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An Approach to Quality Improvement Projects in Cytopathology

Gavin Giles (Coordinator Cytopathology, SRA and Autopsy)

Quality Improvement

What is QI?

- ❑ Systematic, data guided, activities designed to bring about immediate improvement in health care delivery.
 - More specifically, activities that seek to improve outcomes such as reducing atypical rate, insufficiency rate, diagnostic error rates or shortening turnaround time.

LHSC PaLM Quality Improvement

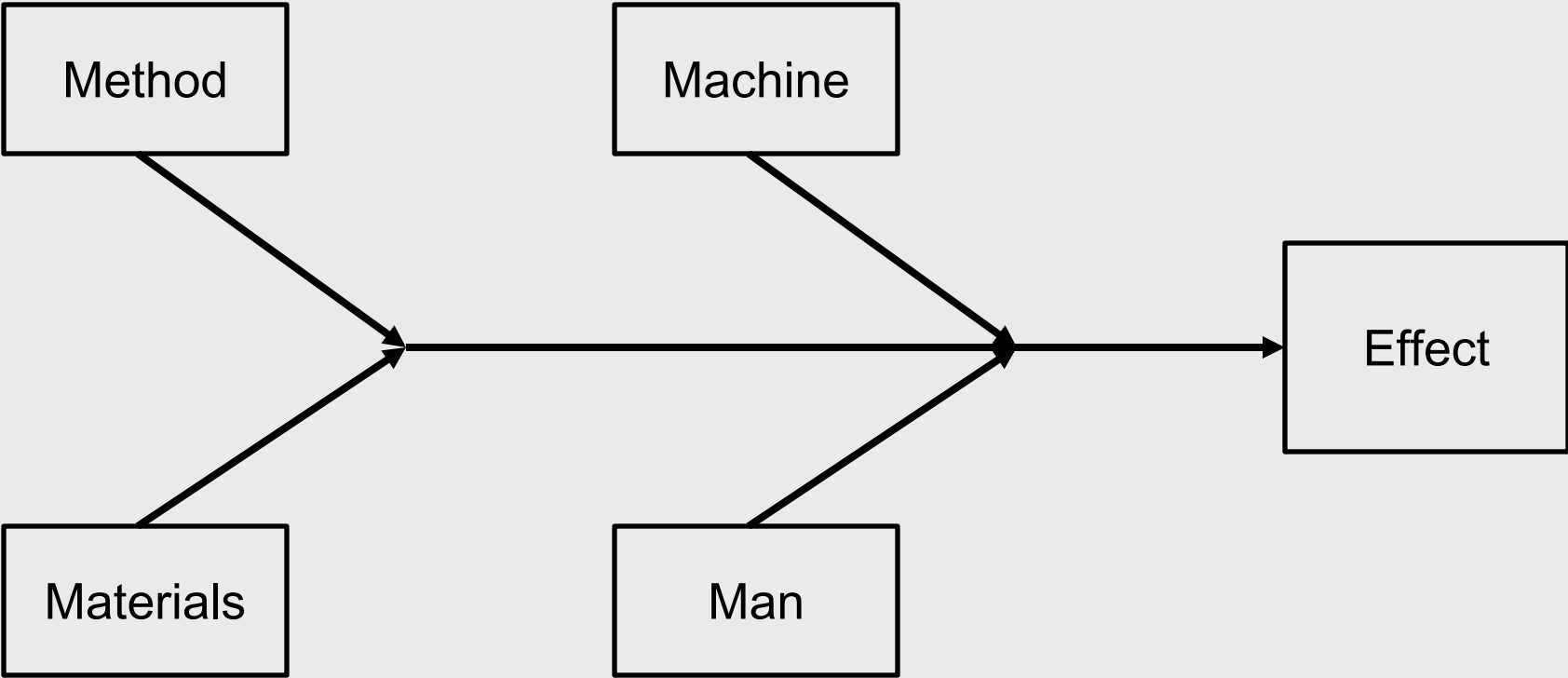
Approach:

- ❑ Focused on solving everyday problems
 - Simple problems that are observed at Gemba
 - Large scale issues are addressed via special projects
- ❑ Team-based
 - Representation from all relevant sub-teams and leadership
 - Resources to gather and summary data from issue forms
 - Expert resource to guide use of quality improvement tools
 - Meetings to ensure discussion of all perspectives

Method

- ❑ Process Review
 - Map current state
- ❑ Problem Description
 - What + How + Which + When + Where + Who = Problem Description (Problem Statement)
- ❑ Determination of Root Cause
 - Categorize or group potential causes in a clear and consistent manner
 - E.g. Fishbone Diagram to explore 6Ms (Man, Machine/Tools, Materials/Inputs, Methods, Measures, Environment)

Fishbone Diagram



5 Whys

Possible Cause	Why?	Why?	Why?	Why?	Why?
	???	???	???	???	???
	Because ↗	Because ↗	Because ↗	Because ↗	Because ↗

- 5 Whys are typically required to dive deep enough to get to the root cause

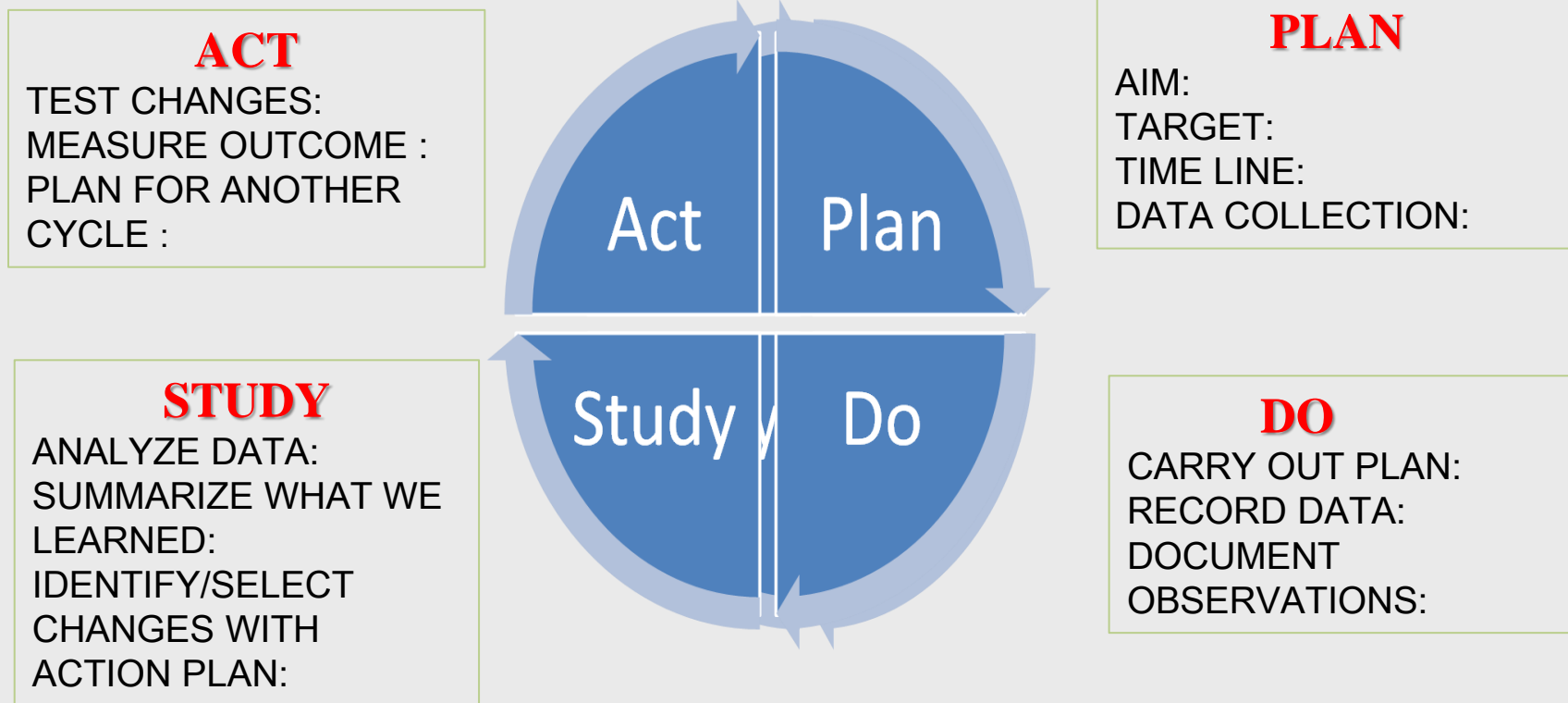
Method

□ Action Plan Development

- What will we do?
- Who will do it?
- When will we do it? Progress/Status?
 - Clarity on the root cause and change required to fix it = make the change (e.g. fix the equipment, update the SOP, etc.)
 - Clear on root cause but unclear of change required to fix it = test a change via a PDSA cycle (e.g. process change to address root cause)

PDSA Cycle

- Simple, powerful, action oriented tool for testing change in the work setting



Method

- ❑ Verify outcomes
 - Confirm expected outcome was achieved via data
- ❑ Standardize and Spread
 - Ensure all relevant documentation and processes are updated to reflect the change
 - Ensure training and communication has occurred to all relevant parties to ensure the change is supported, spread, and sustained



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Maximizing Diagnostic Yield in Biliary Brush Cytology:

A QI Project

Susan McRae (Senior Cytotechnologist)

Biliary Brush Cytology

Problem Raised by Clinicians
(Early 2016)



High Atypical rate

Poses Difficulty for Clinical Management

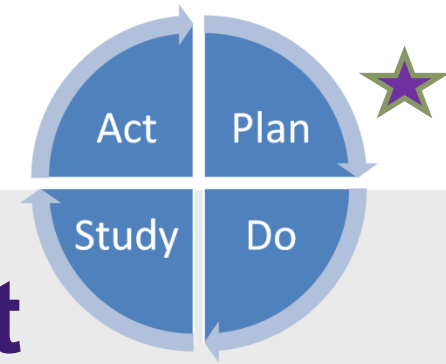
Objectives:

- ❑ **Evaluate the current performance characteristics of ERCP biliary brush cytology service at LHSC**
- ❑ **Design a QI project to improve the diagnostic accuracy of this test**

PDSA cycle



PLAN



Time Line for QI Project

Early 2016

- **IDENTIFY PROBLEM:** Conversation between ERCP physician and Dr. Joseph - discussed high proportion of atypical diagnosis

March 2016

- **PLAN:** Initial data collection to identify specific issues

October 2016

- **DO:** Design the QI project with an AIM statement
- Slide review with cytotechs and cytopaths

April 2017

- **STUDY:** Analyze data
- Review by statistician, summarize what we learned

October 2017

- **ACTION:** :Initiate action plan,
- Move to next cycle

DO

Biliary Cytology QI Project

Aim Statement

**Reduce atypical diagnoses from
36% to 25%
in one year**



Analyzed Possible Factors

Pre-analytical

- Brush collection techniques (sampling issue)
- Cytopreparation techniques

Analytical

- Diagnostic criteria (interpretation)
- Use of 2014 Pap Society Pancreaticobiliary guidelines
- Interpretative variations amongst CT & CP

Post-analytical

- No site specific retro review of atyp/susp

PAP Society 2014:

Pancreaticobiliary Cytology

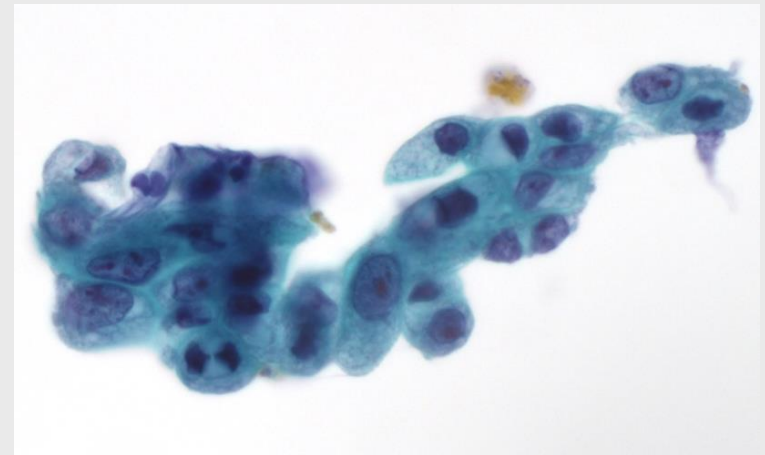
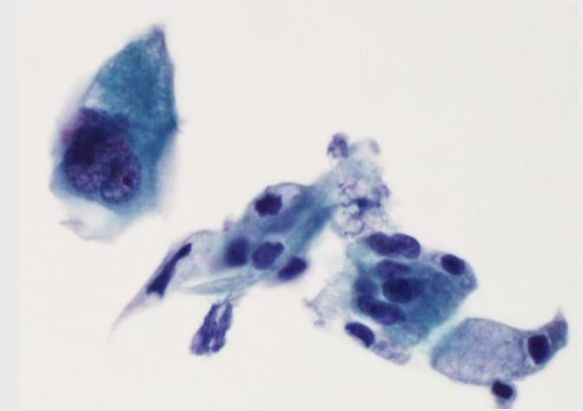
- ❑ The category of **atypical** should be applied when there are cells present with **cytoplasmic, nuclear, or architectural features** that are not consistent with normal or reactive cellular changes of the pancreas or bile ducts, and are insufficient to classify them as a neoplasm or suspicious for a high-grade malignancy. The findings are insufficient to establish an abnormality explaining the lesion seen on imaging. Follow-up evaluation is warranted
- ❑ Heterogeneous category, multiple scenarios

Martha Pittman, Lester Layfield: Cytopathol. 2014;42:338–350

Slide review

Cytomorphology

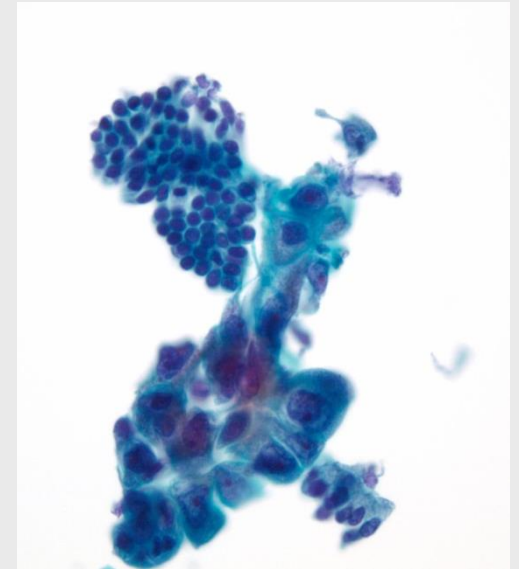
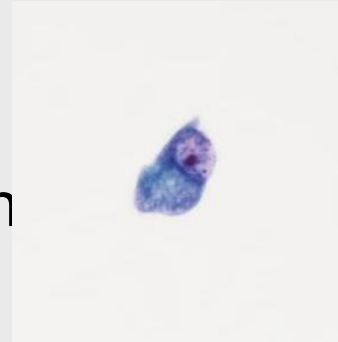
- ❑ Overall cellularity
- ❑ Abnormal group cellularity
- ❑ Atypical single cells
- ❑ Loss of polarity
- ❑ Nuclear features
 - Nuclear enlargement, N/C ratio
 - Anisonucleosis
 - Hyperchromasia
 - Chromatin clumping
 - Chromatin clearing
 - Irregular nuclear contour
- ❑ Cytoplasmic vacuolation



Heath et al: Journal of the American Society of Cytopathology 2015, 4: 282-289

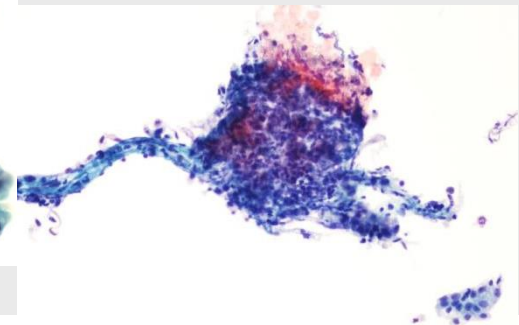
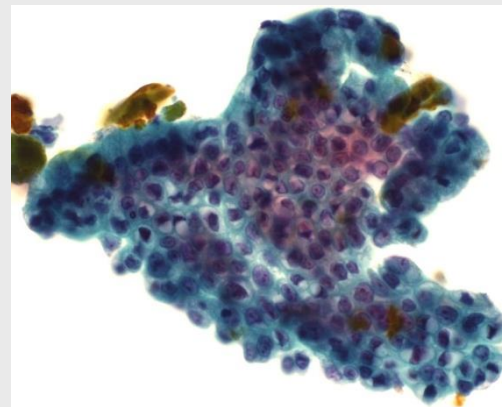
Features favour malignancy

- Atypical single cells
- Two distinct cell population
- Anisonucleosis



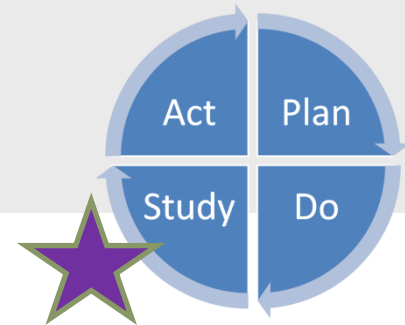
Features favour benign

- Distinct cell borders
- Acute inflammation



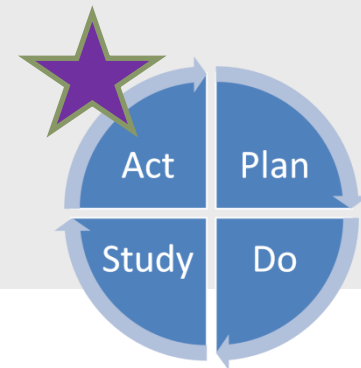
Identify Action Plan

- ❑ Review and reclassify all atypical cases using newly defined criteria (Heath and ours)
- ❑ Stratify “atypical category” into
 - favour benign
 - NOS
 - favour suspicious for malignancy
- ❑ Reanalyze data to determine whether the above approach has an impact on 1) reducing atypical rate and 2) improving diagnostic accuracy



Future Implementation

- ❑ Provide in service to CTs and CPs
- ❑ Encourage peer internal consultation of atypical cases
- ❑ Ongoing QA monitoring of atypical rate



Move to Next Cycle

PLAN

- Discuss sampling technique with **clinical colleagues** in an attempt to improve sample cellularity
- Evaluate role of ancillary technique (FISH) for atypical cases – expensive test



- ✓ Design and implement a QI project for cytology using PDSA cycle model
- ✓ Address strategies that may reduce atypical rate and improve diagnostic accuracy of biliary brush cytology
- ✓ Implement and evaluate these strategies in future

Improve Patient Care

Acknowledgements

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