Prostate Cancer Screening

Ahmad Shabsigh, MD, FACS
Assistant Professor
Department of Urology
The Ohio State University Wexner Medical Center

The Committee:

U.S. Preventive Services Task Force

The Date: August 2008

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The issue:

Prostate Cancer Screening



SCREENING FOR PROSTATE CANCER CLINICAL SUMMARY OF U.S. PREVENTIVE SERVICES TASK FORCE RECOMMENDATION

Population	Adult males
Recommendation	Do not use prostate-specific antigen (PSA)-based screening for prostate cancer. Grade: D



SCREENING FOR PROSTATE CANCER CLINICAL SUMMARY OF U.S. PREVENTIVE SERVICES TASK FORCE RECOMMENDATION

Population	Adult males
Recommendation	Do not use prostate-specific antigen (PSA)-based screening for prostate cancer. Grade: D

The reduction in prostate cancer mortality 10 to 14 years after PSA-based screening is, at most, very small, even for men in the optimal age range of 55 to 69 years.

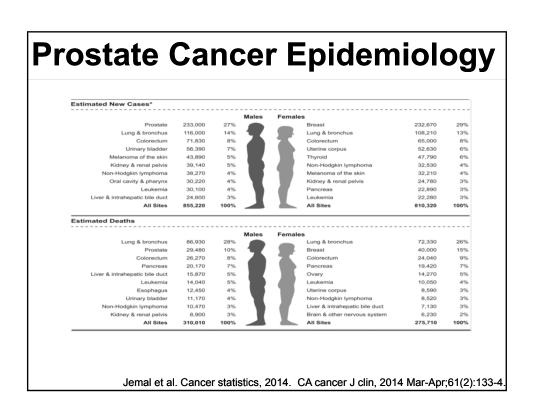
The harms of screening include pain, fever, bleeding, infection, and transient urinary difficulties associated with prostate biopsy, psychological harm of false-positive test results, and overdiagnosis.

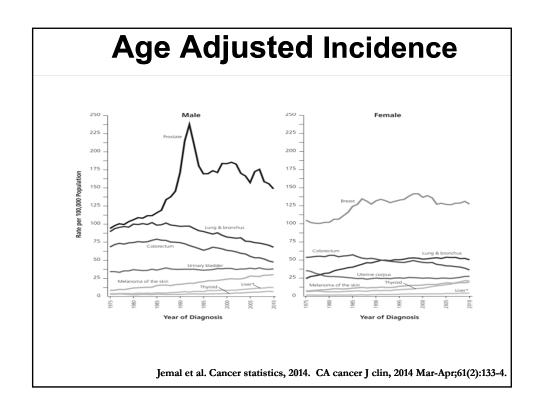
Harms of treatment include erectile dysfunction, urinary incontinence, bowel dysfunction, and a small risk for premature death.

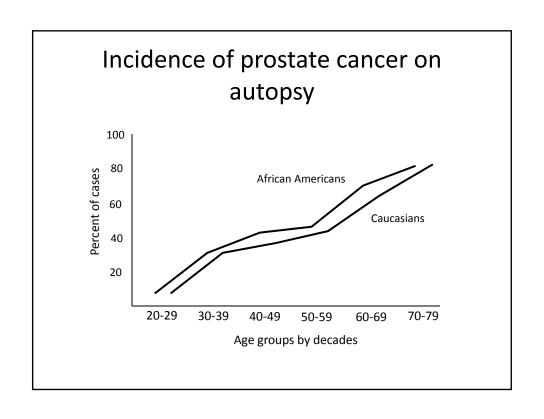
Because of the current inability to reliably distinguish tumors that will remain indolent from those destined to be lethal, many men are being subjected to the harms of treatment for prostate cancer that will never become symptomatic.

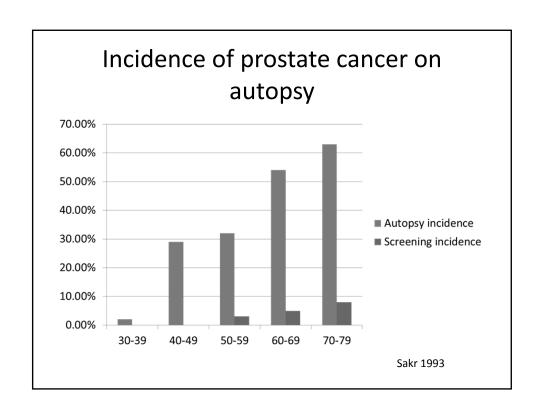
The benefits of PSA-based screening for prostate cancer do not outweigh the harms.

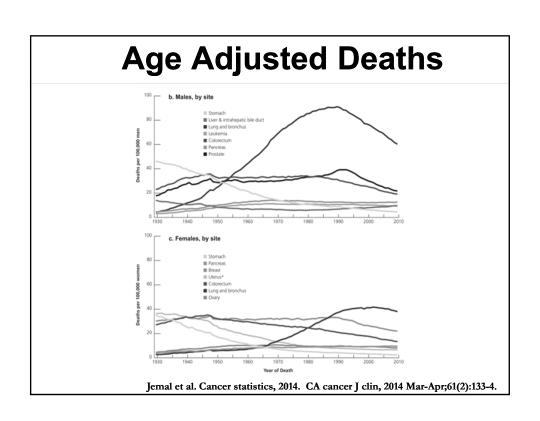
The Impact of the Disease











Lifetime Risk of Dying from CaP

- Risk of dying from prostate cancer is ~3%
- Once metastatic disease develops there is no cure
- Prior to PSA screening only 25% of CaP presented confined to prostate vs. 91% since
- 5 year CSS rates increased from ~70% to 100% (from 1980s to early 2000s)

Jemal et al. Cancer statistics, 2010. *CA cancer J clin, 2011 Mar-Apr;61(2):133-4.* Comprehensive Textbook of Genitourinary Oncology, 3rd edition Catalona et al. Detection of organ-confined prostate cancer is increased through prostate-specific antigen-based screening. *JAMA 1993; 270(8):948*

What is Cancer Screening?

- Checking for disease when there are no symptoms. Since screening may find diseases at an early stage, there may be a better chance of curing the disease.
- · The source: NCI

What Is Prostate Cancer Screening?

- HPI
- DRE
- PSA

Prostate Specific Antigen

- Discovered in 1979 by Wang et al
- Approved by FDA in 1986
- Produced by prostate and periuretheral glands epithelial cells
- Liquefaction of seminal coagulum
- Serine protease from the kallikrein family
- · In serum, most is bound

Prostate Specific Antigen

- Inflammation, hyperplasia, neoplasia lead to disruption of physiological barriers and increased serum PSA levels
- Half life is 2-3 days
- Used for
 - > Initial diagnosis of disease and screening
 - Monitor for recurrence after initial therapy
 - > Prognosis of outcomes after therapy

Prostate Cancer Screening

- Controversial:
 - Prostate cancer has a relatively slow course, Long term follow up is needed (>15 years).
 - > Patient's age
 - > Comorbidities
 - > Treatments are associated with significant morbidity
 - No comparisons of efficacy between therapeutic options

Screening for Prostate Cancer: *Potential Harms*

- · Additional medical visits
- Adverse effects of prostate biopsies
- Anxiety
- Over diagnosis
- Over treatment
- Morbidity and mortality associated with treatment
- Financial burden

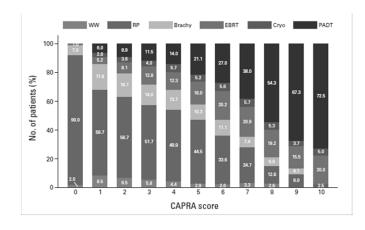
Complications of TRUS Prostate Biopsy

Complications	%
Hematospermia	37.4
Hematuria > 1 d	14.5
Rectal bleeding <2 d	2.2
Prostatitis	1.0
Fever > 101.3°F, epididymitis, rectal bleeding >2 d, retention	<1.0
Other complications requiring hospitalization	0.3

EAU Guidelines

Time Trends and Local Variation in Primary Treatment of Localized Prostate Cancer

Matthew R. Cooperberg, Jeanette M. Broering, and Peter R. Carroll



J Clin Oncol 28:1117-1123.

Screening for Prostate Cancer: *Potential Benefits*

- Early detection and treatment
- Improve cancer Specific Survival
- Alleviate symptoms of locally advanced disease

The Evidence

ERSPC

The European Randomized study of Screening for Prostate Cancer

ERSPC

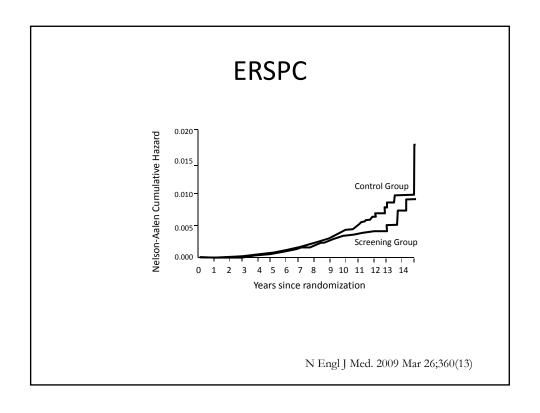
- Primary objective: PC mortality
- Ages 50-74
- 162.387 men
- Screen interval every 4 yrs (87%) Sweeden every 2 yrs (13%)
- Sextant TURS Bx for PSA ≥ 3.0 ng/ml, abnormal DRE, F/T ratio 3-4 ng/ml

N Engl J Med. 2009 Mar 26;360(13)

ERSPC

- Screen 72.890
- Control 89.353
- 85.8% biopsied of the positive tests PPV 24.1
- Median F/U 9 years
- Screen arm: 5990 PC (8.2%), that is 71% higher, 214 deaths
- Control arm: 4.307 PC (4.8%), 326 deaths

N Engl J Med. 2009 Mar 26;360(13)



ERSPC

- 20% fewer men die of PC in the screen group (p=0.04)
- Adjustment for non compliance, 27% fewer deaths in the screened men
- Absolute risk reduction 7 per 10.000 screened men
- NNS: 1.410, NNT: 48 in excess of the control arm.
- NNT to prevent mets 24
- All centers showed the same outcome (16-26%)

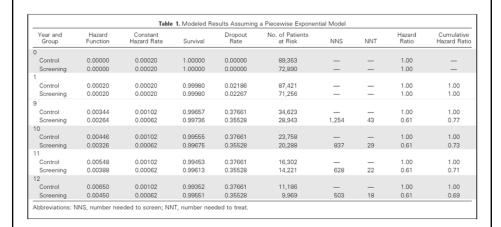
N Engl J Med. 2009 Mar 26;360(13)

Number Needed to Treat (NNT)

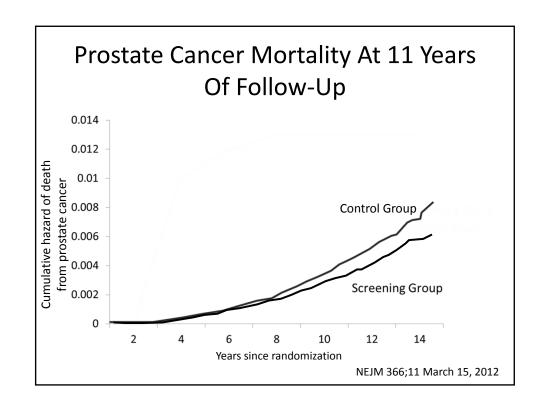
- Estimates Will Decrease
- In Northern Ireland (with very little screening), the NNT to prevent 1 case of metastatic prostate cancer was only 15
- THAT IS similar to the NNT to prevent 1 breast cancer death through mammography screening and follow-up surgery
- The number needed to treat to save 1 life with prostate cancer screening will decrease with correction for compliance and longer follow-up

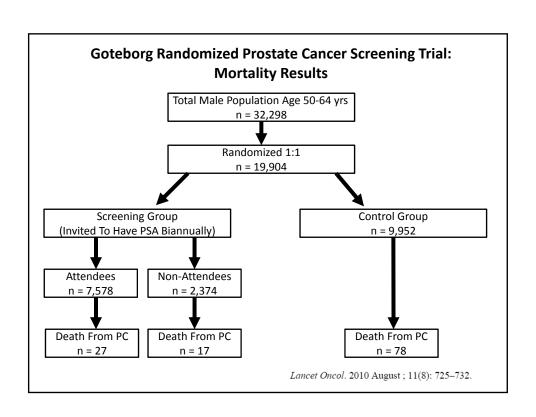
Roobol MJ et al, Eur Urol 56: 592, 2009

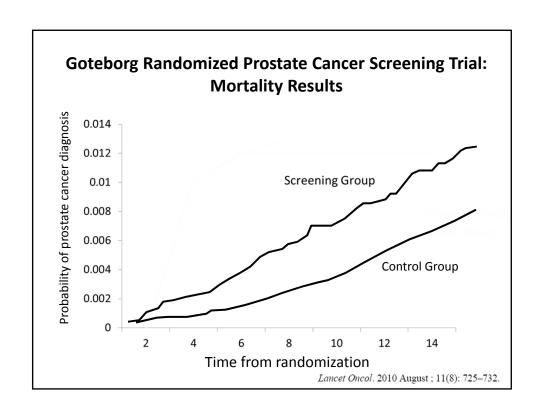
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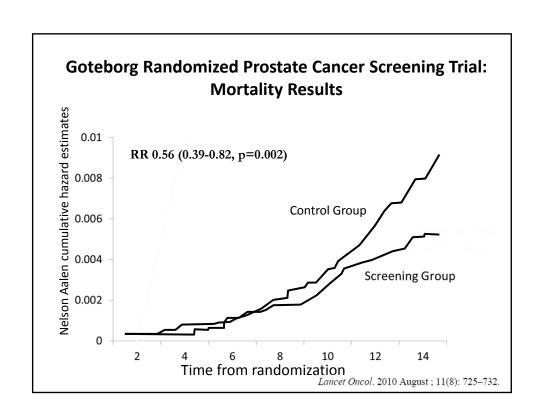


Catalona, J Clin Oncol 2011, 29:464-467









PLCO

The US Prostate, Lung, Colorectal, and **Ovarian Cancer Screening Trial**

PLCO

- 74,000 ages 55 to 74 years
- 1:1 randomization to receive annual PSA and DRE screening to a total of 4 screens vs usual care in the community
- PSA cut-off is 4 ng/mL
- Follow-up of abnormal screening results was at the discretion of physicians
- In the screening group, rates of compliance were 85%
- Rate of screening in control arm 40\$ in first year
- and 52% in sixth yrs
 Rates of screening in the control group
 increased from 40% in the first year to 52% in the sixth year
- 7 years of follow-up

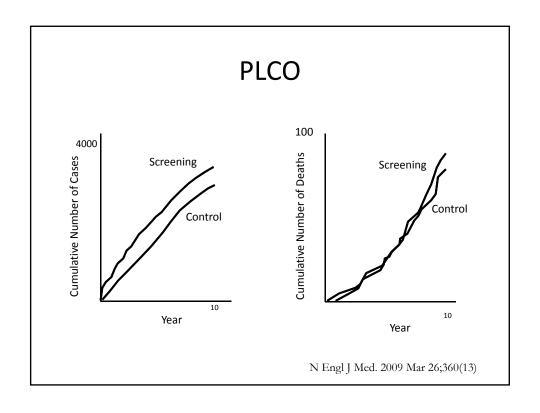
N Engl J Med. 2009 Mar 26;360(13)

Variable	Screening Group (N=38,343)	Control group (N=38,350)
	Pe	rcent
Age		
55-59 yr	32.3	32.3
60-64 yr	31.3	31.3
65-69 yr	23.2	23.2
70-74 yr	13.2	13.2
Race or ethnic group†		
Non-Hispanic white	86.2	83.8
Non-Hispanic black	4.5	4.3
Hispanic	2.1	2.1
Asian	4.0	3.9
Other	0.8	0.9
Missing data	2.4	5.0
Enlarged prostate or benign prostatic hyperplasia	21.4	20.5
Previous prostate biopsy	4.3	4.3
Family history of prostate cancer	7.1	6.7
PSA test within past 3 yr		
Once	34.6	34.3
Two or more times	9.4	9.8
Digital rectal examination within past 3 yr		
Once	32.8	31.9
Two or more times	22.2	22.0

PLCO

- Screen: PC 116 per 10,000 person-years (2820), 50 deaths
- Control: PC 95 per 10,000 person-years (2322), 44 deaths
- Rate ratio, 1.13; 95% CI, 0.75 to 1.70). The data at 10 years were 67% complete and consistent with these overall findings.

N Engl J Med. 2009 Mar 26;360(13)



PLCO

- 40%-52% of controls were screened during the study (contamination) thus, comparing 85% vs 52% screened
- Poor prompt Bx compliance for PSA > 4
- Reported PCa mortality at 7-10 yr (med 11.5) but f/u was only 5.3 to 6.2 years for PCa patients
- 10-year prostate cancer detection rate was only
 15% higher in screened men 9.0% vs 7.8%
- PCa death rate = 2.0 screened vs 1.7 control /104 per-yr
- Authors conclude: no mortality benefit from screening

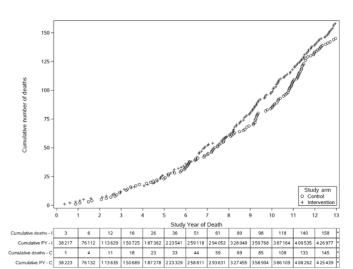
Assessing contamination and compliance in the prostate component of the prostate, lung, colorectal, and ovarian (PLCO) cancer screening trial

			Time Period	Of Latest Test		
		< 1 year	1-2 years	2-3 years	> 3 years	
PSA	# Men Surveyed		Routine Use (%)			
0	181	33	15	3	2	38
1	422	31	14	6	5	34
2	385	41	17	5	4	24
3	410	39	16	8	5	21
4	435	46	15	7	3	17
5	392	46	18	5	3	15
0-5	2225	40	16	6	4	23
0-5 adjusted		46	14	5	4	21
0-5 screened arm		78	8	3	2	9
DRE						
0-5	2336	28	17	17	9	28
PSA or DRE						
0	196	39	16	6	10	20
1	454	37	20	8	10	15
2	415	49	17	7	6	13
3	450	43	20	10	7	12
4	466	49	17	7	6	12
5	418	52	22	5	5	8
0-5	2399	46	19	7	7	13
0-5 adjusted		51	17	6	6	12

Clinical Trials 2010; 7: 303-311

Prostate Cancer Screening in the Randomized Prostate, Lung, Colorectal, and Ovarian Cancer Screening Trial: Mortality Results after 13 Years of Follow-up

Gerald L. Andriole, E. David Crawford, Robert L. Grubb III, Saundra S. Buys, David Chia, Timothy R. Church, Mona N. Fouad,



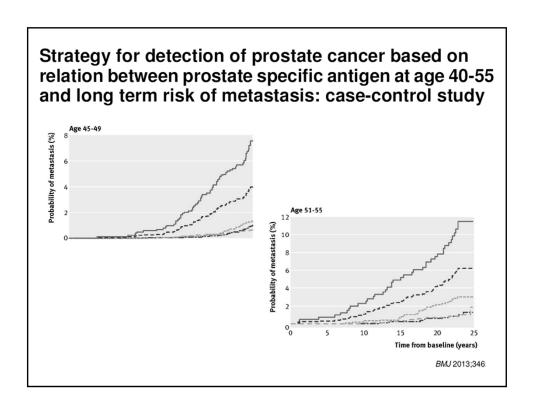
A Smarter Way to Screen for Prostate Cancer

Smarter Screening

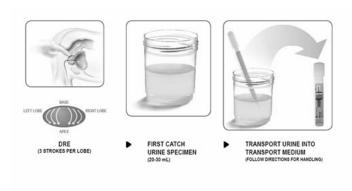
- Risk-adjust screening by age, comorbidities, family history, ethnicity and PSA (reduce false positives)
- Reduce false positive PSA results by repeating (verifying) positives and by adding additional markers (reduce indications for biopsy)
- Active surveillance for low-risk cancers (reduce harms of unnecessary therapy)
- Refer patients who need treatment to experienced high-volume physicians or centers (reduce harm of necessary therapy)

		Proporti	ion (95% CI)	
PSA concentrat	ion (µg/L)	Deaths	Metastases	
Age 45-49 at ba	seline scree	n		
Highest 10 th	<u>></u> 1.6	44 (34 to 53)	40 (33 to 48)	
Highest quarter	<u>></u> 1.06	54 (45 to 63)	51 (44 to 59)	
Below median	<0.68	28 (20 to 37)	28 (22 to 35)	
Age 51-55 at se	cond screen			
Highest 10 th	<u>≥</u> 2.4	44 (32 to 56)	42 (32 to 52)	
Highest quarter	<u>></u> 1.4	59 (47 to 71)	56 (46 to 66)	
Below median	<0.85	16 (7 to 25)	18 (10 to 26)	

BMJ 2013;346



PCA3 Screening



PCA3 Screening

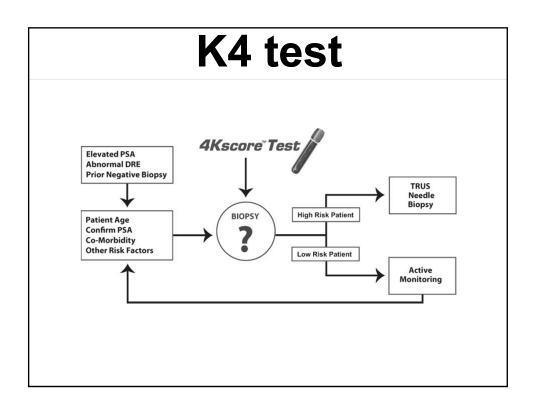
- PCA3 is a non-coding mRNA molecule that is believed to be prostate specific.
 - It is highly over-expressed in cancerous prostate cells relative to benign tissue
 - Present in urine (no blood test necessary)
- Potential to be used as supplement for PSA testing
 - > PSA has a 21% specificity but a 87% sensitivity for prostate cancer
 - Conversely, a test for PCA3 was reported to have a sensitivity of only 49%, but a specificity of 78%
 - Additional studies are needed

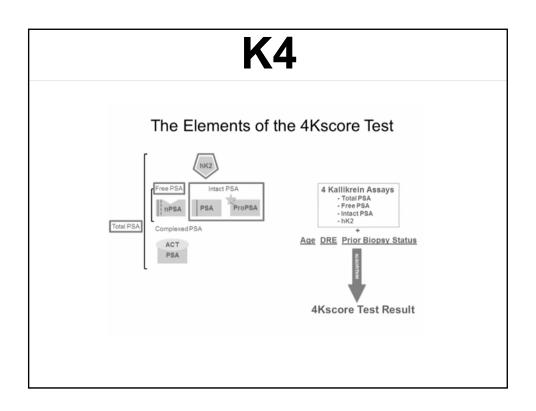
PCA3 Screening

Table 2: Operating Characteristics of PCA3 vs. PSA in 225 Men Undergoing Prostate Re-Biopsy

PCA3/PSA mRNA ratio vs. Serum PSA: Previous negative biopsy group

	PCA3 Assay	Serum PSA
Cutoff	PCA3/PSA = 35 x 10 ⁻³	4.0 ng/mL
Sensitivity	58%	83%
Specificity	74%	17%
*ROC AUC	0.680	0.506
Odds ratio	3.6	1.2
*P = 0.002		
		•





A panel of kallikrein marker predicts prostate cancer in a large, population-based cohort followed for 15 years without screening

Andrew Vickers, PhD 1 , Amit Gupta, MD 2 , Caroline J. Savage, MPH 1 , Kim Pettersson, PhD 5 , Anders Dahlin, PhD 6 , Anders Bjartell, MD, PhD 7 , Jonas Manjer, MD, PhD 8 , Peter T. Scardino, MD 3 , David Ulmert, MD, PhD 3 , 6 , and Hans Lilja, MD, PhD 2 ,3,4,6

	Biopsies		Prostate cancers		Palpable prostate cancers		Advanced prostate cancer	
	Performed	Avoided (%)	Found	Delayed	Found	Delayed	Found	Delayed
Cancers diagnosed within 5 years from baseline								
Biopsy all (i.e., PSA ≥3.0 ng/ml)	1000	0	152	-	112	-	49	-
Biopsy based on age-specific PSA threshold*	539	461(46%)	112	40	88	24	40	9
Biopsy those with PSA ≥4.0 ng/ml	637	363(36%)	128	23	100	12	46	4
Biopsy if risk on full kallikrein panel is ≥20%	579	421(42%)	131	21	102	9	47	2
_								

Cancer Epidemiol Biomarkers Prev. Author manuscript; available in PMC 2012 March 1.

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	Biopsies		Prostate cancers		Palpable prostate cancers		Advanced prostate cancer	
	Performed	Avoided (%)	Found	Delayed	Found	Delayed	Found	Delayed
Cancers diagnosed	seline							
Biopsy all (i.e., PSA ≥3.0 ng/ml)	1000	0 (0%)	367	-	242	-	143	-
Biopsy based on age-specific PSA threshold*	539	461(46%)	265	103	192	50	115	28
Biopsy those with PSA ≥4.0 ng/ml	637	363(36%)	285	82	202	40	120	23
Biopsy if risk on full kallikrein panel is ≥20%	579	421(42%)	299	68	210	31	129	14

 ${\it Cancer\ Epidemiol\ Biomarkers\ Prev.}\ {\it Author\ manuscript;\ available\ in\ PMC\ 2012\ March\ 1}.$

A panel of kallikrein marker predicts prostate cancer in a large, population-based cohort followed for 15 years without screening

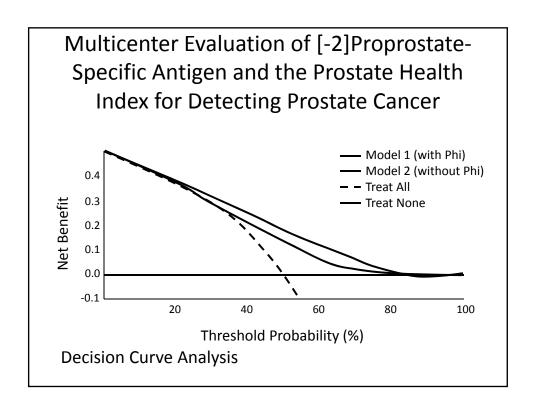
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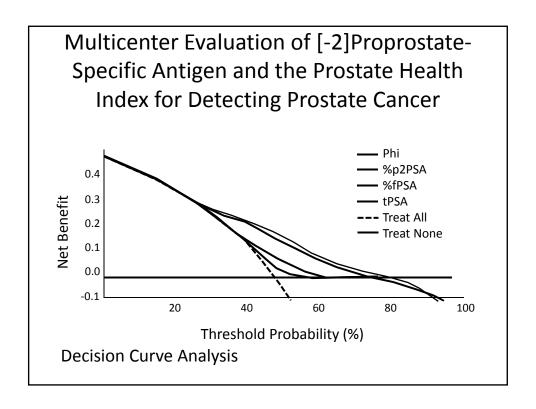
	Any Prostate Cancer	Palpable Prostate Cancer (clinical stage T2 or higher at diagnosis	Advanced Prostate Cancer (clinical stage T3 or higher or evidence of metastasis at diagnosis
Base Model	0.654 (0.621, 0.683)	0.708 (0.671, 0.741)	0.716 (0.664, 0.762)
Full model	0.751 (0.726, 0.777)	0.803 (0.774, 0.831)	0.824 (0.785, 0.858)
Full model without hK2	0.752 (0.728, 0.782)	0.803 (0.777, 0.832)	0.825 (0.784, 0.855)
Full model without intact PSA	0.711 (0.680, 0.746)	0.746 (0.706, 0.779)	0.754 (0.698, 0.801)
Full model without free PSA	0.654 (0.619, 0.689)	0.698 (0.650, 0.731)	0.695 (0.638, 0.751)
Model including total PSA, free PSA and age only	0.692 (0.664, 0.719)	0.723 (0.686, 0.764)	0.720 (0.658, 0.771)

Cancer Epidemiol Biomarkers Prev. Author manuscript; available in PMC 2012 March 1.

Prostate Health Index (PHI)

• ([-2]proPSA/free PSA) $\times \sqrt{PSA}$.

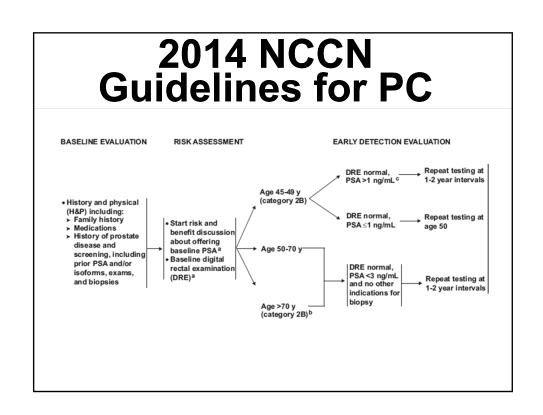




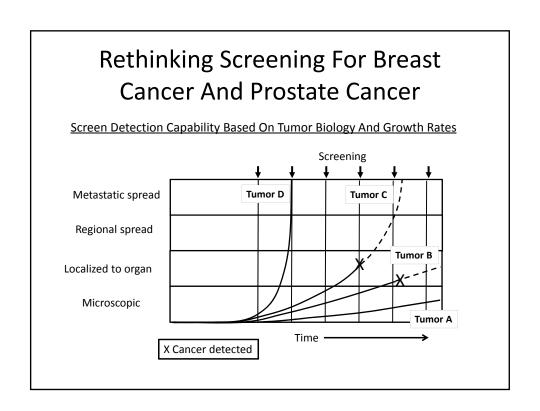
Where do we stand?

AUA

- No screening < 40 yrs.
- No routine screening in men 40 to 54 yrs at average risk.
- Individualized for high risk < 55 yrs
- shared decision-making for 55 to 69 yrs
- Every 2 or more yrs according to baseline PSA
- No screening for >70 yrs or any man with less than a 10 to 15 year life expectancy.
- Some men age 70+ years who are in excellent health may benefit from prostate cancer screening.



Rethinking Screening for Cancer



Benefit and Burden of Mammographic Screening and Prostate-Specific Antigen Screening in the United States and Europe

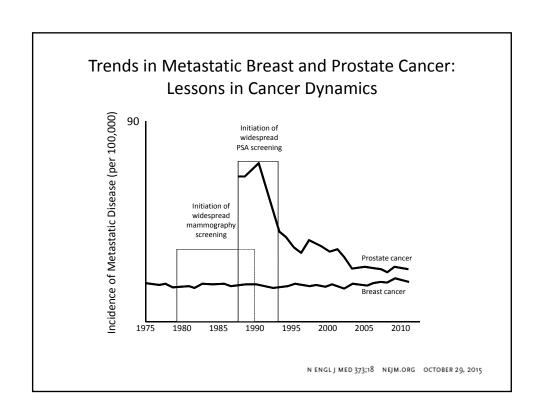
Breast Cancer

Region	Deaths Averted	Cancers Detected, Treated	Biopsies/ Recalls	Screening Visits	Individuals Screened (#)	Years Of Screening (#)
U.S.	1	18 Invasive 6 DCS	90/535	5866	838	6
Europe	1	15 Invasive 5 DCS	41/162	3352	838	6

Prostate Cancer

Region	Deaths Averted	Cancers Detected, Treated	Biopsies/ Recalls	Screening Visits	Individuals Screened (#)	Years Of Screening (#)
U.S.	0					
Europe	1	48		2397	1410	9

JAMA 2009; 302:1685



Conclusions

• PSA is not a perfect screening test

Conclusions

 PSA is not a perfect screening test (But it is the best we have)

Conclusions

PSA is not a perfect screening test

(But it is the best we have)

 Yes most men will have PC and most will not die from it

Conclusions

PSA is not a perfect screening test

(But it is the best we have)

- Yes most men will have PC and most will not die from it
- Tens of thousands die from the disease, and the numbers will increase with increased life expectancy

Conclusions

PSA is not a perfect screening test

(But it is the best we have)

- Yes most men will have PC and most will not die from it
- Tens of thousands die from the disease, and the numbers will increase with increased life expectancy
- PSA screening for PC detects cancers earlier and at a lower stage where curative therapies more effective

Conclusions

PSA is not a perfect screening test

(But it is the best we have)

- Yes most men will have PC and most will not die from it
- Tens of thousands die from the disease, and the numbers will increase with increased life expectancy
- PSA screening for PC detects cancers earlier and at a lower stage where curative therapies more effective
- PC screening saves lives



Cracks on Airbus A380 Wings

- January 2012: Qantas A380 plane encounters severe turbulence on London-Singapore flight
 - Aircraft checked and cleared to fly on to Sydney
- <u>February 5, 2012</u>: Plane grounded in Sydney after further precautionary inspection finds 36 hairline cracks on the wing rib brackets similar to "Type 1" cracks found on previous A380 checks



When It Comes to Prostate Cancer:

"Diagnostically aggressive"

Peter T. Scardino, MD

When It Comes to Prostate Cancer:

"Diagnostically aggressive"

"Therapeutically conservative"

Peter T. Scardino, MD