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Spring 2-8-2023

# PROTOCOL: New York State Race, Ethnicity, and Insurance Disparities in Follow-up Prostate Cancer Screening

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#### **Recommended Citation**

Greenspan, Seth; Chandra, Mansi; Joo, Hyun Woo; Sapir, Netanel; Gorman, Jonathan; Yang, Jie; Li, Xiaoning; Cavale, Barghav; Fierro, Allegra; Shroyer, Annie Laurie Laurie; and Fitzgerald, John P., "PROTOCOL: New York State Race, Ethnicity, and Insurance Disparities in Follow-up Prostate Cancer Screening" (2023). *Department of Urology Faculty Publications*. 3. https://commons.library.stonybrook.edu/dou-articles/3

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arghav Cavale,	Allegra Fierro, A	Annie Laurie L	aurie Shroyer	and John P. F	itzgerald	

<u>TITLE</u>: New York State Race, Ethnicity, and Insurance Disparities in Follow-up Prostate Cancer Screening

**Principal Investigator:** Dr. John Fitzgerald, MD, MSCI, and Dr. A. Laurie Shroyer, PhD, MSHA

Co-Principal Investigators: Hyun Woo Joo, Seth Greenspan, Mansi Chandra and Netanel Sapir

Biostatistician: Jonathan Gorman

### **PURPOSE AND SPECIFIC AIMS:**

Using de-identified reports from the Statewide Planning and Research Cooperative System (SPARCS) data, this descriptive study will identify the impact of socioeconomic status (SES) metrics on the follow-up prostate cancer screening care within 3 years of index prostate cancer screening test in NYS. The socioeconomic status metrics will be subclassified into race, insurance, and ethnicity and each of these sub-components will be evaluated for its impact on the follow-up cancer screening care. The exclusion criteria for this study includes patients records with unknown age, age <55 or >75, previous history of prostate cancer or radical prostatectomy, previous prostate biopsy, female sex, lives outside NYS, unknown or missing data on race, ethnicity, or insurance status, or multi-ethnic patients. For the included patients, initial prostate cancer screening, follow-up screening, characteristics (e.g., age, SES), and risk profiles will be evaluated. Moreover, patients diagnosed with prostate cancer or receiving prostatectomy will be reported.

Additionally, the following hypotheses will be tested:

H(0): Among patients with a baseline PSA test, socioeconomic status (SES) metrics (i.e., vulnerability based upon race/insurance/ethnicity) may pose as barriers to follow-up prostate cancer screening care within 3 years of index prostate cancer screening test (e.g., Vulnerability = V = Black, Hispanic, and Self-pay Insurance)

- H(0): Among patients with a baseline PSA test, race does not impact the likelihood of follow-up prostate cancer screening care within 3 years of index prostate cancer screening test (e.g., R-FC)
- H(0): Among patients with a baseline PSA test, insurance does not impact the likelihood of follow-up prostate cancer screening care within 3 years of index prostate cancer screening test (e.g., I-FC)
- H(0): Among patients with a baseline PSA test, ethnicity does not impact the likelihood of follow-up prostate cancer screening care within 3 years of index prostate cancer screening test (e.g., E-FC)

Please note, the SPARCS database de-identified reports will be used. Additionally, a not human subject's research (NHSR) determination is requested.

#### **BACKGROUND AND SIGNIFICANCE:**

For the period from January 2010 to December 2018, this analysis will use the NYS SPARCS database trends to evaluate the influence of race and insurance on follow-up prostate cancer screening. Disparities in prostate cancer survival based on patient's race has been well documented by previous research. A 2018 retrospective propensity analysis of prostate cancer patients diagnosed between 2004-2010 found that white men had a greater overall survival than black men, and that this survival difference was eliminated when their model simulated equal access to prostate cancer care.<sup>2</sup> A 2017 study found that African American and Hispanic men are less likely to receive definitive treatment for prostate cancer than white men, and the rates of treatment declined throughout their study period from 2004-2011.<sup>3</sup> Additionally, while Black men have a greater lifetime incidence of prostate cancer, a recent study found that they are less likely to receive PSA-testing.<sup>4</sup> Moreover, a 2013 post-hoc study of patients enrolled in the Prostate, Lung, Colorectal, and Ovarian (PLCO) trial found that among patients screened with a PSA test, Black patients were less likely than non-Hispanic Whites to receive follow-up PSA testing or prostate biopsies.<sup>5</sup>

In addition to the racial disparities in prostate cancer outcomes, previous research has exhibited that there are disparities in prostate cancer care based on the patient's insurance status. A 2018 study on the association between expanded insurance coverage and prostate cancer screening found that the gap in PSA test utilization between the higher and lower income male patients in Medicaid early expansion states was significantly reduced.<sup>6</sup> A 2019 study conducted on patients diagnosed with prostate cancer in Florida found that patients without insurance or with Medicaid were more likely to be diagnosed with late-stage prostate cancer.<sup>7</sup> Furthermore, a 2019 retrospective cohort study found that patients with private and public insurance were more likely to receive favorable treatment.<sup>8</sup> Bledsoe et al. (2018) found that insurance has an effect on treatment modality, as patients with private insurance were more likely to receive minimally invasive surgery and less likely to receive external beam radiotherapy than patients without insurance.<sup>9</sup>

Our study intends to further explore the relationship between socioeconomic factors and access to prostate cancer care, by examining how race and insurance affects patient's access follow up prostate cancer screening with PSA or biopsy. While this disparity has been studied with respect to Black and White race,<sup>5</sup> there are no previous studies examining the effect of insurance and ethnicity on follow-up screening. All results will be adjusted for patient comorbidities and age. Lastly, these variables will be stratified by year, so that trends over time can be evaluated.

#### **RESEARCH DESIGN AND METHODS:**

This retrospective observational cohort study will be done using the SPARCS Health Facts dataset. With the help of the SBU SOM Bioinformatics Department and Biostatistics Core Lab, the SPARCS database will be matched/merged to the enclosed coding listings to create a study-specific de-identified prostate cancer screening database. Furthermore, the Bioinformatics and Biostatistics team members will be responsible for providing the descriptive statistics listed below as well as providing a study-database for future analyses. For this study's primary hypothesis, a p-value of <0.001 will be used (however, all p-values will be reported by separate interpretation by readers). All secondary and tertiary analyses, as well as all exploratory analyses, will use a p-value of < 0.01. SAS version 9.4 will be used to complete all the

necessary statistical tests. More detailed methods including ICD-10, ICD-9, and CPT codes used in this study are described in the protocol below.

## **REFERENCES**

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- 7. Ramirez E, Morano J, Beguiristain T, et al. Insurance status as a modifier of the association between race and stage of prostate cancer diagnosis in Florida during 1995 and 2013. Cancer Epidemiol 2019;59:104-8.
- 8. Awasthi S, Gerke T, Williams VL, et al. Interrelationship Between Health Insurance Status and Prostate Cancer Grade Can Have Critical Impact on Prostate Cancer Disease Control: A Retrospective Cohort Study. Cancer Control 2019;26:1073274819837184.
- 9. Bledsoe TJ, Park HS, Rutter CE, Aneja S, Nguyen PL, Yu JB. Impact of Health Insurance Status on Prostate Cancer Treatment Modality Selection in the United States. Am J Clin Oncol 2018;41:1062-8.

## **Protocol:**

**Population:** The population of this study is NYS males from the ages of 55-75 who received a prostate cancer screening test from 2010-2018 as defined in **Table 1**. Each patient will be sorted by the year of the initial screening test performed, in order to track trends over time.

**Table 1: Prostate Cancer Screening Test Codes** 

Prostate Cancer S	Prostate Cancer Screening Codes						
Procedure	ICD-9	ICD-10	СРТ	Notes			
Elevated	790.93,	R97.20, Elevated	NA				
Prostate Specific	Elevated	PSA					
Antigen (PSA)	prostate						
	specific						
	antigen						
Prostate Specific	V76.44	Z12.5	84152, Complex PSA				
Antigen (PSA)							
			84153, Total PSA				
			84154, Free PSA				
			81593, Panel PSA				
Prostate Needle	60.11,	0VB03ZX	55700, 55705,	Any ONE ICD-9, ICD-10, or			
Biopsy (Open or	60.12	0VB04ZX	55706	CPT satisfies criteria			
Closed)		0VB07ZX					
		OVB08ZX,					
		0VB00ZX					

#### **Inclusion Criteria:**

- -PSA test, elevated PSA test, or prostate biopsy between 2010-2018
- -Male sex
- -Age 55-75
- -NYS Resident

## **Exclusion Criteria:**

- -In-hospital death in initial record
- -With prostate cancer diagnosis prior to or at the time of initial encounter (**Table 2a** or **2b** codes)
- -With prostatectomy or prostatectomy prior to or at the time of initial encounter (**Table 3** codes)
- -With prostate biopsy prior to or at the time of initial encounter
- -With unknown or missing data on race, ethnicity, or insurance or multi-ethnic patients

**Step 1:** The population will be divided into **three** groups based on the **first** screening test recorded in SPARCS. They are listed in order of priority below (Group 1 = highest priority)

Group 1: Elevated PSA test from diagnostic code

Group 2: Prostate Specific Antigen (PSA) test, results unspecified (outpatient data only)

Group 3: Prostate biopsy

**Table 2a:** Prostate Cancer Diagnosis Codes:

Diagnosis	ICD-9	ICD-10	Notes
Prostate Cancer	185	C61	
Carcinoma in	233.4	D07.5	
situ of prostate			

**Table 2b:** Prostate Cancer Metastasis Codes

Diagnosis	ICD-9	ICD-10	Notes
Prostate Cancer	198.5	C79.51 C79.82	
Metastasis	198.82	C77.2 C77.5	
	196.2		
	196.6		

**Table 3:** Radical Prostatectomy Codes

Procedure	ICD-9	ICD-10	СРТ	Notes
Radical	60.3, 60.4,	OVTOOZZ,	55810, 55812,	Any ONE ICD-9, ICD-10,
Prostatectomy	60.5,	0VT04ZZ,	55815, 55840,	or CPT satisfies criteria
(Laparoscopic/robotic	60.62, or	OVTO7ZZ,	55842, 55845,	
or open)	60.69	or 0VT08ZZ	55866	

Table 4: Radiation Procedure Codes

Procedure	ICD-9	ICD-10 PCS	СРТ	Notes
Radiation	92.30, 92.31,	DV20DZZ,	77373, 77385, 77386,	Any ONE ICD-9, ICD-
(Stereotactic	92.32, 92.33,	DV20HZZ,	77424, 77425, 77520,	10, or CPT satisfies
Radiosurgery,	92.39, 92.20,	DV20JZZ,	77522, 77523,	criteria
Brachytherapy,	92,23, 92. 28,	DV1097Z,	77525,77600, 77605,	
IMRT, Beam	92.27, 92.22,	DV1098Z,	77610, 77615, 77620,	
Radiation)	92.24, 92.25,	DV1099Z,	77770, 77771, 77772,	
	92.26, 92.21,	DV109BZ,	77778, 77371, 77372,	
	99.85, 92.29,	DV109CZ,	77373, 7740177402,	
	17.69, 92.41	DV109YZ,	77403, 77404, 77406,	
		DV10372,	77407, 77408, 77409,	
		DV10B72,	77411, 77412, 77413,	
		·	77414, 77416, 77418,	
		DV10B9Z,	77423, 77424, 77425,	
		DV10BBZ,	77520, 77522, 77523,	

	DV10BB1, DV10BCZ, DV10BYZ, 3E0N304, 3E0N704, 3E0N804, DV000ZZ, DV001ZZ, DV002ZZ, DV003ZO, DV003ZZ, DV004ZZ, DV005ZZ, DV006ZZ, DVY07ZZ, DVY08ZZ, DVY0CZZ, DVY0FZZ, DVY0KZZ	77525, 77781, 77782, 77783, 77784, 77785,77786, 77787 0395T, G0173, G0251, G0339, G0340, G6003, G6004, G6005, G6009, G6010, G6011, G6012, G6013, G6014, G6015, G6016	
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**Step 2:** For each group of patients, the number/percentage that had each of the following follow-up scenarios will be recorded as outcomes.

#### **Screening Outcomes:**

Outcome 1A: Repeat/Follow-up PSA, results unspecified, within 3 years

Outcome 1B: Repeat/Follow-up elevated PSA within 3 years

Outcome 1C: Follow up biopsy (Table 2) within 3 years

#### No follow-up

Outcome 2: No repeat screening tests within 3 years

**Step 3:** Initially, a univariate analysis will be performed to determine the marginal association between the following variables/exposures and the above outcomes using either Chi-Square tests and/or other statistical tests. Additionally, the specific vulnerable groups listed below will be compared to others who are outside that group.

## Variables/Exposures:

Age: Yearly

Race: Black, Non-Hispanic White, other races

Ethnicity: Hispanic, Non-Hispanic

Insurance: Medicare, Medicaid, Commercial, Self-pay

Vulnerable groups: Black race, Hispanic ethnicity, Self-pay insurance status

Additional note: The following variables will be adjusted for in the analysis including comorbidities, risk factors for prostate cancer, and subsequent prostate cancer diagnosis/treatment after initial treatment. To clarify, our study will **exclude** patients who are diagnosed/treated with prostate cancer before their initial PSA test or biopsy but will **adjust for** patients in the analysis who received a prostate cancer diagnosis or treatment after their initial screening test but before their subsequent screening test.

Comorbidities: Charlson or Elixhauser Comorbidity Score
Risk Factors: Smoking/Tobacco, Morbid Obesity, Family History of Prostate Cancer, Personal
History of Irradiation
Prostate Cancer Diagnosis or Treatment w/ prostatectomy or radiation (Tables 2a, 2b, 3, and 4)

## **SAMPLE TABLES**

Table 1: Descriptive table of patients' characteristics, risk factors, comorbidity score, index of comorbidities and outcomes by vulnerability status

Variable	Level	Total (N =)	V (N =)	NOT V (N =)	P-value*
Patients' characte	eristics at the ti	ime of Initial P	C Screening		
Abnormal PSA					
Age	Yearly				
Age Categories	40-55				
	55-69				
	70 - 75				
Year of prostate cancer screening	2011				
cancer screening	2012				
	2013				
	2014				

	2015						
Risk factors at the time of Initial PC Screening							
Smoking/Tobacc o	No						
	Yes						
Morbid Obesity	No						
	Yes						
Family History of Prostate	No						
Cancer	Yes						
	Yes						
Personal History of Irradiation	No						
	Yes						
Comorbidities' sc	ore at the time	e of Initial PC	Screening				
Elixhauser Score							
<b>Charlson Score</b>							
Elixhauser Comorbidities at the time of Initial PC Screening							
Congestive heart failure	No						
	Yes						
Cardiac arrhythmias	No						
·	Yes						
	1						

Valvular disease	No				
	Yes				
Pulmonary circulation	No				
disease	Yes				
Peripheral vascular disease	No				
	Yes				
Hypertension, uncomplicated	No				
	Yes				
Hypertension, complicated	No				
	Yes				
Paralysis	No				
	Yes				
Other neurological	No				
disorders	Yes				
Chronic pulmonary disease	No				
	Yes				
Diabetes w/o chronic	No				
complications	Yes				
<u> </u>	1	1	ı	ı	1

Diabetes w/ chronic complications	No			
	Yes			
Hypothyroidism	No			
	Yes			
Renal failure	No	2		
	Yes			
Liver disease	No			
	Yes			
Peptic ulcer Disease, excluding	No			
bleeding	Yes	2		
HIV and AIDS	No			
Lymphoma	No			
	Yes			
Metastatic cancer	No			
	Yes			
Solid tumor w/out metastasis	No			
	Yes			
	No			
	•		 -	-

		Т			
Rheumatoid arthritis/collage n vascular d	Yes				
Coagulopathy	No				
	Yes	3			
Obesity	No				
	Yes	2			
Weight loss	No				
	Yes				
Fluid and electrolyte	No				
disorders	Yes	2			
Blood loss anemia	No				
	Yes	2			
Deficiency Anemia	No				
	Yes				
Alcohol abuse	No				
	Yes				
Drug abuse	No	3			
	Yes				
Psychoses	No	9			
	Yes	Þ			
L			l .	l .	

Depression	No									
	Yes	3								
Charlson Comork	Charlson Comorbidities at the time of Initial PC Screening									
Congestive heart failure	No									
lanure	Yes	2								
Chronic	No									
pulmonary disease	Yes									
Diabetes w/o chronic complications	No									
complications	Yes	2								
Diabetes w/ chronic	No	5								
complications	Yes									
Myocardial infarction	No	2								
	Yes									
Peripheral vascular	No									
disorder	Yes									
Cerebrovascular disease	No									
	Yes									
Dementia	No									

	Yes	3		
Rheumatic disease	No	9		
	Yes			
Peptic ulcer disease	No			
	Yes			
Mild liver disease	No			
	Yes			
Hemiplegia or paraplegia	No			
	Yes			
Renal disease	No			
	Yes			
Any malignancy w lymphoma and leukemia	No			
w/o malignant neoplasm of skin	Yes	9		
Moderate or severe liver	No	3		
disease	Yes			
Metastatic solid tumor	No			
	Yes			

AIDS/HIV	No				
Coronary Artery Disease	No				
	Yes				
Connective Tissue Disorder	No				
	Yes				
Moderate-Severe Renal Disease	No				
I	Yes				
ANY Follow-up Screening	Yes				
	No				
Follow-up Biopsy	Yes				
	No				
Follow-up Abnormal PSA	Yes				
	No				
Follow-up PSA/ PC Screening (not abnormal PSA) NEW CODES+	Yes				
CPT CODES	No				
*. F4:- 1	. 11 1	1 1 (1)	1	, 1 C	M + C 1

<sup>\*:</sup> For categorical variables, p-values were based on Chi-squared test with exact p-value from Monte Carlo simulation; for continuous variable, p-value was based on Wilcoxon rank sum test.

Table 2: Descriptive table of patients' characteristics, risk factors, comorbidity score, index of comorbidities and outcomes by race

Variable	Level	Total (N =)	Black (N =)	NOT Black (N =)	P-value*
Patients' characte	eristics at the t	ime of Initial l	PC Screening		
Abnormal PSA					
Age	Yearly				
Age Categories	40-55				
I	55-69				I
	70 - 75				
Year of prostate cancer screening	2011				
realised screening	2012				I
	2013				
	2014				
	2015				
Risk factors at th	e time of Initia	l PC Screenin	g g	<u> </u>	
Smoking/Tobacc o	No				
1	Yes				1
Morbid Obesity	No				

	Yes				
	103				
Family History of Prostate Cancer	No				
	Yes				
	Yes				
Personal History of Irradiation	No				
	Yes				
Comorbidities' sc	ore at the time	of Initial PC	Screening		
Elixhauser Score					
Charlson Score					
Elixhauser Como	rbidities at the	time of Initia	PC Screening	,	
Congestive heart failure	No				
	Yes				
Cardiac arrhythmias	No				
•	Yes				•
Valvular disease	No				
ı	Yes				l
Pulmonary circulation	No				
disease	Yes				-
	No				

Peripheral vascular disease	Yes		
Hypertension, uncomplicated	No		
	Yes		
Hypertension, complicated	No		
	Yes		
Paralysis	No		
•	Yes		•
Other neurological	No		
disorders	Yes		
Chronic pulmonary	No		
disease	Yes		
Diabetes w/o chronic	No		
<sup>1</sup> complications	Yes		•
Diabetes w/ chronic	No		
complications	Yes		'
Hypothyroidism	No		
	Yes		
Renal failure	No		

	Yes		
Liver disease	No		
	Yes		
Peptic ulcer Disease, excluding	No		
bleeding	Yes		
HIV and AIDS	No		
Lymphoma	No		
1	Yes		'
Metastatic cancer	No		
'	Yes		'
Solid tumor w/out metastasis	No		
•	Yes		•
Rheumatoid arthritis/collagen	No		
vascular d	Yes		•
Coagulopathy	No		
1	Yes		
Obesity	No		
	Yes		

Weight loss	No			
ı	Yes			
Fluid and electrolyte	No			
disorders	Yes			'
Blood loss anemia	No			
•	Yes			
Deficiency Anemia	No			
	Yes			
Alcohol abuse	No			
	Yes			
Drug abuse	No			
	Yes			
Psychoses	No			
•	Yes			
Depression	No			
•	Yes			
<b>Charlson Comor</b>	bidities at the t	ime of Initial	PC Screening	
Congestive heart failure	No			
	Yes			

Chronic pulmonary	No		
disease	Yes		•
Diabetes w/o chronic	No		
complications	Yes		
Diabetes w/ chronic	No		
complications	Yes		
Myocardial infarction	No		
	Yes		
Peripheral vascular disorder	No		
	Yes		
Cerebrovascular disease	No		
	Yes		
Dementia	No		
	Yes		
Rheumatic disease	No		
	Yes		
Peptic ulcer disease	No		
	Yes		

Mild liver disease	No		
1	Yes		ı
Hemiplegia or paraplegia	No		
	Yes		•
Renal disease	No		
•	Yes		
Any malignancy w lymphoma and leukemia w/o	No		
malignant neoplasm of skin	Yes		
Moderate or severe liver	No		
<sup>1</sup> disease	Yes		
Metastatic solid tumor	No		
	Yes		
AIDS/HIV	No		
Coronary Artery Disease	No		
	Yes		
Connective Tissue Disorder	No		
	Yes		

Moderate- Severe Renal Disease	No Yes						
ANY Follow- up Screening	Yes						
	No						
Follow-up Biopsy	Yes						
	No						
Follow-up Abnormal	Yes						
PSA	No						
Follow-up PSA/ PC Screening (not abnormal PSA) NEW CODES+ CPT CODE	Yes						
	No						
	*: For categorical variables, p-values were based on Chi-squared test with exact p-value from Monte Carlo simulation; for continuous variable, p-value was based on Wilcoxon rank sum test.						

Note: For continuous variable, median+/-IQR were reported.

Table 3: Descriptive table of patients' characteristics, risk factors, comorbidity score, index of comorbidities and outcomes by insurance status

Variable	Level	Total (N =)	Self-Pay (N =)	NOT Self- Pay (N =)	P-value*				
Patients' characteristics at the time of Initial PC Screening									
Abnormal PSA									

Age	Yearly			
Age Categories	40-55			
ı	55-69			I
	70 - 75			
Year of prostate cancer screening	2011			
	2012			•
	2013			•
	2014			•
	2015			•
Risk factors at the	time of Initial	PC Screening		
Smoking/Tobacco	No			
'	Yes			•
Morbid Obesity	No			
ı	Yes			I
Family History of Prostate Cancer	No			
1	Yes			
	Yes			•
Personal History of Irradiation	No			
	Yes			

Comorbidities' score at the time of Initial PC Screening							
Elixhauser Score							
Charlson Score							
Elixhauser Comorl	bidities at the t	time of Initial	PC Screening				
Congestive heart failure	No						
'	Yes				•		
Cardiac arrhythmias	No						
,	Yes				1		
Valvular disease	No						
I	Yes				1		
Pulmonary circulation disease	No						
I	Yes				I		
Peripheral vascular disease	No						
•	Yes				1		
Hypertension, uncomplicated	No						
<b></b>	Yes				'		
Hypertension, complicated	No						
-	Yes						
Paralysis	No						

	Yes		
Other neurological disorders	No		
	Yes		
Chronic pulmonary disease	No		
	Yes		
Diabetes w/o chronic	No		
complications	Yes		
Diabetes w/ chronic complications	No		
	Yes		
Hypothyroidism	No		
	Yes		
Renal failure	No		
	Yes		
Liver disease	No		
	Yes		
Peptic ulcer Disease, excluding bleeding	No		
	Yes		

HIV and AIDS	No		
Lymphoma	No		
1	Yes		I
Metastatic cancer	No		
ı	Yes		1
Solid tumor w/out metastasis	No		
1	Yes		'
Rheumatoid arthritis/collagen	No		
<sup>1</sup> vascular d	Yes		•
Coagulopathy	No		
1	Yes		1
Obesity	No		
1	Yes		1
Weight loss	No		
	Yes		'
Fluid and electrolyte	No		
disorders	Yes		
Blood loss anemia	No		
	Yes		

Deficiency Anemia	No				
	V		li .	li e	
	Yes				
Alcohol abuse	No				
	Yes				
Drug abuse	No				
•	Yes				'
Psychoses	No				
•	Yes				
Depression	No				
•	Yes				
Charlson Comorbio	dities at the tir	ne of Initial Po	C Screening		
Congestive heart failure	No				
•	Yes				
Chronic pulmonary disease	No				
I	Yes				'
Diabetes w/o chronic	No				
complications	Yes				
Diabetes w/ chronic complications	No				

	Yes		
Myocardial infarction	No		
	Yes		_
Peripheral vascular disorder	No		
	Yes		
Cerebrovascular disease	No		
	Yes		
Dementia	No		
	Yes		
Rheumatic disease	No		
	Yes		
Peptic ulcer disease	No		
	Yes		
Mild liver disease	No		
	Yes		
Hemiplegia or paraplegia	No		
	Yes		
Renal disease	No		
	Yes		

Any malignancy w lymphoma and leukemia w/o malignant neoplasm of skin	No Yes		
Moderate or severe liver disease	No		
'	Yes		'
Metastatic solid tumor	No		
'	Yes		'
AIDS/HIV	No		
Coronary Artery Disease	No		
	Yes		
Connective Tissue Disorder	No		
	Yes		
Moderate-Severe Renal Disease	No		
	Yes		
ANY Follow-up Screening	Yes		
	No		
Follow-up Biopsy	Yes		
	No		

Follow-up Abnormal PSA	Yes		
	No		
Follow-up PSA/ PC Screening (not abnormal PSA) NEW CODES+ CPT CODE	Yes		
	No		

<sup>\*:</sup> For categorical variables, p-values were based on Chi-squared test with exact p-value from Monte Carlo simulation; for continuous variable, p-value was based on Wilcoxon rank sum test.

Note: For continuous variable, median+/-IQR were reported.

Table 4: Descriptive table of patients' characteristics, risk factors, comorbidity score, index of comorbidities and outcomes by ethnicity

NOT Hispanic (N Hispanic (N **Total** (N =)Variable Level P-value\* =) =) Patients' characteristics at the time of Initial PC Screening Abnormal PSA Yearly Age 40-55 Age Categories 55-69 70 - 75 2011 Year of prostate cancer screening 2012 2013

	2014								
	2015								
Risk factors at the time of Initial PC Screening									
Smoking/Tobacc o	No								
	Yes								
Morbid Obesity	No								
ı	Yes				'				
Family History of Prostate Cancer	No								
	Yes								
	Yes								
Personal History of Irradiation	No								
	Yes								
Comorbidities' sc	ore at the time	e of Initial PC	Screening						
Elixhauser Score									
Charlson Score									
Elixhauser Comorbidities at the time of Initial PC Screening									
Congestive heart failure	No								
	Yes								
	No								

Cardiac arrhythmias	Yes		
Valvular disease	No		
•	Yes		•
Pulmonary circulation	No		
<sup>1</sup> disease	Yes		
Peripheral vascular disease	No		
•	Yes		•
Hypertension, uncomplicated	No		
	Yes		
Hypertension, complicated	No		
	Yes		
Paralysis	No		
	Yes		
Other neurological	No		
disorders	Yes		
Chronic pulmonary	No		
disease	Yes		•
	No		

	I		ĺ
Diabetes w/o chronic complications	Yes		
Diabetes w/ chronic	No		
complications	Yes		•
Hypothyroidism	No		
•	Yes		•
Renal failure	No		
•	Yes		•
Liver disease	No		
•	Yes		•
Peptic ulcer Disease, excluding	No		
bleeding	Yes		
HIV and AIDS	No		
Lymphoma	No		
•	Yes		•
Metastatic cancer	No		
•	Yes		•
Solid tumor w/out metastasis	No		
	Yes		

Rheumatoid arthritis/collagen	No		
<sup>1</sup> vascular d	Yes		•
Coagulopathy	No		
•	Yes		
Obesity	No		
•	Yes		'
Weight loss	No		
•	Yes		•
Fluid and electrolyte disorders	No		
disorders	Yes		
Blood loss anemia	No		
•	Yes		
Deficiency Anemia	No		
•	Yes		
Alcohol abuse	No		
•	Yes		•
Drug abuse	No		
•	Yes		•
Psychoses	No		

	Yes				
Depression	No				
	Yes				
Charlson Comor	bidities at the t	time of Initial	PC Screening	<u>I</u>	
Congestive heart failure	No				
	Yes				
Chronic pulmonary disease	No				
	Yes				'
Diabetes w/o chronic complications	No				
	Yes				
Diabetes w/ chronic complications	No				
	Yes				
Myocardial infarction	No				
	Yes				
Peripheral vascular disorder	No				
	Yes				
Cerebrovascular disease	No				
	Yes				-

Dementia	No		
1	Yes		
Rheumatic disease	No		
	Yes		•
Peptic ulcer disease	No		
	Yes		
Mild liver disease	No		
	Yes		•
Hemiplegia or paraplegia	No		
	Yes		
Renal disease	No		
1	Yes		•
Any malignancy w lymphoma and leukemia w/o malignant neoplasm of skin	No		
	Yes		
Moderate or severe liver disease	No		
	Yes		
Metastatic solid tumor	No		
	Yes		

AIDS/HIV		No						
Coronary Artery Disease		No						
		Yes						-
Connective Tissue Disorder		No						
		Yes						
Moderate- Severe Renal Disease	No							
	Yes	S						
ANY Follow- up Screening	Yes	es						
	No							
Follow-up Biopsy	Yes	S						
	No							
Follow-up Abnormal PSA	Yes	S						
	No							
Follow-up PSA/ PC Screening (not abnormal PSA) NEW CODES+ CPT CODE	Yes	S						
	No							
*· For categorica	1 var	iables n-value	25 V	vere based on	Chi-squared	l test wi	th exact n-value f	om Monte Carlo

<sup>\*:</sup> For categorical variables, p-values were based on Chi-squared test with exact p-value from Monte Carlo simulation; for continuous variable, p-value was based on Wilcoxon rank sum test.

Note: For continuous variable, median+/-IQR were reported.