Anatomic Pathology Process Improvement Stories

From the Simple (Tissue Storage Process) to the Complex (Placental/Fetal Tissue Process)

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KEY LEARNING OBJECTIVES

After this program you will be able to:

Describe a basic improvement approach in handling nonconformities within an anatomic pathology laboratory.

Identify potential opportunities for improvement within the participant's own laboratory based on the three different case studies.

Apply the common lessons learned from this session to the participants own laboratory.





Agenda

Background

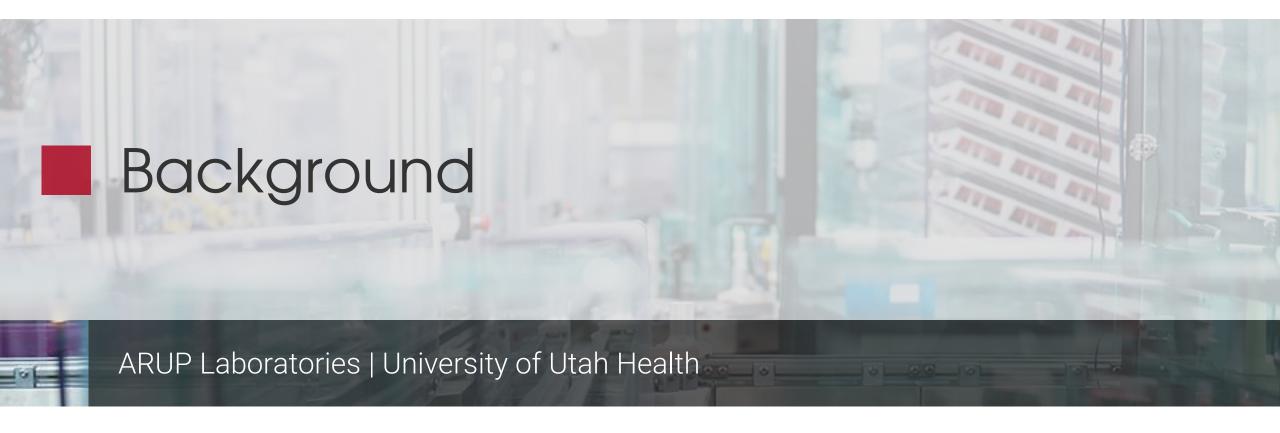
Case Studies

Outcomes

Lessons Learned

What's Next?







1 location

90+

medical directors

>99%

of testing performed in-house

>6.5M patients supported each year

A nonprofit enterprise of the University of Utah and its Department of Pathology





BACKGROUND

Nonprofit, academic affiliate



3,500+ employees

including 90+ medical directors

Provides testing for:

- Genetics
- Immunology
- Oncology
- Pediatrics
- Pain management

Broad test menu

3,000+ tests and test combinations

Clients include:

- university teaching hospitals
- children's hospitals
- multihospital groups
- commercial laboratories
- group purchasing organizations

>10M

specimens/year

>6.5M

patients affected/year







PLACENTA/FETAL TISSUE HANDLING PROCESS IMPROVEMENT

Problem

- APP Staff identified a problem with the Placenta/Fetal Process:
 - » Average 10% of pathology orders placed on placental tissue required follow up.
 - Tissue often not submitted to pathology
 - Necessary information frequently not included
 - > i.e. gestational age
 - » Tissue loss had occurred.



PLACENTA/FETAL TISSUE HANDLING PROCESS IMPROVEMENT

Goals

- Eliminate loss outcomes.
- Improve shared understanding of expectations for hand-off between UH and ARUP Pathology.
 - » Distinguish between L&D issues coming from OR/Delivery rooms

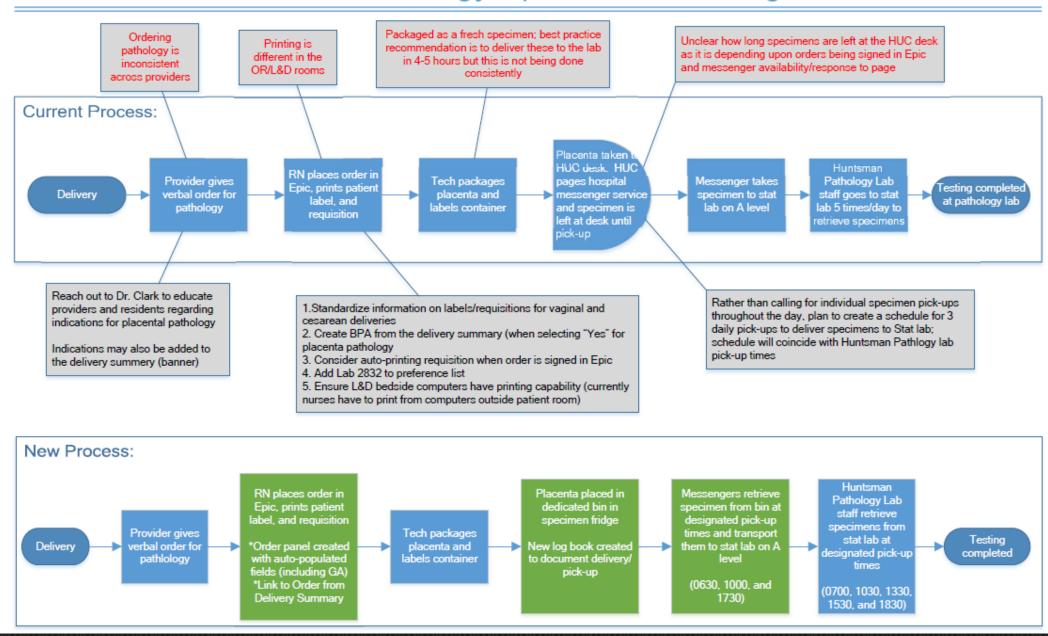


How did we do it?

- Assembled team/identified stake holders
- Mapped Current State and Future State
 - » Identified problem areas
 - » Vetted with Stake holders
 - Tissue stability
 - Indications list
- Developed improvements with full team participation



L&D Pathology Specimen Handling



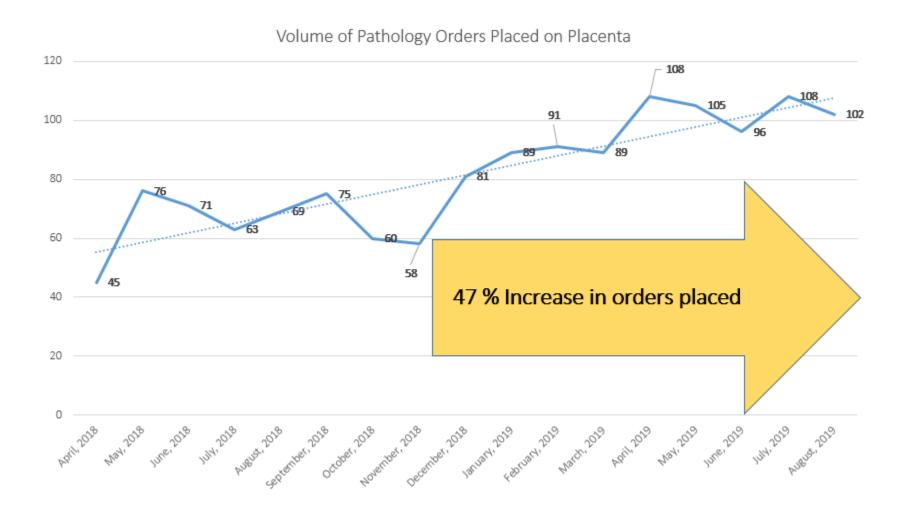
PLACENTA/FETAL TISSUE HANDLING PROCESS IMPROVEMENT

Outcome

- Process is understood and improved between organizations.
 - » Indications for ordering pathology on placental tissue are now known
 - » Designated storage locations in L&D "Pick up for Pathology"
 - » Messengers scheduled delivery to ARUP pick-up sites 3X daily
 - » EPIC order panel built with required fields and process guidance
 - » Both OR and delivery rooms produce barcoded requisitions when orders are placed. (Reduces mislabel opportunities)
- Collaborative relationship developed. (Trust)



Outcome Measure





Future plans / Transferability



- Next Project with OB/GYN team
 » Fetal Demise Process
- Development of shared expectations with additional clinical areas
- Barcoded requisitions from all clinics
- Ordering requisition auto-printing
- Designated specimen locations





Problem

- The intraoperative process is complex, fast-passed and crosses organizational lines which resulted in a significant error.
 - » Ownership of process tasks are not explicitly defined
 - Lack of associated training/competency
 - » Gross description was not documented: residual tissue was missed during clean-up step



FROZENS—INTRAOPERATIVE CONSULTATION PROCESS IMPROVEMENT

Goals

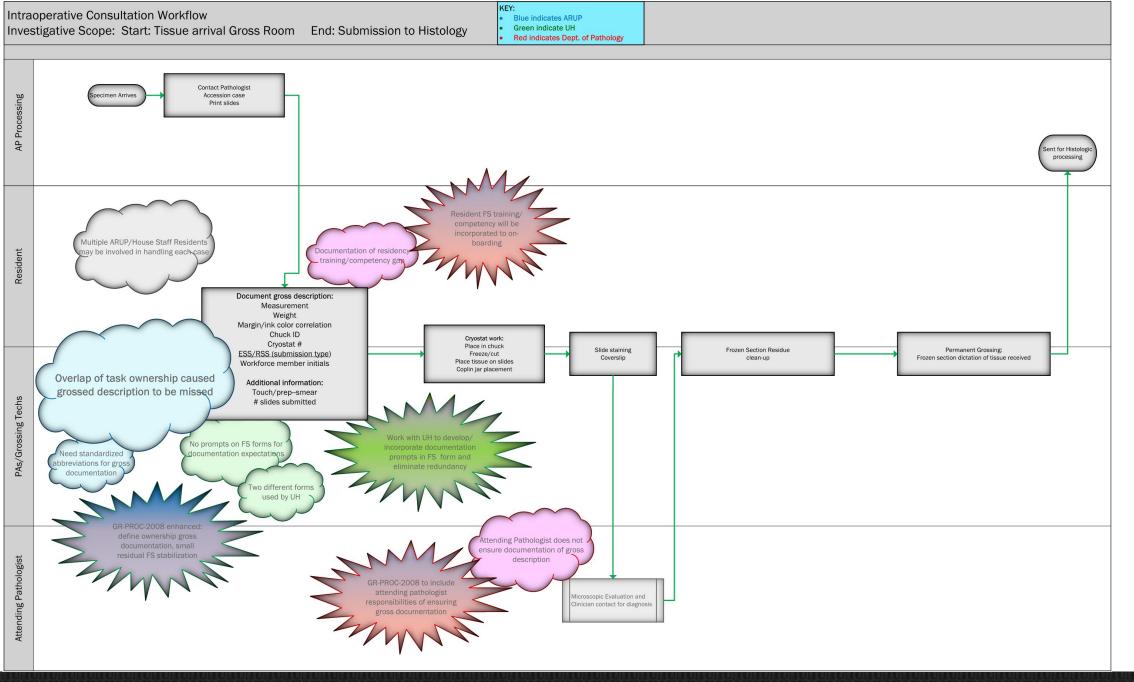
- Eliminate loss outcomes.
- Understand ownership of steps.



How did we do it?

- Assembled team and identified stake holders
- Mapped current state and identified problem areas.
 - » Vetted with Stake holders/team
 - Dept. of Pathology/Residency program
 - Gross Dissection/AP Processing
- Developed improvements with full team participation







Corrective Actions, Implementation, Outcomes

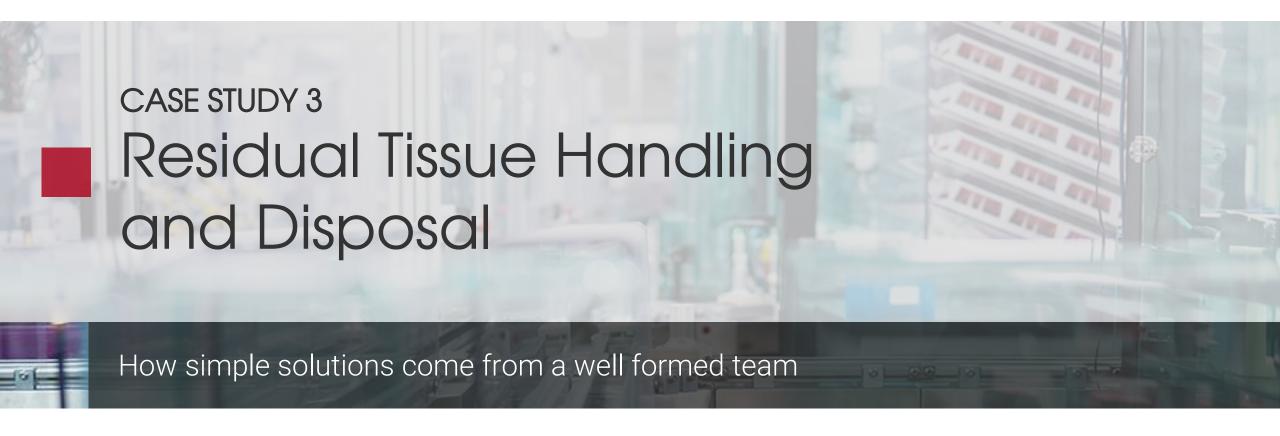
Department of Pathology/ Residency program

- Enhanced procedural steps
 - » Tasks associated with Intraoperative process are better understood
- Training further developed for on-boarding residents
- No tissue loss to date

ARUP: Gross Dissection/AP Processing

- Standing meeting developed between ARUP and Dept. of Pathology residency team
- Enhanced procedural steps
 - » Development of prompts on Intraoperative form
- No tissue loss to date







Problem

- The tissue disposal process was taking two people, 3–4 hours to complete.
- It was a labor intensive, time consuming task that was prone to error.



RESIDUAL TISSUE HANDLING AND DISPOSAL

Goals

- Automate the reconciliation of the specimen in the disposal process
 - » Save time
 - » Increase accuracy



Volume of Containers Evaluated for Disposal Weekly

Tissue Containers



Jake discarding specimens



RESIDUAL TISSUE HANDLING AND DISPOSAL

Old Process

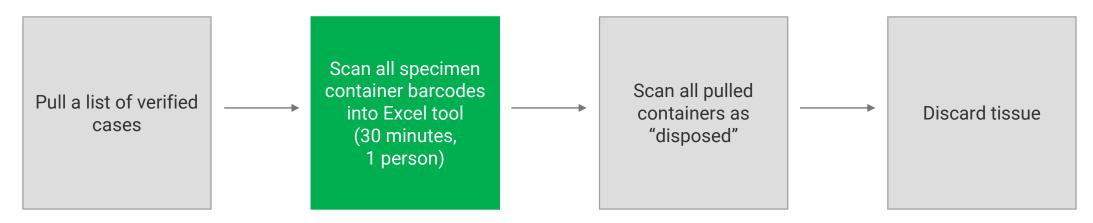
Pull a list of verified cases

One person reads case # from specimen container, another checks list (2-3 hours, 2 people)

Scan all pulled containers as "disposed"

Discard tissue

New Process



How did we do it?

- Team worked with IT
 Analyst to develop an excel tool for us to utilize.
- Transferability:

 Could be considered for any lab that performs disposal of specimens



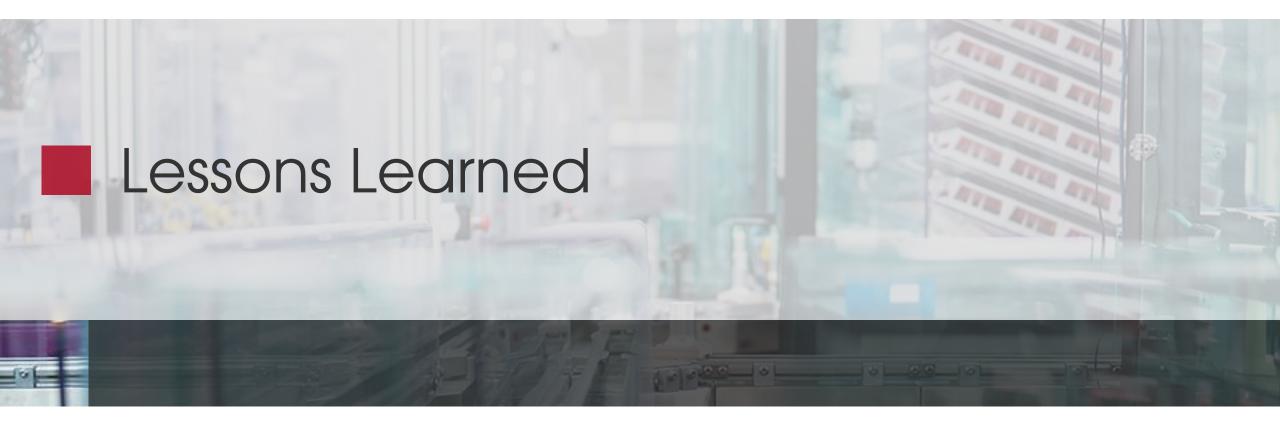
Outcome

Outcome

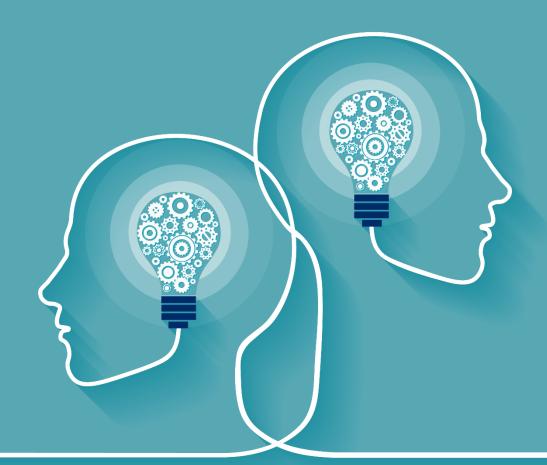
- Time saved:3-4 hours per week
- Financial benefit: approx. \$3000 per year
- Increased accuracy:
 automated a portion of a
 manual process, reducing the
 potential for human error.

Outcome Measure

 Audit in 6 months to demonstrate sustained improvements.



Lessons Learned



- Trust is the foundation of successful team work
- A well-developed team can solve any problem
- Remove "blame" by vetting with your team and then confirming with data.
- Never forget who the "customer" is.



Using Data to Identify Opportunities

- Specimen Intake Process
 Improvement Project
- Gross Room TAT Improvement Project
- Pathology Review and Concordance: PaRC—Development of a digital tool to assist pathologist with overall quality and competency assessments





Shout Outs

Labor and Delivery Department University of Utah Health

Nikole Ihler, RN, Clinical Nurse Coordinator

Erin Clark, MD Division Director, Maternal-Fetal Medicine

PCH attending Pathologists

Sarah Lauer, Quality

Jennifer Spackman, BSN RN Application systems Analyst III

Labor and Delivery Registered Nurses

Julie O'Neil/Erica Cuvelier, RN

Breann Hilton/Maria May, RN

Teressa Gilbert, RN

Daniel Albertson, MD, Section Chief, Surgical Pathology

Erika Prince, AP Processing, Supervisor

AP Processing group

Adnan Milicevic, ARUP AP IT Analyst

Douglas Pulvirenti, Gross Dissection, Supervisor

Gross Dissection group (GTs and PAs)

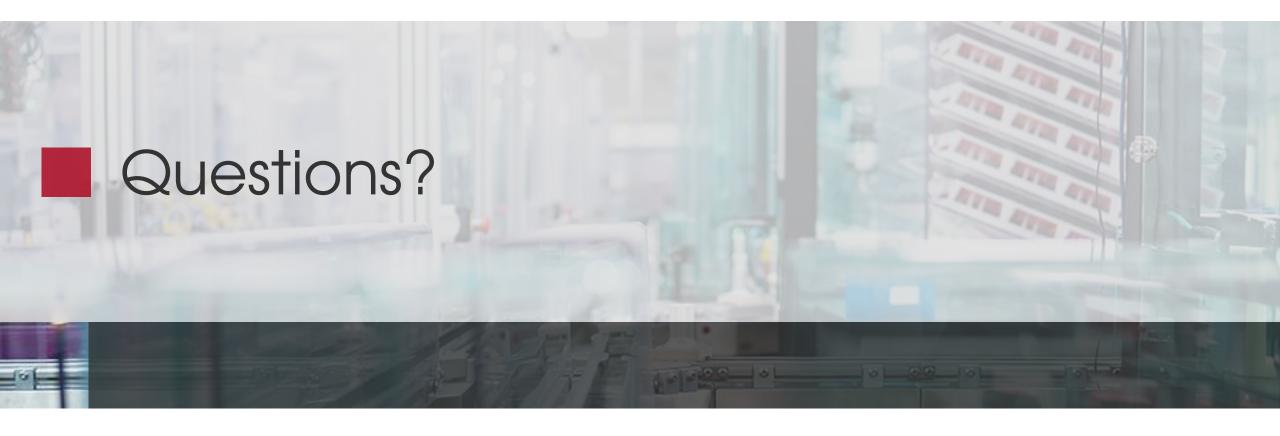
Christian Davidson MD, Gross Dissection Section Chief

Kristi Smock, MD, Residency Coordinator

Maria Pletneva, MD, Surgical Pathology Residency, Liaison

Margaret Coppin, Operations Director, Pathology







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