Feinberg School of Medicine MP21 #18-4062

Black Race Predicts Significant Prostate Cancer Independent of Clinical Setting and Clinical and Socioeconomic Risk Factors

Oluwarotimi S. Nettey MD, MHS¹, Austin J. Walker BS¹, Mary Kate Keeter MPH¹, Aishwarya Nugooru¹, Iman C. Martin PhD², Maria Ruden MS³, Pooja Gogana BS¹, Michael A. Dixon BS¹, Tijani Osuma MD⁴, Courtney M.P. Hollowell MD⁵, Roohollah Sharifi MD^{6,7}, Marin Sekosan MD⁹, Ximing Yang MD, PhD¹⁰, William J. Catalona MD¹, Joshua J. Meeks MD, PhD¹, Andre Kajdacsy-Balla MD, PhD⁸, Virgilia Macias MD⁸, Rick A. Kittles PhD¹¹, Adam B. Murphy MD, MSCI^{1,6}

¹Department of Urology, Northwestern University Feinberg School of Medicine, Chicago, IL, ²Community Oncology and Prevention Trials Research Group, National Cancer Institute, Bethesda, MD, ³Department of Medicine, University of Illinois at Chicago, IL, ⁴Ross University School of Medicine, Miramar, FL, ⁵Division of Urology, Cook County Health and Hospitals System, Chicago, IL, ⁶Section of Urology, Jesse Brown VA Medical Center, Chicago, IL, ⁹Department of Medicine, Chicago, IL, ⁸Department of Pathology, University of Illinois at Chicago, IL, ¹⁰Department of Pathology, Northwestern University Feinberg School of Medicine, Chicago, IL, ¹¹Division of Health Equities, Department of Population Sciences, City of Hope Cancer Center, Duarte, CA

Background

- Black men have 1.6 fold higher prostate cancer (PCa) incidence and 2-3 times the mortality rate compared to White men
- Studies have linked Black race to PCa risk but most fail to account for established risk factors such as 5-ARI use, prostate volume, socioeconomic status, and clinical setting

Research Objectives

- To assess whether Black race independently predicts overall and significant Pca diagnosis on initial biopsy when controlling for established clinical, behavioral and socioeconomic risk factors, and hospital funding type in a multi-racial cohort
- To examine changes in the effect size of Black race in men ages 40-54, who are excluded from US Preventive Services Task Force (USPSTF) PCa screening recommendations

Methods

- Recruited 564 men over age 40 undergoing initial prostate biopsy for abnormal PSA or digital rectal examination (DRE) from three publicly funded and two private hospitals in Chicago from 2009-2014
- Genetic West African ancestry (WAA) estimated using panel of 105 ancestry informative markers
- Multivariate analyses examined the associations between clinical setting, race, WAA, clinical and sociodemographic risk factors, PCa diagnosis and Gleason ≥3+4 PCa
- Subgroup analysis performed for men age 40-54

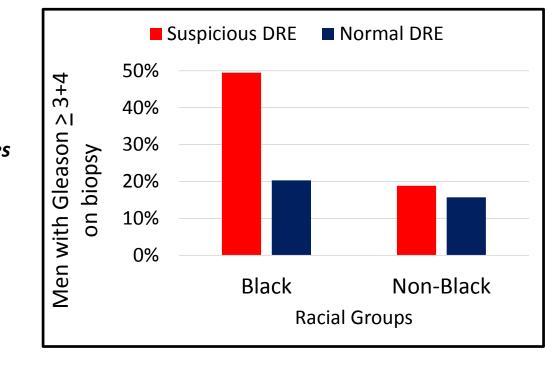
	Black	Non-Black	
	(N =287)	(N =277)	p value
	N (%)	N (%)	
Biopsy Outcomes			
Cancer on Biopsy	181 (63.1)	115 (41.5)	<0.001
<u>></u> Gleason 3+4	86 (47.5)	46 (40.0)	<0.001
Selection Selection	26 (14.4)	11 (9.6)	0.02

Figure 1: Race and abnormal rectal exam and frequency of Gleason >3=4 PCa

Results

- Black men had higher median PSA (8.1 vs 5.6 ng/ml), PSAD (0.22 v 0.15 ng/ml/cm³) compared to non-Blacks (all p<0.05)
- Blacks had lower frequency of marriage (39.0% vs 72.2%), higher rates of poverty (61.7% vs 43.3%), were more likely to have smoked (64.8% vs 56.0%) and more likely to be recruited from public hospitals (89.2% vs 51.3%, all p<0.05)
- Blacks had increased rates of Gleason ≥3+4 PCa relative to non-Blacks in both public (27.7% vs 11.6%, p<0.001) and private (48.4% vs 21.6%, p=0.002) settings
- WAA was not predictive of overall PCa diagnosis in Blacks either as a continuous variable (p=0.71) or in quartiles (Q1-Q3, p=0.17, 0.86, 0.13 respectively)
- For men aged <55, Black race (OR 5.66, 95% CI: 1.39-23.16, p=0.02) and family history (OR 4.98, 95% CI: 1.39-17.87, p=0.01) were independently positively associated with overall PCa diagnosis

Table 1: Biopsy outcomes stratified by race



Limitations

- Central pathologic review was not performed across sites
- Referred population
- Race was self-reported, used as a proxy for genetics and environmental exposures

Multivariable Logistic Regression for Cancer on Biopsy vs. Negative Biopsy		Multivariable Logistic Regression for Gleason ≥3+4 Prostate Ca vs. Gleason 3+3/Negative Biopsy			
Covariates	Odds Ratio (95% C.I.)	p value	Covariates	Odds Ratio (95% C.I.)	p value
Black race	2.13	0.002	Black race x	2.93	0.009
	(1.33-3.40)		Abnormal DRE	(1.31-6.53)	
Abnormal DRE (yes)	1.14 (0.74-1.76)	0.57	Black race x Normal DRE ^a	1.14 (0.56-2.32)	0.72
	,		Non-Black race x Abnormal DRE ^a	0.86 (0.39-1.94)	0.72
			Non-Black race x Normal DRE ^{a (ref)}	1	-
1st degree Fam Hx	1.74	0.02	1st degree Fam Hx	1.48	0.16
(yes)	(1.07-2.83)		(yes)	(0.86-2.56)	
Age, years	1.04	0.02	Age, years	1.05	0.02
	(1.01-1.07)			(1.01-1.08)	
Log(PSA), ng/ml	4.32	<0.001	Log(PSA), ng/ml	13.09	<0.001
	(2.29-8.14)			(6.06-28.27)	
Publicly funded site	0.53	0.04	Publicly funded site	0.29	0.001
(yes)	(0.28-0.98)		(yes)	(0.14-0.60)	
High School	1.17	0.49	High School	0.84	0.55
completion (yes)	(0.74-1.84)		completion (yes) (0.48-1.48)		
Prostate volume,	0.98	<0.001	Prostate volume, cm ³ 0.98		< 0.001
cm ̃	(0.97-0.99)			(0.97-0.99)	
5-ARI use (yes, ≥6	0.38	0.004	5-ARI use (yes, ≥6	0.32	0.02
months)	(0.19-0.74)		months)	(0.12-0.85)	
Married (yes)	0.86	0.81	Married (yes)	0.61	0.06
	(0.56-1.33)			(0.36-1.03)	
Annual Income <	1.1	0.72	Annual Income <	1.15	0.63
\$30K/year	(0.67-1.78)		\$30K/year	(0.64-2.06)	

Table 3: Binary logistic regressions for Black race versus overall prostate cancer and Gleason ≥3+4 prostate cancer diagnosis

Conclusions

- Black race remains associated with PCa after adjusting for clinical setting, clinical and socioeconomic risk factors
- Black race is the strongest risk factor of PCa for men under 55 years
- West African Ancestry does not predict overall or significant PCa in models substituting WAA in place of Black race