

Collaborative-Wide Meeting

October 18, 2024



Nonprofit corporations and independent licensees of the Blue Cross and Blue Shield Association



Welcome

Khurshid Ghani, MD, MS, FRCS



A community that partners to improve patients' lives by inspiring high-quality care through data-driven best practices, education, and innovation

Agenda



- Welcome & General Updates
- Prostate Persistent and Biochemically Recurrent Cancer after Prostatectomy
- KIDNEY Initial Dive into Cancer-Specific Outcomes
- Keynote Physician Wellness and.
 WellPrept Pilot

- Lunch
- BPH QI Opportunities
- ROCKS Improving URS Practice: Lessons from an Ongoing Clinical Trial
 - Closing Remarks

Data Abstractor Session – Arbor Research Training





10 International Webinars





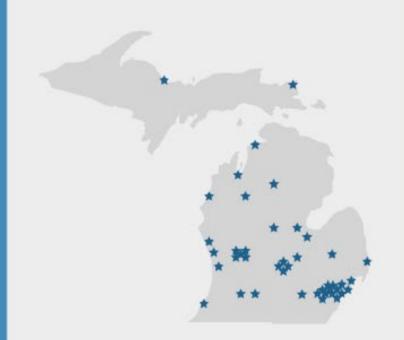
4 Clinical Trials >\$6M Funding

>\$100M Heathcare Cost Savings



Impact of





43 MI Practices
3 Non-MI Practices





>270 Urologists

15 Patient Advocates





>25,000 Patients' Lives Improved

Thank You! COORDINATING CENTER STATISTICIANS





Rod Dunn



Stephanie Daignault-Newton



Sabir Meah



Junzhi Sun



Caitlin Seibel

Welcome! MEMBERS and GUESTS



Guest Speakers



David Canes
Lahey Hospital and Medical Center



Daniel Krauss
Corewell Health

MUSIC Urologist



Henry Rosevear
Michigan Institute of Urology

Guest

Lei Wang, MD

Patient Advocates

Doug Adams

BCBSM Partners

- Marc Cohen
- Emily Santer
- Monica Whitted



Updates

Congrats Dr. Miller!





September 19, 2024

Share on: X f



David Miller to become EVP for medical affairs, Michigan Medicine CEO

Known for a spirit of continuous improvement, he'll start July 2025



Clinical Registry Transition





 Moving to Arbor Research to improve data collection

Abstractor training begins today

Go-live 11/11

In God we Trust.

All others must

bring Data.

-Dr. Robert Fletcher,

Patient Advocate, quoting
W. Edwards Deming



BLUES Clinical Trial Completes Enrollment!



Thank you very much!



















NCT#05026710

PCP Engagement & Working Group



Payment

BCBSM uplift payments for PCPs involved in specific aspects of urologic care

Templating

EHR templating for urologists in their communications to PCPs

Educational Materials

PCP education around specific urologic guidelines

Central Hub

Physician and patient education available via a single website



Spanish Resources Now Available







Value Based Reimbursement

2025 (payout) VBR Metrics



Performance Measure	Baseline Performance	Target Performance	Current Performance
PSA testing within 90 days of radical prostatectomy	92%	≥ 95%	93%
Opioid-limited partial and radical nephrectomy	59%	≥ 60%	52%
Ureteral stenting following URS in pre-stented patients	63%	≤ 62%	54%
Radical nephrectomy for benign renal masses	8%	≤ 6%	7%
Renal mass surveillance follow up	45%	≥ 50%	58%
Proportion of smokers who receive smoking cessation counseling	82%	≥ 85%	81%
Proportion of smokers who quit smoking at 3 months post-RP	27	≥ 30%	31%

MUSIC members will be eligible for a 2% VBR uplift in 2025

2026 Standard VBR (3%): Collaborative-Wide



Population-based Performance Measure	Baseline Performance	Target Performance
Prostate: Active Surveillance Follow-Up	87%	≥ 89%
ROCKS: Post-URS Ureteral Stent Duration	15%	≤ 13%
KIDNEY : Active Surveillance Follow-Up	29%	≥ 35%

COLLABORATIVE must meet 2 of 3 metrics

2026 Additional VBR (2%): Practice-Level



Population-based Performance Measure*	Baseline Performance	Target Performance
Prostate: Post-RP PSA	89%	≥ 92%
ROCKS: PRO Enrollment	54%	Current practices: Maintain or improve by 5% New Practices: ≥ 30%
KIDNEY: Opioid-limited Partial and Radical Nephrectomy	52%	Maintain or improve by 5%

PRACTICES must meet 1 of 3 metrics

2026 Smoking Cessation VBR (2%):

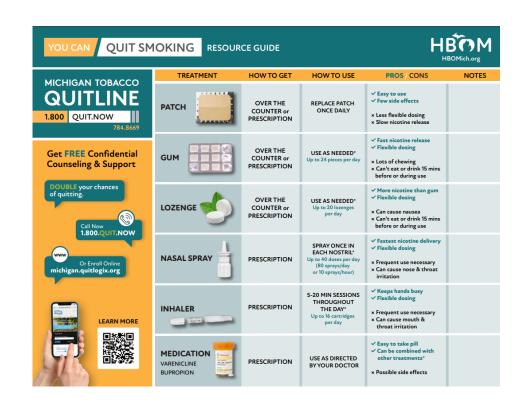
TYUSIC Michigan Urotopical Surgery

Collaborative-Wide

Data collected via RP PRO surveys

- Metrics
 - Pre-op cessation counseling for smokers
 - Smokers who quit by 3 months post-op

Resources available on MUSIC website



COLLABORATIVE must meet 2 of 2 metrics

2026 VBR Participation: Physician-Level —



Each **physician** must do at least 1 of the following from July 1, 2024 – June 30, 2025 to earn any VBR:

- 1) Attend a collaborative-wide meeting
- 2) Attend a skills workshop
- 3) Attend your MUSIC site visit
- 4) View your reports in the registry (coming in 2025)

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Persistent and Biochemically Recurrent Cancer after Prostatectomy

Understanding and Managing the Challenges



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Two patients undergo prostatectomy for localized disease

Patient 1

Pathology shows T3bN0, GG4, multifocal positive margin PSA is 0.3 six weeks after RP

Patient 2

Pathology shows T2Nx, GG2
Undetectable PSA for 3 years, rises
from 0.12 to 0.3 over next 2 years

Should they be treated differently?

More Than Words: Defining Adjuvant, Consolidative, EUROPEAN and Salvage Treatment after RP



Clinical state	Definition of clinical state	Treatment recommendation
Undetectable PSA after RP: Absence of disease with low clinical suspicion	Undetectable PSA Favorable pathology	Surveillance
Undetectable PSA after RP: Absence of disease with high clinical suspicion	Undetectable PSA Concerning pathology o pN+ o ISUP grade group 4-5 AND pT3b-4 ± positive margin	Surveillance or adjuvant therapy
PSA persistence after RP : Persistence of disease with low clinical suspicion	Detectable PSA immediately after RP Favorable pathology Absence of radiographic evidence of disease	Surveillance or consolidative therapy
PSA persistence after RP: Persistence of disease with high clinical suspicion	Detectable PSA immediately after prostatectomy Concerning pathology o pN+ o pT3-4 ± positive margin o ISUP grade group 3-5Radiographic evidence of disease on molecular imaging	Consolidative therapy
PSA recurrence after RP: Biochemical recurrence	Period of undetectable PSA followed by detectable PSA	Salvage therapy (early) or surveillance
Metastatic disease	Evidence on conventional imaging or pathology	Management of metastatic prostate cancer
ISUP = International Society of Urological pathology;	RP = radical prostatectomy; PSA = prostate-specific antigen.	

Tailoring Treatment for Similar Patient Populations—



Persistently Positive (PP)

Initial PSA 0.14 or higher

Biochemical Recurrence (BCR)

PSA rise to 0.2 from < 0.1



Metastatic Cancer and Death Occur in Patients with PSA Recurrence





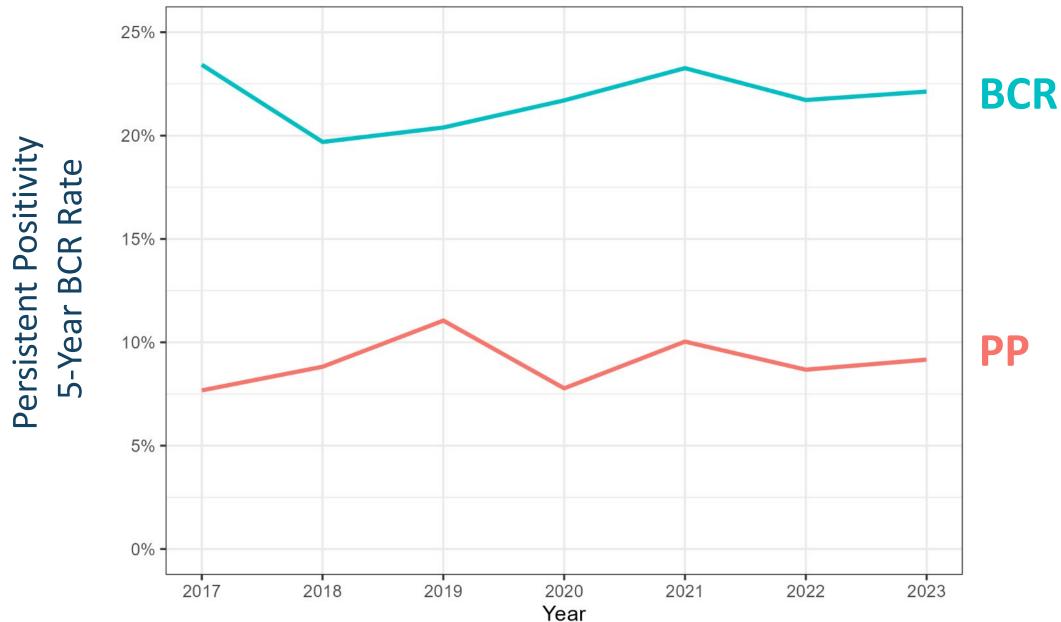
2023 AUA Guideline:

Patients should be informed that the development of a PSA recurrence after surgery is associated with a <a href="https://www.night.n

Congruent with this clinical principle, physicians should <u>regularly</u> <u>monitor PSA</u> after radical prostatectomy to enable early administration of salvage therapies if appropriate. (Clinical Principle)

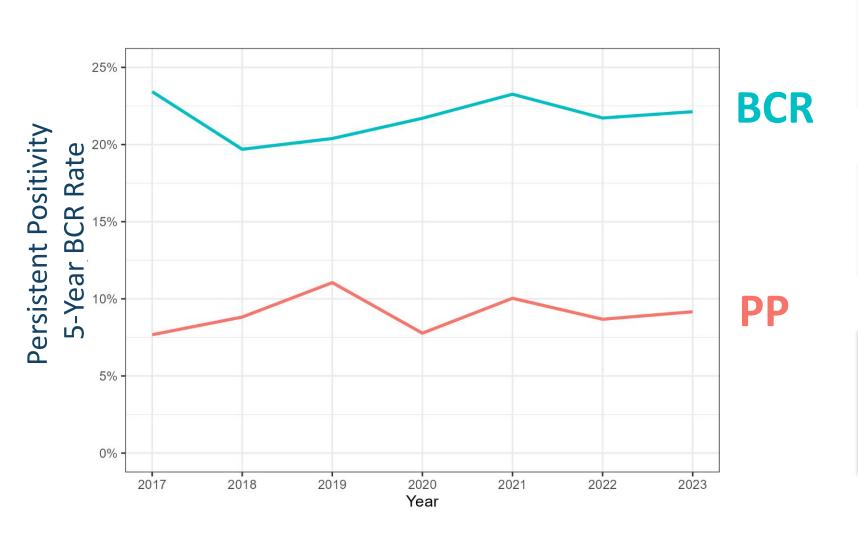
Between 30 and 40% of Patients Have Residual Cancer (PP or BCR) -





One Third of Patients Have Cancer Post RP





MUSIC Rate of PP and BCR is 20%

Predicted Rate is around 33%

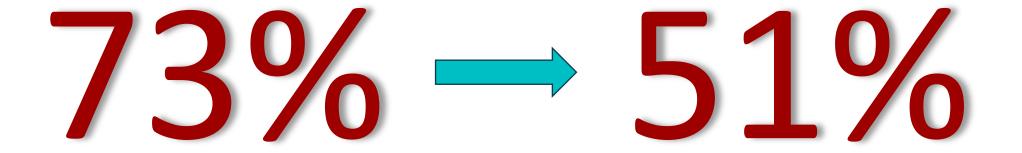
Missing ~10% of patients





Missing ~10% of RP patients with BCR

because we only have post-RP PSAs on

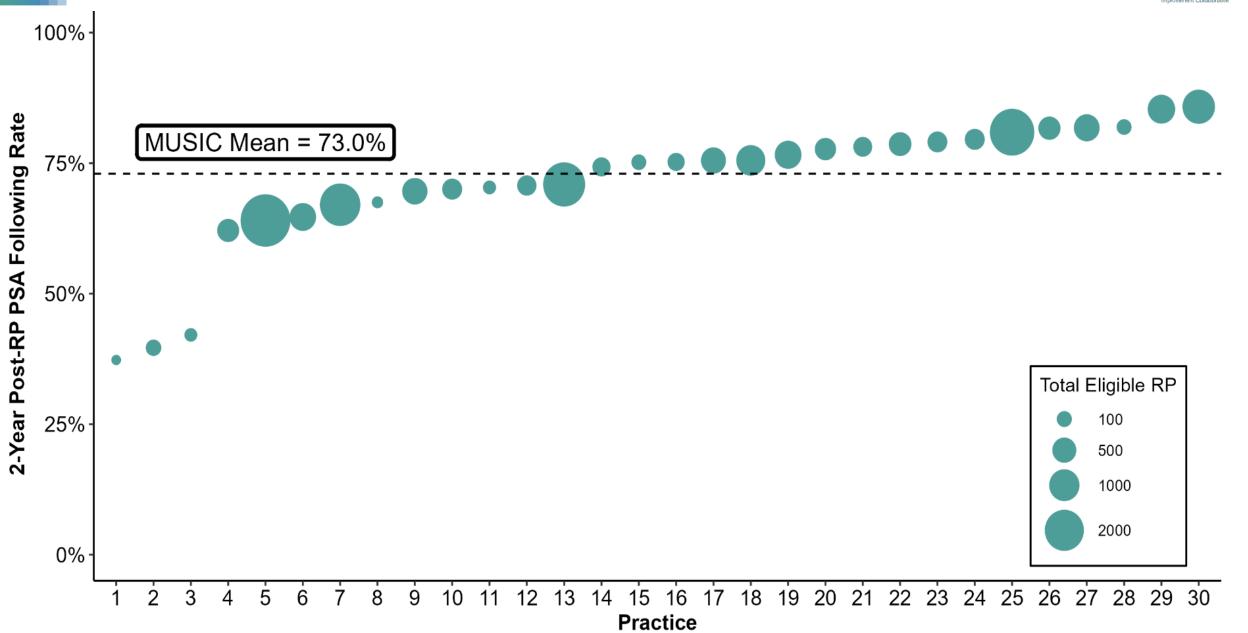


of patients at 2 years

of patients at 4 years

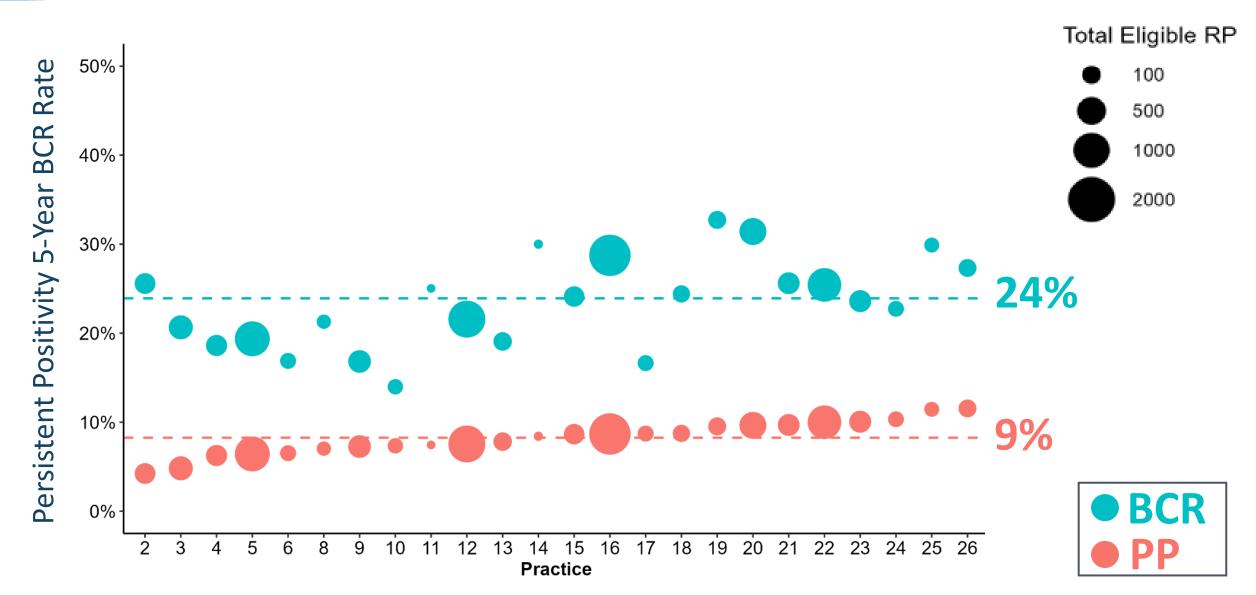
Practice Variation: 2 Year Post-RP PSA Rate





Practice Level Variation: PP and 5-Year BCR







PP and BCR Patients Differ in Pre-Operative Risk



Characteristic	PP N = 1,720	BCR N = 2,617 ¹	p-value ²
Clinical T-Stage			<0.001
T1	938 (55%)	1,595 (61%)	
T2	540 (31%)	757 (29%)	
T3	70 (4.1%)	47 (1.8%)	
Tx	167 (9.7%)	211 (8.1%)	
Pre-Operative PSA	9 (6, 16)	7 (5, 10)	<0.001
Pre-Operative PSA			<0.001
<10	890 (55%)	1,876 (75%)	
10-20	449 (28%)	464 (19%)	
20-50	230 (14%)	150 (6.0%)	
>50	59 (3.6%)	17 (0.7%)	
NCCN Risk Group			<0.001
Very Low	12 (0.7%)	24 (0.9%)	
Low	66 (3.9%)	161 (6.2%)	
Favorable Intermediate	153 (8.9%)	408 (16%)	
Unfavorable Intermediate	583 (34%)	1,160 (45%)	
High	318 (19%)	314 (12%)	
Very High	582 (34%)	539 (21%)	

Persistently Positive more likely with

Stage cT3

PSA >20

High or Very High Risk

PP and BCR Patients Differ in Pathologic Risk



DD	DCD	
N = 2,715 ¹	N = 2,118 ¹	p-value ²
		<0.001
41 (2.4%)	92 (3.5%)	
385 (22%)	1,016 (39%)	
567 (33%)	889 (34%)	
206 (12%)	205 (7.9%)	
513 (30%)	397 (15%)	
436 (25%)	1,144 (44%)	
600 (35%)	976 (37%)	
654 (38%)	491 (19%)	
27 (1.6%)	5 (0.2%)	
1,208 (70%)	1,384 (53%)	<0.001
683 (41%)	499 (19%)	<0.001
1,026 (60%)	1,315 (50%)	< 0.001
		<0.001
1,205 (70%)	2,088 (80%)	
329 (19%)	150 (5.7%)	
186 (11%)	379 (14%)	
627 (36%)	439 (17%)	<0.001
	385 (22%) 567 (33%) 206 (12%) 513 (30%) 436 (25%) 600 (35%) 654 (38%) 27 (1.6%) 1,208 (70%) 683 (41%) 1,026 (60%) 1,205 (70%) 329 (19%) 186 (11%)	N = 2,715¹ N = 2,118¹ 41 (2.4%) 92 (3.5%) 385 (22%) 1,016 (39%) 567 (33%) 889 (34%) 206 (12%) 205 (7.9%) 513 (30%) 397 (15%) 436 (25%) 1,144 (44%) 600 (35%) 976 (37%) 654 (38%) 491 (19%) 27 (1.6%) 5 (0.2%) 1,208 (70%) 1,384 (53%) 683 (41%) 499 (19%) 1,026 (60%) 1,315 (50%) 1,205 (70%) 2,088 (80%) 329 (19%) 150 (5.7%) 186 (11%) 379 (14%)

Persistently Positive patients more commonly have

RP Grade Group 4-5

Stage pT3b - 4

Margin Status

Nodal Disease

Metastatic Cancer and Death Occur in Patients with PSA Recurrence





2023 AUA Guideline:

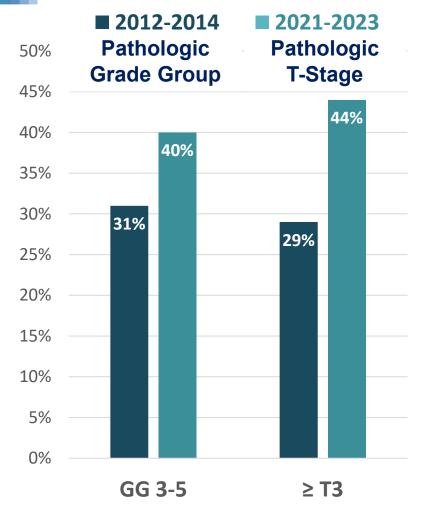
Patients should be informed that the development of a PSA recurrence after surgery is associated with a <u>higher risk</u> of development of <u>metastatic prostate cancer or death</u> from the disease.

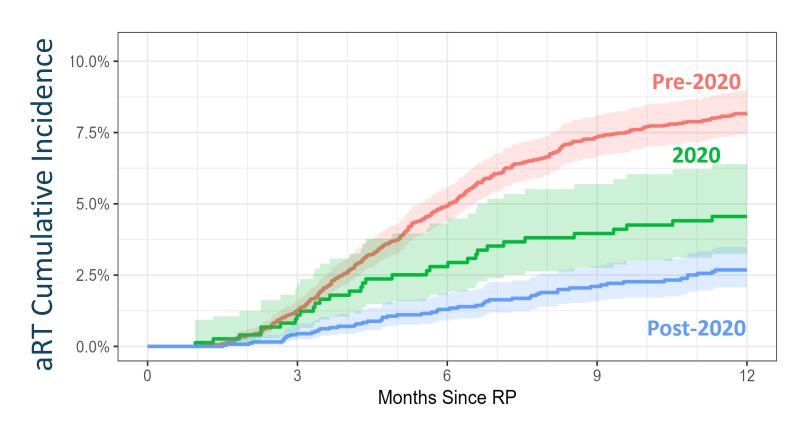
Congruent with this clinical principle, physicians should <u>regularly</u> <u>monitor PSA</u> after radical prostatectomy to enable early administration of salvage therapies if appropriate. (Clinical Principle)

Metastatic Cancer and Death Occur in Patients with PSA Persistence

Post-RP Cancer Becoming a Worse Problem





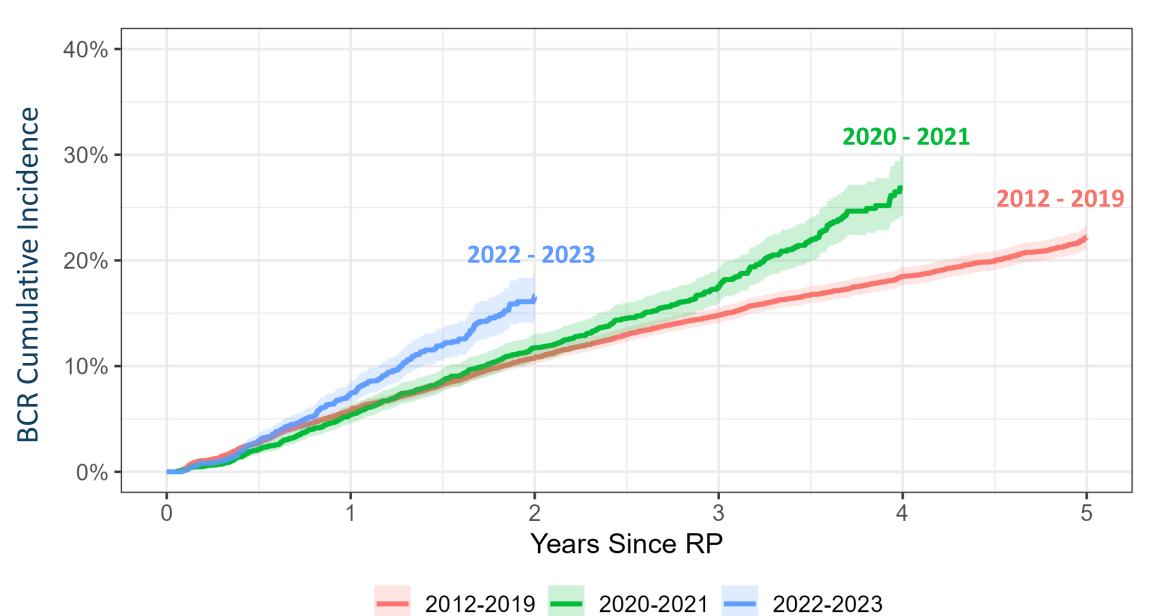


9% increase in Grade Group 3-5 and 15% increase in ≥ T3 Disease

Less adjuvant treatment since 2020

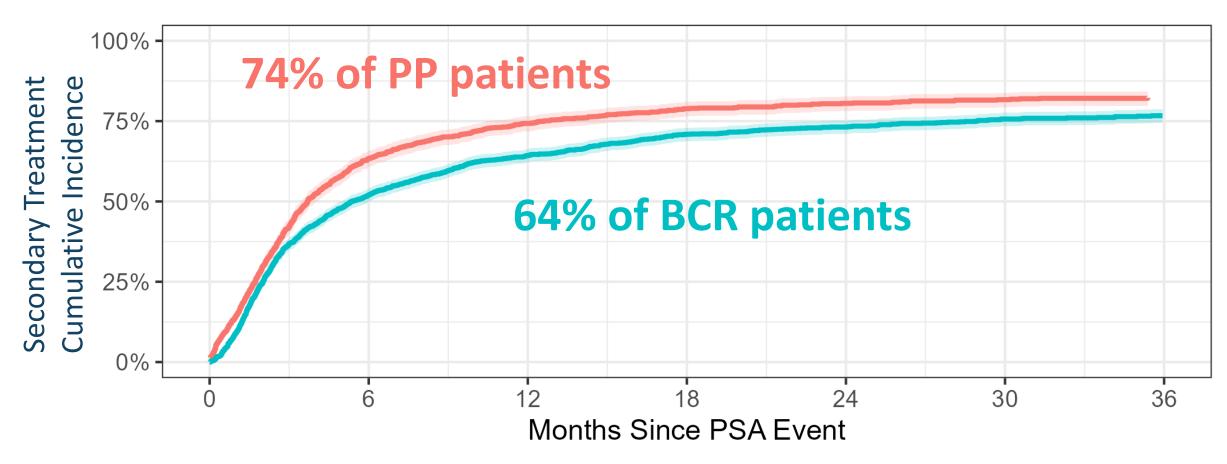
BCR Increasing in Recent Years -





Treatment at 1-Year from PSA Event

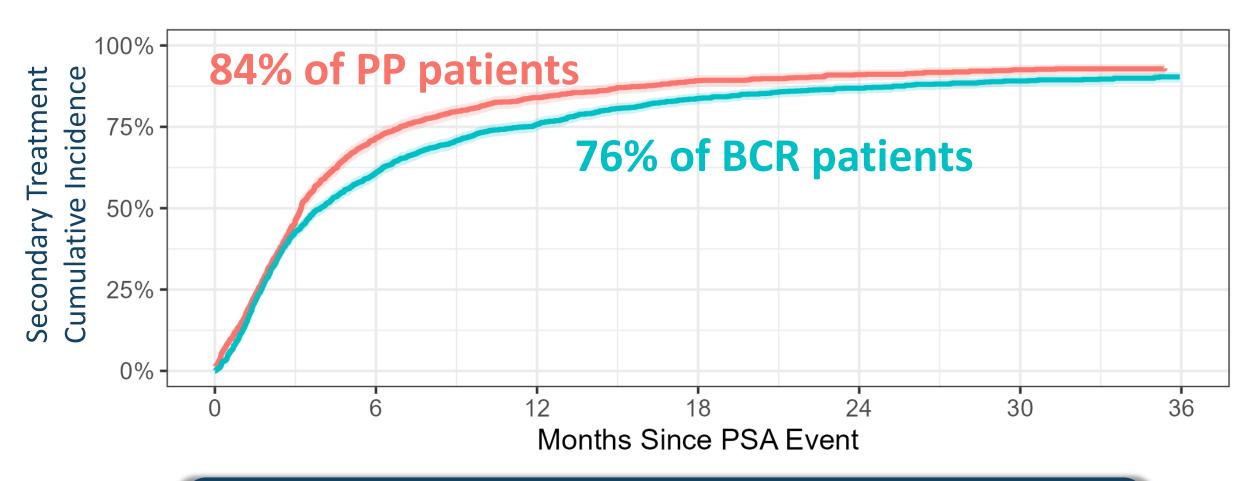




BUT, we know we under capture treatment

Treatment at 1-Year from PSA Event



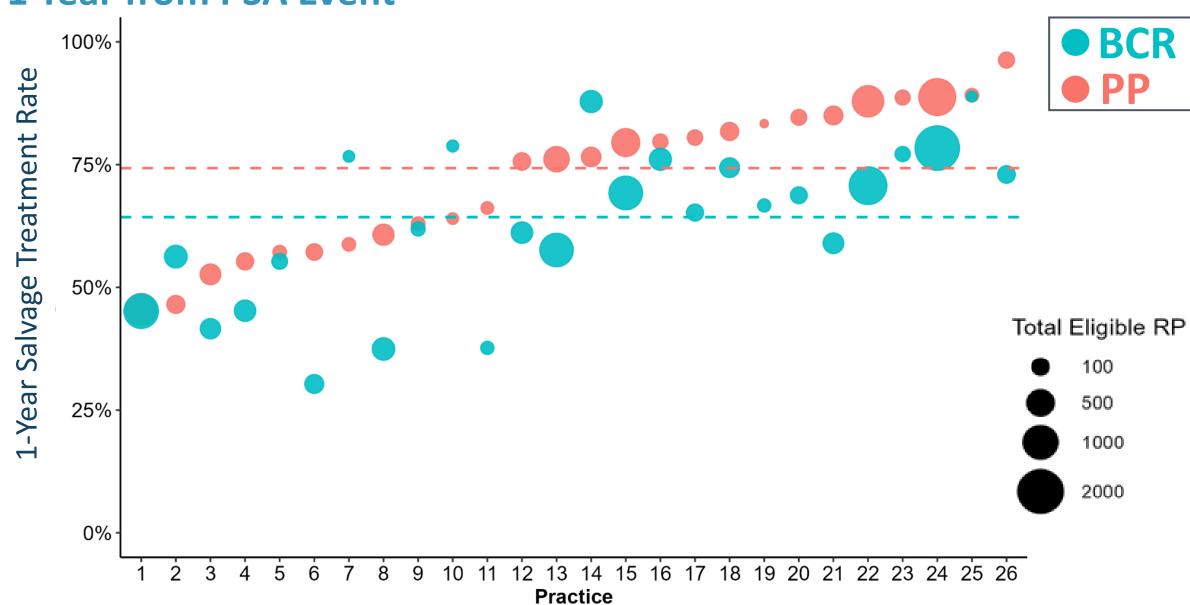


Including patients whose **PSA became undetectable** without recorded treatment

Rates of Consolidative or Salvage Treatment at

IN USIC Michigan University (California)

1 Year from PSA Event





PP and BCR Patients Differ in Post-Operative Risk —



Characteristic	PP N = 1,159	BCR N = 1,579	
Number of PSA Tests Between PP/BCR and Secondary Treatment	2 (1,3)	2 (1,3)	
PSA at PP/BCR Event	0.64 (0.30, 2.20)	0.20 (0.14, 0.27)	
Highest PSA Pre-Secondary Trt (Post-RP)	1.00 (0.40, 2.84)	0.23 (0.17, 0.40)	
PSA at PP/BCR Event			
< 0.1	0 (0%)	0 (0%)	
[0.1, 0.2)	75 (6.5%)	475 (30%)	
[0.2, 0.3)	213 (18%)	727 (46%)	
[0.3, 0.4)	112 (9.7%)	185 (12%)	
[0.4, 0.5)	83 (7.2%)	54 (3.4%)	
[0.5, 0.6)	54 (4.7%)	40 (2.5%)	
[0.6, 0.7)	56 (4.8%)	26 (1.6%)	
[0.7, 0.8)	35 (3.0%)	10 (0.6%)	
[0.8, 0.9)	29 (2.5%)	8 (0.5%)	
[0.9, 1)	31 (2.7%)	7 (0.4%)	
>= 1	471 (41%)	47 (3.0%)	

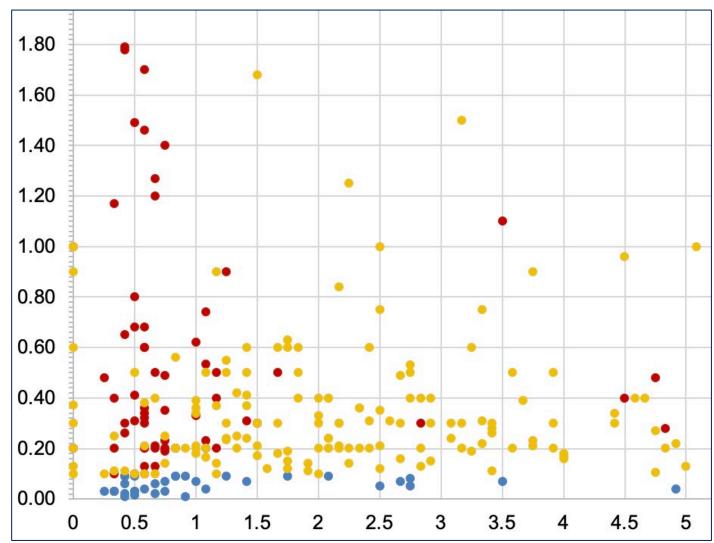
91% BCR patients receiving early salvage (PSA<0.5)

41% PP patients treated at PSA >1

PSA and Time to Treatment for Patients Receiving XRT







One quarter of patients receiving XRT after RP are persistently positive

PSA data consistent with MUSIC

AdjuvantConsolidativeSalvage

Years from Radical Prostatectomy to Initiation of Radiation

Key Takeaways



- One third of patients undergoing prostatectomy have cancer post-op
 - Disease aggressiveness INCREASING over recent years
 - High quality PSA surveillance imperative
 - MUSIC likely missing 10% of patients with BCR
 - Patients with more advanced disease at substantially higher risk
- Residual cancer either persistently positive or biochemically recurrent
 - Different disease characteristics → different approach to management
 - Risk adapted timely subsequent treatment critical
- Shared management of patients with Radiation Oncology colleagues
 - Mutual understanding of treatment goals and potential morbidity



Radiation Oncology: Key Player in Managing Biochemical Recurrence

Daniel Krauss, MD
Radiation Oncologist at Corewell Health





Post-Prostatectomy Radiation Therapy: Approaches to Varying Clinical Presentations

Daniel J. Krauss, M.D.

Professor of Radiation Oncology

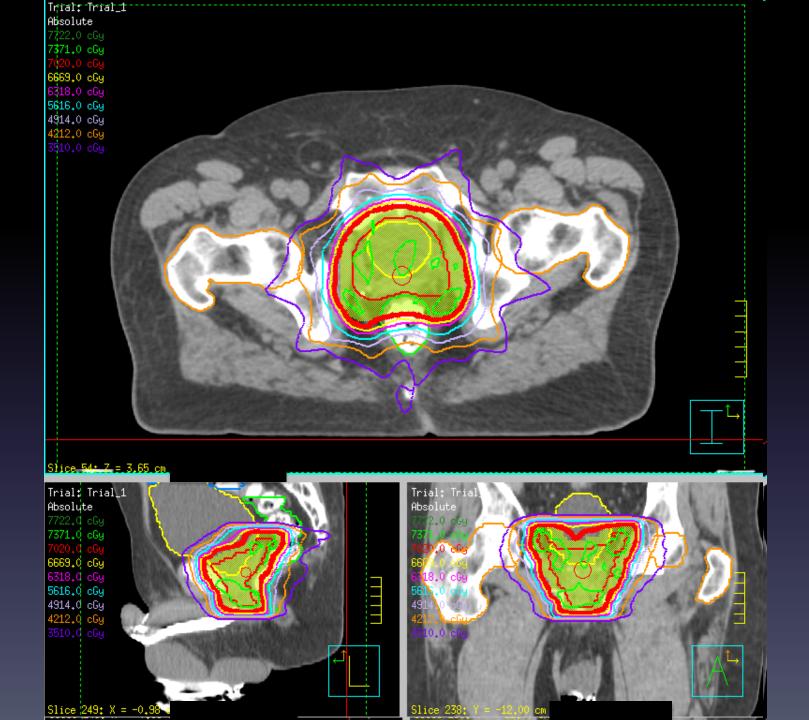
Oakland University William Beaumont School of Medicine

Disclosures

None

Radiotherapy After Prostatectomy What do we treat?

- No tumor
- No target organ
- Standardization of target volume through multiple prospective trials.
 - Inferior: 0.5 1.0 cm inferior to VUA
 - Lateral: obturator internus m.
 - Anterior: pubic symphysis
 - Posterior: rectum
 - Superior: seminal vesicle remnant/2-3 cm superior to top of pubic symphysis



Background: How did we get to where we are now?

Adjuvant or Salvage Therapy? (Undetectable Post-Op PSA Nadir)

- Prospective randomized trials evaluating adjuvant radiotherapy following prostatectomy with high-risk pathologic features identified:
 - **SWOG** Thompson et al. *J Urol* 2009;181:956-62.
 - EORTC Bolla et al. *Lancet* 2012:380:2018-27.
 - ARO (German) Wiegel et al. Eur Urol 2014;66:243-50.

SWOG 8794

Thompson et al. *J Urol* 2009.

- 1988-1997: 431 patients with \geq 1 of the following
 - Extracapsular extension
 - Seminal vesicle invasion
 - Positive surgical margin
- Randomization: "Adjuvant" RT (60-64 Gy) vs. Observation
- Negative pelvic nodes (lymphadenectomy for all but low-risk disease patients)
- Undetectable post-op PSA NOT required
 - ~1/3 of patients had PSA \geq 0.2 ng/mL (i.e. were "salvage" cases)

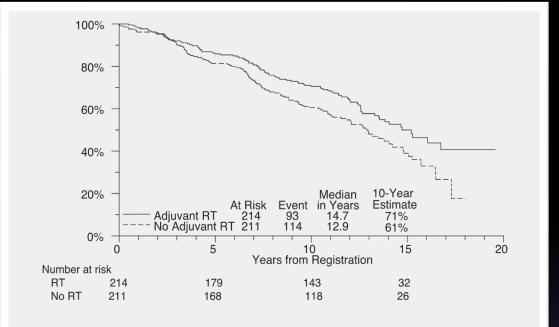
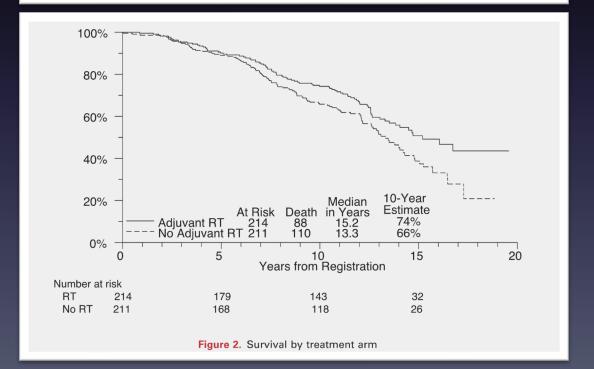
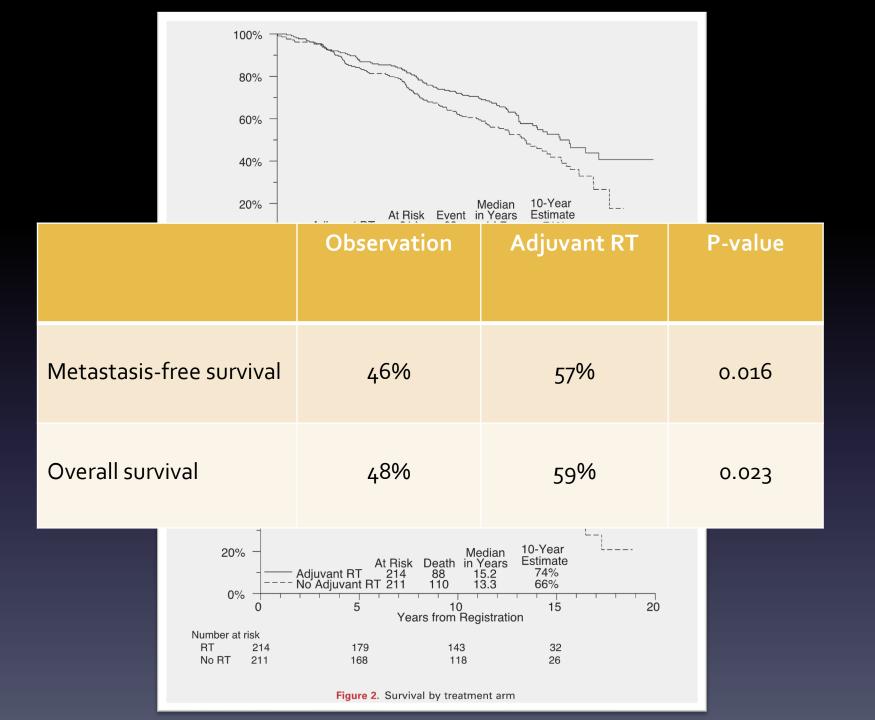


Figure 1. Metastasis-free survival by treatment arm

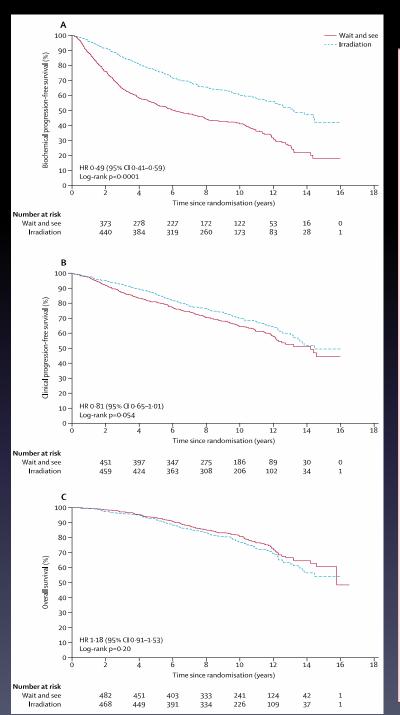




EORTC 22911

Bolla et al. *Lancet* 2012.

- 1992-2001: 1005 patients
 - Node-negative and \geq 1 of the following:
 - + ECE
 - + SVI
 - + margin
- Randomization: Immediate post-op RT (60 Gy) vs. Observation
 - 113 patients in observation arm eventually received salvage RT for relapse
- 10.7% of patients had detectable (> 0.2 ng/mL) PSA
- Median follow-up: 10.6 years



	Wait and see (n=503)	Irradiation (n=502)	Total (n=1005)
Biochemical or clinical progression or death	311 (61.8%)	198 (39·4%)	509 (50.6%)
Treated for prostate cancer, without biochemical progression	10 (2.0%)	7 (1.4%)	17 (1.7%)
Biochemical progression only	238 (47·3%)	105 (20.9%)	343 (34·1%)
Biochemical progression and locoregional failure	20 (4.0%)	11 (2.2%)	31 (3·1%)
Biochemical progression and distant failure	2 (0.4%)	1 (0.2%)	3 (0.3%)
Locoregional failure	2 (0.4%)	3 (0.6%)	5 (0.5%)
Distant failure	3 (0.6%)	5 (1.0%)	8 (0.8%)
Death without biochemical or clinical progression	36 (7·2%)	66 (13·1%)	102 (10·1%)
Clinical progression or death	101 (36 0%)	157 (31.3%)	338 (33.6%)
Locoregional failure	83 (16·5%)	35 (7.0%)	118 (11.7%)
Distant fallure	36 (7 2%)	3 0 (7·2%)	72 (7·2%)
Death without clinical progression	62 (12·3%)	86 (17·1%)	148 (14·7%)
Death	115 (22.9%)	130 (25.9%)	245 (24·4%)
Prostate cancer	34 (6.8%)	25 (5.0%)	59 (5.9%)
Cardiovascular disease	27 (5·4%)	33 (6.6%)	60 (6.0%)
Other cancer	29 (5.8%)	33 (6.6%)	62 (6.2%)
Alzheimer's disease	2 (0.4%)	3 (0.6%)	5 (0.5%)
General deterioration/ageing/sudden death at home	3 (0.6%)	1 (0.2%)	4 (0.4%)
Anaemia	1 (0.2%)	1 (0.2%)	2 (0·2%)
Renal insufficiency	0	2 (0.4%)	2 (0·2%)
COPD/embolism/pulmonary failure	6 (1.2%)	5 (1.0%)	11 (1·1%)
Infection not further specified	3 (0.6%)	10 (2.0%)	13 (1.3%)
Complication of surgery during follow-up	0	1 (0.2%)	1 (0.1%)
Accident/suicide	0	3 (0.6%)	3 (0.3%)
Unspecified, not prostate cancer	3 (0.6%)	3 (0.6%)	6 (0.6%)
Unknown	7 (1.4%)	10 (2.0%)	17 (1.7%)
Second cancer	69 (13.7%)	68 (13.5%)	137 (13.6%)

Data are number (%). Biochemical progression specifically refers to protocol-defined biochemical progression (defined in Methods section). COPD=chronic obstructive pulmonary disease.

Table 2: Events at long-term follow-up

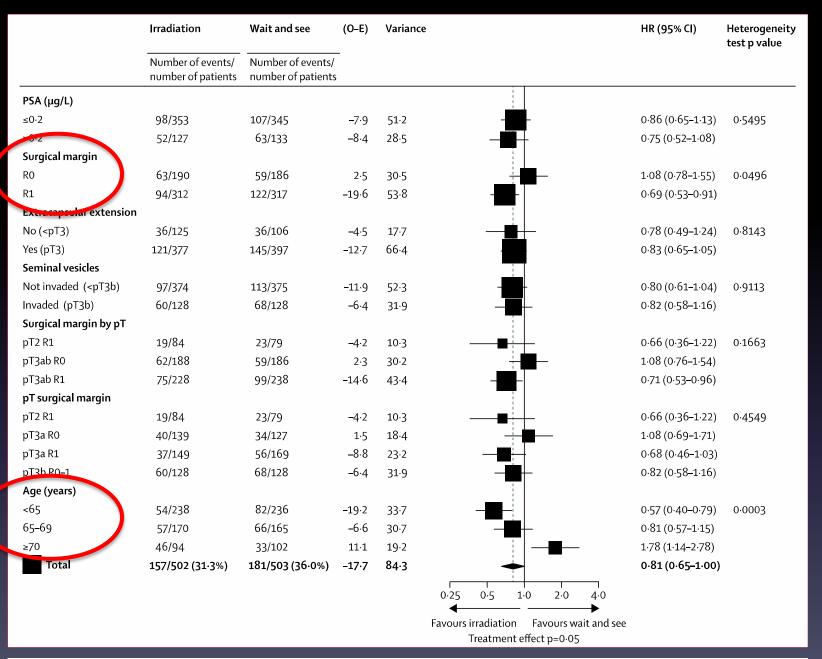
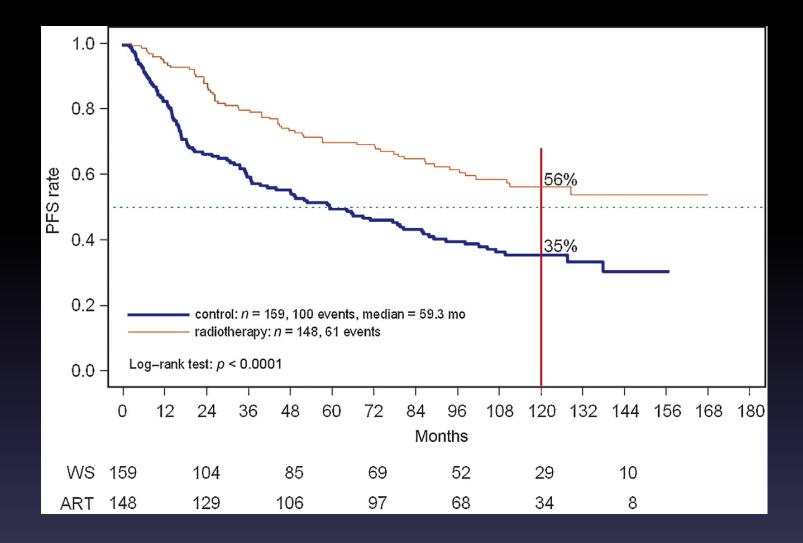


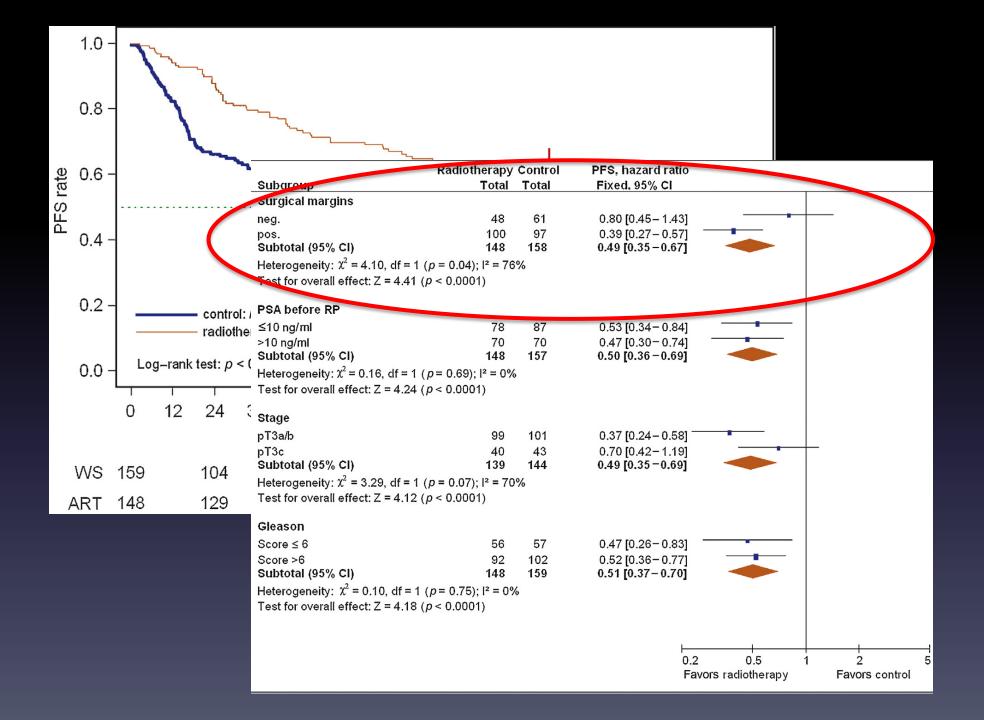
Figure 3: Effects of baseline factors on clinical progression-free survival O=observed. E=expected. HR=hazard ratio. PSA=prostate-specific antigen.

ARO 96-02/AUO AP 09/95

Wiegel et al. Eur Urol. 2014.

- 1997-2004: 307 patients with undetectable PSA post-op randomized to:
 - adjuvant RT (6o Gy)
 - "wait-and-see" approach
- 34 patients in RT arm refused treatment
- 5 patients in "wait-and-see" arm got RT



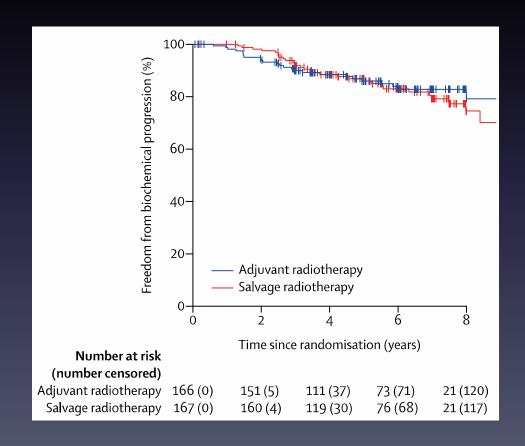


Take-Home Messages

- Adjuvant RT for men with pT₃ disease and/or positive margins reduces biochemical relapse rates.
 - Younger patients (< age 70) and those with positive surgical margins most likely to benefit
- Questions remain regarding clinical relapse/survival benefit.
- Left unclear what advantages adjuvant RT holds over early salvage.
 - Can potentially spare significant proportion of patients unnecessary treatment
 - Published while awaiting results of prospective studies evaluating this question
 - RAVES, GETUG-17, RADICALS

TROG o8-o3 (RAVES)

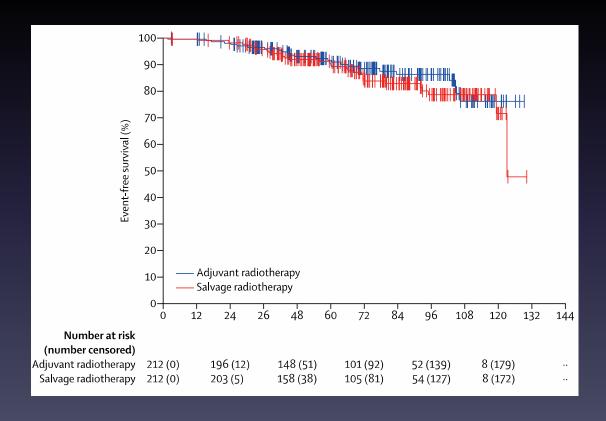
 Randomized 333 patients s/p prostatectomy with undetectable PSA to immediate (adjuvant) RT or "early salvage" – started once PSA was > 0.2.



• 5-yr FFP 86% vs. 85% with reduced Gr 2 GU toxicity 54% vs. 70% for salvage arm

GETUG-AFU 17

• 424 patients randomized to adjuvant vs. early salvage RT post RP with high risk features



- 5-yr EFS 92% vs. 90%
- 54% of patients on salvage arm eventually required RT

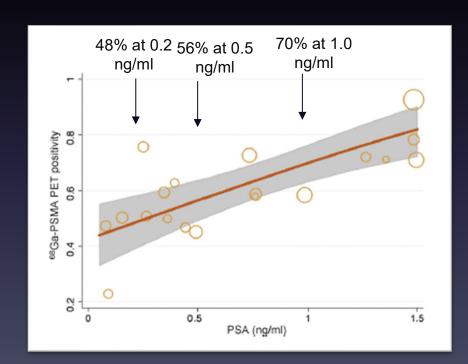
But things change...

Approach to Workup and Treatment Considerations

PSMA PET

Very sensitive clinical detection for patients with biochemical failure^{1,2}

Nearly 80% detection of disease in LN's < 8 mm³



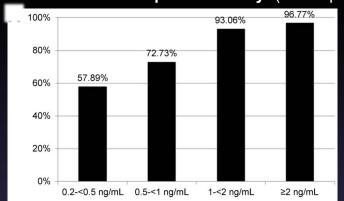
PSA <1ng/ml PSA 1-<2ng/ml PSA 2-<3ng/ml PSA >3ng/ml

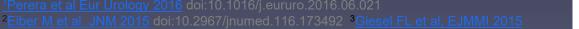
Fig. 1 Detection rate of [11C]Choline-PET/CT plotted against the PSA-value for recurrent prostate cancer

Detection rate after prostatectomy² (n=248 pts)

73%

62%





100

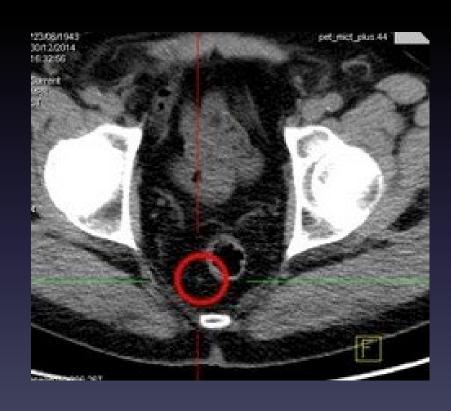
rate [%]

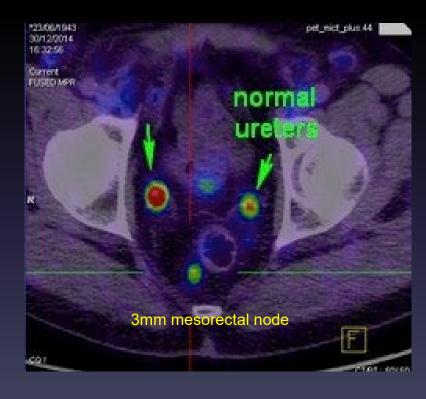
Detection

Pelvic Nodal Recurrence Post-Prostatectomy Undetectable nadir --> PSA rise to 0.23



Peri-Rectal Nodal Recurrence PSA = 0.4 ng/mL

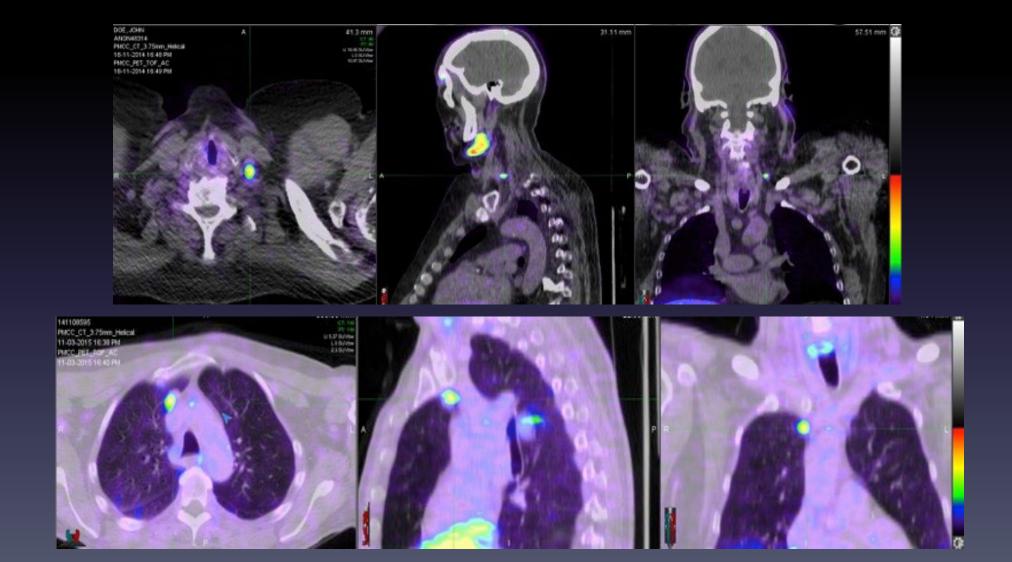




Local Failure Post-op PSA = 0.31 after undetectable nadir



Unusual Recurrence Patterns



Approach to Salvage Radiotherapy

Local Therapy

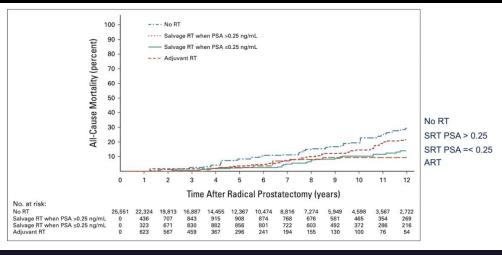
- RT to prostate bed
- Better early than late

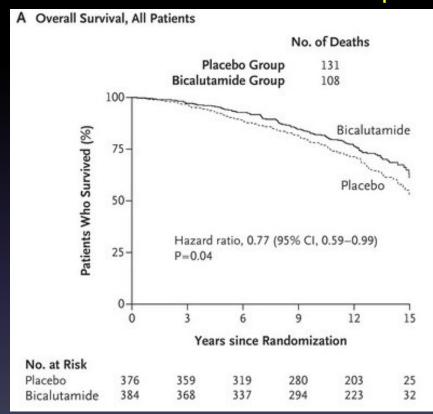
Systemic Therapy

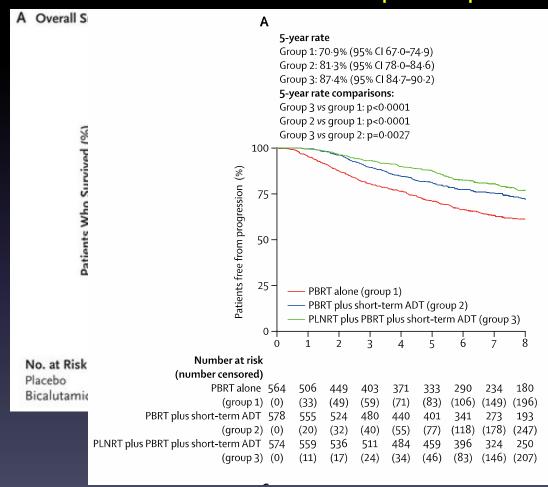
- Endocrine therapy
 - LHRH agonist/antagonist (short- vs. long-term)
 - Abiraterone
 - Anti-androgen

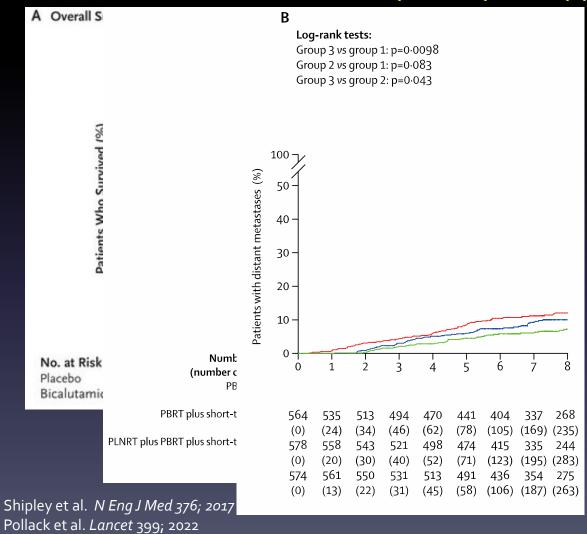
Treat pelvic nodes?

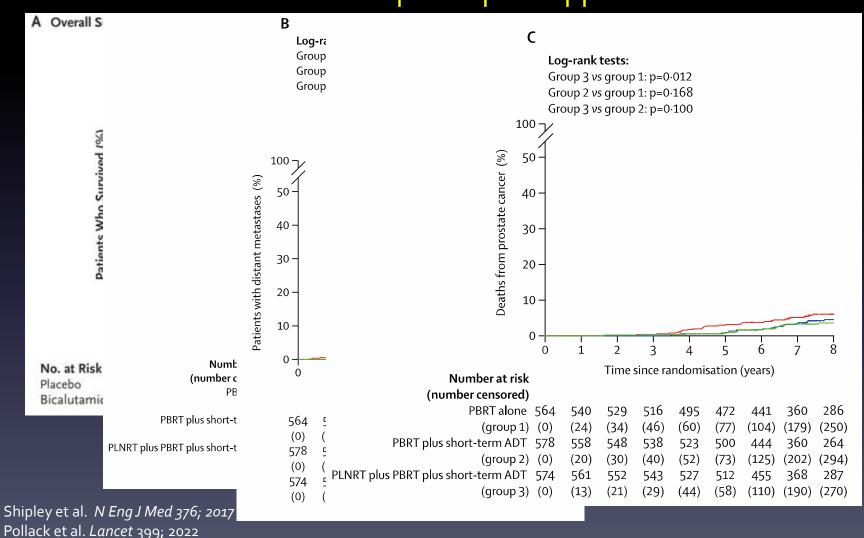
- Local disease stage
- Gleason score
- Secondary pathologic factors (LVSI, PNI)
- PLND performed/extent?
- PSA velocity/doubling time











What about patients with persistently elevated (> 0.1-0.2 ng/mL) post-prostatectomy?

- May occur in up to 10-15% of prostatectomy cases
- More likely to occur with:
 - Higher pre-op PSA values
 - Older patients
 - Pre-op Gleason (≥ 8)
 - Advanced disease stage (≥T3a)
 - + margins or +LN
- Could be representative of variable clinical scenarios
 - Trace amounts of benign prostate tissue left in bed (unprovable)
 - PSA production from locally persistent disease
 - Correlate with path findings (EPE, margin status)
 - Remaining regional or distant metastatic disease
 - Pre-op staging workup?
 - Extent of nodal surgery?

Little prospective data on this group of patients...what do retrospective data suggest?

- Preisser et al. Eur Urol 76(1); 2019.
 - ~1000 patients; ~50% received salvage RT
 - RT administration associated with improved OS and CSS (~50% relative reduction)
- Stish et al. *J Clin Oncol* 34(32); 2016.
 - > 1100 patients treated between 1987-2013 with salvage RT
 - Nearly 2/3 of patients failed biochemically
 - 10-yr rate of DM ~20%
 - Reduced by RT dose > 68 Gy and administration of ADT
 - Each pre-RT PSA doubling increased DM risk by 32% (TREAT EARLY!)

 Table 1
 Retrospective studies on patients treated with salvage radiotherapy for a persistent detectable postoperative PSA level

Studies with pN0 patients	Median follow-up (years)	% patients with detectable PSA	Population characteristics	Salvage RT (% patients, RT dose, concurrent HT	Pre-RT PSA (ng/ml)	Median time from PR to RT	Survival results	Factors predicting for better survival on mul- tivariable analysis
Stish [10]	8.9	1106	pT3a-pT4: 43.1% Positive margins: 48.6% Gleason≥8: 16.2%	100% RT Dose < 66 Gy: 30% Dose > 68 Gy: 51.5% 83,7% without HT 6,5% with HT > 1 year	≤0.5 ng/ml (45%)	33.6 months	5-year BRFS: 50.1% 10-year BRFS: 45.7% 5-year MFS: 89.1% 10-year MFS: 80.1%	pT, Gleason score, pre- RT PSA Dose > 68 Gy, HT use
Preisser [3]	3.9	1025 (8.8%)	pT3a-b: 63.2% Positive margins: 42.9% Gleason≥8: 21.6%	100% RT Dose NR	NR	5.4 months	NR	pT, Gleason score, pre- RT PSA
Ploussard [6]	3.7	201 (100%)	pT3a-b: 54.2% Positive margins: 67.7% Gleason≥8: 14.9%	100% RT Dose NR 0% HT	0.48	7 months	BRFS: 42.8%	pT3b, Gleason score, post-RP PSA, pre-RT PSA > 1 ng/ml, surgi- cal margins, PSA velocity
Ploussard [12]	3.1	496 (5.1%) of 9735 patients	pT3a-4: 49.6% Positive margins: 54% Gleason≥8: 19.1%	40.4% RT 8.9% RT+HT 19,5% HT	0.1–6.1	NR	1-year BRFS: 34.3% 5-year BRFS: 21.5% 5-year OS: 94.7%	pT3b, Gleason score, post-RP PSA, pre-RT PSA > 1 ng/ml, surgi- cal margins, PSA velocity
Gandaglia [9]	9.2	496 (50%) of 982 patients	pT3a-4: 53.4% Positive margins: 50% Gleason≥8: 33.3%	251 (50.6%) RT 50.4 Gy pelvis, 68 Gy fossa 23% HT	NR	NR	10-year CSS: 88%	pT, Gleason score, pre-RT PSA, 10-year metastasis risk > 30%
Fossati [11]	8.0	224 (24%) of 925 patients	pT3a-4: 56% Positive margins: 44% Gleason≥8: 24%	100% RT 68 Gy fossa 30% HT	0.2	1.3 months	8-year MFS: 74% if high risk and 62% if very high risk	pT, Gleason score, pre- RT PSA
Barthowiak [13]	6.1	133 (24%) of 555 patients	pT3a-4: 43% Positive margins: 45% Gleason≥8: 26%	100% RT Median of 66.6 Gy 0% HT	0.56	10 months	5-year RFS: 49%	pT, pre-RT PSA > 0.5 ng/ml

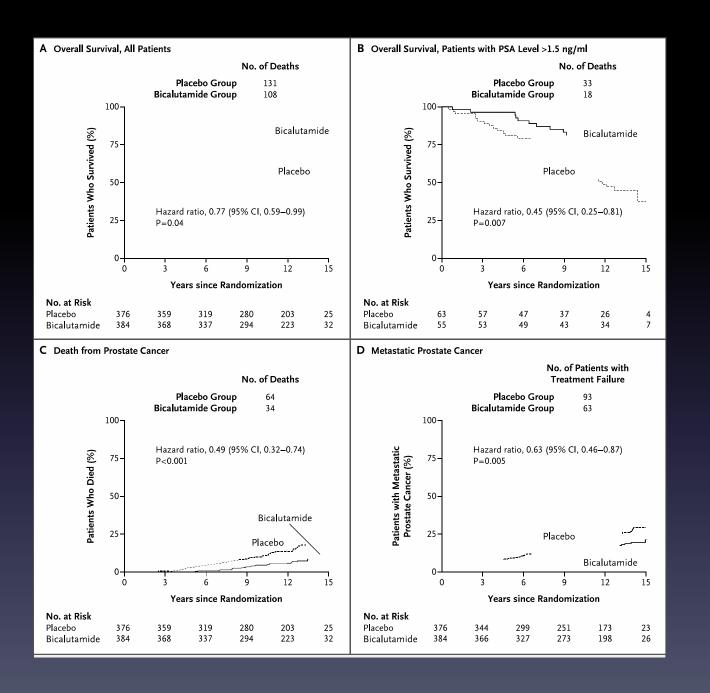
PSA: Prostate-specific antigen); pT: tumor pathological stage; NR: not reported; RP: radical prostatectomy; RT: radiotherapy; HT: hormonotherapy; BRFS: biochemical relapse-free survival; RFS: relapse-free survival; MFS: metastasis-free survival; CSS: cancer-specific survival; OS: overall survival

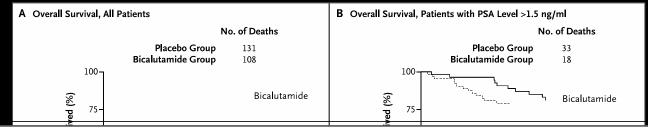
RTOG 9601

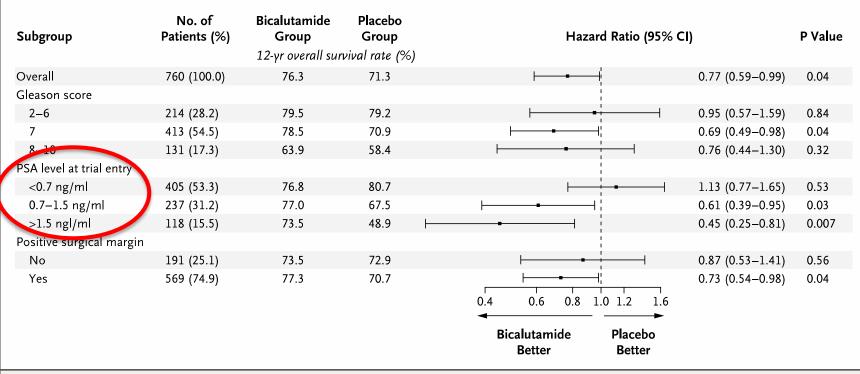
Shipley et al. *NEJM* 2016.

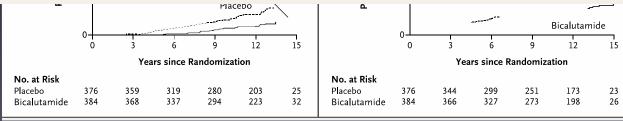
• 1998-2003: 760 patients randomized to post-op RT (64.8 Gy)+/- bicalutamide (150 mg/day) x 2 years

- Eligibility
 - pT₃ OR pT₂ with + margin AND
 - Post-op PSA o.2-4.o ng/mL.
 - Lymph node negative
- Median f/u: 13 years









A Overall Survival, All Patients	B Overall Survival, Patients with PSA Level >1.5 ng/ml				
No. of Deaths	No. of Deaths				
Placebo Group 131	Placebo Group 33				

Table	2. An	titumor Effic	acy with Res	pect to Ke	y Sec	onda	ary End	Points at	12 Yea	rs.
							_			

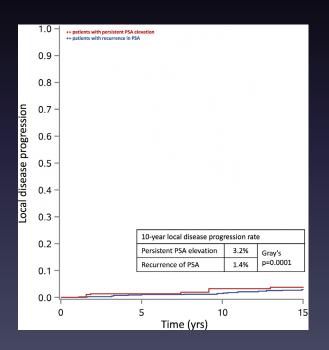
End Point and Subgroup	Bicalutam	Bicalutamide Group		o Group	Hazard Ratio (95% CI)	P Value
	Patients at Risk	Rate of End Point	Patients at Risk	Rate of End Point		
	no.	%	no.	%		
Metastatic prostate cancer						
All patients	384	14.5	376	23.0	0.63 (0.46–0.87)	0.005
Gleason score						
2–6	111	7.8	103	16.5	0.64 (0.30–1.36)	0.25
7	205	15.4	208	19.8	0.80 (0.52–1.22)	0.31
8–10	67	21.2	64	44.7	0.35 (0.18–0.67)	0.001
PSA level at trial entry						
<0.7 ng/ml	210	13.4	195	17.1	0.76 (0.47–1.22)	0.26
0.7–1.5 ng/ml	119	17.4	118	28.4	0.67 (0.40–1.12)	0.13
>1.5 ng/ml	55	13.1	63	31.1	0.36 (0.15–0.84)	0.01
Positive surgical margin						
No	96	22.9	95	31.1	0.79 (0.47–1.32)	0.38
Yes	288	11.8	281	20.3	0.56 (0.38–0.84)	0.005
Death from prostate cancer*	384	5.8	376	13.4	0.49 (0.32–0.74)	< 0.001
Death from other causes	384	17.9	376	15.3	1.10 (0.79–1.53)	0.58

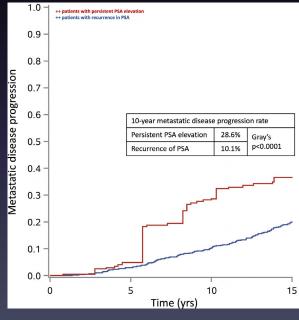
No. at Risk							No. at Risk						
Placebo	376	359	319	280	203	25	Placebo	376	344	299	251	173	23
Bicalutamide	384	368	337	294	223	32	Bicalutamide	384	366	327	273	198	26
Bicalutamide	384	368	33/	294	223	32	Bicalutamide	384	366	327	2/3	198	

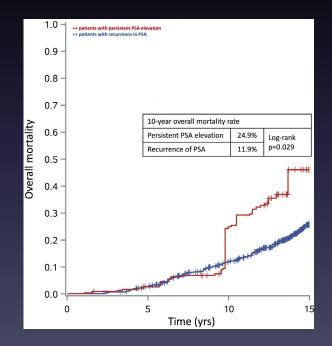
RTOG 9601 Update

Sood et al. *Urol Oncol* 38; 2020

- Post-hoc analysis of patients with persistently PSA (nadir \geq 0.4) vs. recurrent PSA
 - 670 recurrent vs. 90 persistent
 - ~50% of patients received bicalutamide in both recurrent and persistent subgroups

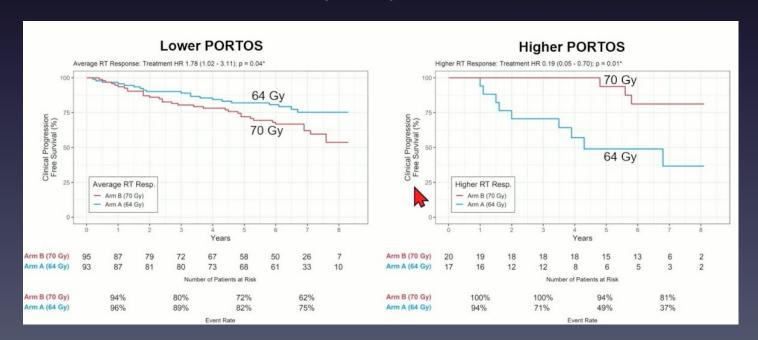






A word about genomic classification...

- Decipher analysis of 352 patients from RTOG 9601
 - Not statistically significant, but suggestive of greater benefit of endocrine Rx for patients with higher Decipher scores
 - 5% vs. 16% benefit for DM
 - 4.5% vs. 12% for PCSM
 - 2.4% vs. 9% for OS
- PORTOS score may be predictive for RT dose escalation
 - Dal Pra data secondary analysis of SAKK 09/10 trial



Additional Considerations

• Treatment intensification?

- LN-directed therapy
- Oligometastatic disease
 - MDT/SBRT (COMET, ORIOLE, NRG GU 011)
- Abi/pred
 - STAMPEDE (LN+)
- Anti-androgen
 - Enzalutamide, Apalutamide, Darolutamide
 - STEEL, NRG GU 006
- Chemotherapy
 - Probably not...
- Immunotherapy
- Radiopharmaceutical

To Conclude...

Most Common Post-op Patient Presentations/Approaches

- High-risk path (EPE/+ margin) with undetectable PSA
 - Surveillance
 - Can consider adjuvant for very HR (+SV, +LN, Gleason 9-10)
- Rising PSA after initially undetectable
 - Restaging PSMA PET negative
 - Salvage RT to prostate bed +/- pelvic nodes +/- ADT
 - PET + prostate bed
 - Salvage RT to prostate bed (consider focal boost to nodule) +/- LN +/- ADT
 - PET +LN
 - Salvage RT to prostate bed + LN + ADT + Abi/pred (STAMPEDE)
 - Escalate RT dose to PET + disease
 - PET + solitary or oligo-metastatic distant disease
 - SBRT to metastatic disease
 - RT to prostate bed/nodes???
 - Systemic therapy?

Most Common Post-op Patient Approaches (cont'd)

- Persistent PSA elevation post-op
 - Restage with PSMA PET if not done pre-op
 - Early treatment initiation with combination local + systemic therapy
 - Consideration and future study of intensification of systemic therapy due to high risk of metastatic progression

ThankYou



Table Discussions

Help us understand relationships with Radiation Oncology across the state

Questions to Consider -

1. What are the most **common barriers** you encounter when considering radiation therapy for a patient following prostatectomy?

2. What do you wish you knew more about to improve your counseling or decision making for patients with a detectable PSA after prostatectomy?

3. What **effective strategies** have you used when co-managing patients with a radiation oncologist?

4. What can MUSIC do to improve the care of patients requiring post-prostatectomy radiation?



Key Takeaways

Tudor Borza, MD



Clinical Stage and Recurrence / Metastases: Need for Improved Documentation

Brian Lane, MD, PhD



Nonprofit corporations and independent licensees of the Blue Cross and Blue Shield Association

Why is Clinical Staging Important?





Without accurate N and M staging, we cannot draw conclusions on oncologic outcomes



MUSIC's contributions are significant, especially within AS

>8000 patients from clinically diverse settings

>2500 T1RM on AS



Who progresses after AS or treatment? And when?

What is the outcome of tiny lung nodules?

Are mets from RCC or other cancer?



Many questions to be answered to improve the quality of care delivered to our patients!





- 57 yo with 5.5 cm right renal mass and 12 mm lung nodule (ddx: primary lung Ca vs. met)
- So cT1bN0M1 (if RCC met) vs.
 cT1bN0M0 (if not met)
- What would you say?





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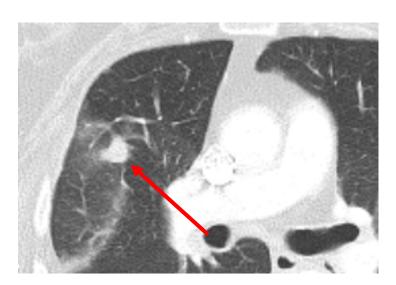


- 57 yo with 5.5 cm right renal mass and 12 mm lung nodule (ddx: primary lung Ca vs. met)
- So cT1bN0M1 (if RCC met) vs.
 cT1bN0M0 (if not met)
- Code as cM1



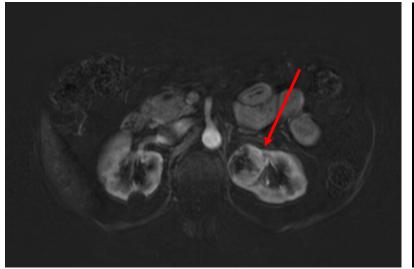
- Path: squamous cell carcinoma (not from kidney)
- Plan: surgical resection of both likely localized cancers: lung cancer and cT1bN0M0 kidney cancer
- Path from MIRNx: ccRCC, 5cm, pT1bN0M0, grade 3



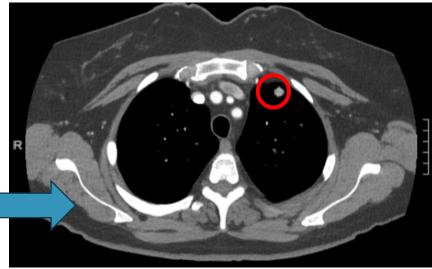




- 76yo woman with 4.5cm left renal mass on imaging done for CKD
- Biopsy done ccRCC grade 3 GFR: 37
- Pre-op chest imaging multiple non-calcified scattered nodules bilaterally, largest 6mm; no comparison study available
- Partial nephrectomy ccRCC grade 3, pT1b with negative margins
- 3-month interval CT chest showed enlarging nodules, now up to 15.1cm; What is the M stage?

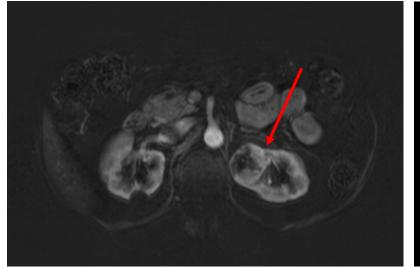


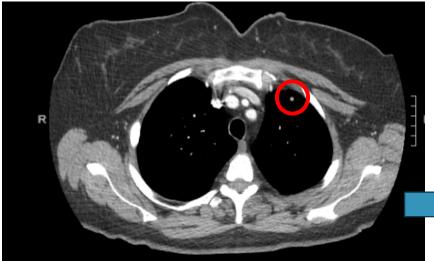


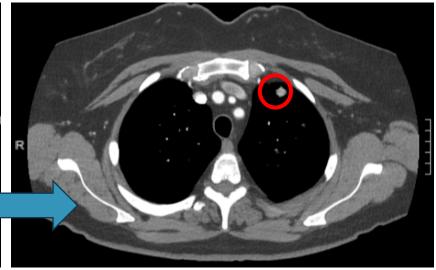




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- Partial nephrectomy ccRCC grade 3, pT1b with negative margins
- 3-month interval CT chest showed enlarging nodules, now up to 15.1cm; biopsy arranged: path:







Primer on Documentation of cTNM Stage



for Clinicians

T stage hasn't changed:

cT1a: localized, ≤4 cm

cT1b: localized, 4.1 - 7 cm

cT2a: localized, 7.1 - 10 cm

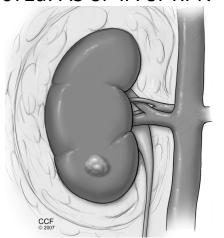
cT2b: localized, >10 cm

cT3a: radiographic suspicion of fat or venous invasion

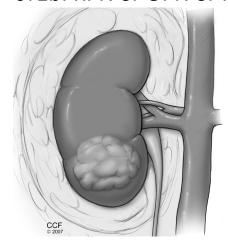
cT3b/c: rad. suspicion of IVC invasion

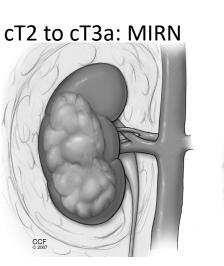
cT4: rad. suspicion of direct invasion into another organ (adrenal, liver, etc.)

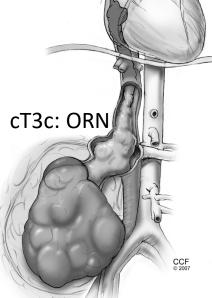




cT1b: RPN or OPN or MIRN or AS







Clinical vs. Pathologic N and M Staging -



Clinical N and M Staging

- Use of cN1 and/or cM1 means that the index cancer is suspected to have spread
 - Lesions may or may not be related to RCC (they may be from a different cancer or not be cancer). Over time, some may be determined to be M0
- cNx and/or cMx relates to lesions that are 'indeterminate' for metastasis
 - RCC met, other cancer, or not cancer'

Pathologic N and M staging

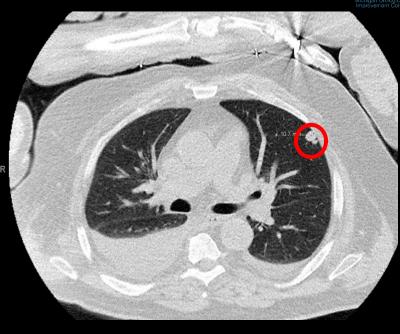
- Frequent use of pNx and pMx is correct
- They are correctly used when there
 has been no pathologic tissue (biopsy
 / surgery) for LN or no distant mets
 (most patients)

Clinical N and M Staging





Mx: Indeterminate lung nodule M1: lung metastasis





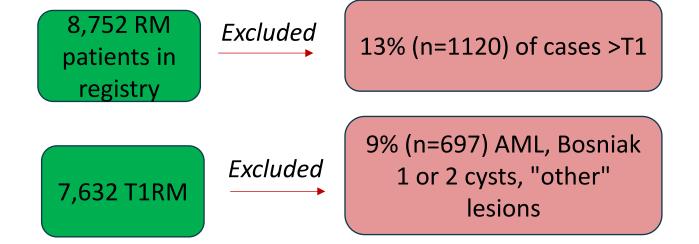
Nx:
Indeterminate
lymph node
(7mm)

N1: Lymph node metastases

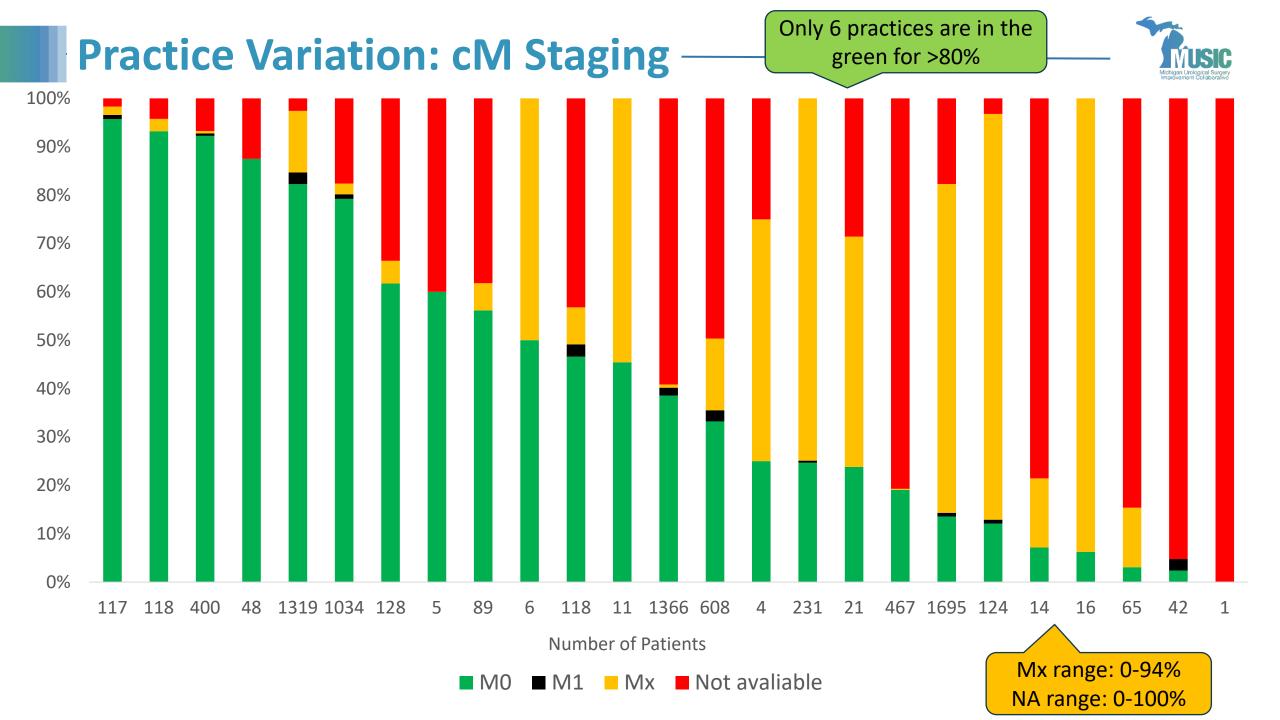


The Problem





23% (n=1577) NX 25% (n=1753) Missing = Staging not available 25% (n=1724) MX 26% (n=1817) Missing = Staging not available

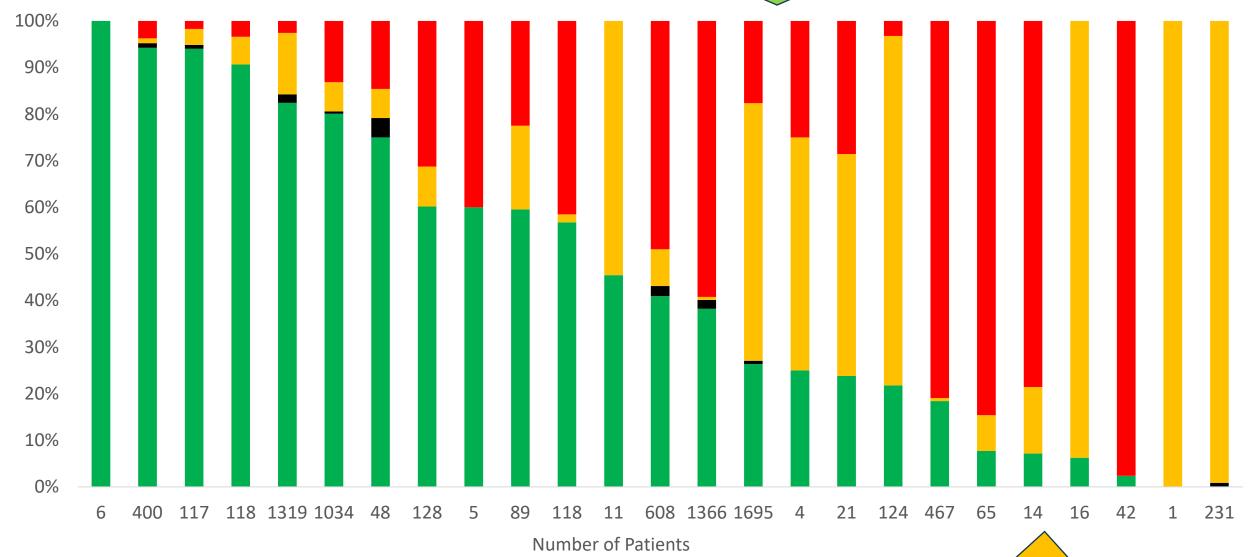




Practice Variation: cN Staging —

Only 6 practices are in the green for >80%





Nx ■ Not avaliable

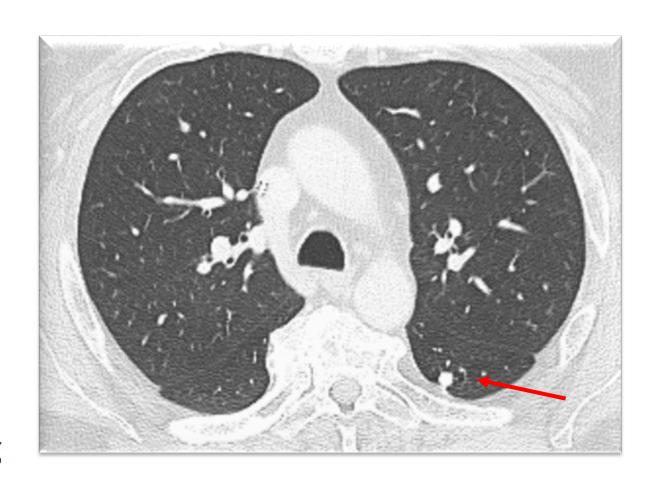
Nx range: 0-99%

NA range: 0-98%





- Clear cell RCC, 3.5 cm, neg margins s/p left RALPN (3/2010) Stage cT1aN0M0 and pT1aNxMx
- F/u with Urology ended after 5 yrs post-op
- Referred back for "Several nonspecific noncalcified nodules within both lung fields not clearly visualized on prior examinations due to difference in technique"
- Shown is a 5 mm indeterminate lung nodule in 2/2023
- What is the M stage?



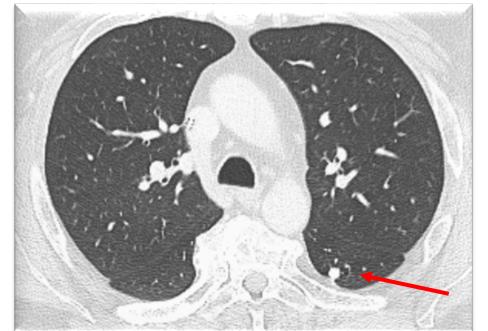


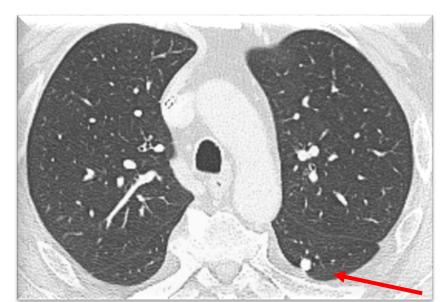
Case Study



- Clear cell RCC, 3.5 cm, neg margins s/p left RALPN (3/2010) Stage cT1aN0M0 and pT1aNxMx
- F/u with Urology ended after 5 yrs post-op
- Referred back for "Several nonspecific noncalcified nodules within both lung fields not clearly visualized on prior examinations due to difference in technique"
- Shown is a 5 mm indeterminate lung nodule in 2/2023, and stable as of 9/2024
- This is cMx

... and now clinically determined to be cM0 with subsequent f/u







TNM Staging Documentation in Prostate

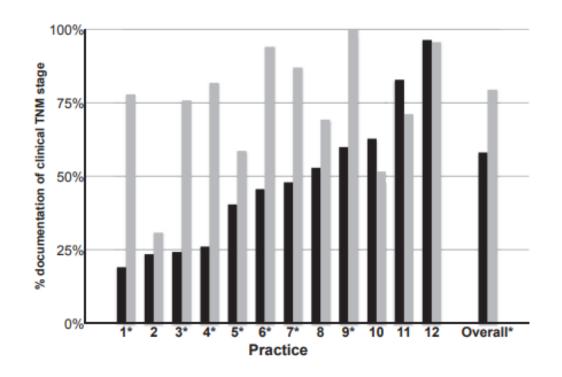


Health Services Research

Improvement in Clinical TNM Staging Documentation Within a Prostate Cancer Quality Improvement Collaborative

Christopher P. Filson, Brooke Boer, Jon Curry, Susan Linsell, Zaojun Ye, James E. Montie, and David C. Miller

- At baseline, 58% of patients had clinical TNM staging in the medical record, ranging from 19% to 96% across 12 practices
- After the intervention, documentation improved to 79% of patients overall



Staging Documentation in KIDNEY



- At baseline, ~70% of patients had clinical staging in the medical record, ranging from 0% to 96% across 25 practices
 - However, ~26% of these cases are incorrectly documented as indeterminate
- We are hopeful to have similar success as we had with PCa (58% to 79%)





KIDNEY Cancer Visit Template

Patient Name:
MRN:
Date of Visit

o M1

o MX

	ological Surgery et Collaborative						D	ate of Visit
/ Pa	tient Vi	sit – Ren	nal Mass					
•		bidities		•	Tumor	Com	nple	exity
	0	Conges	stive heart failure (CHF)		0	R		
	0		c kidney disease (CKD)				•	1
	0	Chronic	c obstructive pulmonary				•	2
			(COPD)				•	3
	0	Cerebr	ovascular disease		0	E		
	0	Periphe	eral vascular disease				•	1
		(PVD)					•	2
•	Labs						•	3
	0	CMP			0	N		
			Date				•	1
			ALT Value				•	2
			AST Value				•	3
			ALP Value		0	Α		Antonion
	0	CBC					:	Anterior
		•	Date				:	Posterior
	0	Urinaly	rsis		0	L	•	Х
		•	Date		0	L		1
		•	Proteinuria Value				:	2
	0	Creatin	ine					3
		•	Date		0	Hila	ar	3
		•	Value		0	11116		Yes
	0	eGFR						No
		•	Date		0	Tot	al s	core:
		•	Value					4
•		Size (cm	1)					5
•	Tumor							6
	0	Right					•	7
	0	Left					•	8
•		T Stage					•	9
	0	T1a T1b					•	10
	0	T2a					•	11
	0	T2b					•	12
	0	T3a		•	Initial (Clinic	al I	mpression
	0	T3b			0		nign	
	0	T3c			0		-	ious
	0	T4			0			rminate
•		N Stage	.	•				ssessment of volume
	0	NO	•		preser	vatio	n (F	PAVP): %
	0	N1		•	Pre-op	erati	ive a	assessment:
	0	NX			0			rd PN
•		M Stage	e		0	Tec	hni	cally challenging PN
	0	мо			0	No	t an	nenable to PN

• Treatment Recommendation

o Surveillance



KIDNEY Cancer Visit Template

Patient Name:	
MRN:	
Date of Visit	

- Timing of next imaging
- o Ablation
- o Partial Nephrectomy
- o Radical Nephrectomy
- o No Treatment Needed
- Chest Imaging
 - Not ordered
 - o CT thorax
 - o Chest X-Ray

Key Takeaways





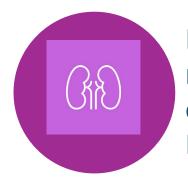
Use cNx and cMx only when you really mean it (indeterminate lesions or unclear if metastasis are from RCC)



Abstractors should record staging as best as they are able; add text notes for cases where they are unclear



Clinicians should use cN0 and cM0



Properly identified cases = reporting of ACCURATE oncologic outcomes across MUSIC-KIDNEY

We have a large (>8000) series of patients from clinically diverse settings; We are providing important information for patients and providers about the safety of AS and Rx

Collecting additional, accurate data is essential!



Physician Wellness: Harvesting the High Hanging Fruit

David Canes, MD



Nonprofit corporations and independent licensees of the Blue Cross and Blue Shield Association

Physician Wellness:
Harvesting the
High-Hanging Fruit

David Canes, MD October 18, 2024





Disclosures:

Founder, WellPrept

Patient care is the most fulfilling thing.



Wellness = thriving (not just surviving)

Manage stress of being urologist

Work - life balance

+ Relationships, pts & colleagues

Sense of pers & prof fulfillment



Be the doctor you set out to be.



Wellness = burnout shield/antidote



Review > PLoS One. 2016 Jul 8;11(7):e0159015. doi: 10.1371/journal.pone.0159015. eCollection 2016.

Healthcare Staff Wellbeing, Burnout, and Patient Safety: A Systematic Review

Louise H Hall ^{1 2}, Judith Johnson ^{1 2}, Ian Watt ³, Anastasia Tsipa ^{1 4}, Daryl B O'Connor ¹

Affiliations + expand

PMID: 27391946 PMCID: PMC4938539 DOI: 10.1371/journal.pone.0159015

Abstract

Objective: To determine whether there is an association between healthcare professionals' wellbeing and burnout, with patient safety.

Design: Systematic research review.

Data sources: Psychlnfo (1806 to July 2015), Medline (1946 to July 2015), Embase (1947 to July 2015) and Scopus (1823 to July 2015) were searched, along with reference lists of eligible articles.

Eligibility criteria for selecting studies: Quantitative, empirical studies that included i) either a measure of wellbeing or burnout, and ii) patient safety, in healthcare staff populations.

Results: Forty-six studies were identified. Sixteen out of the 27 studies that measured wellbeing found a significant correlation between poor wellbeing and worse patient safety, with six additional studies finding an association with some but not all scales used, and one study finding a significant association but in the opposite direction to the majority of studies. Twenty-one out of the 30 studies that measured burnout found a significant association between burnout and patient safety, whilst a further four studies found an association between one or more (but not all) subscales of the burnout measures employed, and patient safety.

Conclusions: Poor wellbeing and moderate to high levels of burnout are associated, in the majority of studies reviewed, with poor patient safety outcomes such as medical errors, however the lack of prospective studies reduces the ability to determine causality. Further prospective studies, research in primary care, conducted within the UK, and a clearer definition of healthcare staff wellbeing are

High quality care

Burnout is associated with worse outcomes in MANY domains

Stanford model of professional fulfillment

2016



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1

Culture of wellness

2

Personal resilience

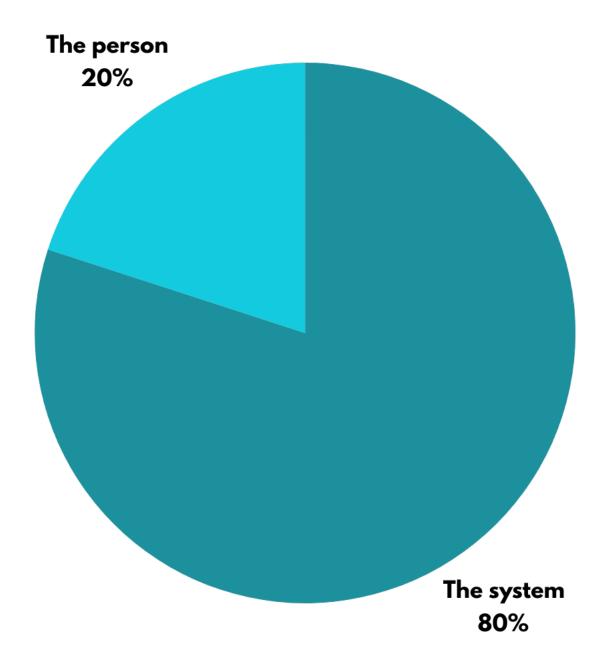
3

Efficiency of practice



2

Personal resilience





Dr. Tait Shanafelt

Nation's first ever Chief Wellness Officer

Ann Intern Med. 2002 Mar 5;136(5):358-67. doi: 10.7326/0003-4819-136-5-200203050-00008.

Burnout and self-reported patient care in an internal medicine residency program

Tait D Shanafelt 1, Katharine A Bradley, Joyce E Wipf, Anthony L Back

Affiliations + expand

PMID: 11874308 DOI: 10.7326/0003-4819-136-5-200203050-00008

Abstract

Background: Burnout is a syndrome of depersonalization, emotional exhaustion, and a sense of low personal accomplishment. Little is known about burnout in residents or its relationship to patient care.

Objective: To determine the prevalence of burnout in medical residents and explore its relationship to self-reported patient care practices.

Design: Cross-sectional study using an anonymous, mailed survey.

Setting: University-based residency program in Seattle, Washington.

Participants: 115 internal medicine residents.

Measurements: Burnout was measured by using the Maslach Burnout Inventory and was defined as scores in the high range for medical professionals on the depersonalization or emotional exhaustion subscales. Five questions developed for this study assessed self-reported patient care practices that suggested suboptimal care (for example, "I did not fully discuss treatment options or answer a patient's questions" or "I made...errors that were not due to a lack of knowledge or inexperience"). Depression and at-risk alcohol use were assessed by using validated screening questionnaires.

Results: Of 115 (76%) responding residents, 87 (76%) met the criteria for burnout. Compared with non-burned-out residents, burned-out residents were significantly more likely to self-report providing at least one type of suboptimal patient care at least monthly (53% vs. 21%; P = 0.004). In multivariate analyses, burnout--but not sex, depression, or at-risk alcohol use--was strongly associated with self-report of one or more suboptimal patient care practices at least monthly (odds ratio, 8.3 [95% CI, 2.6 to 26.5]). When each domain of burnout was evaluated separately, only a high score for depersonalization was associated with self-reported suboptimal patient care practices (in a dose-response relationship).

Conclusion: Burnout was common among resident physicians and was associated with self-reported suboptimal patient care practices.

First quantitative association between burnout and poor care

2002



It's about organizational change, systems change, and culture change, not tips and tricks for personal resilience.

Tait Shanafelt, MD



66

When organizational wellness efforts are either lip service, or manifest as yoga and granola and learn how to practice mindfulness... they will fall flat.

Tait Shanafelt, MD



66

A bad system will beat good people every single time.

Tait Shanafelt, MD



Personal Resilience

Healthy habits

Time for recovery

Wellbeing in the face of adversity

Safety net systems

Culture of Wellness

Effective leadership

Respect & inclusivity

Recognition

Regular measurement

Efficiency of practice



Unnecessary admin burden

Optimize workflows

Redesign of inefficient work

Streamline EHR and IT interfaces

Why is efficiency high-hanging fruit?



Humans get used to inefficiencies.





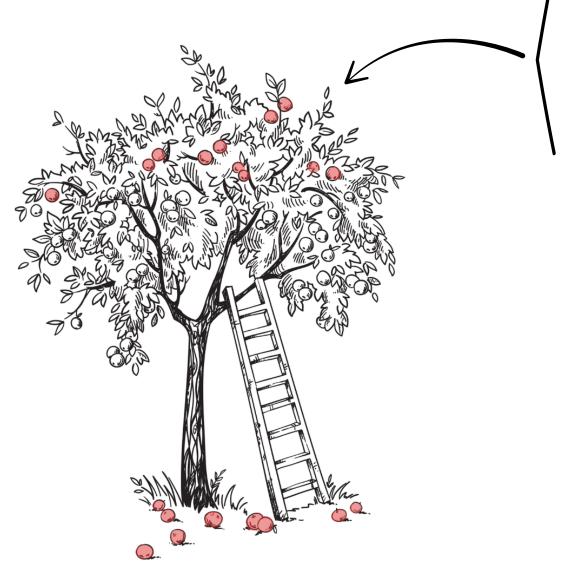
No training re: operational expertise

"Just the way things are"

ROI may not be instantaneous

Unclear metrics





Can we control what a patient knows and when they know it?

Can we control what a patient knows and when they know it?

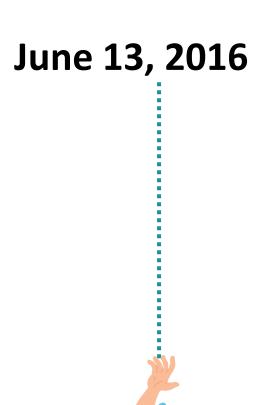




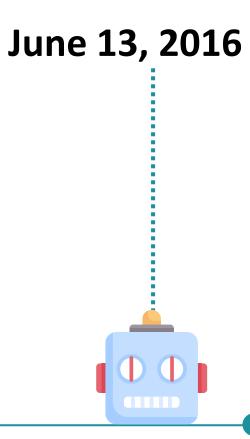


Why address patient pre-education?



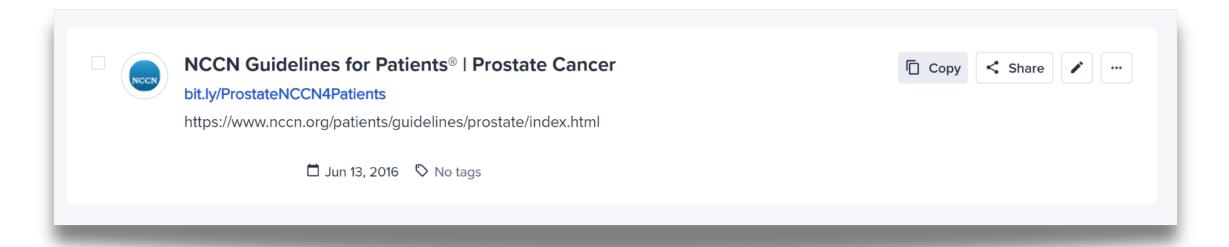


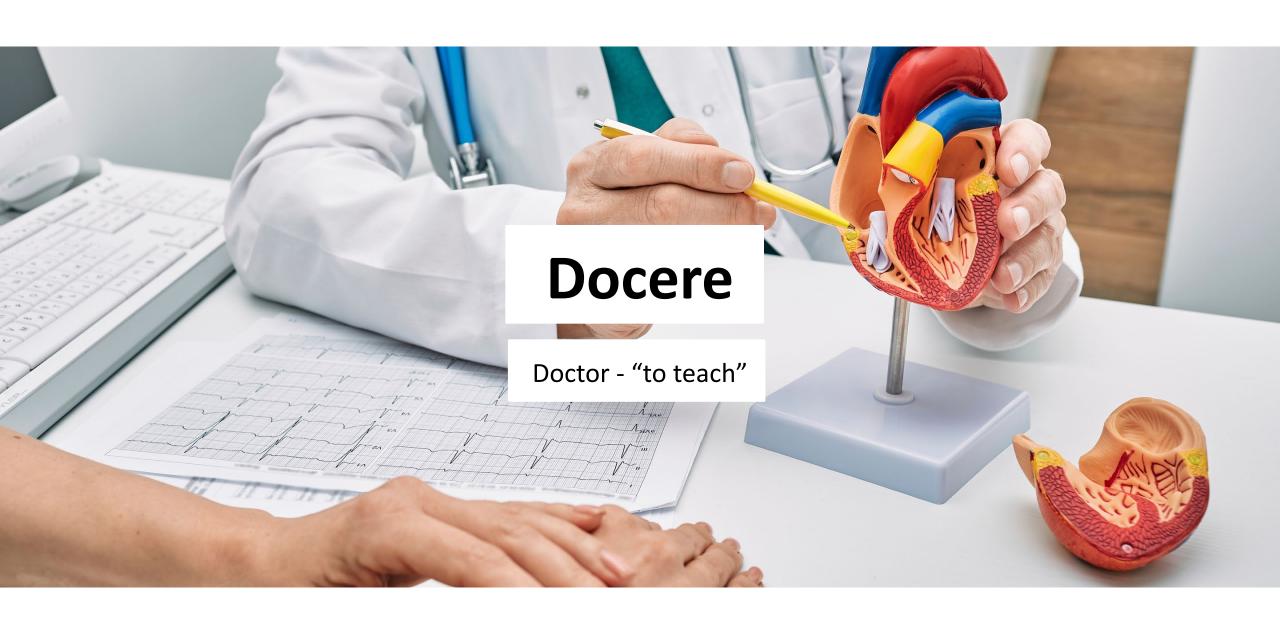
Born 1975 Today October 18, 2024



Born 1975 Today October 18, 2024







Would teachers have their students show up cold?













Does it work? (Arleeta's story)





Patient satisfaction n = 1750

96%

More likely to recommend

83%

Less anxious

97%

Better understanding

-66

"Now I don't have to search Google. What a huge relief! And this is way better than anything I would have found on my own."

-Actual WellPrept patient quote

— 66 ———

"This is so great. I wish all my doctors did this."

-Actual WellPrept patient quote

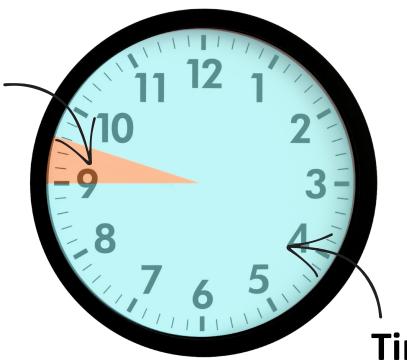


"I find it nice to see what is presented by the physician. Makes me more at ease to see him."

-Actual WellPrept patient quote

What do doctors think?

Time spent on basic spiels

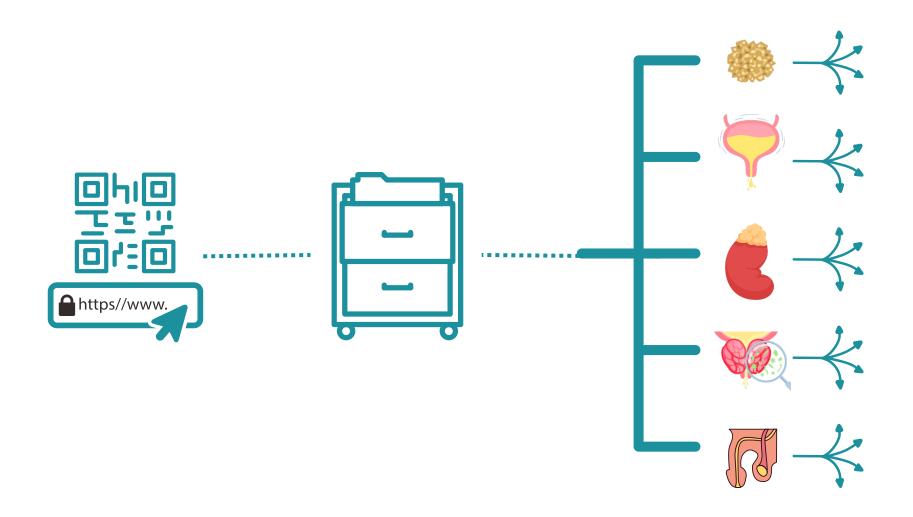


Time spent doing literally anything else

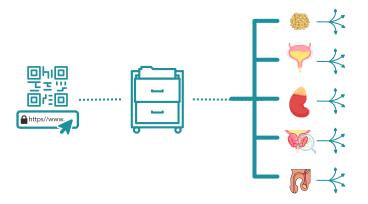
How does this actually work?



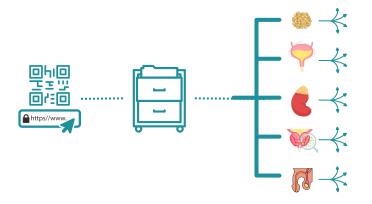
It starts with one link



Vetted
resources &
your favorite
things



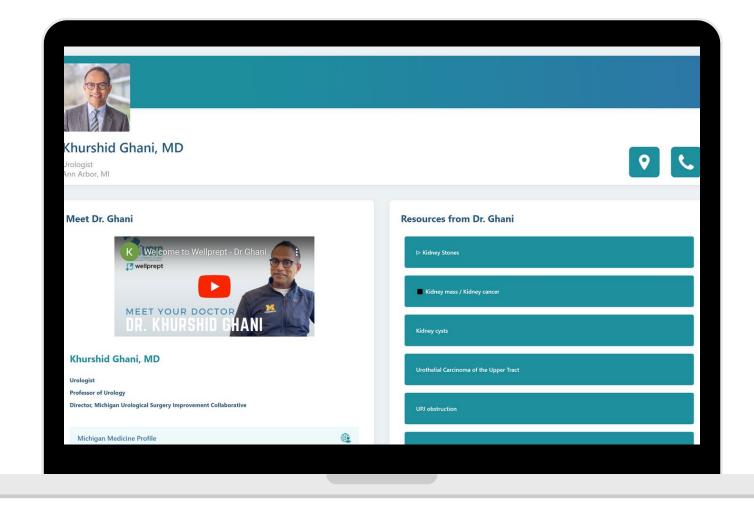
Conversations you and the patient actually want to have

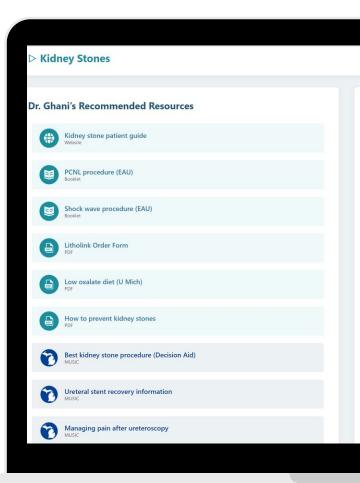


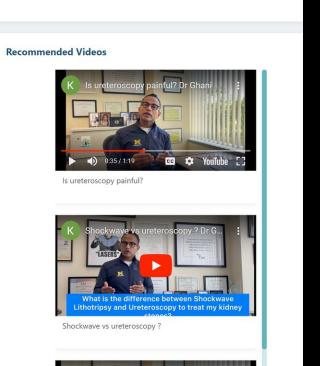
Better outcomes?
(more on how MUSIC can help later)



What does it look like to a patient?

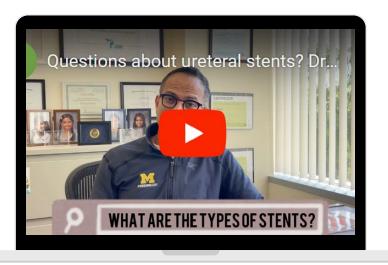








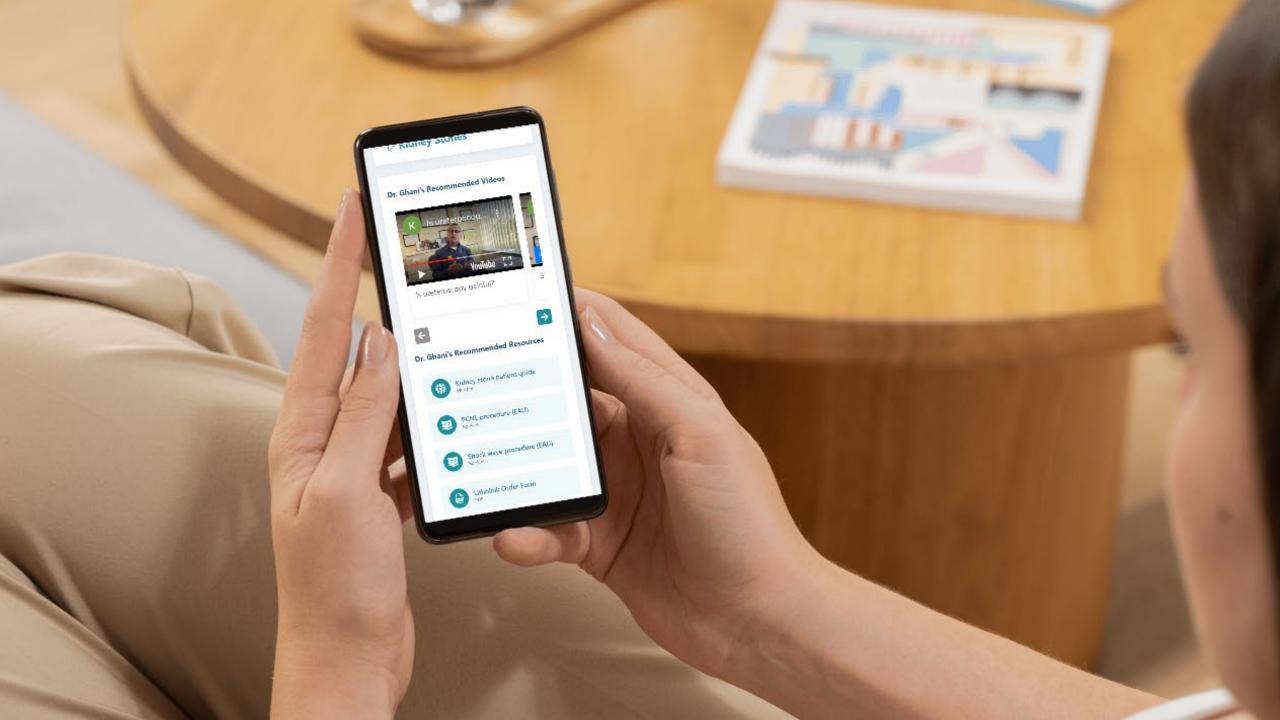


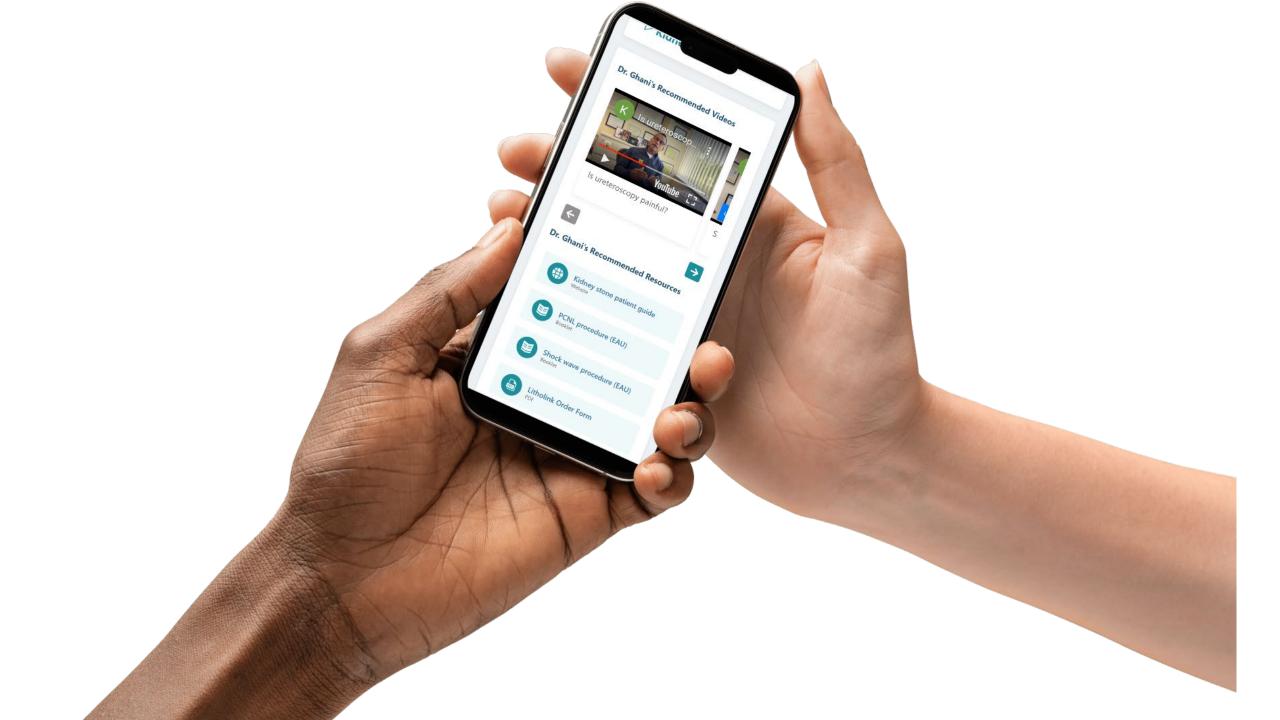


Teach it once. Share it forever.









How do you share it?

2

3

Pre-education

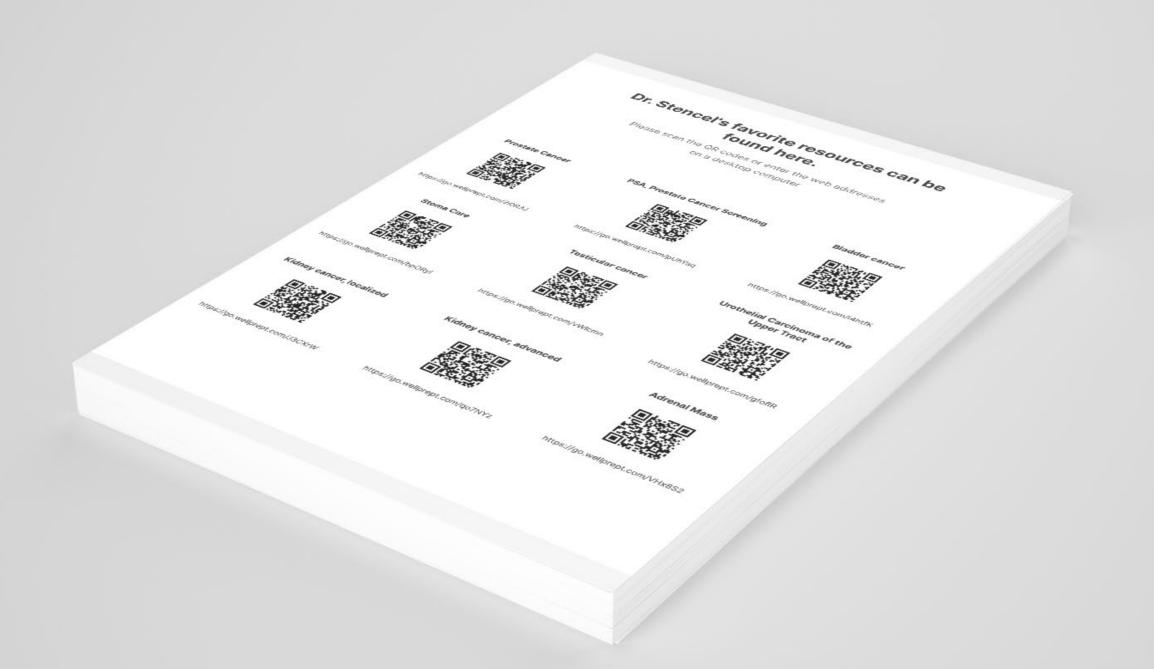
Face to face

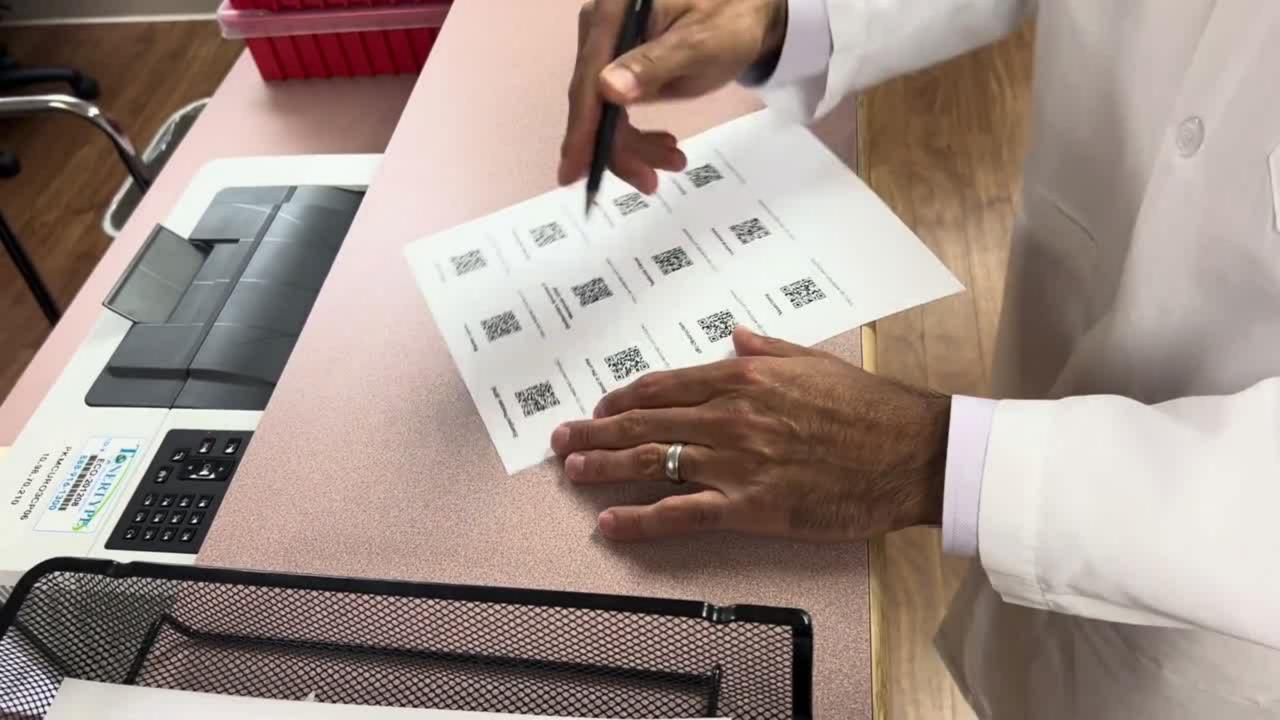
Parting gift

Face to face



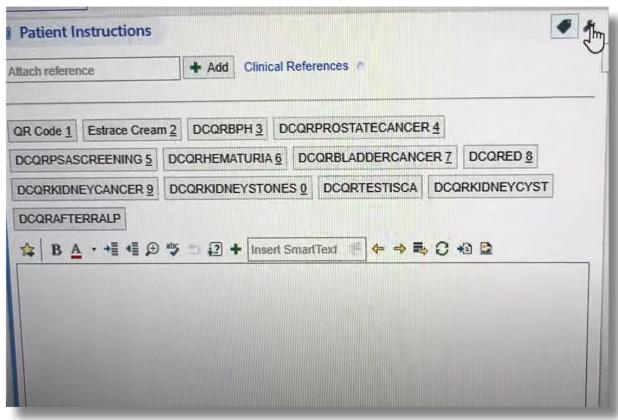
Parting gift

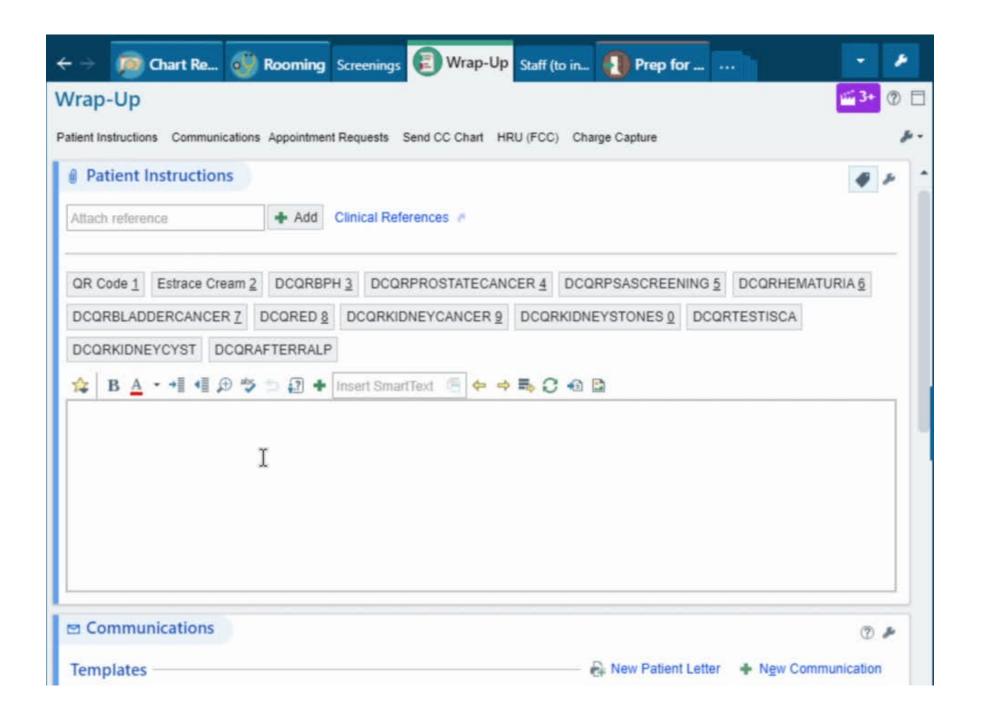




After visit summary / MyChart message



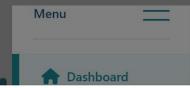




Pre-education

WellPrept Notify

You (when precharting), and/or your team (admin assist, MA, scribe, care coordinator, appointment scheduler)



Send Condition Information

Please remember to not send PHI in your default message



What's next

Improve your wellness

Incorporate into your workflow

Study it

Disease state Patient education Outcome

Disease state Patient education Outcome

Small renal masses - does it allay anxiety on surveillance?

Low risk CaP - does it increase adherence to surveillance?

BPH - improve decision making among surgical choices?

Post prostatectomy incontinence - does targeted edu improve coping/anxiety?

Does edu on dietary and lifestyle modifications improve adherence and reduce stone recurrence?

Does stent education via WellPrept reduce post procedure phone calls?





Let's start with your wellness & your patients' satisfaction

And then tackle high hanging fruit together





































Electronic Delivery of Patient Education: WellPrept Pilot Results

Tudor Borza, MD



Nonprofit corporations and independent licensee of the Blue Cross and Blue Shield Association



A community that partners to improve patients' lives by inspiring high-quality care through data-driven best practices, education, and innovation

Patient Educational Materials

. With time, many men learn more about the disease and find

· Most patients report significantly lower levels of anxiety

ways to deal with their stress

within 2 years

· Maintaining normal routines

· Meditation and yoga

· Spending quality time with loved ones

· Getting the proper amount of sleep

0:00 / 6:02





□ ❖ YouTube []

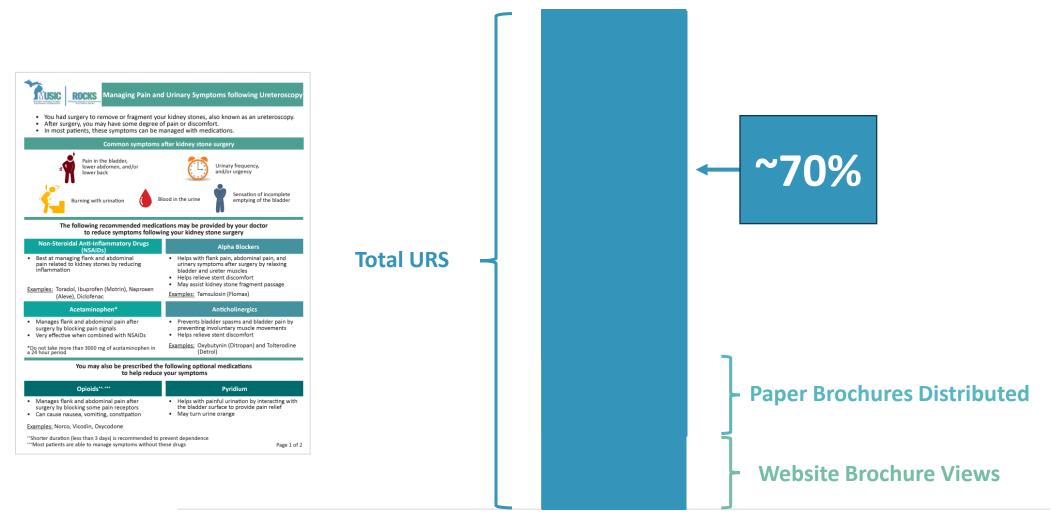
- Recent data suggests that giving radiation at a level of ≥0.2ng/mL (ideally before 1.0 ng/mL) is as effective as giving radiation therapy while the PSA is still undetectable.
- A PSA of ≥0.2ng/mL is typically when doctors and patients consider giving salvage radiation
- The exact value that is best for you may vary and you should discuss this further with your urologist or radiation oncologist.

Page 1 of 2

Resources are Underutilized



Resources Distributed vs Procedures per Month





A community that partners to improve patients' lives by inspiring high-quality care through data-driven best practices, education, and innovation

Can We Increase Distribution?

TYUSIC Michigan Livological Surgery

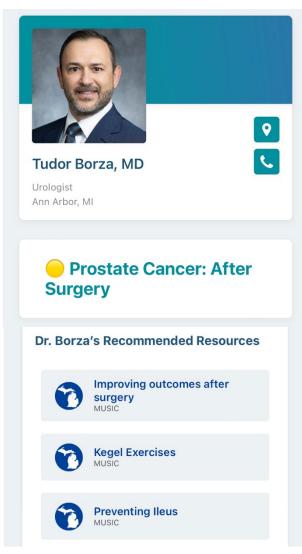
 Doctor to patient content distribution system

Pilot ran July - Sept

Personalized pages for MUSIC physicians

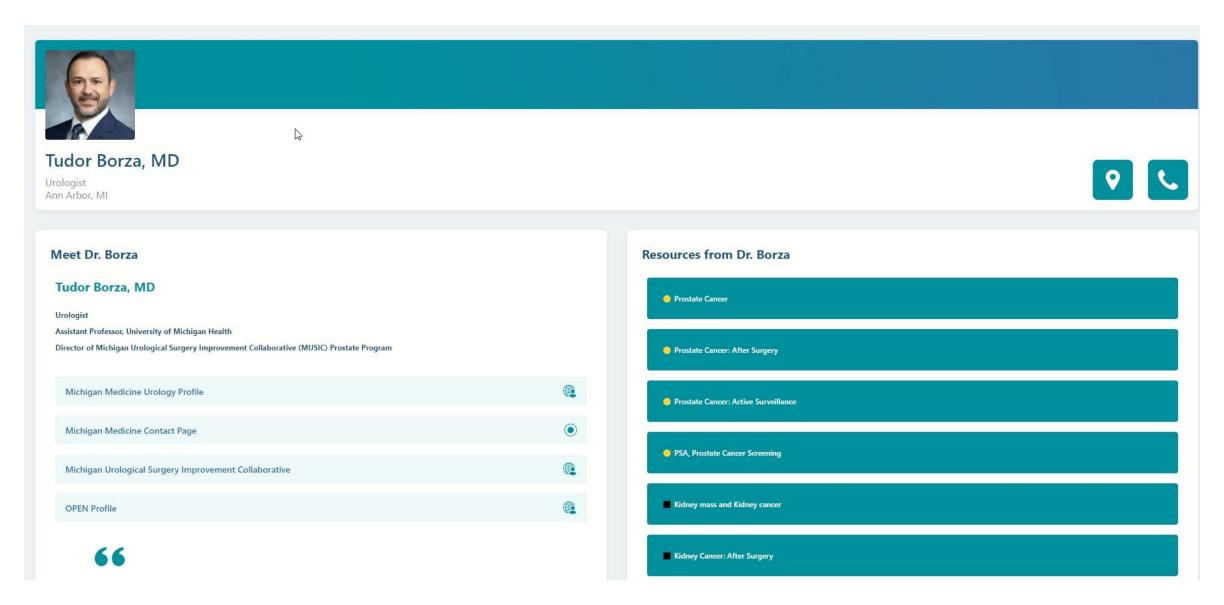
MUSIC materials pre-loaded





How It Works: WellPrept Page





Goals of the Pilot —





#1 Evaluate implementation strategies



Explore provider perceptions

How It Works: Implementation



QR Code/Link

- Flyers, business cards, table tents
- Give to patients during appointment

EHR Integration

- Ad hoc integration
- SmartPhrases added to AVS/discharge summary

Patient Portal/Email

- Added to appt reminders
- In response to questions

WellPrept Notify

- Patient info into WellPrept database
- Links auto sent before appt

П

WellPrept Pilot by the Numbers -





13 MUSIC Urologists

- 11 distributed materials to patients
- 11 responded to pilot survey



3 Months

Most took a month to implement

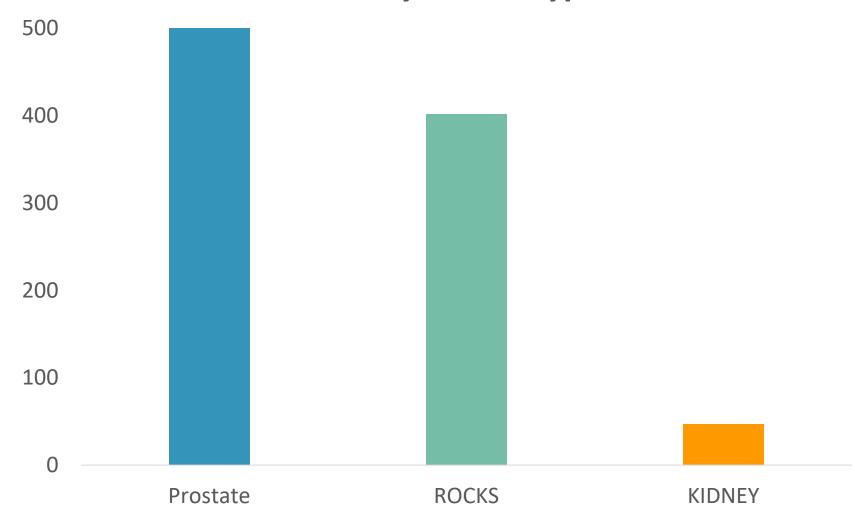


- >1,300 page views, 1400 resource clicks
- Top pilot provider had 354 views!

WellPrept Pilot by the Numbers -



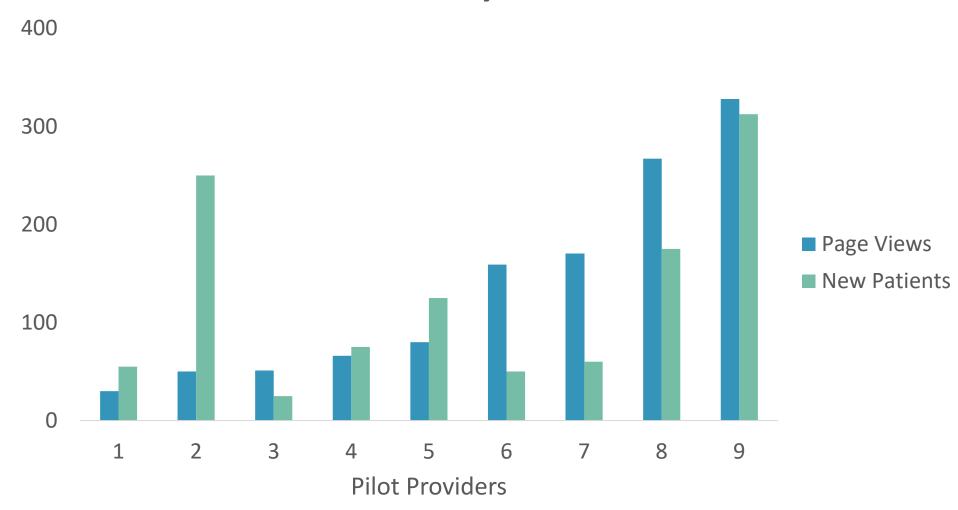




WellPrept Pilot by the Numbers -





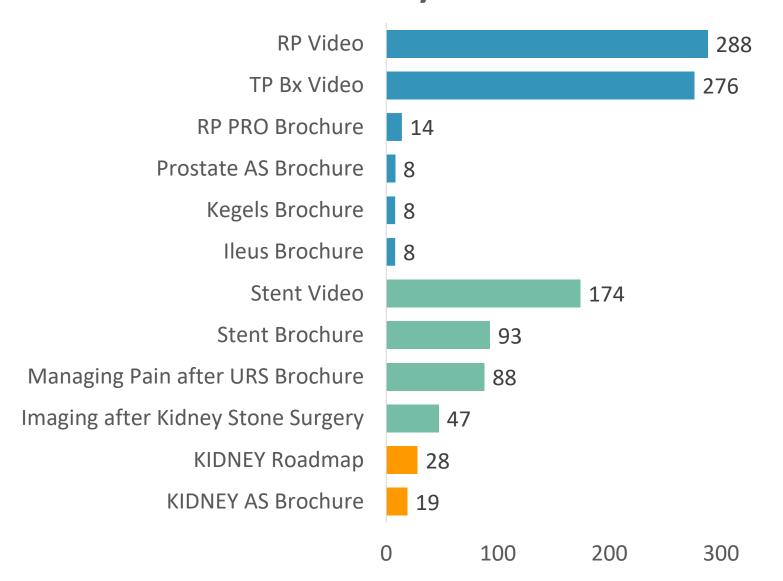


WellPrept Pilot by the Numbers



400

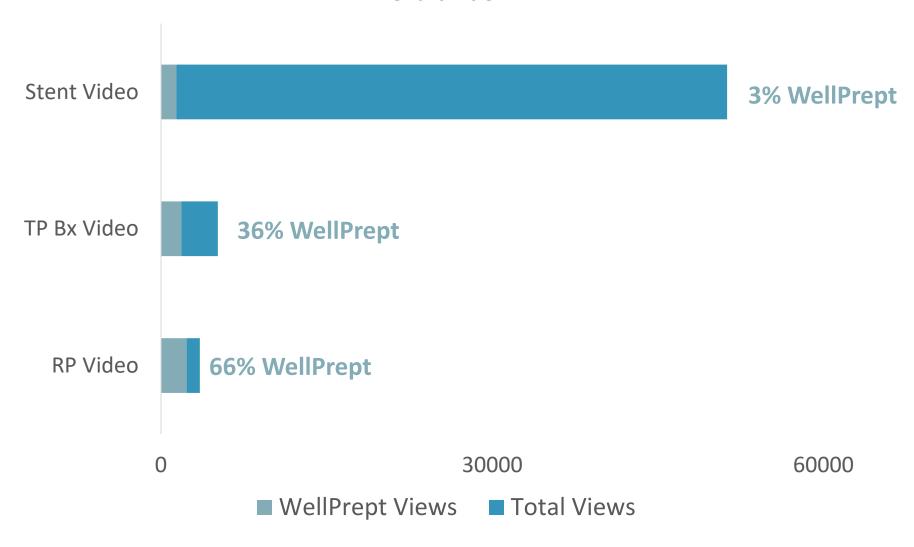
Views by Resource



Beyond the Pilot: MUSIC Resources on WellPrept —



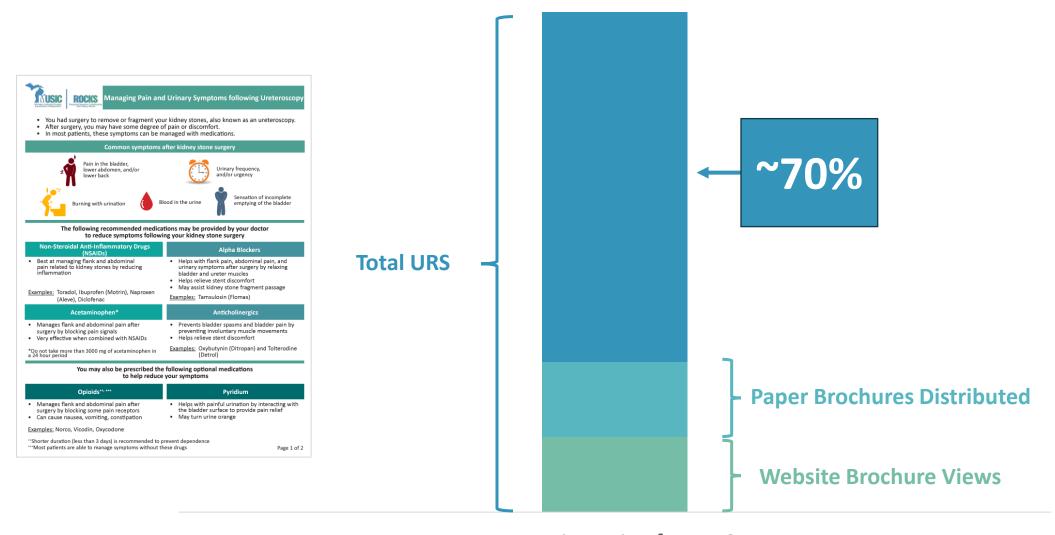
Chart Title



Resources Utilization: Did WellPrept Help?



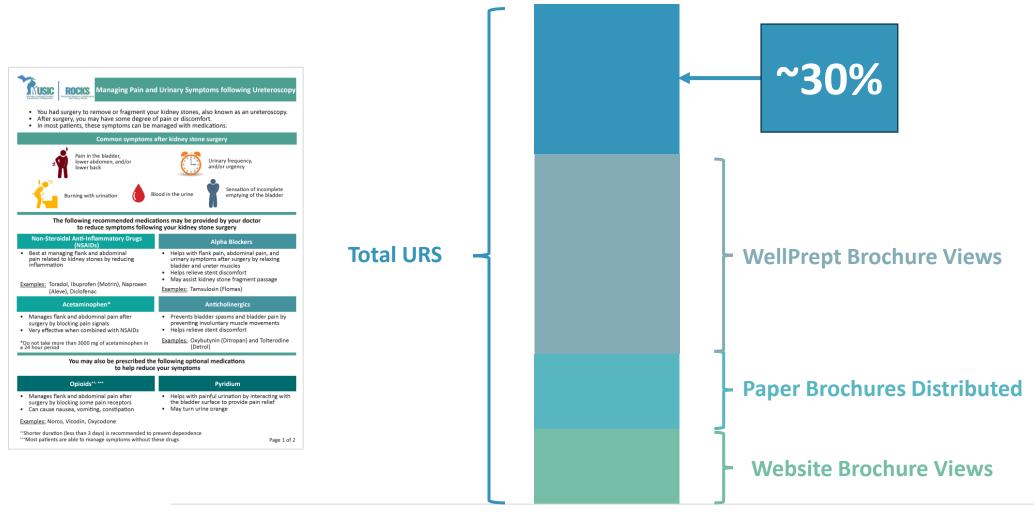
Resources Distributed vs Procedures per Month



Resources Utilization: Did WellPrept Help?



Resources Distributed vs Procedures per Month



WellPrept Views by Implementation Strategy



QR Code/Link

- 4 Providers
- 615 page views
- ~2000 new and return patients

EHR Integration

- 2 Providers
- 329 page views
- ~250 new and return patients

Patient Portal/Email/Text

- 5 Providers
- 370 page views
- ~2000 new and return patients

WellPrept Notify

- 2 Providers
- 267 page views
- ~300 new and return patients

Provider and Patient Experience



Providers

All found WellPrept useful

All reported WellPrept improved patient understanding

20% reported WellPrept shortened office visits

"Patients had more specific questions, especially about surgery"

What could be improved?

- Integration with EHR
- Easier editing
- Searchable on Google

Patients

90% of providers received positive feedback on WellPrept from patients

- "Liked surgical videos"
- "All felt this was helpful"
- "Happy to have a resource"

What are Patients Saying?



"This is insanely useful to me"

-Patient of Dr. Rogers

"These documents are very helpful during a very stressful time. Thanks"

-Patient of Dr. Palka

"Frankly I'm BLOWN AWAY that my doctor would organize all of this for me. There is good in the world after all"

-Patient of Dr. Rogers

WellPrept Beyond the Pilot -













LUNCH



Growing MUSIC: Benign Prostatic Hyperplasia (BPH)

Khurshid Ghani, MD, MS



Nonprofit corporations and independent licensees of the Blue Cross and Blue Shield Association



Harvard Business Review



Spotlight Article

Managing People

The Business Case For Curiosity

by Francesca Gino





 At December 2023 MUSIC Strategic Retreat and in follow-up conversations with current MUSIC members, general urology and specifically BPH was noted as a key area of interest

- BPH is a common condition and the management as a urologic condition is variable and costly
 - > 100,000+ BPH-related procedures performed annually
 - > 12 treatment options!
 - ➤ Nearly \$4 Billion spent on the management of BPH per year

BPH as a QI Opportunity -





П

Development Process



- MUSIC providers interested in serving in a program leadership role invited to submit a BPH Program Letter of Intent (LOI) describing their general aims
- Proposed projects outlined at today's collaborative-wide meeting
- Formal project proposal detailing the project and initiative including patient care benefits and potential ROI submitted by November 18th
- Proposals reviewed by Executive Committee and Coordinating Center
- Decision on successful proposal/leadership team by end of 2024
- BPH summit meeting to be hosted Q1/Q2 in 2025
- Targeting BPH program pilot go-live in Q3 Q4 2025



BPH Letters of Intent (LOIs) Received





Jay Lonsway, MD

Western Michigan Urological Associates

- Revise or create new instruments to adequately assess patient feelings of anxiety and depression post treatment
- PROs to compare multiple surgical intervention outcomes
- Development of quality measures to accurately assess the quality of BPH care
- Share decision making including the creation of a decision aid



Wilson Sui, MD



John DiBianco, MD

Michigan Medicine Urology University of Florida Urology

- Measure the variation in ED visit rates following outpatient BPH surgery
- Examine post-surgical ED care as a driver of episode payments for BPH
- Identify the processes of care employed by high-performing practices



Sabry Mansour, MD

Urology Specialists of MI

- Preserve Bladder Health: Introduce MIST at an earlier stage to prevent complications related to BPH
- Symptoms: Address the limited efficacy of current medical management strategies
- Enhance Diagnostic Accuracy: Utilize advanced diagnostic modalities to prevent unnecessary delays in treatment

We Want to Hear from YOU



 Break out individual tables to discuss what you see as important quality improvement opportunities as it relates to BPH

Where is the unmet need?

What could we do?

Where can we have IMPACT?



Breakout Session: 15 Minutes



Report Out: 15 Minutes

Next Steps



 We will record the feedback received today as we move toward standing up this new program

 We will communicate the selected proposal including the leadership team and specific aims by the end of the year

 Additional thoughts or feedback as it relates to BPH between now and then, please email Susan @ <u>slinsell@med.umich.edu</u>



Improving Ureteroscopy Practice: Lessons Learned from an Ongoing Stent Omission Clinical Trial

Casey Dauw, MD Khurshid Ghani, MD, MS



Nonprofit corporations and independent licensee of the Blue Cross and Blue Shield Association

Disclosures



- Casey Dauw
 - Boston Scientific Corporation, paid consultant
 - Cook Medical, Inc., paid consultant
 - Ethicon, paid consultant
 - Karl Storz Endoscopy, paid consultant

- Khurshid Ghani
 - Ambu, paid consultant, royalties/patent beneficiary
 - Boston Scientific Corporation, consulting fee, grant or research support
 - Coloplast, grant or research support



Part 1: WHY?

1cm LP stone treated with URS —



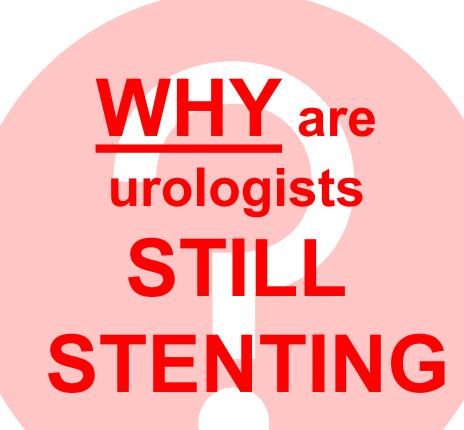




Stenting after Ureteroscopy for Stones

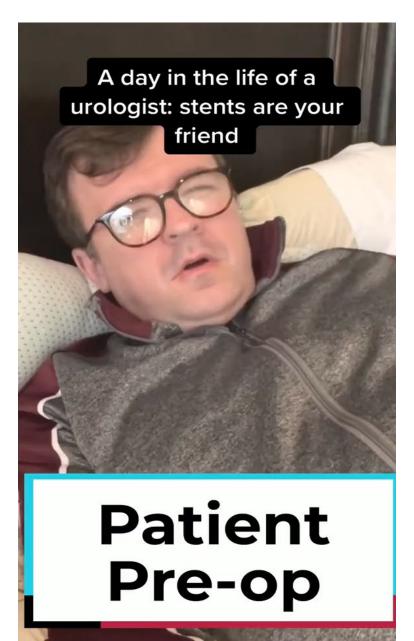


AUA guidelines recommend stent omission after uncomplicated ureteroscopy, but 80% of patients get stented



The Perspective of a Urologist







Strategies to Facilitate Stent Omission: Counselling -





Ureteral Stent Placement following Ureteroscopy Increases Emergency Department Visits in a Statewide Surgical Collaborative

Spencer C. Hiller,* Stephanie Daignault-Newton, † Hector Pimentel, Sapan N. Ambani, John Ludlow, John M. Hollingsworth, Khurshid R. Ghani‡,§ and Casey A. Dauw§

Doc, I really do not want to be stented!



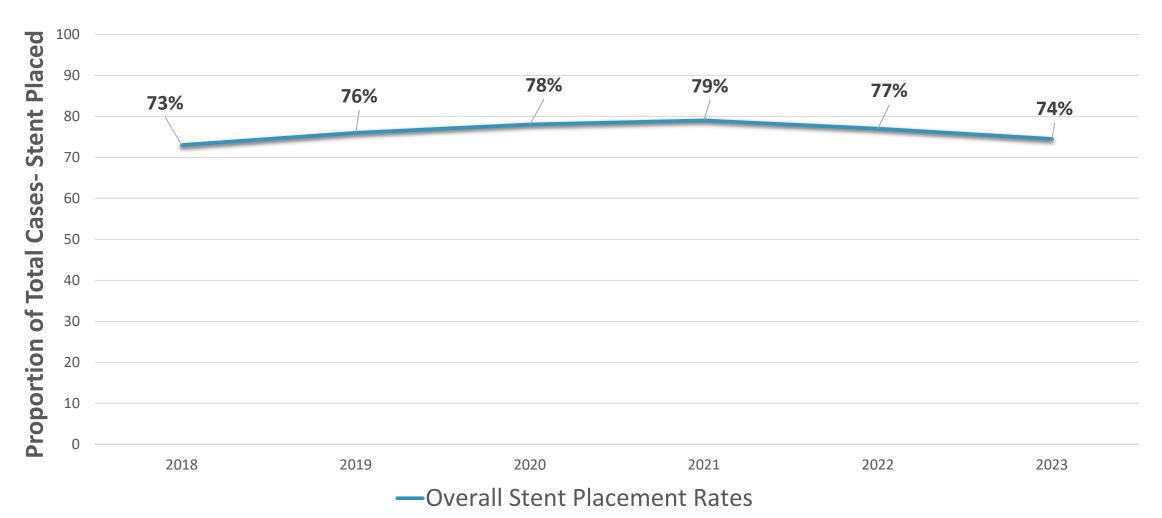
well, 0.5% of cases where a stent is omitted will require urgent intervention postoperatively.

Despite Our Efforts, Stenting Rates Remain Unchanged



Collaborative Wide Stent Placement Rates

(2018 - 2023)



Stent vs No Stent: EVIDENCE IS UNCERTAIN



Ureteral stent versus no ureteral stent for ureteroscopy in the management of renal and ureteral calculi (Review)

Ordonez M, Hwang EC, Borofsky M, Bakker CJ, Gandhi S, Dahm P

2019 Cochrane Review

- Stenting may slightly reduce the number of unplanned return visits
- "But we are very <u>uncertain of this finding</u>"

"Given the importance of this question, higher-quality and sufficiently large trials are needed to better inform decision-making."

We Asked Patients





What is the SOUL MUSIC study?



Hypothesis

Stent omission (vs placement) is associated with improvements in patient reported outcomes (PROs) and 30-day healthcare utilization after ureteroscopy.



Inclusion Criteria

- Small stones <1cm
- Not prestented
- Uncomplicated URS

SOUL Endpoints



Co-Primary Endpoints

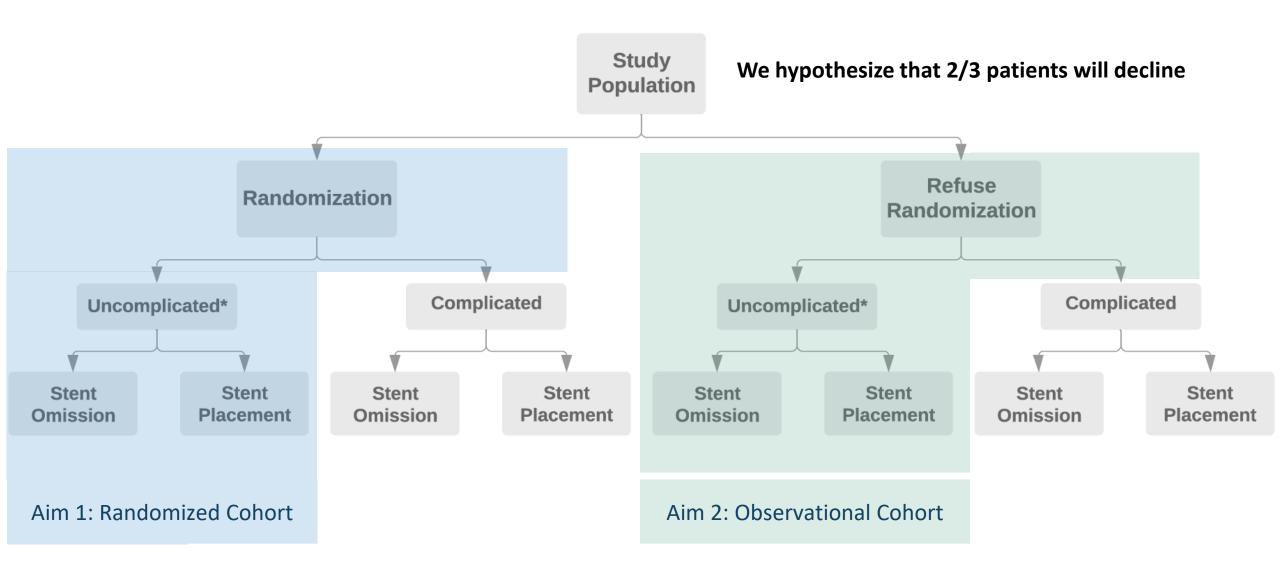
- PROMIS Pain Interference at 7-10 days
- Unplanned healthcare utilization within 30 days

Secondary Endpoints

- Compare/assess the following between treatment arms:
 - Healthcare utilization at each level of composite score
 - Pain and health-related quality of life.
 - Urinary symptoms
 - Treatment satisfaction
 - Time off work for patients and caregivers

Combined Randomized & Observational Design







SOUL is Unique

Michigan Urciogical Surgery Improvement Collaborative

There are **very few** federally funded kidney stone surgical trials...







"Successful completion of this scientifically rigorous study will likely positively impact the field of urology."

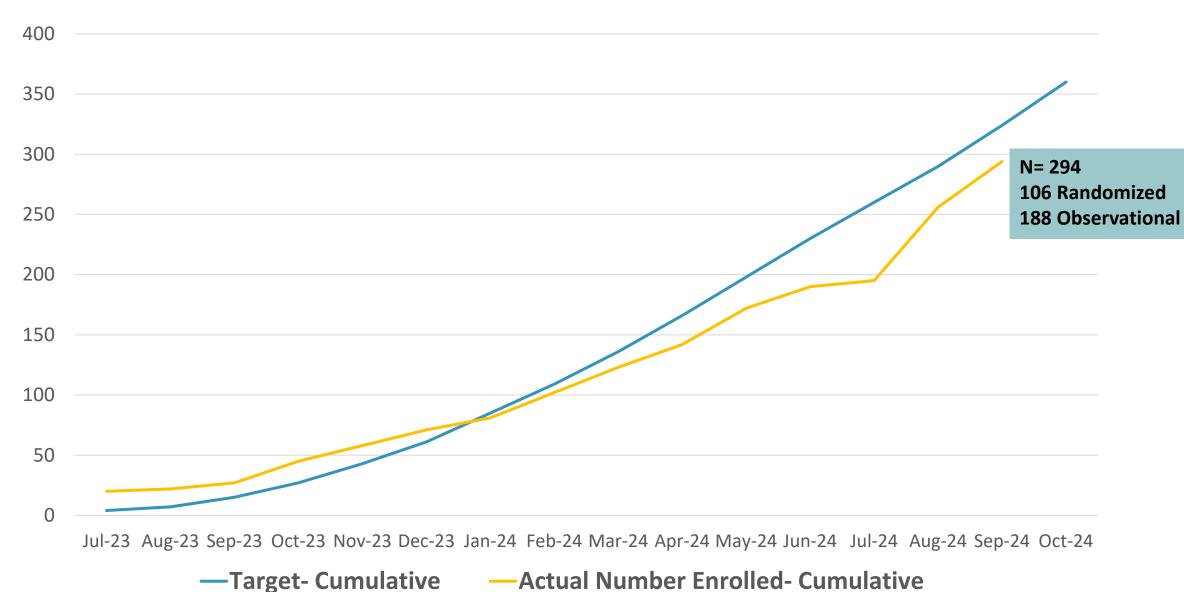
- Peer reviewer



Part 2: Current State of the SOUL Trial

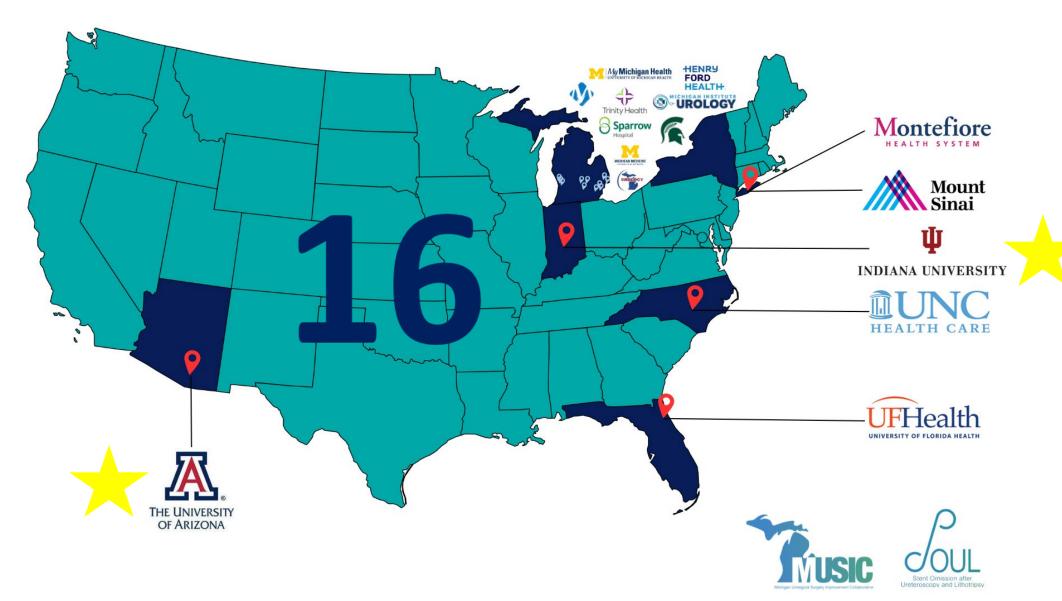
Combined Cohort: Cumulative Patient Enrollment —





2 New Sites to Come Onboard





Thank You for Participating! -





Dr. Mantu Gupta



Dr. Casey Dauw



Dr. Khurshid Ghani



Dr. Karla Witzke



Dr. Jeremy Konheim



Dr. William Roberts



Dr. Sapan Ambani



Dr. Andre King



Dr. Eduardo Kleer



Dr. Dima Raskolnikov



Dr. William Atallah



Dr. Russell Becker



Dr. Henry Rosevear



Dr. Alexander Small



Dr. Neil Pugashetti



Dr. Elena Gimenez



Dr. David Wenzler



Dr. Kara Watts



Dr. Joseph Haddad



Dr. Christina Fox



Dr. Anthony Bonzagni



Dr. Arvin George



Dr. Andrew Schwinn



Dr. Davis Viprakasit



Dr. Suprita Krishna



Dr. Roy Miller



Dr. Wilson Sui



Dr. Dave Friedlander



Dr. Kristian Stensland



Dr. Laris Galejs



Dr. David Leavitt



Dr. Andrew Higgins



Part 3: What have we learned so far?

What Have We Learned so Far? —





Perspectives on stent omission: Patients and Physicians



Incidence of negative ureteroscopy



Preop Alpha-Blockers in ureteroscopy

Guest Surgeons





Dr. Henry Rosevear, MD





Dr. Eduardo Kleer, MD



Integrated Health
Associates



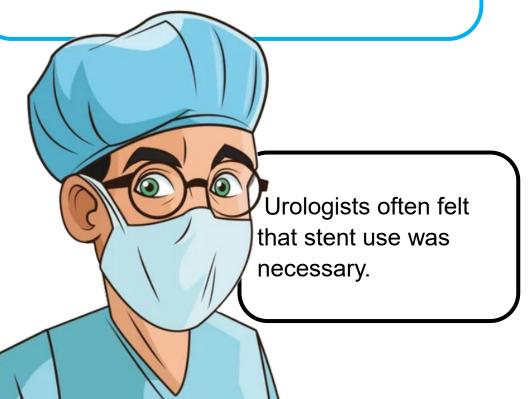
Perspectives on Stent Omission

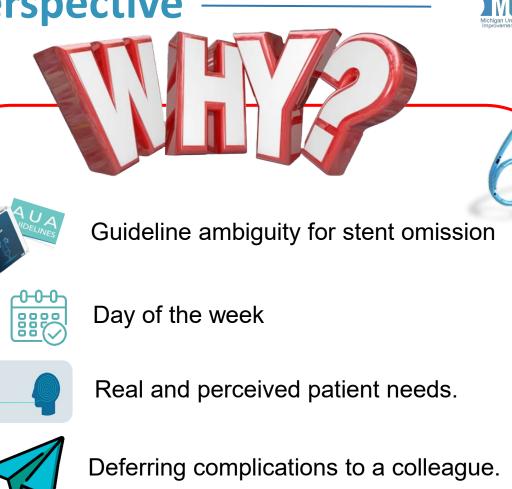
Panel Discussion

Understanding the Surgeon Perspective



Physicians acknowledged that patients prefer stent omission, but **optimal kidney** stone treatment outcomes were prioritized.



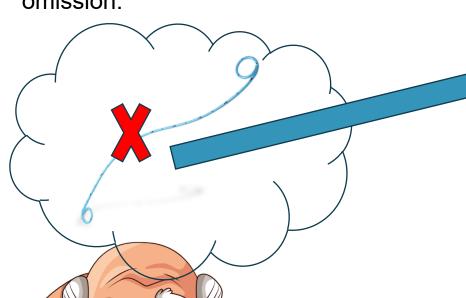


Financial Incentives

Understanding the Patient Perspective



Prior experience led patients towards a preference of stent omission.



Patients strongly preferred stent omission

With stent omission, patients reported less pain, faster recovery and return to work, and a higher quality of life

Patients
emphasized the
need for more education,
especially on stent removal.



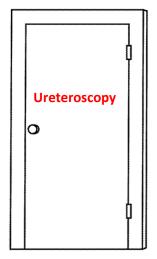
Negative Ureteroscopy

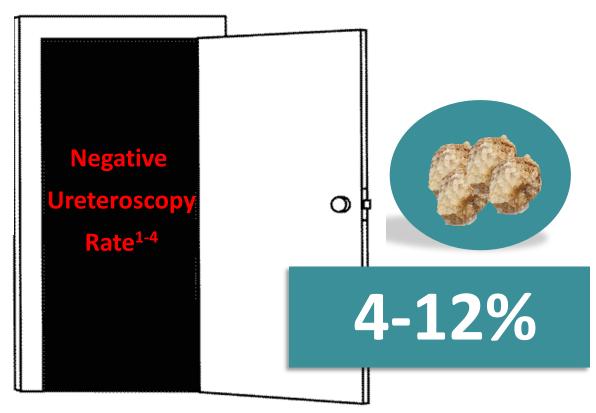
Panel Discussion

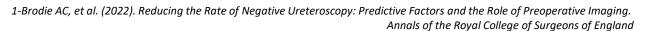


Negative Ureteroscopy: Identifying a Quality Gap









2- Katafigiotis I, et al. (2018). "Stoneless" or Negative Ureteroscopy: A Reality in the Endourologic Routine or Avoidable Source of Frustration?

Estimating the Risk Factors for a Negative Ureteroscopy. Journal of Endourology

3- Lamberts, R. W., et al. (2017). Defining the Rate of Negative Ureteroscopy in the General Population Treated for Upper Tract Urinary Stone Disease. Journal of Endourology





<7mm **ureteral** stones

Cost Implications of Negative Ureteroscopy





Variation in Spending around Surgical Episodes of Urinary Stone Disease: Findings from Michigan

Juan San Juan, Hechuan Hou, Khurshid R. Ghani,* James M. Dupree† and John M. Hollingsworth‡



~\$11,000

Each

Ureteroscopy Procedure



56Cases in Michigan
Each Year



~\$600,000 Estimated Annual Cost

Reducing negative ureteroscopy cases can lead to substantial savings



Preoperative Alpha Blockers

Panel Discussion

Failed Ureteroscopy







Effect of preoperative alpha-blockers on ureteroscopy outcomes: A meta-analysis of randomised trials

First published: 03 April 2024 | https://doi.org/10.1002/bco2.358 | Citations: 1

7% Failed URS



Strategies to Avoid Failed URS: Preop Alpha Blockers





Effect of preoperative alpha-blockers on ureteroscopy outcomes: A meta-analysis of randomised trials

Naeem Bhojani, Ben H. Chew, Samir Bhattacharyya, Amy E. Krambeck, Khurshid R. Ghani, Larry E. Miller



Shorter procedure time



Reduced need for mechanical Ureteral Dilation



Fewer complications



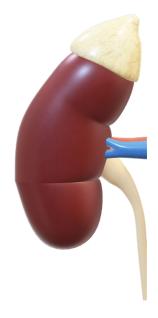
Decreased Access Failure Rate

Preoperative Alpha Blocker Use in MUSIC —



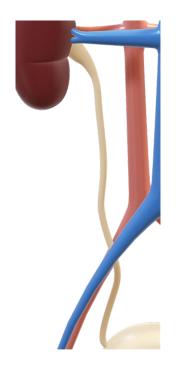
KIDNEY

35%



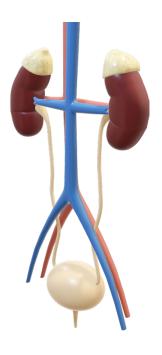
URETER/URETER + KIDNEY

45%



OVERALL

40%



Key Takeaways



MUSIC

Coordinating Center

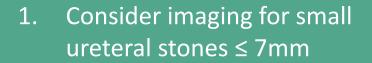


Patient educational video/leaflet on stent removal strategies



Results from the ongoing SOUL Clinical Trial







- Reduce failed ureteroscopy
- Facilitate stent omission?

Participating Practices/ Urologists

Potential Impact of SOUL MUSIC





Change Urological Norms

Routine stenting may not be necessary post-ureteroscopy.

June 2025

Patient-Centric Approach

Improved patient outcomes by reducing unnecessary stent placements.

Practice Changing

Influencing future clinical guidelines based on trial results.

Long-Term Patient Benefits

Improving recovery and lowering complication rates for patients.



Closing Remarks

Tudor Borza, MD



Nonprofit corporations and independent licensees of the Blue Cross and Blue Shield Association

Prostate Key Takeaways



• 1/3 of prostatectomy patients have cancer after surgery

Persistently positive and biochemically recurrent cancer are different

Risk adapted management and collaboration with radiation oncology are key

KIDNEY Key Takeaways



Accurate clinical stage is needed to study oncologic outcomes

26% of MUSIC KIDNEY cases have incorrect clinical stage documented

- Clinicians should
 - Use cNx and cMx only for indeterminate lesions
 - Use cN0 and cM0 more frequently

ROCKS Key Takeaways



SOUL clinical trial is underway in 13 centers throughout MUSIC

- Interviews have found that patients strongly prefer stent omission
- While physicians often feel stents are needed

There is a ~4% negative URS rate in SOUL

Consider re-imaging for small ureteral stones to reduce negative surgery

Consider pre-op alpha-blockers to reduce failed URS AND facilitate stent omission



Save the Date



MUSIC Nationwide Webinar

Beyond the Operating Room:
Tools and Techniques for Managing Adverse Events



Wednesday
April 16th
6:00-7:45PM ET



