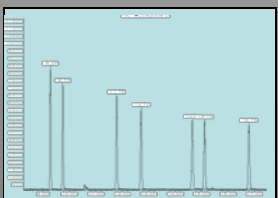
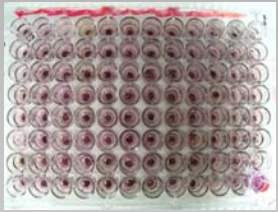


Development & Implementation of a Lean Process Implementation Plan

Jeffrey P. Massey, Dr.P.H., HCLD(ABB)
Quality Manager
Michigan Department of Community
Health Bureau of Laboratories



Lean Waste

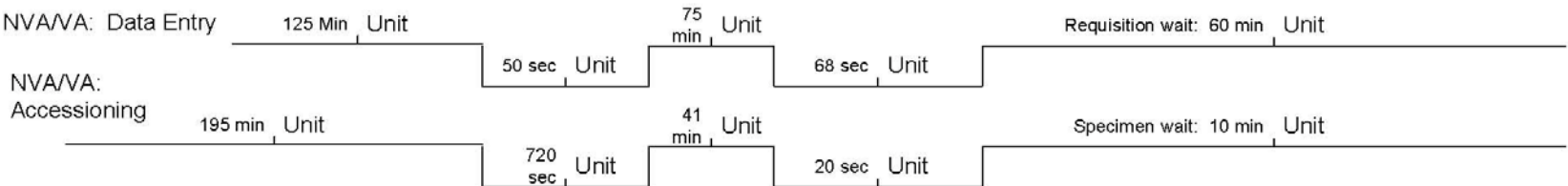
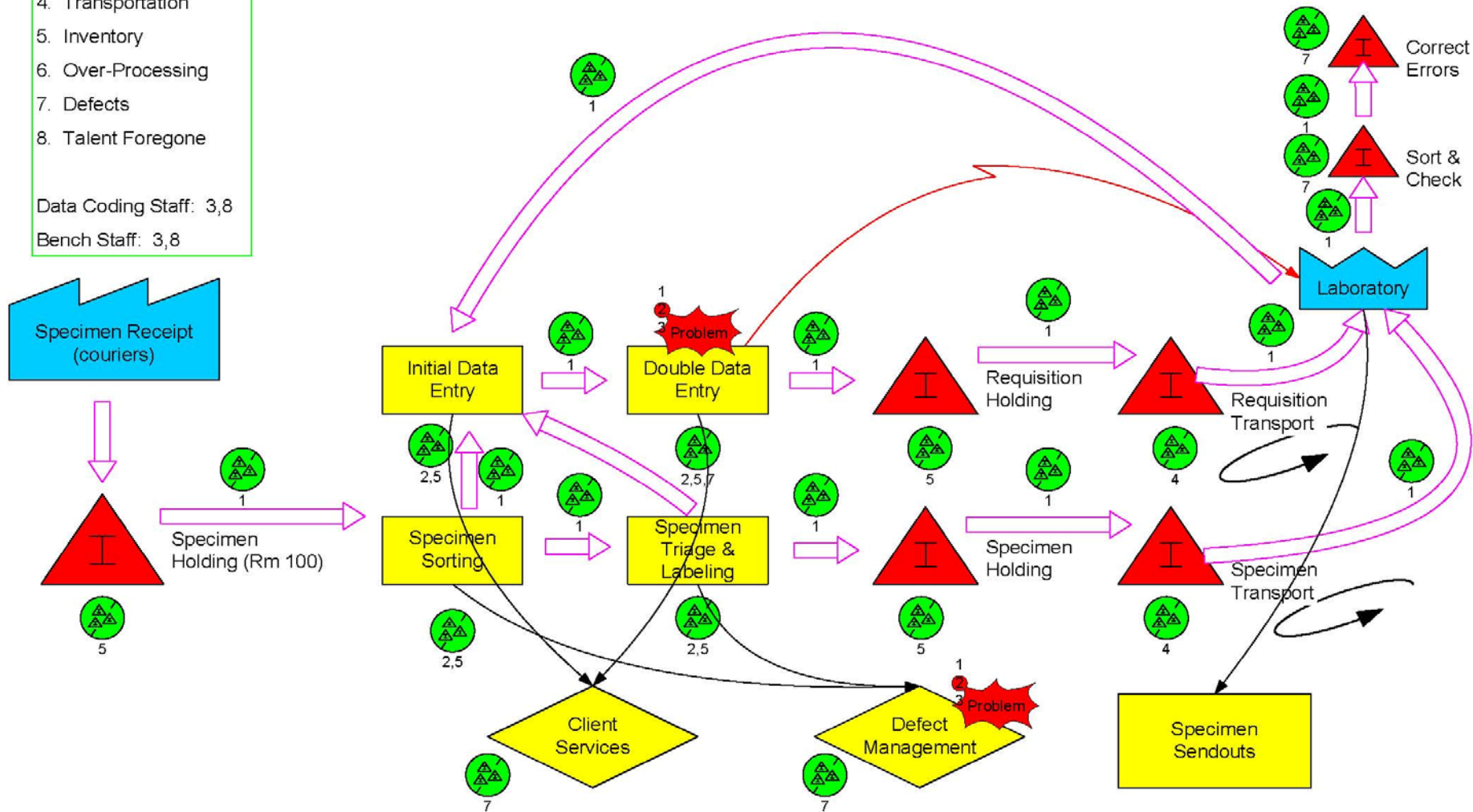
1. Overproduction
2. Waiting
3. Motion
4. Transportation
5. Inventory
6. Over-Processing
7. Defects
8. Talent Foregone

Data Coding Staff: 3,8

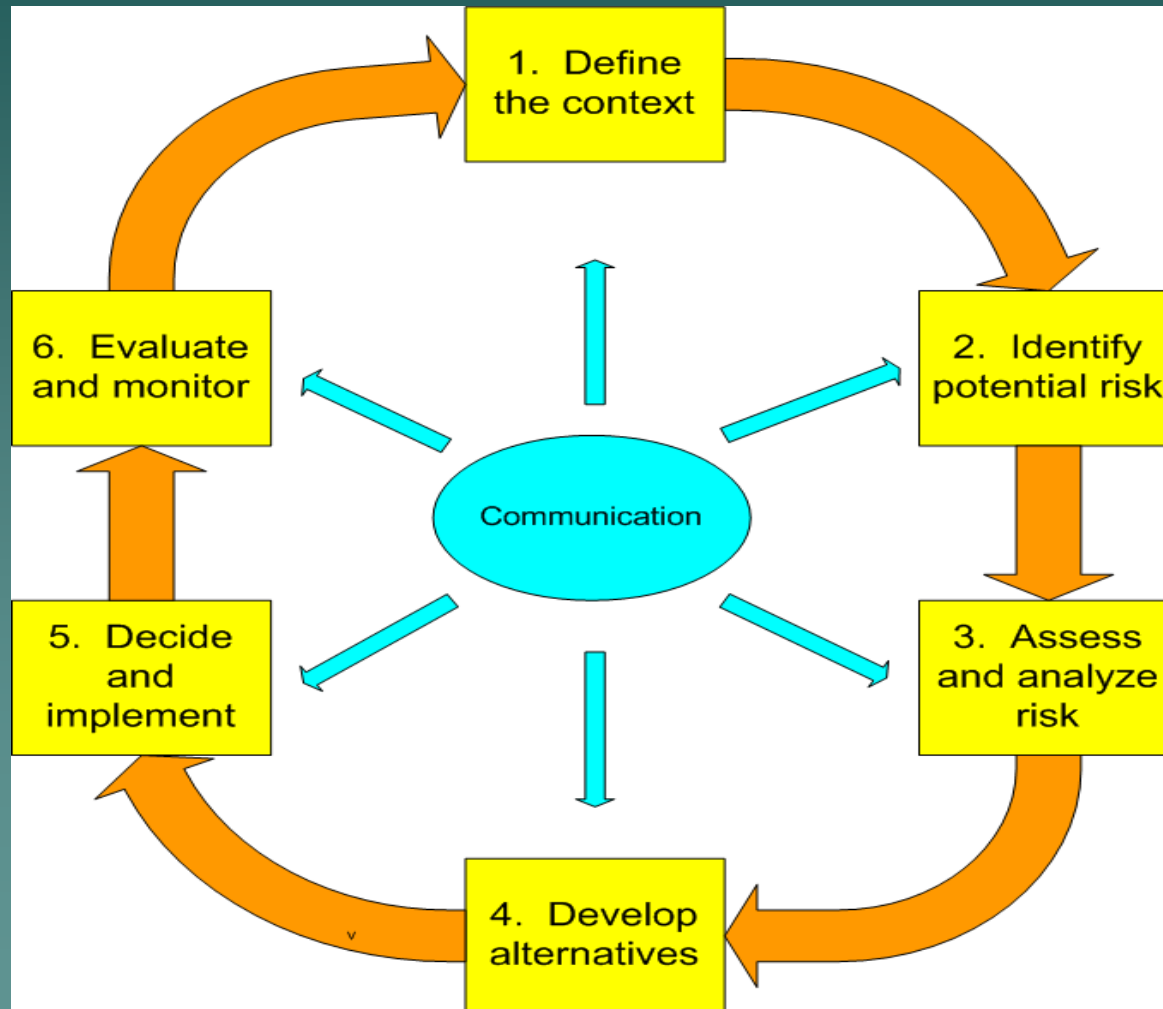
Bench Staff: 3,8

Current Value Stream Map:

DASH Unit – Specimen Accessioning & Data Entry



Risk Management Cycle

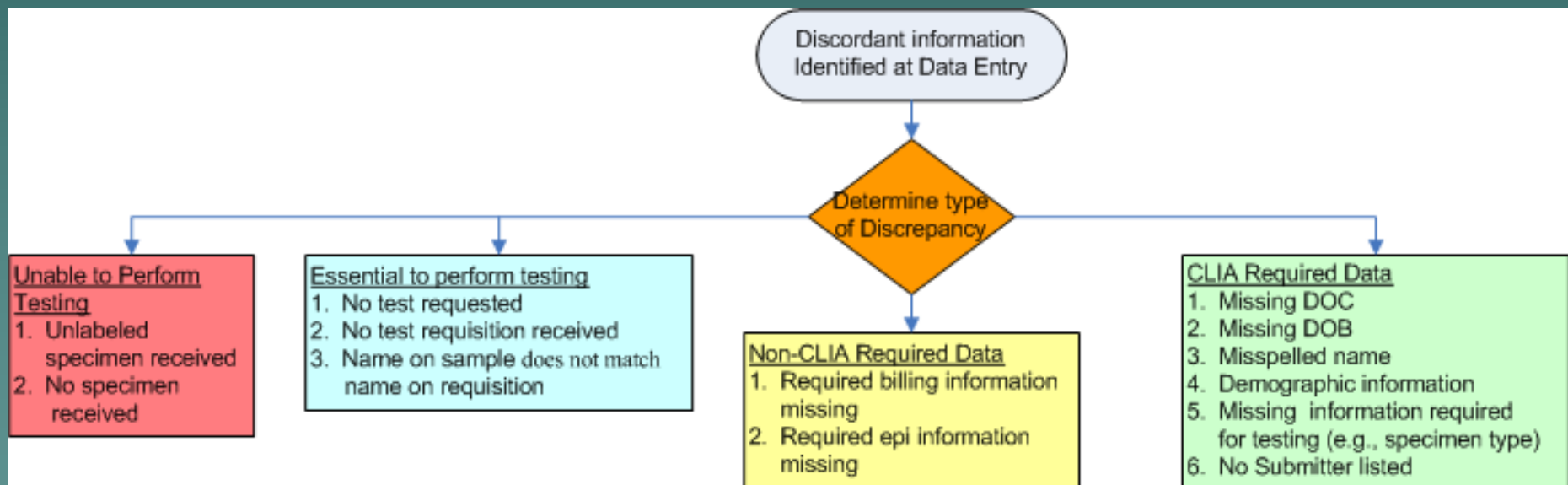


Problem Identification

- Specimens tested – reports held if discordant information present
- No definitive lab policy for reporting results with discordant information
- Regulatory deficiency – reports held past established TAT

Brainstorm: Define Scope of the Problem

- Define type of discordant information
 - What type of error will suppress reports?



Type of Mismatches

- Missing information
- Discrepancy between specimen & requisition
- Wrong or no specimen received
- Unauthorized specimen received
- Test not available
- Specimen improperly submitted or leaking

Will report be suppressed?

- Missing information (YES)
- Discrepancy between specimen & requisition (YES)
- Wrong or no specimen received (NO)
- Unauthorized specimen received (NO)
- Test not available (NO)
- Specimen improperly submitted or leaking (NO)

Define Current Process

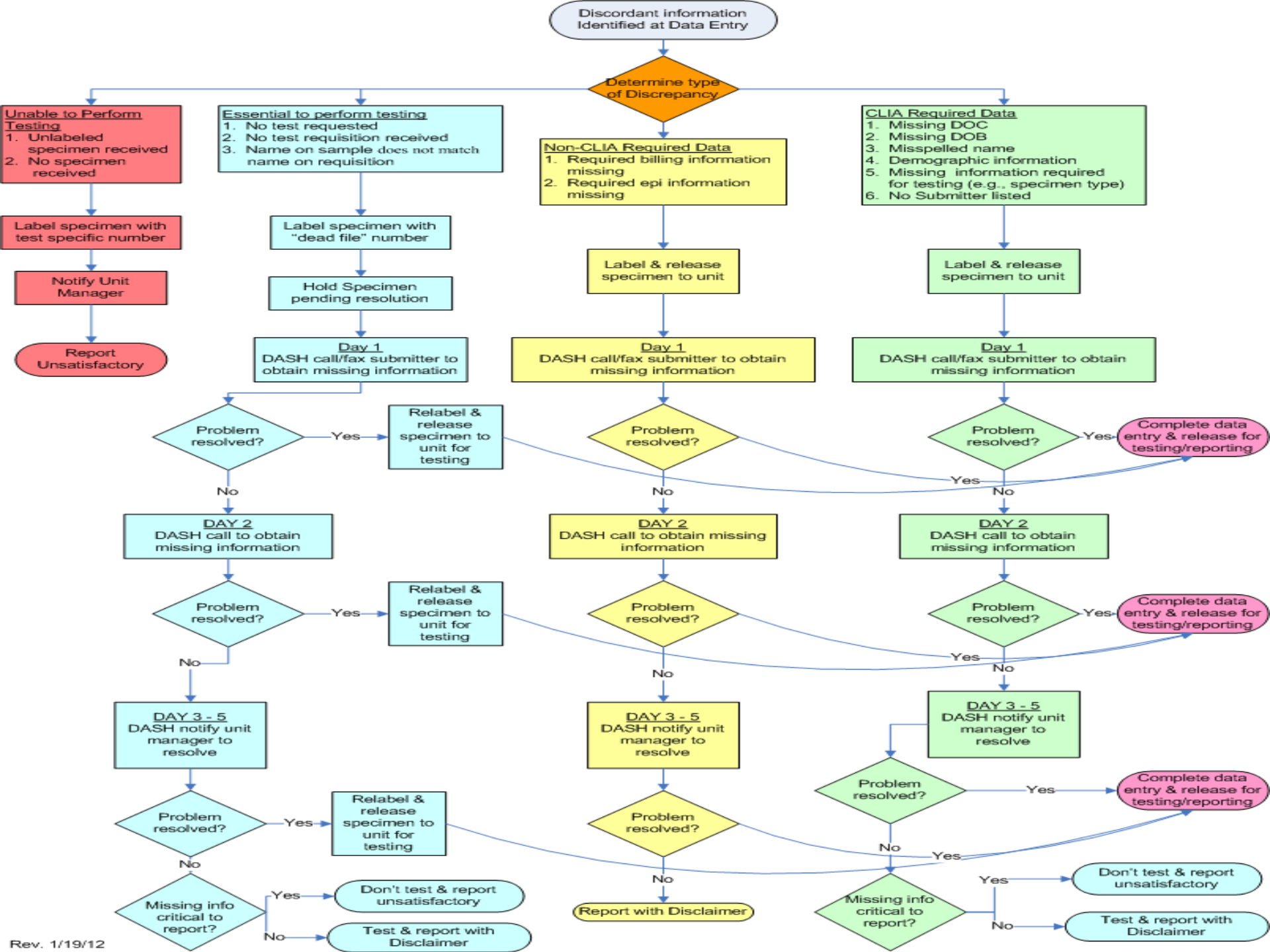
Error identification during specimen triage

- Day 1: fax correction form
- Day 7: initial follow-up via fax
- Day 14: telephone follow-up via phone
- Day 30: final follow-up
- Day 30: report unsatisfactory if not resolved

Define Future State

- Day 1: DASH detects error during specimen triage
- Day 1: DASH calls submitter to resolve. Release specimen for testing/reporting if resolved.
 - Fax requisition if change must be made in writing.
- Day 2: If unresolved, DASH makes repeat contact.
- Day 5: If still unresolved, manager contacts submitter to resolve.

If still unresolved, manager reports with disclaimer or reports as unsatisfactory.



- CLIA Required Data**
1. Missing DOC
 2. Missing DOB
 3. Misspelled name
 4. Demographic information
 5. Missing information required for testing (e.g., specimen type)
 6. No Submitter listed

Label & release specimen to unit

Day 1
DASH call/fax submitter to obtain missing information

Problem resolved?

Yes - Complete data entry & release for testing/reporting

No

DAY 2
DASH call to obtain missing information

Problem resolved?

Yes - Complete data entry & release for testing/reporting

No

DAY 3 - 5
DASH notify unit manager to resolve

Problem resolved?

Yes - Complete data entry & release for testing/reporting

No

Missing info critical to report?

Yes - Don't test & report unsatisfactory

No - Test & report with Disclaimer



Pilot Study

One week period, document:

and type of mismatches

resolved at Day 1

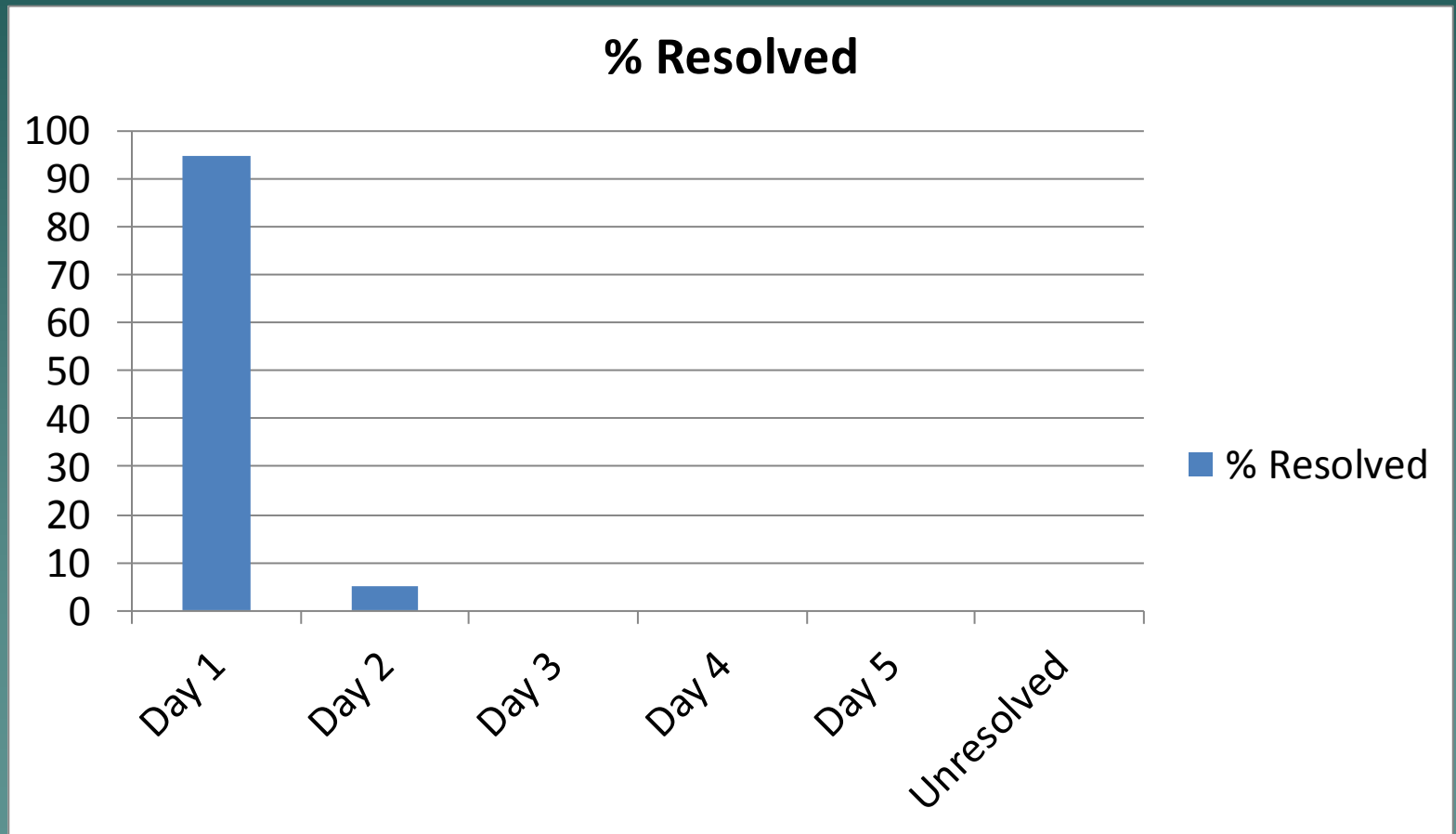
resolved at Day 2

resolved by Manager

unresolved & reported with disclaimer

unresolved & reported as unsatisfactory

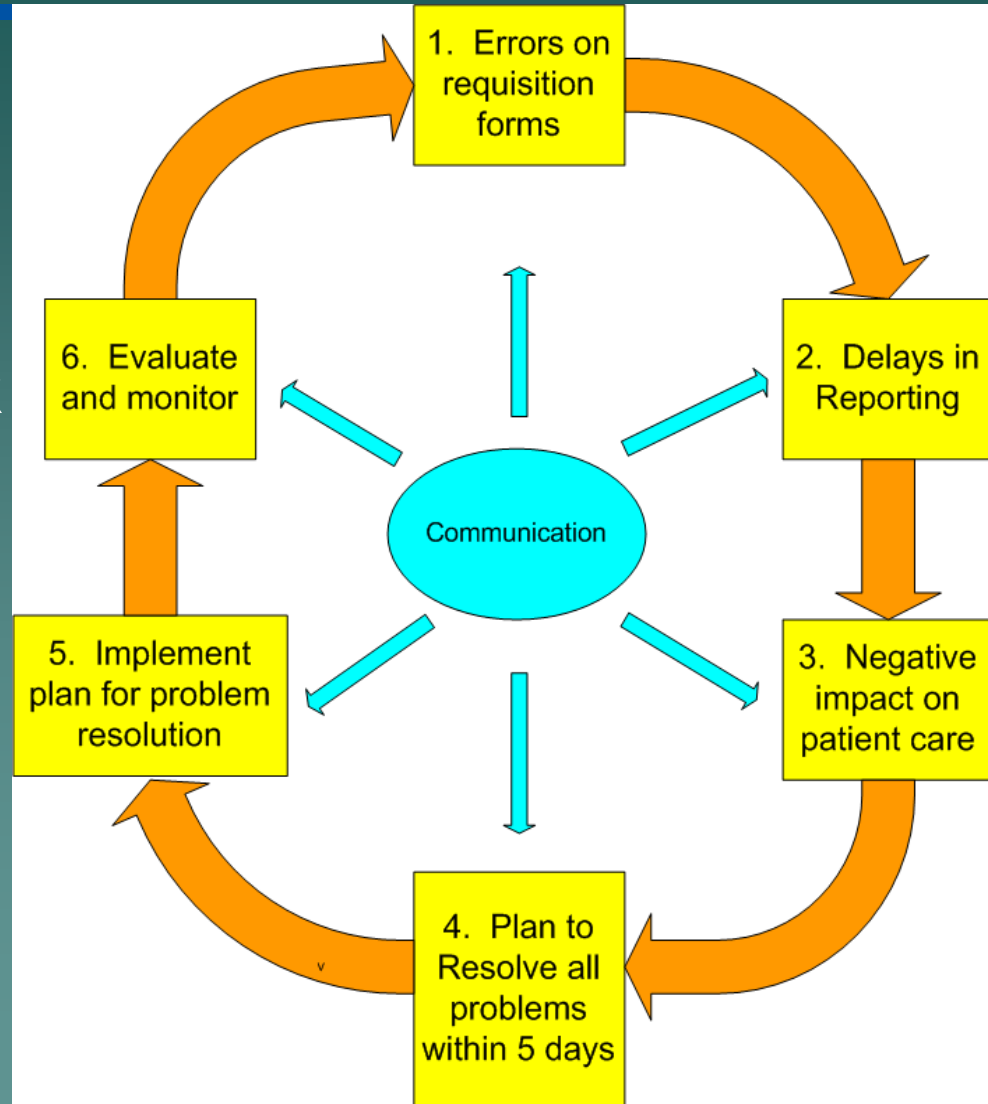
Assessment of Effectiveness (following implementation)



N = 40

Risk Management Cycle

- 1. Define the context
- 2. Identify potential risk
- 3. Assess & analyze risk
- 4. Develop alternatives
- 5. Decide & implement
- 6. Evaluate & monitor



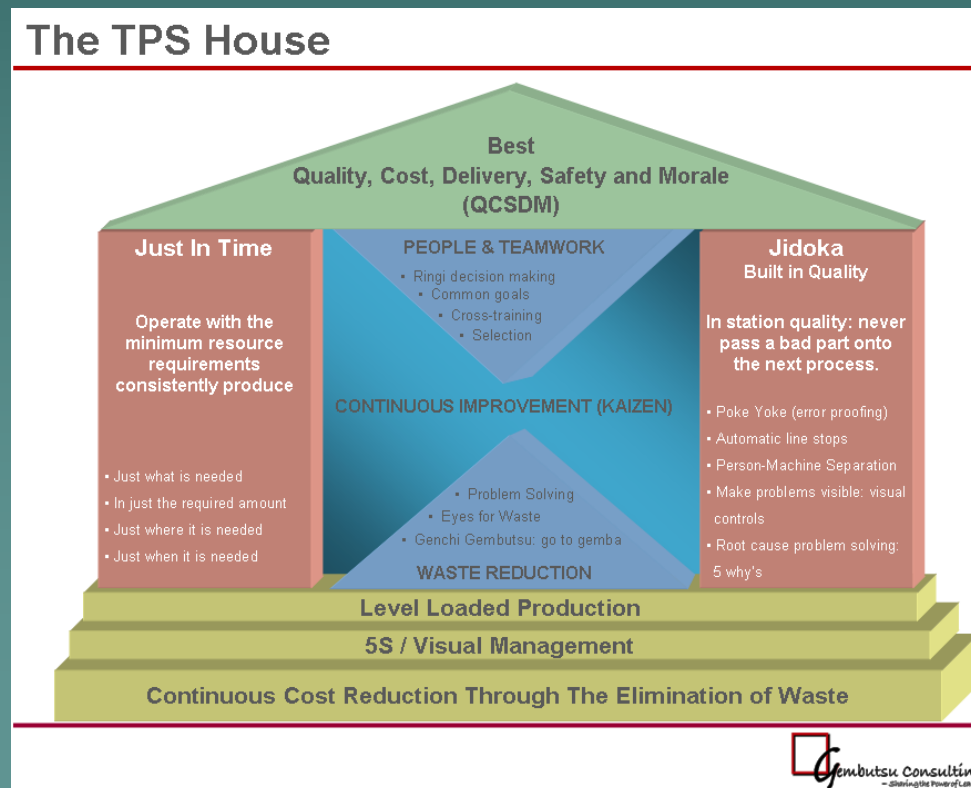
Lessons Learned

- Understand current process before implementing a change
- Get input from staff doing the work before implementing a change
- Communication essential
 - Get input
 - Share ideas
 - Convey results & educate

Lean is Great When it Works, What Could Possibly Go Wrong???????

Problems

- Make changes without prior planning or establishing a foundation of quality



Questions???

