

Welcome!

Before we begin...

Today's session will be recorded

Please add your name and health system in the chat







April 16, 2024, 4:30 PM ET • 3:30 PM CT • 2:30 PM MT • 1:30 PM PT

Prostate Cancer Screening IMPACT ECHO

Session 3: Utilizing Data for Pre-biopsy Risk Calculation and/or Referral to Urology **The PSA is Elevated:** *Now What?*

Welcome to Session 3 of the **Prostate Cancer Screening IMPACT ECHO**



Each ECHO session will be recorded and *will* be posted to a publicly-facing website. Chat content, attendance, and poll responses are also recorded



Please update your Zoom Participant Name to First Last, Org (Molly Black, ACS).



Type your full name, the full name of your organization, and e-mail in the chat box.



You will be muted with your video turned off when you join the call. Use the buttons in the black menu bar to unmute your line and to turn on your video.



Today's materials will be made available on our <u>ACS ECHO website</u>.



All ECHO sessions take place on the <u>iECHO</u> & Zoom platforms. <u>iECHO Terms of Use</u> & <u>Zoom Privacy Policy</u>.



Questions about Zoom during the call? Find **@Beth Graham** in the chat.

This project is being funded by





Every cancer. Every life.







Have a question? Don't wait to ask! Feel free to enter in the **Chat** at any time.





Together with our Primary Care Participant Sites and Subject Matter Experts using the ECHO's all-teach, all-learn approach, we aim to:

Increase appropriate prostate cancer screening.

- 2 **Increase** utilization of prostate cancer shared decision-making tools by primary care teams.
- 3 **Increase** relationships and streamline referral pathways with urologists/other specialty providers within the medical neighborhood.
- **4 Increase** data capacity to:
 - risk stratify patients,
 - track shared decision-making process,
 - track PSA results and identify trends, and
 - collect and utilize data for pre-biopsy risk calculator and/or referral to specialty care.

Today's Agenda



- 1. Welcome, Housekeeping & Data | 7 minutes
- 2. Didactic Presentation & Discussion: Utilizing Data for Prebiopsy Risk Calculation and/or Referral to Urology **The PSA is Elevated: Now What?** 25 minutes Presented by: Andrew M.D. Wolf, MD, MACP
- 3. Participant Site Introduction: Greater Baden Medical Services | 3 minutes
- 4. Case Presentation & Recommendations | 20 minutes Presented by: Debbie Apperson, CPRN | Greater Baden Medical Services, Inc.
- 5. Survey, Schedule, Reminders, & Wrap-Up | 5 minutes



Your ACS ECHO Team



Molly Black Director, Screening American Cancer Society ACS ECHO Program Lead & ECHO Facilitator



Mindi Odom Director, Project ECHO Your ECHO Co-Lead



Beth Graham, MPH, CHES Program Mgr., Project ECHO Your Program Support



Jennifer McBride, PhD Senior Data & Evaluation Manager

Introductions



Meet Our Prostate Cancer Screening IMPACT ECHO HUB – Subject Matter Experts (SMEs)



Andrew M.D. Wolf, MD, MACP Professor, Internal Medicine University of Virginia, School of Medicine



Quoc-Dien Trinh, MD, MBA Chief of Urology Brigham and Women's Faulkner Hospital



William H. Boykin, Jr, MD

Urology Specialist UK King's Daughters Medical Center



Yaw A. Nyame, MD, MS, MBA Assistant Professor, Director of Urology Fred Hutch at University of Washington

We use your feedback!



What would make these ECHO sessions more impactful in the future?

- So far so good thanks
- need to establish practical way to apply in the complex primary care setting of the FQHC where time is limited and there are many social barriers.
- A little more time
- The SMEs are wonderful.
- No specific suggestion
- More emphasis on best practice workflows and practical tools to use to establish a screening program
- I hope we can create a standard of care on when to screen for prostate cancer and the frequency of screening for prostate cancer.

C-SASI Baseline:







Health systems offer reduced cost prostate cancer screenings Health systems use a shared decision-making tool (Athena Alerts)

2

Health systems use a pre-biopsy risk calculator following an abnormal PSA test

11

C-SASI Baseline: Health systems capacity to report prostate cancer screening data



Health systems were most commonly able to provide and trust prostate cancer screening data on **patient eligibility**, followed by **patient ethnicity** and **patient race**









Session 3 The PSA is Elevated: *Now What?*

Andrew M.D. Wolf, MD, MACP Professor, Internal Medicine University of Virginia School of Medicine

Learning Objectives

- To develop a systematic response to elevated PSA screening tests
- To employ strategies to reduce Urology referrals of patients unlikely to have clinically significant prostate cancer
- To use an on-line risk calculator to inform patient discussions regarding elevated PSA levels
- To stimulate discussion of when Urology referral is appropriate



Case #1

62 yo white man without significant past medical history presents for annual preventive visit. He has no family history of prostate cancer. He has mild urinary hesitancy and his prostate is mildly enlarged without induration or nodules. His PSA has been gradually rising:

- 2015: 2.35 ng/ml
- 2017: 2.17
- 2021: 3.75
- 2023: <mark>4.51</mark>

Where do we go from here?



Case #2

55 yo Black man without significant past medical history presents for annual preventive visit. He has no family history of prostate cancer. He has mild urinary hesitancy and his prostate is mildly enlarged without induration or nodules. He has not been screened before. His PSA today returns at 3.51 ng/ml

Where do we go from here?



So what do we do with an elevated PSA?



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What about the good old digital rectal exam?





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The Digital Rectal Exam

- 2018 meta-analysis: sensitivity 51%, specificity 59% (primary care docs)
 - About the same as flipping a coin
- One survey: ½ of medical school graduates never performed a DRE
- Only ½ of primary care docs are confident in their ability to detect prostate cancer with DRE
- Inter-examiner reliability between urologists to identify suspicious nodules is fair at best ($\kappa = 0.22$)
- Major guidelines now make DRE optional for primary screen
 - Still makes sense to do it for abnormal PSA's, symptoms



Causes of False Positive PSA

- BPH the biggie
- Prostatitis (often asymptomatic) don't treat unless symptoms
- Ejaculation (transient)
- Long bike rides
- Probably *not* the DRE
- Rationale for repeating in 1-6 months before acting on it



Age-Specific PSA Thresholds

- 40's: < 2.5 ng/mL
- 50's: < 3.5 ng/mL
- 60's: < 4.5 ng/mL
- 70's: < 6.5 ng/mL
- Actually increases sensitivity for cancer at younger ages
- Overall, reduces false positive rates and overdiagnosis



Reflex Biomarkers to Reduce Biopsies

- % Free PSA
- Prostate Health Index (PHI)
- 4K Score
- PCA3 (prostate cancer antigen 3)
- ExoDx



% Free PSA (fPSA)

- PSA produced by cancer cells is more likely to be complexed to a glycoprotein than PSA produced by non-cancerous cells
- The higher the % free PSA, the lower the risk of cancer
- FDA approved indication: PSA between 4 & 10 ng/ml with normal DRE
- Using a fPSA threshold of <25% detects 95% of cancers & reduces biopsy rate by 20%
 - i.e., only 20% have fPSA levels above 25% & might avoid biopsy



Prostate Health Index (PHI)

- Combination of total PSA, free PSA, and proPSA tests
- Discriminates between high-grade cancer vs low-grade or no cancer
 - Higher score \approx higher cancer risk
- A PHI score cut-off of 24 reduces biopsies by 36-41% at a cost of missing 2.5-5% of high-grade cancers (Gleason <u>></u>7)
- FDA approved for PSA values between 4 & 10 ng/dl
- Mayo sendout (at UVA) & expensive



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4K Score

- Another combo test: total PSA, fPSA, 'intact' PSA & human kallikrein 2
- Also factors in age, DRE result, and prior biopsy status (if done)
- Significantly improve accuracy c/w compared with PSA
 - AUC for 4K compared to PSA alone was 0.82 vs. 0.74 for detection of high-grade cancers
- Can reduce unnecessary biopsies by ~40% depending on threshold used
- Comparable to PHI (& also expensive)



The NEW JOURNAL	ENGLAND of MEDICIN	NE
ESTABLISHED IN 1812	MAY 10, 2018 VOL. 37	8 NO. 19

MRI-Targeted or Standard Biopsy for Prostate-Cancer Diagnosis

V. Kasivisvanathan, A.S. Rannikko, M. Borghi, V. Panebianco, L.A. Mynderse, M.H. Vaarala, A. Briganti, L. Budäus, G. Hellawell, R.G. Hindley, M.J. Roobol, S. Eggener, M. Ghei, A. Villers, F. Bladou, G.M. Villeirs, J. Virdi, S. Boxler, G. Robert, P.B. Singh, W. Venderink, B.A. Hadaschik, A. Ruffion, J.C. Hu, D. Margolis, S. Crouzet, L. Klotz, S.S. Taneja, P. Pinto, I. Gill, C. Allen, F. Giganti, A. Freeman, S. Morris, S. Punwani, N.R. Williams, C. Brew-Graves, J. Deeks, Y. Takwoingi, M. Emberton, and C.M. Moore, for the PRECISION Study Group Collaborators*

Kasivisvanathan V et al. NEJM 2018;378:1767-77.



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MRI vs Standard Biopsy

Table 2. Comparison of Cancer Detection between Groups.*					
Outcome	MRI-Targeted Biopsy Group (N = 252)	Standard-Biopsy Group (N = 248)	Difference†	P Value	
Biopsy outcome — no. (%)			—	—	
No biopsy because of negative result on MRI	71 (28)	0	>		
Benign tissue	52 (21)	98 (40)	>		
Atypical small acinar proliferation	0	5 (2)			
High-grade prostatic intraepithelial neoplasia	4 (2)	10 (4)			
Gleason score					
3+3	23 (9)	55 (22)			
3+4	52 (21)	35 (14)			
3+5	2 (1)	1 (<1)			
4+3	18 (7)	19 (8)			
4+4	13 (5)	6 (2)			
4+5	7 (3)	2 (1)			
5+5	3 (1)	1 (<1)			
No biopsy‡	4 (2)	3 (1)			
Withdrawal from trial§	3 (1)	13 (5)			
Clinically significant cancer¶					
Intention-to-treat analysis — no. (%)	95 (38)	64 (26)	12 (4 to 20)	0.005	
Modified intention-to-treat analysis — no./total no. (%)	95/245 (39)	64/235 (27)	12 (3 to 20)	0.007	
Per-protocol analysis — no./total no. (%)	92/235 (39)	62/227 (27)	12 (3 to 20)	0.007	
Clinically insignificant cancer — no. (%)	23 (9)	55 (22)	-13 (-19 to -7)	< 0.001	
Maximum cancer core length — mm	7.8±4.1	6.5±4.5	1.0 (0.0 to 2.1)	0.053	
Core positive for cancer — no./total no. of cores (%)	422/967 (44)	515/2788 (18)	ss	25 12	
Men who did not undergo biopsy — no. (%) $\ $	78 (31)	16 (6)		53 <u></u> 13	

Kasivisvanathan V et al. NEJM 2018;378:1767-77.



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MRI vs Standard Biopsy Complications

	MRI-Guided Biopsy	Standard Biopsy
Median # of core biopsies	4*	12
Blood in urine (%)	30%	63%
Blood in semen (%)	32%	60%
Post-procedural pain (%)	13%	23%
Rectal bleeding (%)	14%	22%

*Among men who underwent biopsy



Case #1

62 yo white man without significant past medical history presents for annual preventive visit. He has no family history of prostate cancer. He has mild urinary hesitancy and his prostate is mildly enlarged without induration or nodules. His PSA has been gradually rising:

- 2015: 2.35 ng/ml
- 2017: 2.17
- 2021: 3.75
- 2023: <mark>4.51</mark>

Where do we go from here?



Management Options

- Repeat always repeat an elevated PSA in 1-6 months given variability
- DRE if not yet done
- Consider other tests to refine risk assessment, eg % free, PHI, 4K
- Consider using a prostate cancer risk calculator
- Prostate MRI
- Biopsy



riskcalc.org/PCPTRC/



Risk of prostate cancer if biopsy were to be performed

Based on the provided risk factors a prostate biopsy performed would have a:

8% chance of high-grade prostate cancer,

24% chance of low-grade cancer,

68% chance that the biopsy is negative for cancer.

About 2 to 4% of men undergoing biopsy will have an infection that may require hospitalization.

Please consult your physician concerning these results.



If you are Caucasian, click here for a new update to the PCPTRC that incorporates detailed family history into a risk of prostate cancer calculation.

Case #2

55 yo Black man without significant past medical history presents for annual preventive visit. He has no family history of prostate cancer. He has mild urinary hesitancy and his prostate is mildly enlarged without induration or nodules. He has not been screened before. His PSA today returns at 3.51 ng/ml

Where do we go from here?

riskcalc.org/PCPTRC/

Characteristics

Race	
African American	~
Age	
55	
PSA [ng/ml]	
3.51	
Family History of Prostate Cancer	
No	~
Digital rectal examination	
Normal	~
Prior biopsy	
Never had a prior biopsy	~
Percent free PSA available?	
Percent free PSA	

Risk of prostate cancer if biopsy were to be performed

Based on the provided risk factors a prostate biopsy performed would have a:



8% chance of high-grade prostate cancer,

15% chance of low-grade cancer,

77% chance that the biopsy is negative for cancer.

About 2 to 4% of men undergoing biopsy will have an infection that may require hospitalization.

Please consult your physician concerning these results.



Take-Home Points: Managing the Elevated PSA

- Repeat the PSA within 1-6 months given fluctuation.
- Consider DRE if not yet done.
- Consider % free PSA, Prostate Health Index, or 4K score to reduce unnecessary biopsies.
- Consider using prostate cancer risk calculator to help your patients decide next steps if PSA elevated.
- Although usually ordered by urologist, MRI has become a valuable tool to reduce biopsies, overdiagnosis, overtreatment.





Open Discussion: Questions & Answers

Welcome Greater Baden Medical Services, Inc.

Brandywine, MD











Session 3 Patient Case Presentation

Debbie Apperson, CPRN Greater Baden Medical Services

Patient Case Presentation

Presented By: Debbie Apperson, CPRN | Greater Baden Medical Services

Patient-Related Case Presentation

Patient Hx Current Strategies 4/2022: presented w/ swelling in right scrotum- sent to urology 72-year-old, African American/Black male 4/15/2022: PSA at urology office- 203 4/20/2022: repeat PSA at urology office 236 **Past Medical/Surgical History** 4/2022: PET CT scan: metabolic activity in the prostate HTN T2DM CKD3a Inguinal hernia repair 6/2022: inguinal hernia repair **Past Cancer Screening History** 12/12/2022: PSA 244; did not follow up with urologist prostate: 4/15/22 4/2023: came to GBMS office to discuss need for f/u for elevated PSA colon cancer: 11/15/2023 - FIT negative 6/2023: GBMS- again, referral made for urology **Medications** 7/19/2023: new appt w/ urologist - bx done G G3 (G4+3=7) in 7/12 cores 7/2023: repeat PET CT completed – unknown results Irbesartan/HCTZ 300-12.5mg daily, Lovastatin 20mg, qHS 8/16/2023: scan showing prostate cancer w/o evidence of metastasis; Metformin 1000mg, BID Glipizide 5mg daily, Basaglar 20U daily SC shared decision making made w/ patient and urologist - active **Family History** surveillance/radiation therapy/androgen deprivation father: hx of colon cancer dx at 83 yo; currently living mother: hx of therapy/cryotherapy; patient opted for radiation therapy XRTL+ADT X2 diabetes and cardiovascular disease; currently living vears 8/30/23: began ADT therapy w/ urologist 8/31/23: radiation oncologist appt- National Cooperative Cancer Guidelines as a 'High Risk Patient' - 'excellent candidate for external beam radiation therapy and long course of ADT therapy'; given Lupron 11/2023: finished radiation. PSA 9.8; admits s/e hot flashes 2/29/24: PSA 1.2; second Lupron treatment



Prostate Cancer Screening IMPACT ECHO



Presented By: Debbie Apperson, CPRN | Greater Baden Medical Services

Patient-Related Case Presentation

Questions for Discussion:

1) What symptoms are common for checking PSA? We were more using PSA as screening asymptomatic patients.

2) How would the urology team on this panel have handled lack of followup in the setting of an elevated PSA?







Open Discussion: Questions & Answers

Survey Time! Participant Site Team Members Only





How to Use a QR Code



- 1. Turn on your phone camera
- 2. Aim the camera at the code
- 3. A link will show up
- 4. **Tap** the link to go to the survey

Take it now! Survey closes next Tuesday, April 23rd





Session 3 Slides, Recordings, & Resources will be made available within one week on the <u>ACS ECHO Website.</u>





Is **Session 4** in your calendar? **Tuesday, May 21, 2024** 4:30 PM ET • 3:30 PM CT • 2:30 PM MT • 1:30 PM PT

Topic: Increasing relationships and streamlined referral pathways with urologists within the medical neighborhoodCase Presentation: Family Circle of Care Health Centers

Thank You!

See you again

Tuesday, May 21st at 4:30 PM ET • 3:30 PM CT • 2:30 PM MT • 1:30 PM PT in iECHO Zoom



Post-Session 3 Survey