

COLLEGE of AMERICAN PATHOLOGISTS

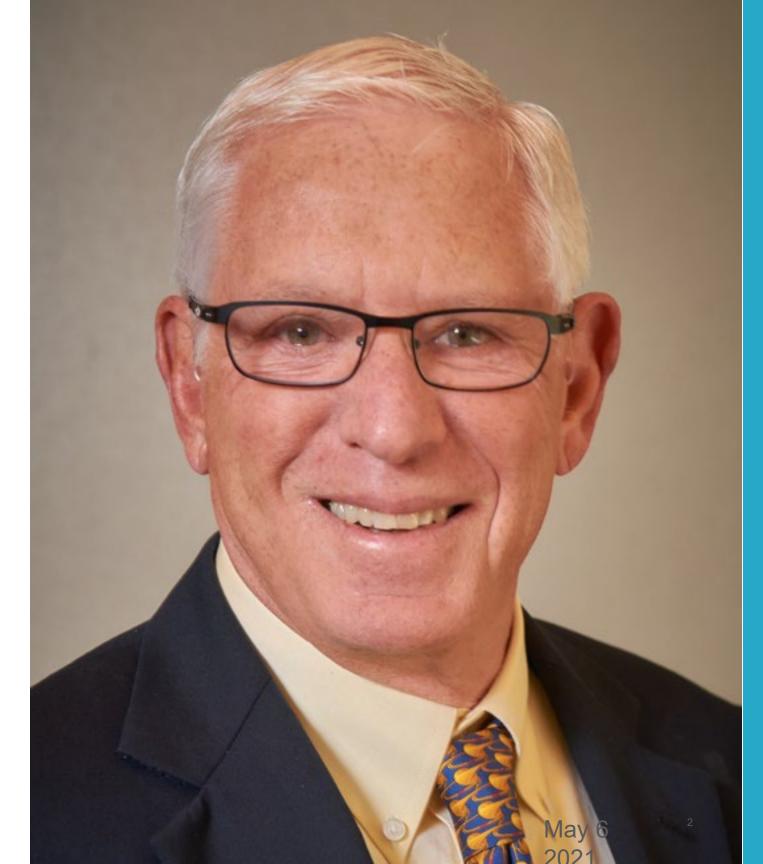
Best Practices for Error Reduction in Anatomic Pathology

David Novis MD, FCAP Stephen Raab MD, FCAP **Esther Yoon MD, FCAP**



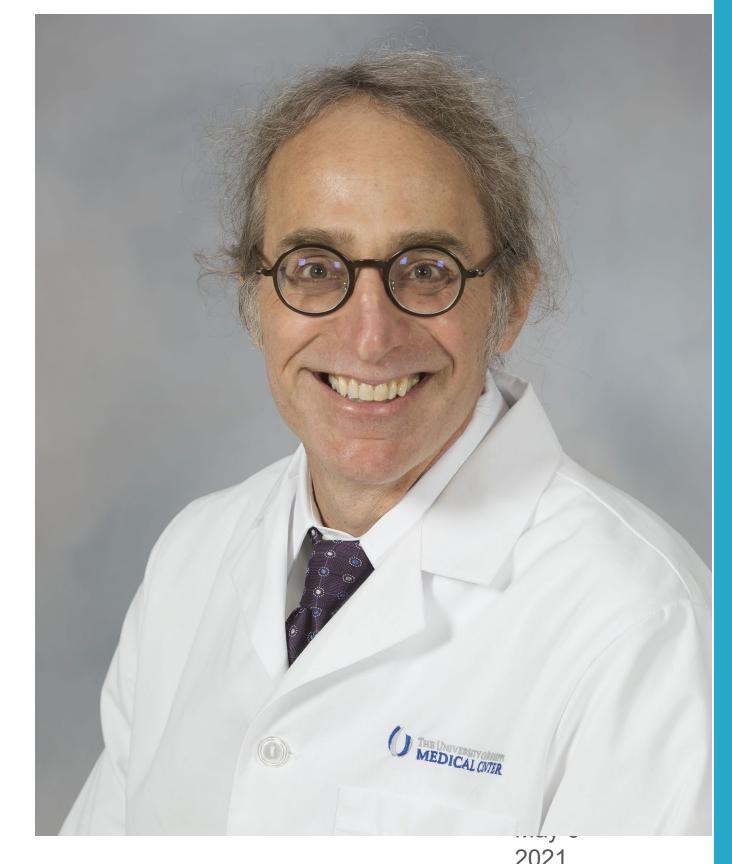
David Novis, MD FCAP

- Owner, CEO Novis Consulting LLC.
- Managing Partner (Ret., Young Novis PA
- Entrepreneur, Business Developer
- Lean Certification, University of Pittsburgh and Henry Ford Hospital
- Past CAP Positions:
 - **o** Speaker of the House of Delegates
 - Member of CAP Board of Governors
 - Vice Chair Quality Practice Committee



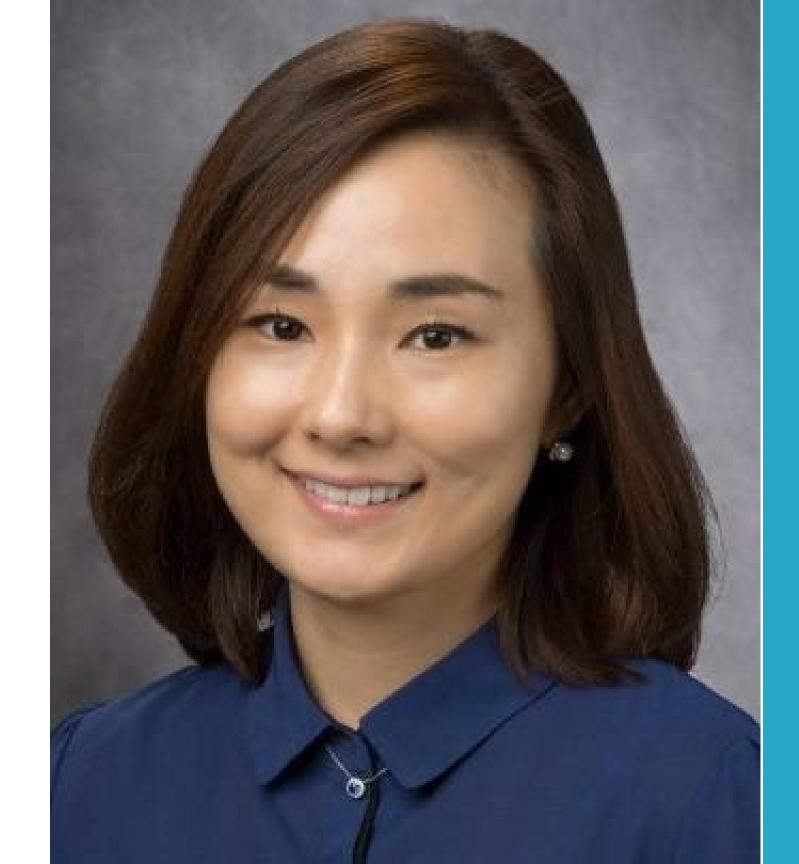
Stephen Raab, MD CAP

- AP/CP Pathologist >30 years
- Medical Officer Agency for HealthCare Research and Quality
- Co-Chair, CAP Interpretive Diagnostic Error Committee.
- Researcher, educator, author in quality and error reduction.
- Recipient CAP Humanitarian Award and Lansky Award (leadership)



Esther Yoon, M.D., FCAP

- Section Head in Surgical Pathology, Florida Region, Cleveland Clinic
- New In Practice Committee, CAP
- Member CAP, USCAP, ASCP
- Board certified AP/CP
- Fellowships: Breast & GYN (2018) and Cytopathology (2019)



Disclaimer

The information presented today represents the opinions of the panelists and does not represent the opinion or position of the CAP.

This should not be used as a substitute for professional assistance.

The information in this presentation is provided for educational purposes only and is not legal advice.

Errors in Anatomic Pathology The Current State

Esther Yoon, M.D. FCAP

Errors = Amendments ?

- How are we doing now?
- Effect of practice setting on error
- Effect of case volume on error
- Can we do better than counting amendments?





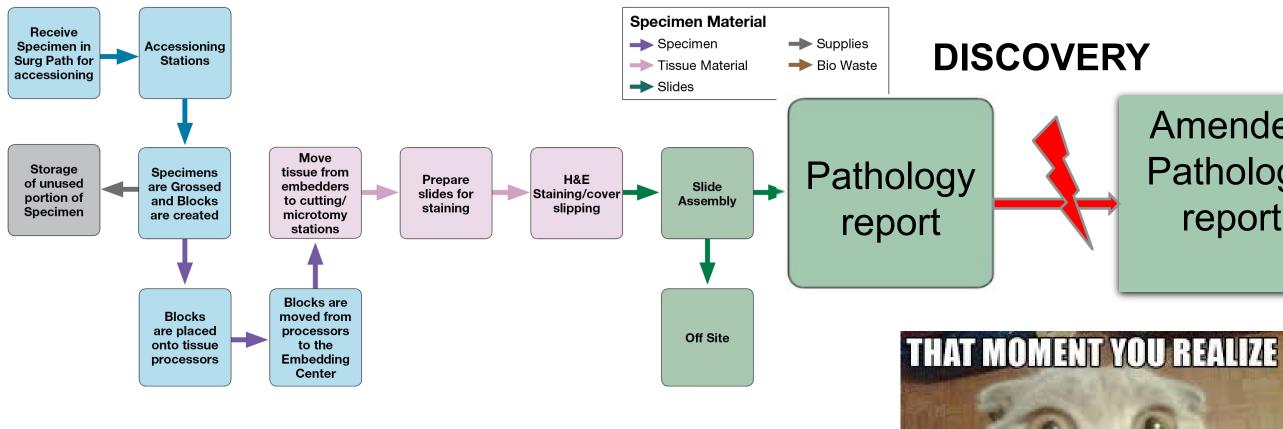
in 2022 Issues, August 2022, In Every Issue, Q&A column

Q. Every month our anatomic pathology laboratory amends patient reports. Does the CAP have a benchmark for amended reports, such as how many are acceptable per month?

Amendment rates range from 0.1 to 10 percent o does not provide a benchmark for amended reports.



Pathology Report and Errors



Histology Specimen Workflow Example

Image modified from https://www.medlabmag.com/article/1436

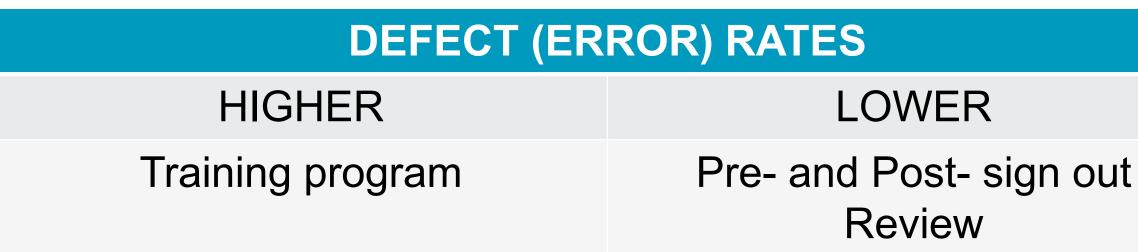
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Amended Pathology report



Surgical pathology report defects: a College of American Pathologists Q-Probes study of 73 institutions.

- 73 participating institutions
- Median defect rate of 5.7/1000 reports



Volmar KE, et al. Surgical pathology report defects: a College of American Pathologists Q-Probes study of 73 institutions. Arch Pathol Lab Med. 2014 May;138(5):602-12.



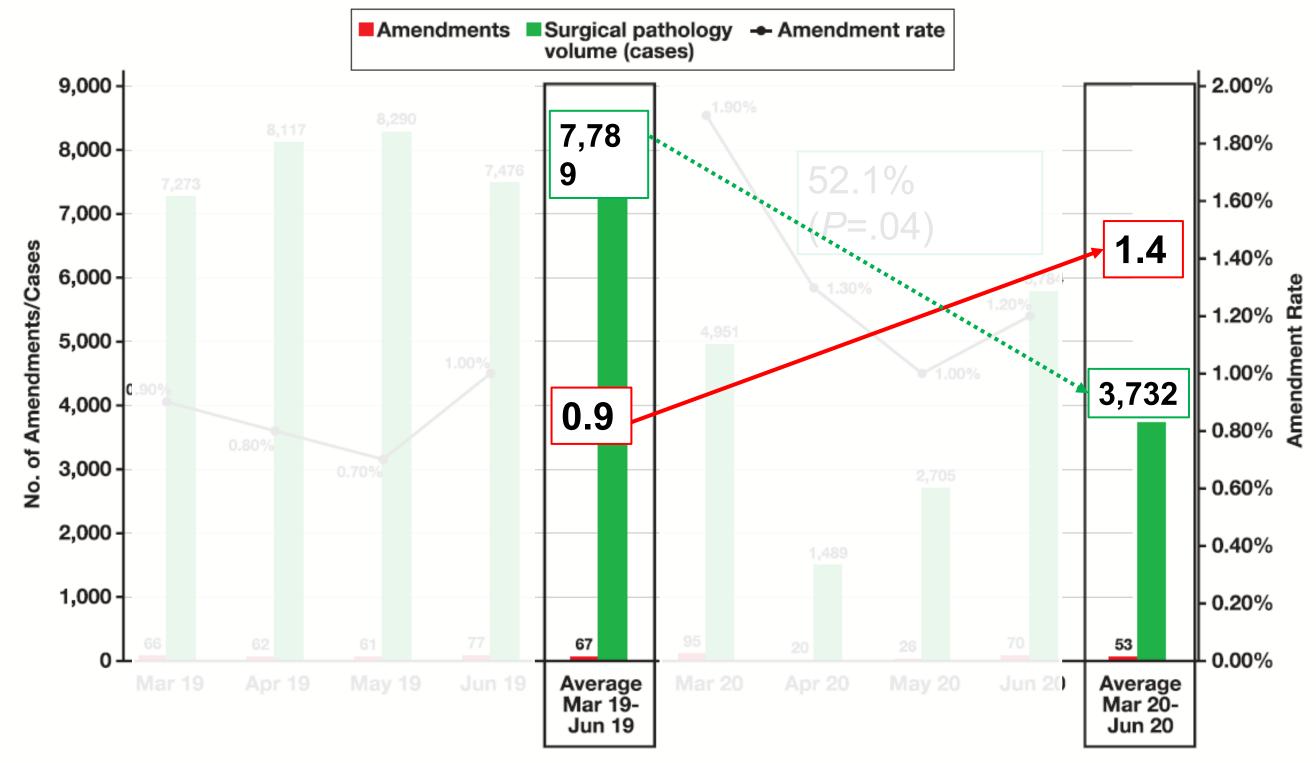


FIGURE 1 Number of amendments and case volume each month (y-axis, left). Rate of amendments each month (y-axis, right).

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Harris CK, et al. Changes in Surgical Pathology Case Volume and Amendment Rates During the COVID-19 Pandemic. Am J Clin Pathol. 2022 Jul 1;158(1):142-147.

% Change in amendments

Types of	Identification	Report defect	D
"Amendments"	(%)	(%)	in
Total change (<i>P</i> =.46)	-53.3	-3.8	



Diagnostic nformation (%)

23.2

Tracking Errors

- Amended Reports
 - Underestimation of magnitude: Follow up
 - Underestimation of severity: Who Decides?
 - ***** Retrospective: Too little too late
- Revised Reports:
 - Accurate estimate
 - Prospective: mitigates risk
 - Promotes intradepartmental standardization



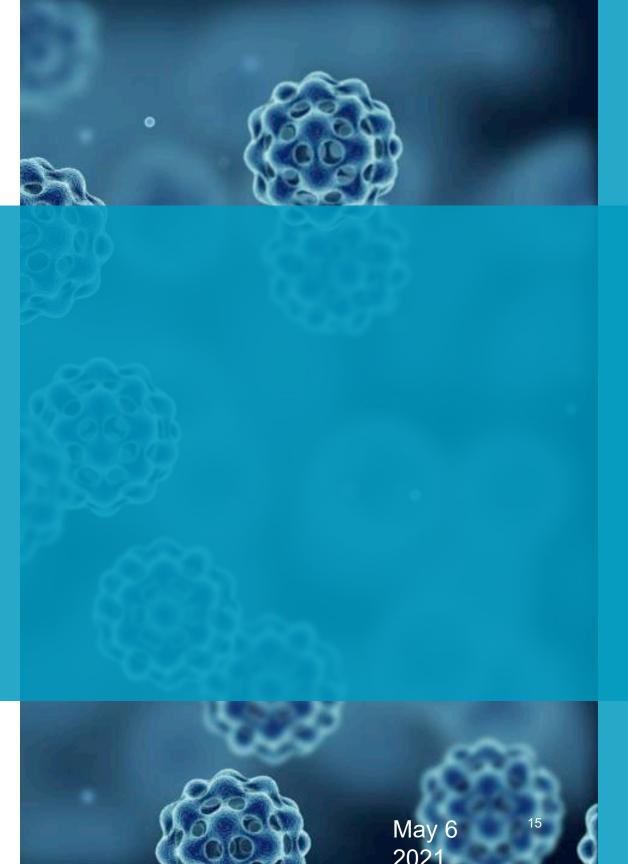


Reducing AP Errors What Works?

Stephen Raab, M.D.

Why Do a Secondary Review

- 1. Types and methods of secondary review
- 2. The benefits of disagreement
- 3. Standardization of reports





What does a blinded retrospective secondary review of a cohort of surgical pathology and cytopathology specimens show?

The College of American Pathologists and the Association of Directors of Anatomic and Surgical Pathology expert panel (2015)

Evidence-based guidelines



What does a blinded retrospective secondary review of a cohort of specimens show?

Guideline Statement 1 – Summary of Studies

Discrepancy rates (%)		Major Discrepancy ra		
No. of studies	Median (25 th -75 th percentile)	No. of studies	Me (25 th – 75 ^t	
116	18.3 (7.5-34.5)	78	5.9 (2	

2.1-10.5)

ledian 5th percentile)

ates (%)

Science Behind Discrepancies— Sources of Variations

- 1. Processes occurring in the patient
- 2. Diagnostic pathways
- 3. Pathologist observers





Science Behind Discrepancies-**Pathologist Judgements of Diagnoses**

Imprecise judgment (noise or repeatability)

Inaccurate judgment (absence of truth or bias)

You are uncertain this tumor is A or B, but do not consider the possibility of C

You are confidant that this tumor is reactive for S100, when it is not

Standardization

We don't know what we don't know

Case Review

Best Detected and Resolved By...



Agreement for Juror First Votes

376 Juror First Votes in Criminal Trials

	Not Guilty	Undecided	Guilty	N	p-Value
African-American juror	46%	13%	41%	743	0.000***
White juror	31%	12%	57%	1,298	0.000***
Hispanic juror	36%	15%	49%	629	0.542
African-American juror-minority defendant	47%	12%	40%	651	0.000***
African-American juror-white defendant	22%	11%	67%	18	0.140
White juror-minority defendant	32%	13%	55%	960	0.008***
White juror-white defendant	30%	10%	60%	136	0.312
Hispanic juror-minority defendant	36%	13%	51%	496	0.955
Hispanic juror-white defendant	33%	9%	58%	33	0.617
Male	36%	11%	53%	1,206	0.311
Female	36%	15%	50%	1,920	0.311

Table 1: Demographics and Juror First Vote

NOTE: Significance levels test the hypothesis that the variables listed in the first column are not associated with a juror's first vote. Significance levels were calculated using ordered logit regression models accounting for the nonindependence of jurors who sat on the same case. The juror's first vote served as the dependent variable. Dummy variables reflecting the juror characteristic or juror characteristic-defendant characteristic combination listed in the first column served as the independent variable.

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Interpretive Summary of Guideline Recommendations



- Implement procedures to detect disagreements and interpretive errors.
- Perform case reviews in manners timely enough to improve patient care.
- Document case reviews and case review procedures.
- Track outcomes of case reviews.
- Implement procedures to maximize diagnostic agreement.



ve errors. tient care.

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Retrospective vs Prospective Review

Retrospective	Prospective
Fixed baseline error rate	Dynamic evolving error ra
Delayed error correction	Immediate error correction
Individual or team activity	Team activity

Nair R, Aggarwal R, Khanna D. Methods of formal consensus in classification/diagnostic criteria and guideline development. Semin Arthritis Rheum. 2011 October ; 41(2):95–10 doi:10.1016/j.semarthrit.2010.12.001.

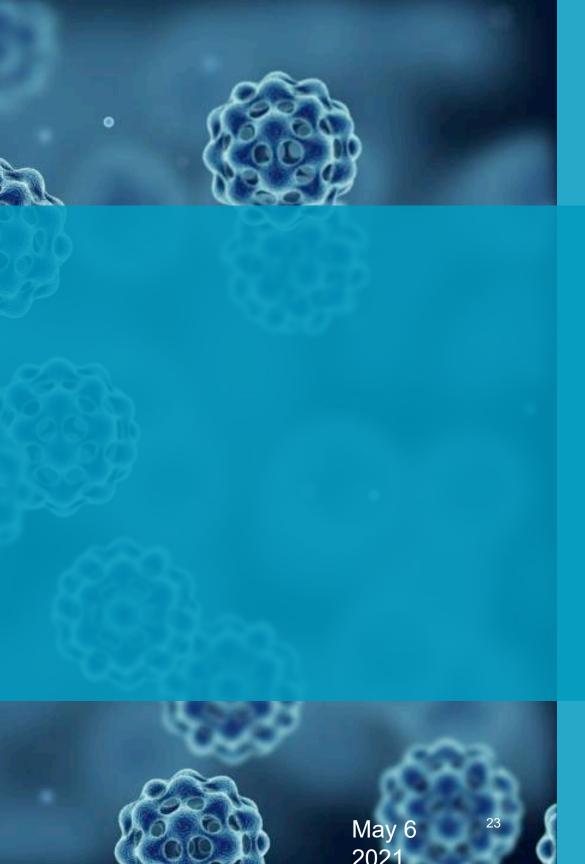
rate

ion

Models of Retrospective Case Review

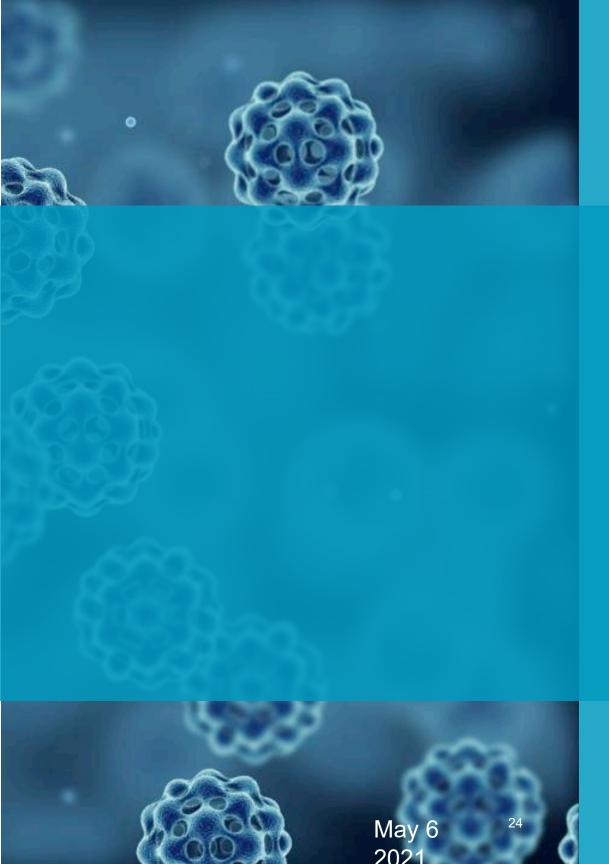
- Formal Model*
- Difficult case conference review,
- Curbside consults
- Secondary opinion (pre-sign-out)
- Tumor boards
- Send outs
- Others.

Nair R, Aggarwal R, Khanna D. Methods of formal consensus in classification/diagnostic criteria and guideline development. Semin Arthritis Rheum. 2011 October ; 41(2):95–10 doi:10.1016/j.semarthrit.2010.12.001



Methods of Prospective Case Review

- Rapid pre-review (e.g., hot seat review)
- Reference class forecasting
- Dyad or team sign-out
- Pilot-Co-pilot diagnosis
- Calibration exercises



Team Signout in a Private Pathology Practice 12K-Accessions

David Novis, MD FCAP

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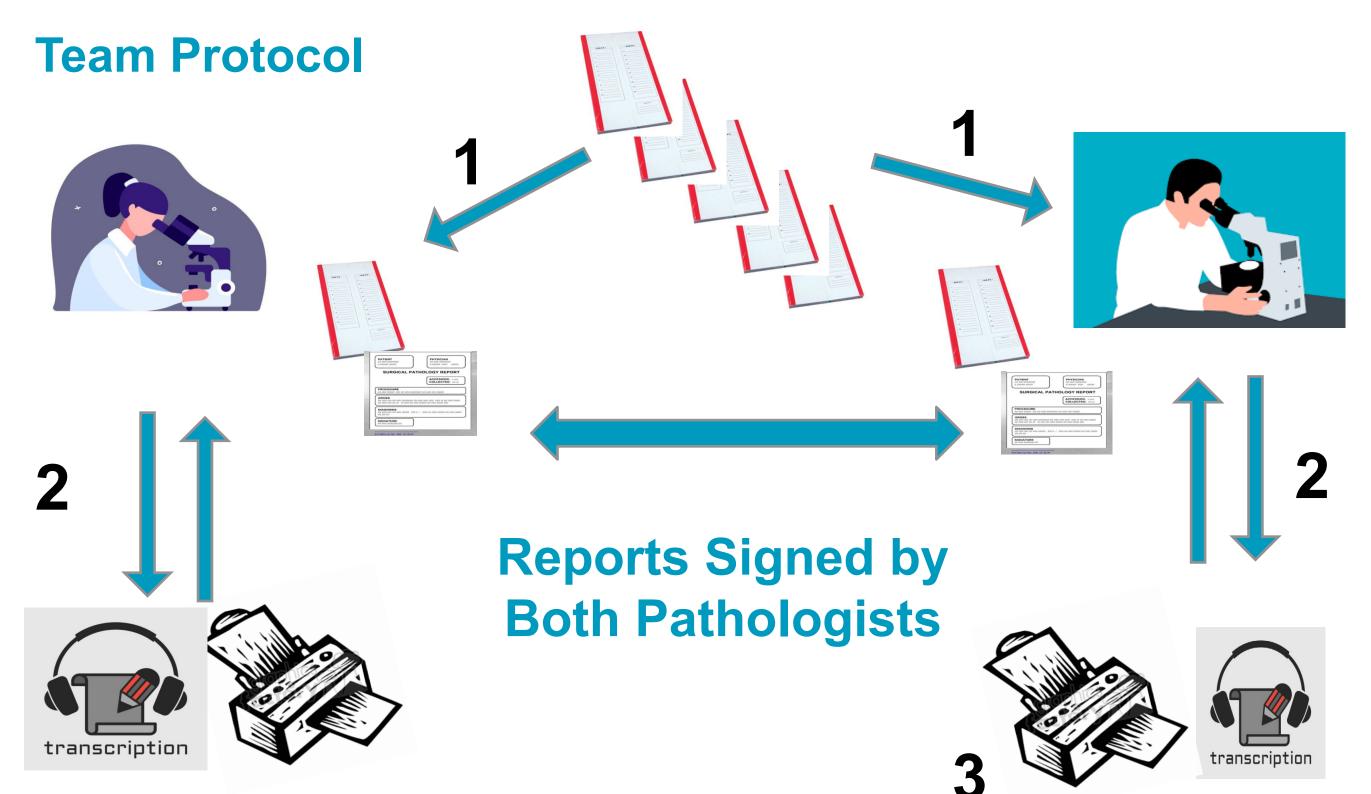


Objectives

- Review Protocol
- Requirements
- Standardization
- Outcomes
- Considerations







Case Review Protocol

- 100% Prospective Review
- *Quality Control*—NOT a double blind read
- Is what's on the report on the slides?
- Is what's on the slides in the report?
- Is the report readable and grammatically correct?
- Does the report address the clinical question?





Requirements of 100% Prospective Review

Culture

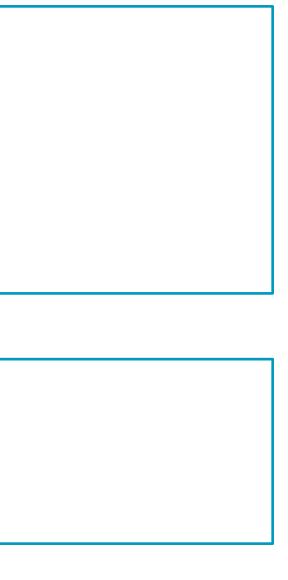
Intolerance of defects Tolerance of work styles

Group rather than individual accountability ✤Trust

Standardized criteria (templates)

ALL diagnoses Diagnostic terms





Standardized Reporting Templates Diagnostic Terms and Criteria

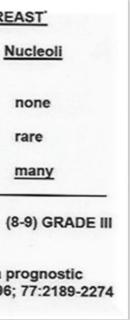
SCORE	TUBULE LUMENS	CYTOLOGY	MITOSES/10 HPF(40X):
1	>75% clear	small, uniform	<u><0-5</u>
2	10-75% clear	some nucleoli	6-10
3	≤10% lumens	anaplastic ^b	>10
TOTAL:	(3-5) GRADE I	(6-7) GRADE II	(8-9) GRADE I

	*Elston CW, Ellis IO. Pathological prognostic factors in breast cancer. I. The value of histological grade in breast cancer:
	experience from a large study with long-term follow-up. Histopathology, 19:403-410, 1991.
	anaplastic: >25% of nuclear diameters > 2rbc; nuclear shape variable; >25% of nuclei possess nucleoli, nuclear chromatic
J	coarea clumped

TOTAL:	(3-5) GRADE I	(6-7) GRADE II	
3	>2 rbc	vesicular	
2	1.5-2 rbc	coarse	
1	1-1.5 rbc	diffuse	
SCORE	Nuclear Size	<u>Nuclear</u> Chromatin	1

* Taken from: Schnittt SJ, Harris JR, Smith BL. Developing a prognostic index for ductal carcinoma in situ of the breast. Cancer 1996; 77:2189-2274

		ENDOMETRIAL	ADENUCARCIN	OMA
7	AR	CHITECTURAL G	RADE	
Architectural	FIC	30	A.	ICC
Growth Pattern	a second and	Glandular Nu	clear Atypia ²	
(% Solid Growth)	Absent	Present	Absent	Present
5%	G1	G2 -	G1	G2
6-50%	G2	G3	G2	
>50%	G3	G3 G3	G3	<u>G3</u> G4
"Glandular nuclear atypia: th mitoses>20/10hpf. NOTE: The grade of the tumo cases	pree of four following or from biopsy specim	features-(1) pleomorphic, tens agrees with the tumo	enlarged (2) coarse chr	tular infiltration associated with omatin, vesicular (3) prominent nucleoil iomy specimens in less than 60% of the metrial Biopsies and Curettings (New

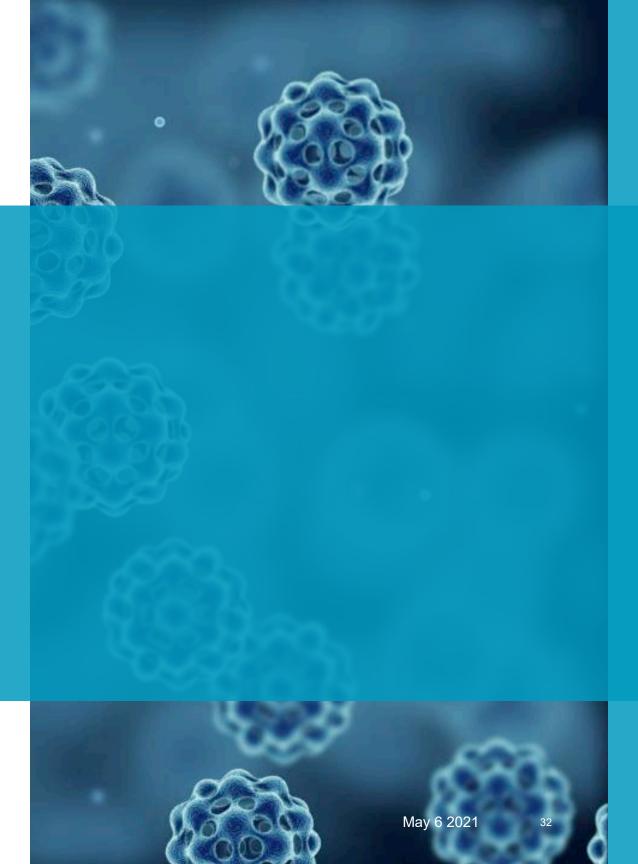


Standardized Reporting Templates

MICROSCOPIC DESCRIPTION B: Squamous mucosa, Squamocolumnar mucosa, glandular mucosa Epithelium Squamous component: acanthosis spongiosis transepithelial migration severity: mild cells: lymphocytes, eos dysplasia: none Glandular Component type: gastric cardia (subsurface mucous glands) gastric body (parietal, chief cells) Specialized (intestinal) type mucosa: not identified inflammation amount: mild distribution: focal type: chronic reactive/inflammatory hypertrophy hyperplasia dysplasia: none Lamina propria/Stroma inflammation amount: severe distribution: diffuse type: chronic

Outcomes

- Reduced errors
- Saved time
- Grew customer satisfaction
- Grew business



Reduced Errors

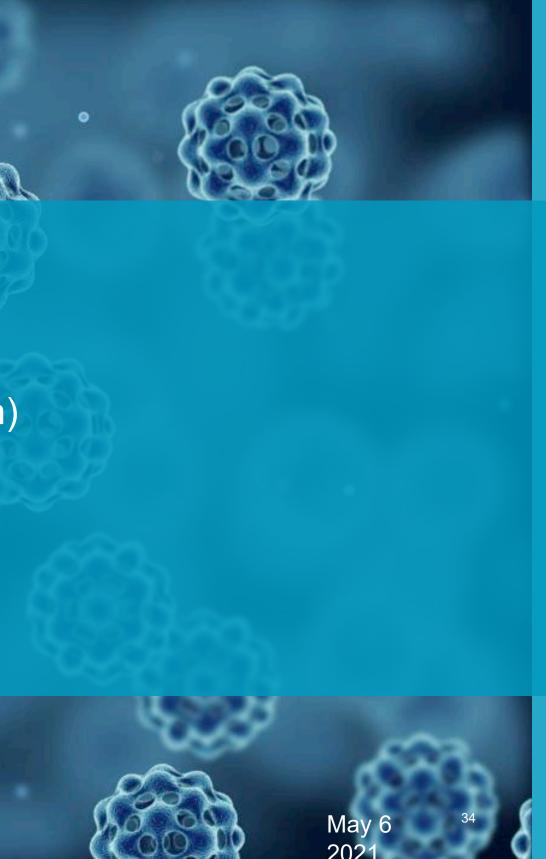
- Amended Reports Rate Decreased From 6 to 1.2 per 10,000*
- Pre-release corrections—Revised Reports: 1/20-1/50

*https://davidnovis.com/wp-content/uploads/2014/03/Doubleread-copy-2.pdf

ber 10,000* /50

Saved Time

- 1. Pareto
- 2. Standardized templates
 - Delegation (conversation not dictation)
 - Clicking on checklist items
- 3. Eliminate disruptive calls for 2nd looks



Customer Satisfaction Business Growth

- Review by second pathologist
- Templates
 - o Customized
 - Complete
 - Customer input...Sense of ownership





Provider Concerns

- Delays turnaround time 1.
- Not paid for QC 2.
- 3. Most errors do not affect patient care
 - Who decides that?
 - Coexisting conditions
 - How long is the follow up?
 - Nightmares

Bottom Line: Comfort level in releasing a defective report with your name on it





May 6 202





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- 2. How Pathology Practices Get Paid
- **3.** Revenue Cycle Management
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- **5.** Basic Practice Cost Analysis
- 6. Capacity Management and Workflow Analysis
- 7. Basic Contracting and Fee Analysis
- 8. Basic Budget Development



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o <u>https://www.cap.org/member-resources/practice-management</u>

Practice Management Articles

https://www.cap.org/member-resources/articles/category/practice-management Ο



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If after attending this discussion and later you applied any of what you learned to your practice, please share your feedback of how it worked for your practice at https://www.cap.org/member-resources/practice-management/practice-managementinquiry-form



Watch for the session evaluation form. Your feedback is important!