

Measuring Sterile Processing Success

High Reliability

## High Reliability: Measuring SPD Success

- Explain the importance of high reliability in sterile processing.
- Describe factors that contribute to people and process failures.
- Explain how a quality assurance program can create a highly reliable SPD.





# **High Reliability**

#### **PEOPLE**

Does everyone have the skills to perform in every area (decon, HLD, Assembly, Sterilization, Case Cartss, ect.)

Does the department suffer when certain people are not there?

#### **PROCESSES**

 Are processes clearly understood?

 Are processes seamless with organized steps and supplies readily available.

## **Definitions:**

• Proficient – Competent or skilled in doing or using something.

• **Perfect** - make (something) completely free from <u>faults</u> or defects.



# **People and Process Failures**



#### Why aren't people proficient?

- No or incomplete training
- No performance accountability



### **PROCESSES**

#### Why aren't processes perfect?

- Disorganized environment
- Inconsistent supply availability
- Location of supplies
- Broken equipment



# Implementing a Continuous Quality Improvement Program

An effective Continuous Quality Improvement (CQI) program requires *education* and *quality audits* that will support people and improve processes continuously.

# **Collect and Review Surgery Tray Error Data**

2025 January		2025 February	
Bioburden	5	Bioburden	3
Missing Locks	6	Missing Locks	1
Missing Filter	2	Missing Filter	0
Missing Required	12	Missing Required	10
Missing Indicator	О	Missing Indicator	4
Misc.	9	Misc.	11
TOTAL ERRORS	34	TOTAL ERRORS	29
Total Sterilized	11,188	Total Sterilized	10,628
Trays	9178	Trays	8864
Peel Pouch	2010	Peel Pouch	1764

## **Conduct and Document Random Audits**

#### **Process Audits**



Follow techs through a process to observe compliance



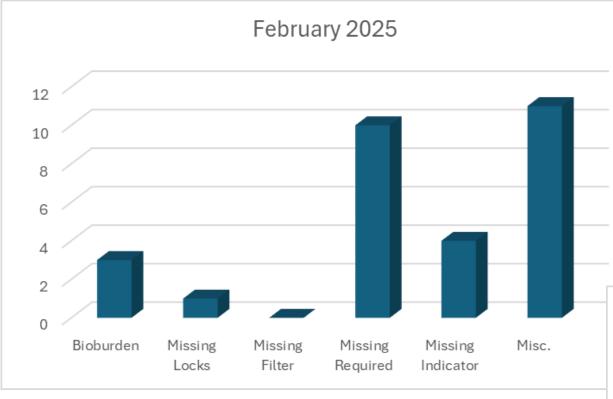
Look for process deviations and ask clarifying questions

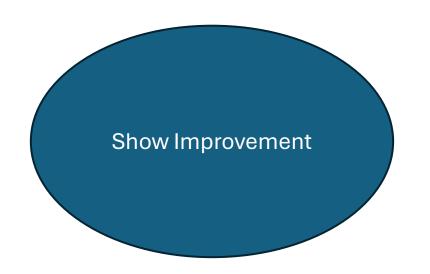


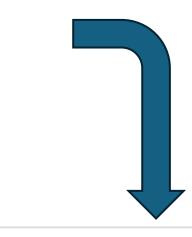
Evaluate process perfection

#### **Product Audits**

- Inspect trays from the washer
- Open and inspect sterile trays
- Inspect completed case cart
- Inspect sterilization and HLD documentation
- Assign education as needed

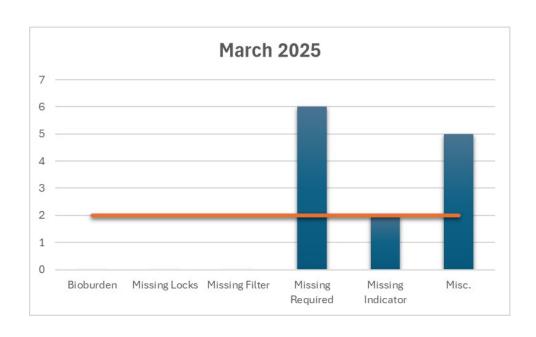








## Prepare to Share Your Reliable Data



- Inspection process created to ensure unsterile loads are inspected for locks and filters prior to sterilization.
- •Corrective action plan developed in partnership with HR to hold technicians accountable for bioburden errors.

Use charts

 Bullet point actions taken to make improvements

Suggest a baseline for your data

# **High Reliability Review**

- Proficient People + Perfect Processes = High Reliability
- Incomplete orientation and or broken processes prevent highly reliable sterile processing service.
- Implementing a continuous quality improvement program can help develop a highly reliable sterile processing service.
- Review errors and allow it to guide audits to identify education and or process requirements
- Document data and share improvements



