- P Chart control limits are based on the Binomial distribution, since pass/fail data is binomial.
 - The standard deviation of the Binomial distribution is:

$$\sqrt{\frac{p(1-p)}{n}}$$

Example of a p Chart (created in JMP)



In this case, there were 50 units in each sample. Overall percent defective was about 5% for this timeframe.

Control Limits for the p Chart

$$UCL = \bar{p} + 3\sqrt{\frac{\bar{p}(1-\bar{p})}{n}}$$
$$CL = \bar{p}$$
$$LCL = \bar{p} - 3\sqrt{\frac{\bar{p}(1-\bar{p})}{n}}$$

 $\overline{p} = rac{\text{Total number of defective units in the samples}}{\text{Total number of units in the samples}}$ n = number of items in each sample

These control limits are the mean +/- *3 sigma for this distribution.*

Exercise 21.4

We want to use a percent defective (p) control chart to monitor the weekly defects per unit occurring during an in-process assembly inspection.

Open Data Sets \rightarrow control chart parts inspected & defective

Use Excel formulas for the following and during calculations, keep the numbers in "fraction defective" form vs percentage:

a) The sample size varies each week, so we'll use an average sample size for calculating control limits. Calculate the average weekly sample size. What concerns might there be about using this number?

b) Calculate the overall percent defective.

This number will be the center line (CL) for the p chart.

c) Use the average sample size and \overline{p} found above to calculate the upper and lower control limits for the p chart.

UCL =

CL =LCL =

d) Optional: Copy the formulas for the control limits down the column for all of the data and use line charts to plot the fraction defective with control chart limits.

Categorical classification:

- np chart: number (count) of defective items per sample with a fixed quantity
- u chart: count of defects per unit
- c chart: count of defects) per sample with a fixed quantity

For np, c and u charts, the control limit calculations and chart appearance are similar to the p chart.

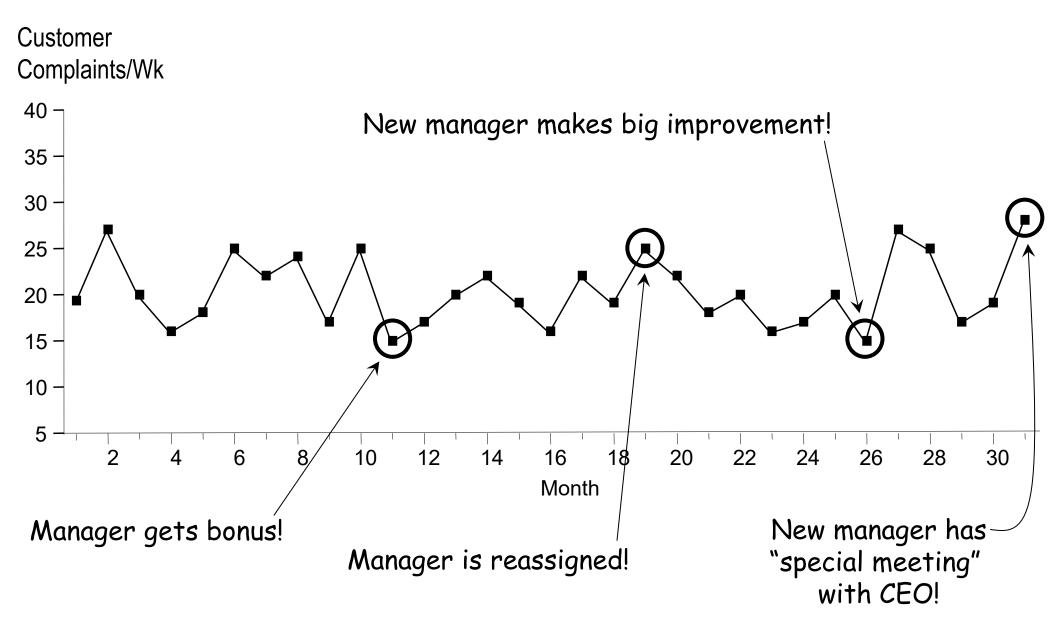
Details of these and other specialized control charts are beyond the scope of this course. More information can be found in any basic statistical process control textbook or reference.

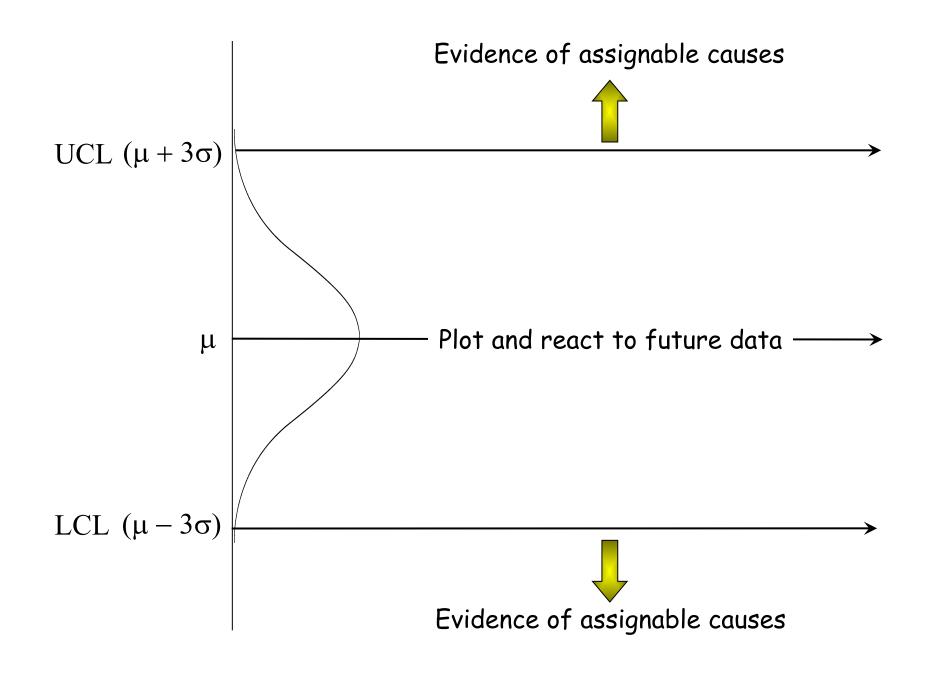
Once the control chart is created, the most valuable work can begin — discerning what the chart is telling us about process variation.

- Is the process "in control" or "out?"
- Are there warning signs that the process may go out of control soon?
- What actions should be take in response to the control chart signals?

The rules we'll discuss for deciding whether a process is in or out of control work only for control limits — *not* for specification limits.

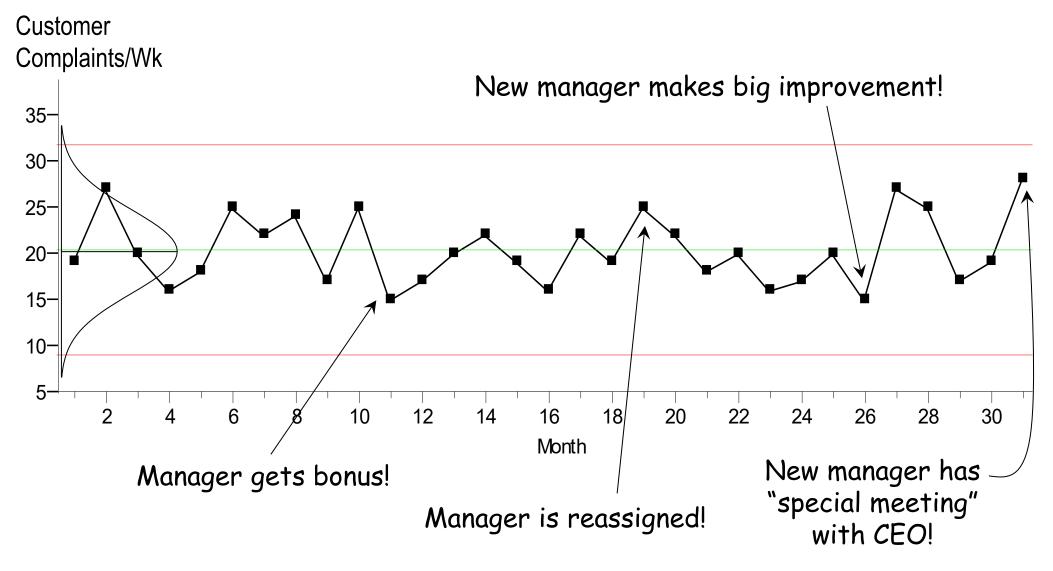
- Our concern with specification limits is whether an item conforms or not.
- Inspection and testing must be used to screen out bad parts, not control limits.





Control Limits show there are no assignable causes!

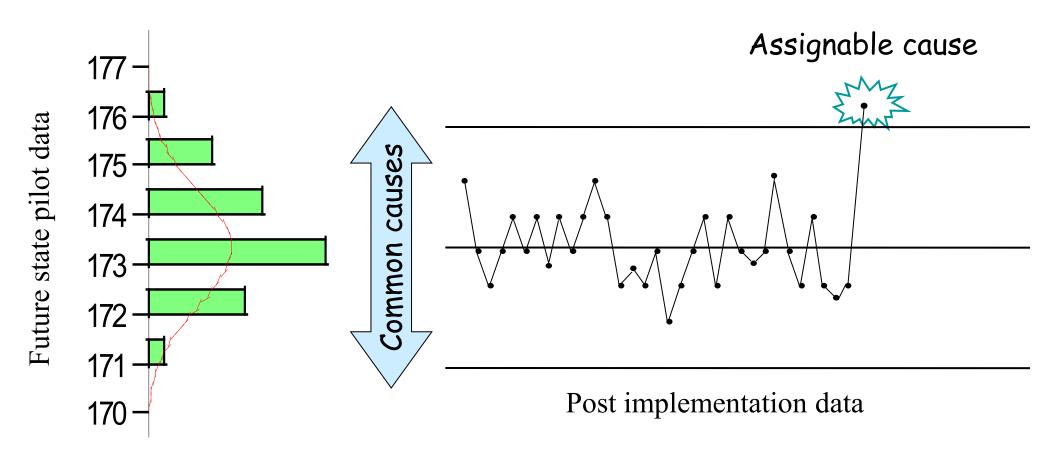
Run charts can cause us to overreact.



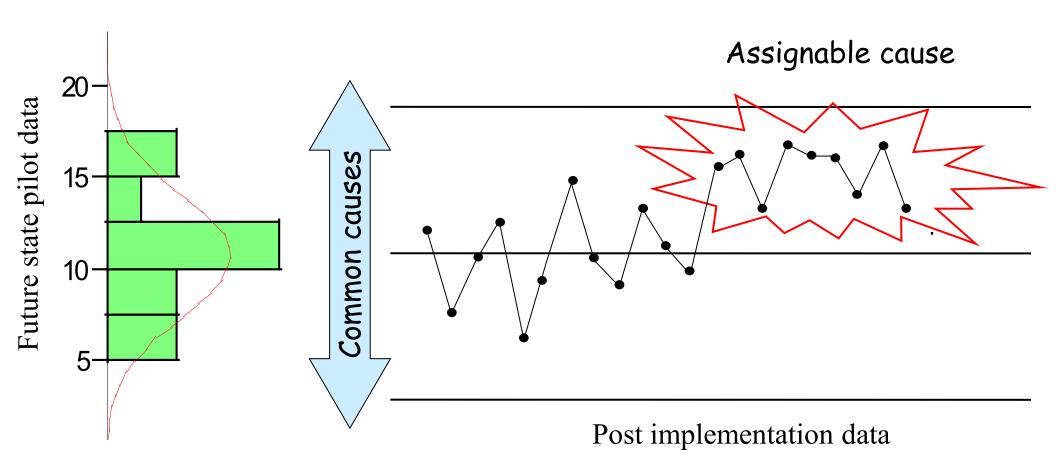
- Control limits provide an operational definition of assignable cause variation
- Simplest rule: points inside the limits are common cause variation, points outside the limits have assignable causes
- 27 in 10,000 common cause data points are expected to fall outside the control limits^{*} this is the nominal *false alarm* rate
- Assignable causes may occur without producing points outside the limits these are *missed signals*
- To reduce missed signals, additional rules are sometimes applied

When monitoring a straightforward KPI, such as number of customer complaints/week or monthly on-time delivery, Management may only want to see a chart of the KPI metric itself.

- In this case, it may be sufficient to use an X-bar or IX chart without the associated standard deviation or range chart.
- Adding control limits to the resulting X-bar or IX chart will provide a statistical basis for action.
- It may also be helpful to add a target or goal line to the chart (aligned with the KPI calculation method).
- An associated variation chart could be created for deeper root cause analysis if necessary. For example:
 - > Are late deliveries "normal" for the organization?
 - > Are there inconsistencies between divisions for global KPI charts?



- 1. This event has probability 0.00135 ($0.0027 \div 2$)
- 2. Investigate to determine the cause
- 3. Take corrective action to eliminate the cause



- 1. This event has probability $0.00195 [(0.5)^9]$
- 2. Investigate to determine the cause
- 3. Take corrective action to eliminate the cause

418

Control chart zones: A, B, and C

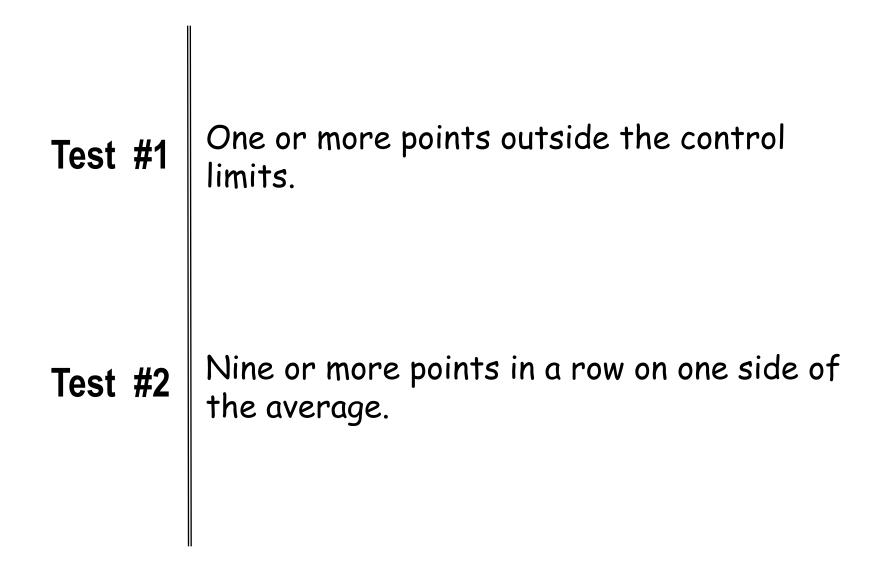
	UCL
A	
B	
С	Avg
С	
В	
A	LCL

Additional tests for assignable causes (cont'd)

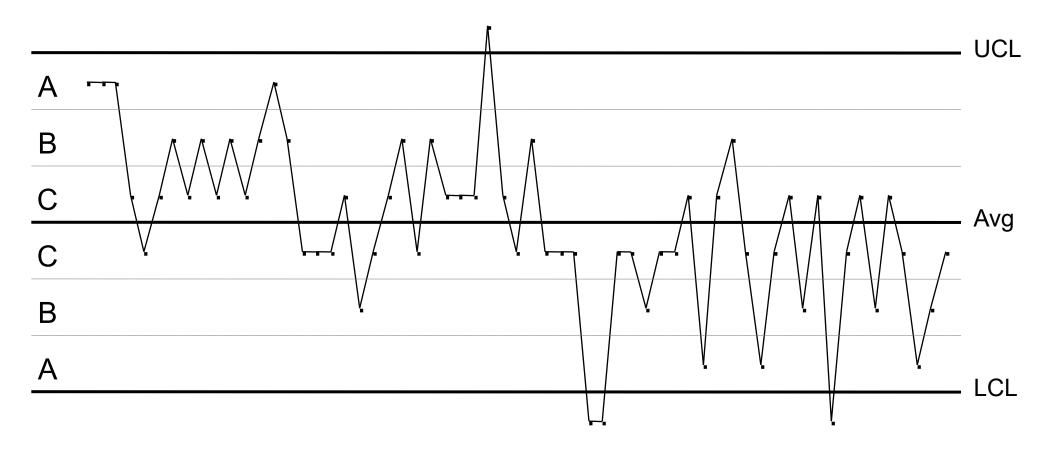
Test 1	One point beyond A (This is the basic test & always used.)
Test 2	9 points in a row on the same side of the average.
Test 3	6 points in a row steadily increasing or decreasing.
Test 4	14 points in a row alternating up and down.
Test 5	Any 2 out of 3 points in a row in A or beyond.
Test 6	Any 4 out of 5 points in B or beyond.
Test 7	15 points in a row in C, above and below the center line.
Test 8	8 points in a row on each side of the average with none in C.

The zone system is based on 3σ limits

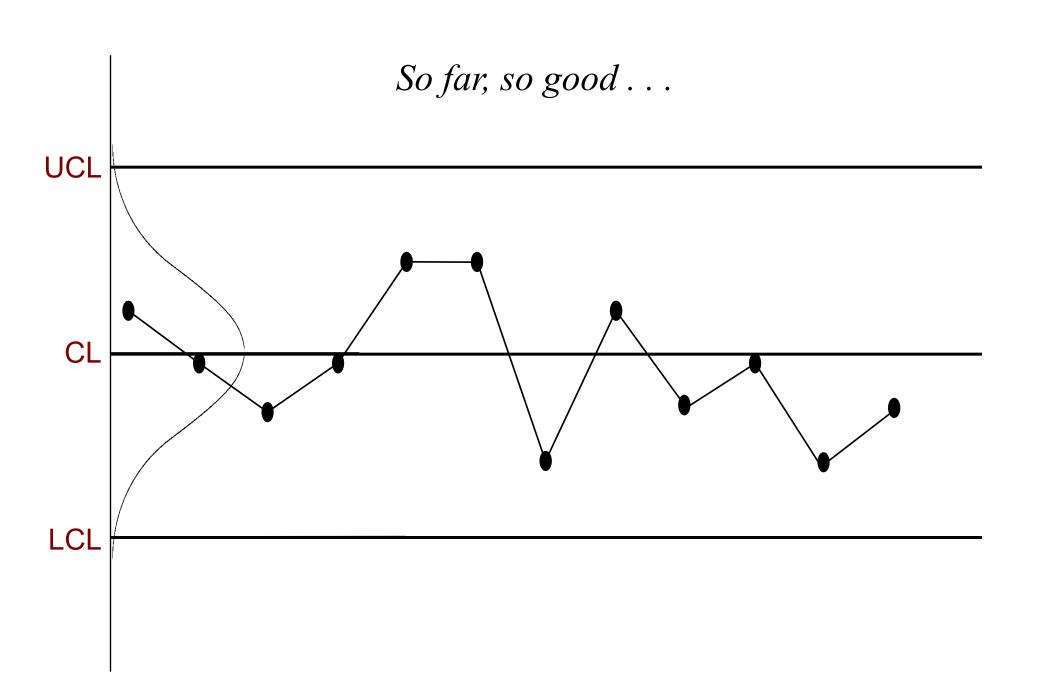
- \cdot C is the region within 1 standard deviation of the mean
- B is the region more than 1 but less than 2 standard deviations from the mean
- A is the region more than 2 but less than 3 standard deviations from the mean



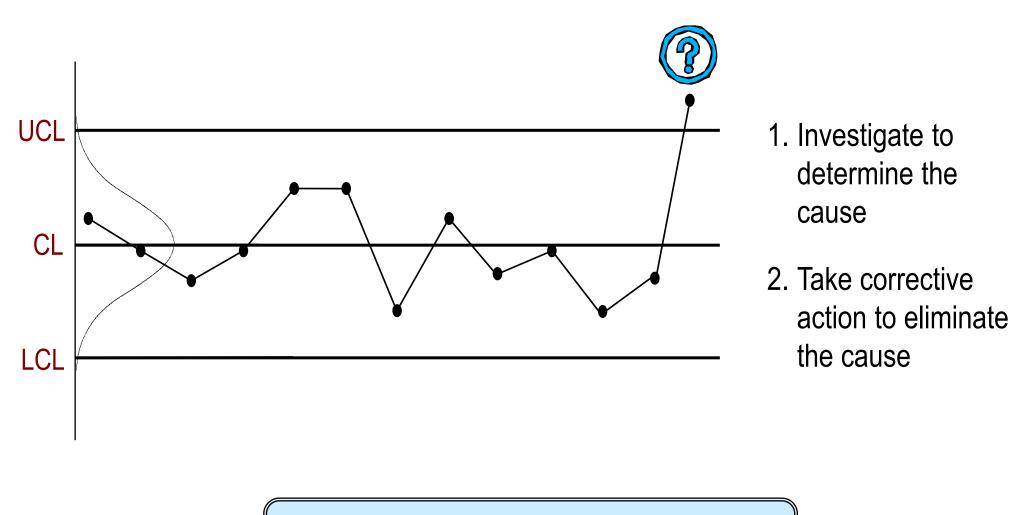
Circle occurrences of Tests 1 and 2 on the control chart shown below. Indicate which is which.



22 Response plans

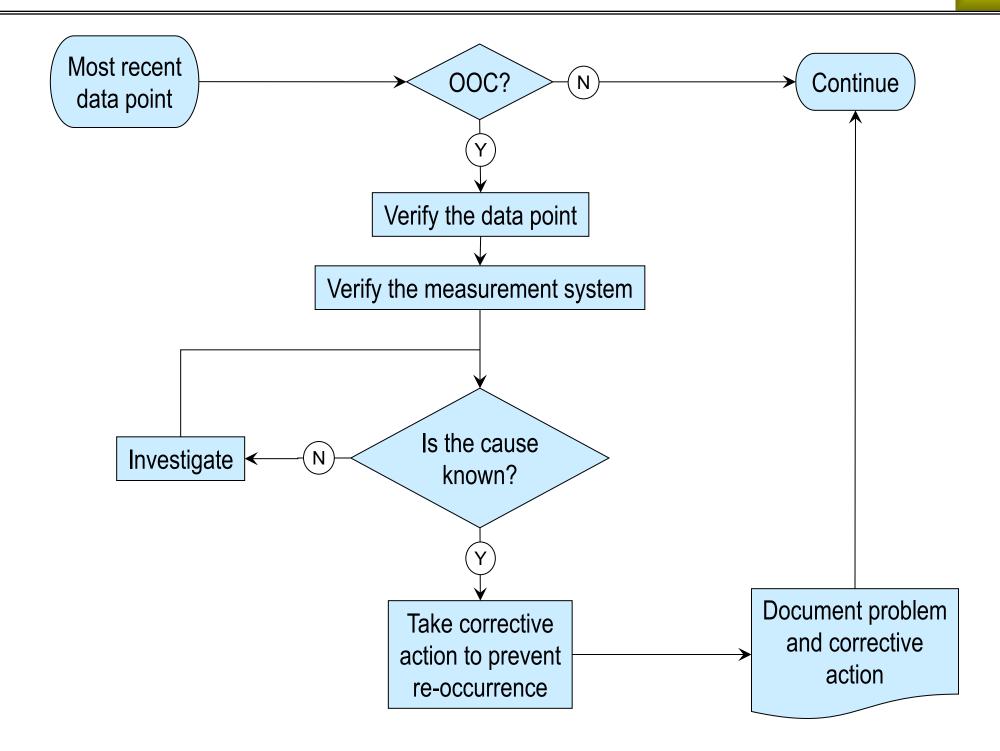


Out-of-control event (OOC)



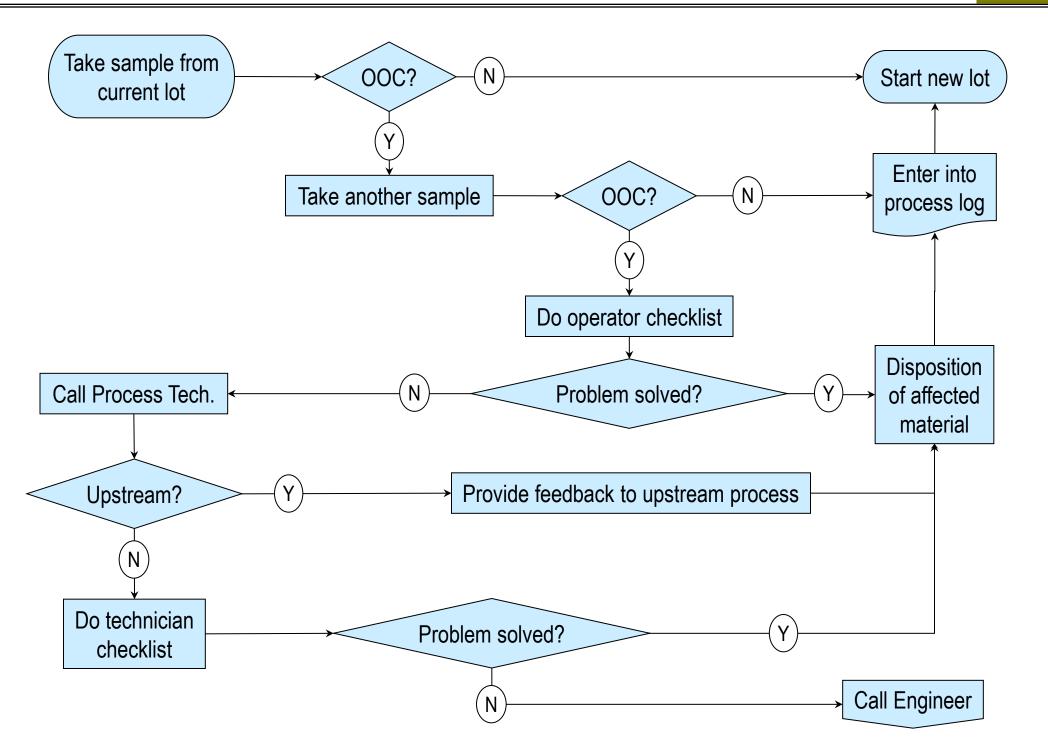
Step 1 requires a response plan

Response plan "skeleton"

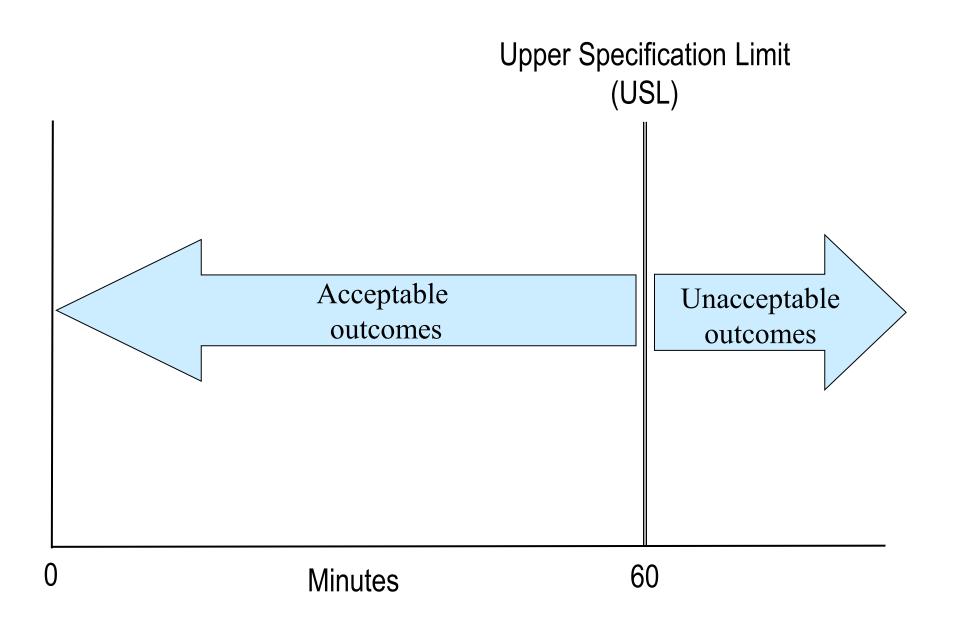


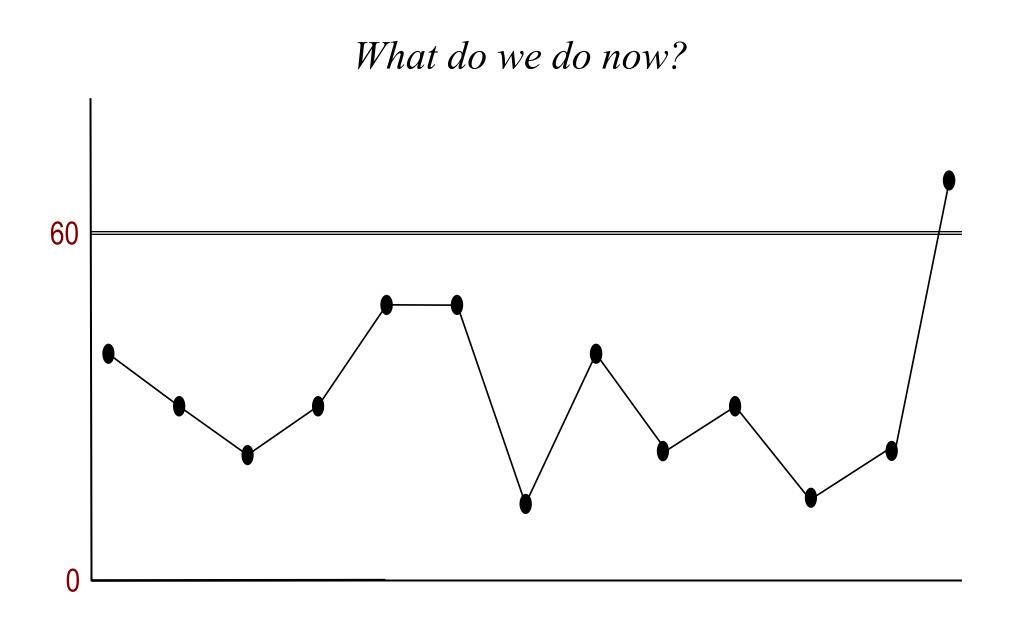
- OOC stands for *out of control*
- This means the control chart indicates an assignable cause according to one or more selected tests
- The success of statistical monitoring depends on having a documented plan for responding to OOCs
- The most effective form of documentation is a process map like the one shown above
- It should posted in a place clearly visible to process participants

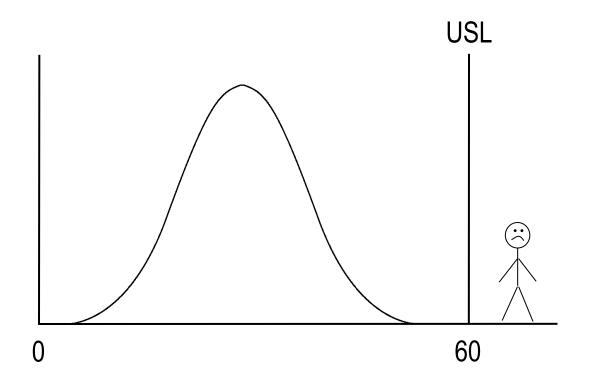
Response plan example



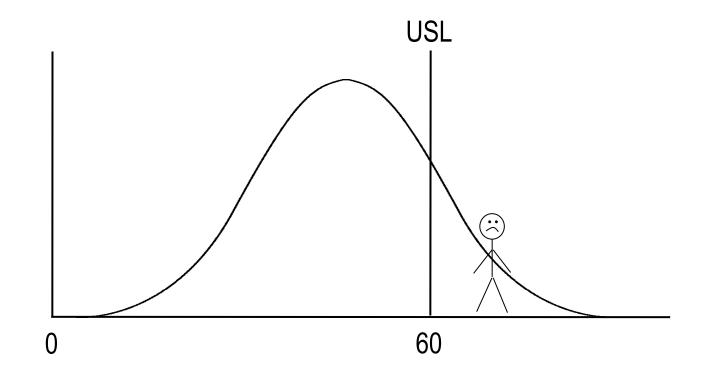
- Example from a high-volume automated assembly process ("sanitized")
- Development team: operators, technicians, engineers, area manager
- Based on experience, they wanted to verify an OOC with a second sample from the same lot
- Note the escalation from Operator to Technician to Engineer.
- When an OOC was confirmed, production was halted
- Within a few months:
 - Chronic equipment and process problems were solved
 - Unplanned downtime and need for Engineering support plummeted
 - Engineers able to focus more on process improvement
 - Productivity increased dramatically



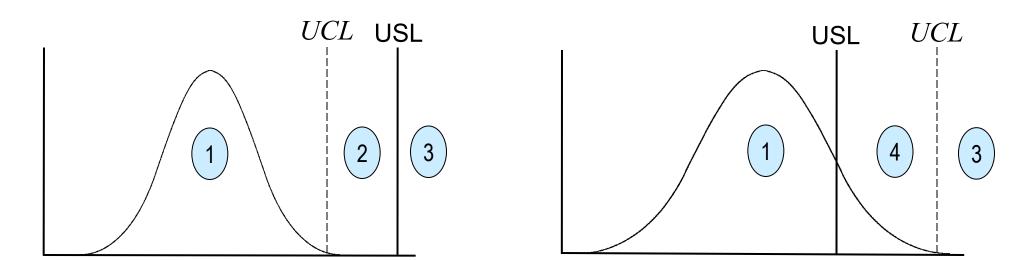




- If the process has good capability, it will virtually never produce a defective outcome, unless there is an assignable cause
- Any OOS point is also OOC
- Any OOS point should trigger the response plan



- If the process has poor capability, there will be OOS outcomes that are not OOC
- These outcomes do not indicate assignable causes
- They should *not* trigger the response plan



Check the appropriate actions for outcomes in each of the 4 zones shown above.

Zone	Initiate OOC response plan	Scrap, rework, do over, etc.	Do nothing
1			
2			
3			
4			