Cognitive Test Score and 25-year Mortality Risk; Does Race Matter?

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Abstract

Objectives: Despite our knowledge of the effect of cognitive test scores on subsequent risk of mortality, few studies have compared Blacks and Whites for this association. The current study was conducted on Black-White differences in the magnitude of the association between baseline cognitive test score and all-cause mortality in a nationally representative sample of adults in the United States over 25 years. Methods: We used data from the Americans' Changing Lives Study, 1986–2011, a national prospective cohort in the U.S. The study followed 3361 adults (2205 White and 1156 Blacks), age 25 and older, for up to 25 years. The independent variable was cognitive test score measured at baseline (1986) using the 4-item version of the Short Portable Mental Status Questionnaire, treated in two different ways (as a dichotomous and as a continuous variable). The dependent variable was time to death (due to all causes) during the follow-up period. Covariates included baseline age, gender, education, income, number of chronic diseases, self-rated health, and depressive symptoms. Race (Black vs. White) was the focal effect modifier. We used a series of Cox proportional hazards models in the total sample, and by race, in the absence and presence of health variables. Results: Overall, cognitive test score predicted mortality risk. A significant interaction was found between race and baseline cognitive test score, suggesting that baseline cognitive test score has a weaker protective effect against all-cause mortality for Blacks un comparison to Whites. In race-stratified models, cognitive test scores at baseline predicted risk of all-cause mortality for Whites but not Blacks, in the absence and presence of baseline socio-economic and health variables. The results were similar regardless of how we treated baseline cognitive test scores. Conclusions: In the United States, baseline cognitive test score has a weaker protective effect against all-cause mortality over a long period of time for Blacks than Whites. The finding is in line with the Minorities Diminished Returns theory and is probably due to structural and interpersonal racism.

Keywords: African Americans, Blacks, Cognition, Cognitive test score, Death, Ethnic groups, Ethnicity, Life expectancy, Mortality, Population groups, Race, Whites

Introduction

It is well established that particular cognitive abilities decline with age, [1] and this decline is associated with morbidity [2] and mortality. [3] Longitudinal studies have shown that cognitive test scores at baseline predicts future morbidity [4] and mortality. [5] Due to the association between baseline cognitive test score and life expectancy, [3] high cognitive function is expected to protect populations against the risk for premature mortality. [6]

Despite the well-defined link between baseline cognitive function and subsequent all-cause

mortality,^[3,7-9] it is not known whether or not diverse population groups differ in this relationship, as very few studies have ever tested predictive ability of cognitive limitation on long-term mortality among racial and ethnic minority populations. Of the few studies that have been conducted among racial and ethnic minorities, most have enrolled Hispanics.^[6,10] In a study among older Mexican Americans, baseline and also 2-year decline in cognitive functioning independently predicted risk of all-cause mortality.^[6] In another study among Mexican Americans, although frailty and cognitive test score were risk factors for mortality separately, the effect





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of cognitive impairment on mortality disappeared by controlling for frailty. ^[10] Unfortunately, compared to Whites or even Hispanics, less is known about this association among Black Americans. ^[11]

Aim

To fill the gap in the literature, and to better understand whether race interferes with the role of cognitive test score on life expectancy in the United States, we conducted this study on Black-White differences in the magnitude of the protective effect of baseline cognitive test score on subsequent all-cause mortality, using a nationally representative sample. In line with the Minorities Diminished returns theory^[85-93] and also some other empirical evidence on ethnic differences in predictive role of baseline risk and protective factors on long-term outcomes,^[12-14] we hypothesized a stronger effect for Whites compared to Blacks.

Methods

Design and setting

The Americans' Changing Lives (ACL) study (ACL; 1986–2011) is a 25-year prospective longitudinal study of American adults (aged 25 years and over). This study recruited a national (representative) sample of U.S. residents. Although more information on their methods and sampling procedure is published elsewhere, we briefly explain some of the aspects of the study here. [15-17]

Ethics

The University of Michigan (UM) Ethical committee (also known as the Institutional Review Board; UM-IRB) approved the study protocol. Written informed consent was received from all participants of the ACL.

Participants and sampling

Using stratified multistage sampling, the ACL recruited a probability sample of U.S. adults who were at least 25 years old. The study participants were 3617 adults who were living in the continental U.S. The only other inclusion criteria for the ACL was being non-institutionalized. The ACL oversampled older adults (age 60 years or more) and African Americans/Blacks. Wave 1 data included 70% and 68% of the sampled households and individuals respectively.



The current analysis is only limited to Europeans/ Whites and African Americans/Blacks (analytic n = 3361 who were either White [n = 2205] or Black [n = 1,156]).

Process

All the baseline data were gathered through faceto-face interviews (first wave; year 1986). Death certificates were used to assess the date of death.

Measures

Baseline demographic factors, socio-economic characteristics, self-rated health (SRH), depressive symptoms, and chronic medical conditions were measured in 1986 to adjust for possible confounding. The main outcome was time to death (due to any causes) during a 25-year follow-up period.

Race/Ethnicity

In this study, race was the focal effect modifier, treated as a dichotomous levels: Black 1 versus White 0 (referent category). Race and ethnicity were self-identified using multiple items in wave 1 of data collection (1986). Responses of these items informed our racial/ethnic categories: "Non-Hispanic White," "Non-Hispanic Black," "Non-Hispanic Native American," "Non-Hispanic Asian," and "Hispanic." The current analysis only included two racial/ethnic groups: Non-Hispanic White and Non-Hispanic Black. Thus, this analysis does not include any Hispanic individuals.

Cognitive test score

The ACL survey adopted the 4-item Short Portable Mental Status Questionnaire (SPMSQ) for the measurement of cognitive test scores. The SPMSQ assesses the following aspects of cognition: knowledge of current events, memory, as well as the ability to perform some basic mathematical tasks. The SPMSQ is designed to measure individuals' cognitive performance from community and institutionalized samples. [18,19] The SPMSQ measure had the following four items: (a) "What is the date today – month, day, and year?" (b) "What day of the week is it?" (c) "What is the name of the president of the United States?" (d) "Subtract 3 from 20 and tell me the number you get. Then, keep subtracting



3 from this number and each new number you get, telling me the results as you go (Stop when the answer is 2 or less)." Each item was coded as 0 "correct" or 1 "incorrect," with a total cognitive score ranging between 0 and 4, with a higher score indicating worse cognitive performance. [20] A score of 2 or more (participants who had 2+ errors) was considered a low cognitive score. [21,22]

Demographics

Demographic characteristics included age (a continuous variable) and gender (a dichotomous variable with male as the referent category) collected in 1986.

Socioeconomic status (SES)

This study collected baseline educational attainment (years of schooling) and income level (an ordinal variable) collected in 1986 as the SES measures.

Number of chronic diseases

The baseline number of chronic medical conditions was measured based on self-reported data. Individuals were asked if a health-care provider had ever informed that they had any of the following seven chronic conditions: Diabetes, hypertension, chronic lung disease, stroke, heart disease, cancer, and arthritis. Responses for whether the individual had a chronic conditions was on a dichotomous scale (yes/no) for each condition. The number of chronic diseases was an interval variable ranging from 0 to 7. [16,17]

SRH

Participants' SRH was measured using a single item with the following five categories: Excellent, very good, good, fair, and poor. In this study, SRH was treated as a dichotomous measure. The five categories were collapsed into the following two categories: Excellent/very good/good 0 versus fair/poor 1. This cutoff is compliant with the literature. The single item SRH measure has shown very high test-retest reliability as well as validity. Poor SRH is a strong predictor of mortality risk net of confounders. The single item SRH measure has shown very high test-retest reliability as well as validity.

Depressive symptoms

Symptoms of depression were captured using the 11-item Center for Epidemiological Studies-Depression scale (CES-D).^[27] CES-D measures the

degree to which the participant felt depressed, sad, lonely, restless, etc. Some of the CES-D items are positively worded, which are reverse-coded. Short versions of the CES-D scale have acceptable reliability and validity. [28-30] Items are on a response scale ranging from 1 to 3. As a result, the CES-D score in this study ranged from 11 to 33, with a higher score indicated more depressive symptoms.

Mortality

Mortality data were primarily gathered either from death certificates or from the National Death Index (NDI). Overall, 1402 deceased participants were detected, while 1959 individuals survived. Collected death data included the date of death, as well as primary and underlying causes of death. In the United States, death certificates are filled out by doctors as soon as possible after individuals are deceased. The cause of death was not used for this study. For the 1402 deceased cases, the month of death was obtained from the NDI or death certificates. [12]

Statistical analysis

Due to the survey design, we used Stata-13 (Stata Corp., College Station, Texas, United States) to calculate design-based standard errors (SE) based on sampling and non-response weights at baseline. Taylor series linearization was used to estimate SEs. Survey linear, logistic, and Cox proportional hazards models were used for data analysis. Hazard ratios (HR), Standardized regression coefficient (B), and odds ratios (OR) with their SEs and 95% confidence intervals are reported.

In the first step, and to test the cross-sectional association between baseline age, gender, education, income, number of chronic diseases, SRH, and symptoms of depression with baseline cognitive test score, we used three logistic regression models (Model 1-3) where the outcomes was cognitive test score treated as a dichotomous measure. For the same aim, we also ran three linear regressions (Model 4-6), where the outcome was cognitive test score treated as a continuous measure. First, we used the pooled sample with the main effect of race in the model (Model 1-4). Then, we ran models that were stratified by race/ethnicity to test if demographic, socio-economic, and health factors predict cognitive test scores (a continuous score) at baseline among Whites (Model 2-5) and Blacks (Model 3-6).



In the next step, and to assess the effect of cognitive test scores on mortality, we used a series of Cox proportional hazards models. Baseline cognitive test score, the independent variable, was measured in 1986, and was treated in two ways: A dichotomous and a continuous measure. The dependent variable was time to death for any cause between 1986 and 2011. Covariates included age, gender, education, income, number of chronic diseases, SRH, and symptoms of depression, which were entered into the model in a step-wise fashion. First, we used the pooled sample of Blacks and White to evaluate the effects of baseline cognitive test scores on mortality, while covariates were being added gradually. Then, we ran models stratified by race to test if a cognitive test score at baseline differently predicts time to death over up to 25 years of follow-up among Whites and Blacks.

Results

Table 1 presents a summary of descriptive statistics for all study variables overall and based on race. Age and gender were not significantly different between racial groups. Blacks had significantly lower education and income and worse SRH, and displayed greater depressive symptoms and more chronic medical conditions. Blacks also had higher cognitive test scores at baseline than Whites.

Table 2 shows three logistic and three linear regressions with binary and continuous cognitive test scores as the outcomes. With a similar pattern among Whites and Blacks, age, education, income, and depressive symptoms, but not gender and number of chronic medical conditions, was associated with baseline cognitive impairment. Whites and Blacks, however, showed a different patterns of associations between SRH and depressive symptoms with cognitive test scores at baseline.

Table 3 summarizes the results of several proportional hazards models that were performed in the pooled sample (Models 1-3, 8-10), as well as Whites (Model 4, 5, 11, 12) and Blacks (Model 6, 7, 13, 14), in the absence of health conditions in the model. According to this table, in the pooled sample, higher baseline cognitive test score was correlated with higher mortality risk. There was also a significant interaction between race and baseline cognitive test scores on the outcome, suggesting that the effect of baseline cognitive test score on mortality is stronger for

Table 1: Descriptive statistics for the analytic sample, stratified by race, and overall

Characteristics	All	95% CI	Whites	95% CI	Blacks	95% CI
	Mean (SE)		Mean (SE)		Mean (SE)	
Age	47.77 (0.534)	46.69–48.84	47.96 (0.601)	46.75–49.17	46.33 (0.717)	44.89–47.78
Education*	12.53 (0.096)	12.34–12.73	12.69 (0.105)	12.48-12.90	11.37 (0.233)	1.90-11.84
Income*	5.41 (0.093)	5.22-5.60	5.57 (0.101)	5.36-5.77	4.25 (0.183)	3.88-4.62
CMC*	0.79 (0.028)	0.74-0.85	0.78 (0.031)	0.71-0.84	0.91 (0.052)	0.81-1.02
SRH*						
Good-Excellent	85.06 (0.008)	83.33–86.64	85.97 (0.009)	84.15-87.60	78.38 (0.017)	74.68–81.68
Poor-Fair	14.94 (0.008)	13.36–16.67	14.03 (0.009)	12.40-15.85	21.62 (0.017)	18.32-25.32
CES-D*	-0.03 (0.025)	-0.08-0.02	-0.07 (0.025)	-0.130.02	0.28 (0.051)	0.18-0.38
Cognitive test score *	0.70 (0.03)	0.63-0.76	0.65 (0.04)	0.58-0.73	1.01 (0.05)	0.91-1.11
	% (SE)	95% CI	% (SE)	95% CI	% (SE)	95% CI
Gender						
Male	47.26 (0.012)	44.86–49.68	47.82 (0.013)	45.12-5.52	43.18 (0.022)	38.79–47.69
Female	52.74 (0.012)	5.32-55.14	52.18 (0.013)	49.48–54.88	56.82 (0.022)	52.31–61.21
Cognitive test score*						
Intact	84.37 (0.01)	82.29–86.25	86.14 (0.01)	83.77–88.21	71.38 (0.02)	67.53-74.95
Impaired	15.63 (0.01)	13.75–17.71	13.86 (0.01)	11.79–16.23	28.62 (0.02)	25.05-32.47



^{*}P<0.05 for all comparisons between Blacks and Whites; CMC: Chronic Medical Conditions; CES-D: Center for Epidemiologic Studies Depression. CI: Confidence interval, SRH: Self-rated health, SE: Standard error

Table 2: Predictors of baseline cognition impairment based on race using logistic and linear regression

Characteristics	All		Whites		Blacks	
	95% CI OR (SE)		95% CI OR (SE)		95% CI OR (SE)	
	Model 1		Model 2		Model 3	
Cognitive function as a continuous measure						
Race (Black)	2.01 (0.27)***	1.52-2.64	-	-	-	-
Gender (Female)	0.87 (0.10)	0.69-1.10	0.81 (0.12)	0.60-1.10	1.16 (0.22)	0.79–1.71
Age	1.01 (0.00)***	1.01-1.02	1.01 (0.00)***	1.01-1.02	1.02 (0.01)*	1.00-1.03
Education	0.49 (0.09)***	0.34-0.70	0.46 (0.10)***	0.30-0.70	0.63 (0.15)#	0.38-1.03
Income	0.92 (0.02)**	0.88-0.97	0.92 (0.03)**	0.87-0.98	0.93 (0.04)	0.86-1.02
СМС	1.01 (0.07)	0.88-1.15	1.00 (0.08)	0.85-1.16	1.08 (0.11)	0.87-1.33
SRH (Poor/Fair)	1.18 (0.18)	0.86-1.61	1.10 (0.22)	0.74-1.64	1.47 (0.30)#	0.98-2.21
CESD-11	1.14 (0.07)*	1.99-1.31	1.14 (.08)#	0.99-1.31	1.15 (0.12)	0.93-1.42
Cognitive	B (SE)	95% CI	B (SE)	95% CI	B (SE)	95% CI
function as a dichotomous measure	Model 4		Model 5		Model 6	
Race (Black)	0.24 (0.06)***	0.12-0.36	-	-	-	-
Gender (Female)	-0.01 (0.04)	-0.09-0.07	-0.03 (0.05)	-0.12-0.06	0.16 (0.07)*	0.01-0.30
Age	0.01 (0.00)***	0.00-0.01	0.01 (0.00)**	0.00-0.01	0.01 (0.00)**	0.00-0.02
Education	-0.38 (0.07)***	-0.520.24	-0.38 (0.08)***	-0.540.22	-0.34 (0.10)**	-0.54 - -0.13
Income	-0.02 (0.01)***	-0.040.01	-0.02 (0.01)**	-0.040.01	-0.04 (0.02)*	-0.07-0.00
СМС	0.00 (0.02)	-0.05-0.04	-0.01 (0.02)	-0.050.04	0.01 (0.05)	-0.09-0.10
SRH (Poor/Fair)	0.09 (0.05)	-0.01-0.19	0.05 (0.06)	-0.07-0.16	0.30 (0.08)***	0.14-0.46
CESD-11	0.06 (0.02)**	0.02-0.09	0.05 (0.02)**	0.01-0.09	0.09 (0.04)*	0.01-0.17

P<0.1, *P<0.05, **P<0.01, ***P<0.001, CMC: Chronic Medical Conditions; CES-D: Center for Epidemiologic Studies Depression. CI: Confidence interval, SRH: Self-rated health, SE: Standard error, OR: Odds ratios

Whites than Blacks, when demographic and baseline socio-economic factors are controlled (Model 4, 5, 11, 12). Race-specific models showed that baseline cognitive test score only predicts mortality among Whites but not Blacks.

Table 4 provides the results of Cox proportional hazards models in the pooled sample (Models 1,2,5,6) and among Whites (Models 3, 7) and Blacks (Models 4, 8), while health variables are also controlled. In this table, in the overall (pooled) sample, a statistically significant interaction effect was found between race and baseline cognitive test scores suggesting a stronger effect of baseline cognitive test score on mortality for Whites relative to Blacks (Models 2,

6). Race-specific models confirmed our finding, as cognitive test scores at baseline predicted mortality among Whites but not Blacks.

Discussion

Our study had three main findings. On the main aim of the study, major Black-White differences were found in the predictive role of baseline cognitive function on long-term mortality. The study also showed lower cognitive function at baseline for Blacks, and Black and White similarities in role of age and socio-economic status on cognitive function. These findings contribute to the existing literature on health disparities in the United States,





Table 3: Effect of baseline cognition test score on all-cause mortality based on race using Cox proportional hazard models in the absence of health variables in the model

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Characteristics	HR (SE)	%56	HR (SE)	826	HR (SE)	%56	HR (SE)	95% CI	HR (SE)	%56	HR (SE)	95% CI	HR (SE)	%56
		ວ		<u></u>	All	ō				ວ		Blacks		ວ
	Model 1		Model 2		Model 3		Model 4		Model 5		Model 6		Model 7	
Race (Black)	1.35	1.18–	1.15 (0.09)#	0.99-	1.30 (0.13)*	1.07–	1	ı	ı		ı	ı	ı	1
Gender (Female)	0.60 (0.04)***	0.53-	0.56	0.49-	0.56	0.49-	0.58	0.51-	0.54	0.47-	0.75	0.61-	0.67	0.54-
Age	1.09	1.09-	1.09	1.08-	1.09	1.08-	1.10 (0.00)***	1.09-	1.09	1.08-	1.08	1.07–	1.07	1.06-
Education			0.82	0.72-	0.82	0.72-			0.81	0.70-			0.88 (0.11)	0.68-
Income			0.92	0.89-	0.92	0.89-			0.92	0.89-			0.91	0.86-
Cognition impairment as a dichotomous measure	0.73	0.54-	0.84	0.65-	0.76	0.55-	0.66	0.43-	0.77	0.55-	0.98 (0.10)	0.75-	0.93 (0.10)	0.84-
Cognition impairment × Black	1	1	1	ı	1.30 (0.10)*	1.07–		1	1	ı	1	1	1	ı
	Model 8		Model 9		Model 10		Model 11		Model 12		Model 13		Model 14	
Race (Black)	1.33 (0.09)***	1.15-	1.14 (0.09)#	0.98-	1.40 (0.16)**	1.11–	1	ı	1		ı	1	1	ı
Gender (Female)	0.60 (0.04)***	0.53-	0.55	0.49-	0.56	0.49-	0.58	0.50-	0.54	0.47-	0.74	0.60-	0.67	0.54-
Age	1.09	1.09-	1.09	1.08-	1.09	1.08-	1.10 (0.00)***	1.09-	1.09	1.08-	1.08	1.07–	1.07 (0.00)***	1.06-
Education			0.83	0.73-	0.83	0.73-			0.82	0.71-			0.89 (0.11)	0.68-
Income			0.92	0.89-	0.92	0.89-			0.92	0.89-			0.91	0.86-
Cognitive test score as a continuous measure	0.87	0.83	0.92	-98.0 0.98	0.88	0.81-	0.83	0.76-	0.88	0.81-	0.97 (0.04)	0.88-	1.02 (0.04)	0.94-
Cognitive test score × Black		'		,	1.15 (0.05)**	1.05-		1	ı	ı		ı	,	
C C C X X X L C C X X X X X X X X X X X			The second second							1		;		

*P<0.05, **P<0.01, ***P<0.001, CMC: Chronic medical conditions; CES-D: Center for Epidemiologic studies depression. CI: Confidence interval, SE: Standard error

Table 4: Effect of baseline cognition impairment on all-cause mortality based on race using Cox proportional hazard models while health is controlled

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Characteristics	HR (SE)	95% CI	HR (SE)	95% CI	95% CI	HR (SE)	12 % S6	HR (SE)
			AII				Blacks	
	Model 1		Model 2		Model 3		Model 4	
Race (Black)	1.08 (0.08)	0.92-1.24	1.27 (0.12)*	1.04-1.55	1		1	
Gender (Female)	0.54 (0.04)***	0.47-0.62	0.54 (0.04)***	0.47-0.62	0.52 (0.04)***	0.45-0.60	0.66 (0.08)**	0.52-0.83
Age	1.08 (0.00)***	1.08–1.09	1.08 (0.00)***	1.08-1.09	1.09 (0.00)***	1.08–1.09	1.06 (0.01)***	1.05-1.07
Education	0.91 (0.06)	0.80-1.03	0.91 (0.06)	0.80-1.03	0.92 (0.06)	0.79-1.04	0.92 (0.12)	0.70-1.19
Income	0.93 (0.01)***	0.91–0.96	0.93 (0.01)***	0.91–0.96	0.94 (0.01)	0.91–0.97	0.91 (0.03)**	0.86-0.97
CMC	1.15 (0.02)***	1.10–1.20	1.15 (0.03)***	1.10–1.20	1.15 (0.03)***	1.10–1.21	1.13 (0.05)*	1.02-1.24
SRH (Poor/Fair)	1.49 (0.11)***	1.28–1.74	1.51 (0.11)***	1.30–1.76	1.59 (0.13)***	1.35–1.88	1.14 (0.11)	0.93-1.38
CESD-11	1.04 (0.04)	0.97-1.12	1.04 (0.04)	0.97-1.12	1.06 (0.05)	0.97–1.15	0.95 (0.04)	0.87-1.04
Cognition impairment as a dichotomous measure	0.86 (0.09)#	0.67–1.11	0.75 (0.10)**	0.54-0.93	0.76 (0.10)***	0.54-0.94	1.10 (0.10)	0.86-1.29
Cognition impairment × Black	-	-	1.39 (0.09)***	1.18–1.54	-	-	-	-
	Model 5		Model 6		Model 7		Model 8	
Race (Black)	1.06 (0.08)	0.90-1.24	1.39 (0.15)**	1.11-1.74	1	-	ı	-
Gender (Female)	0.54 (0.04)***	0.47-0.61	0.54 (0.04)***	0.47–0.62	0.52 (0.04)***	0.45-0.60	0.66 (0.08)**	0.52-0.83
Age	1.08 (0.00)***	1.08-1.09	1.08 (0.00)***	1.08-1.09	1.09 (0.00)***	1.08–1.09	1.06 (0.01)***	1.05-1.07
Education	0.92 (0.06)	0.81–1.04	0.92 (0.06)	0.81–1.04	0.92 (0.06)	0.80–1.06	0.92 (0.12)	0.70-1.19
Income	0.93 (0.01)***	0.91–0.96	0.93 (0.01)***	96.0-06.0	0.93(0.02)	0.90-0.97	0.91 (0.03)**	96.0–98.0
CMC	1.15 (0.02)***	1.10–1.20	1.15 (0.03)***	1.10–1.20	1.15 (0.03)***	1.10–1.21	1.12 (0.05)*	1.02-1.24
SRH (Poor/Fair)	1.49 (0.11)***	1.27–1.73	1.51 (0.11)***	1.29–1.76	1.58 (0.13)***	1.34–1.87	1.14 (0.11)	0.94-1.39
CESD-11	1.04 (0.04)	0.97-1.12	1.04 (0.04)	0.97-1.12	1.06 (0.05)	0.97–1.15	0.95 (0.04)	0.87-1.03
Cognition test score as a continuous measure	0.93 (0.03)*	0.88-0.99	0.88 (0.03)**	0.81–0.95	0.89 (0.03)***	0.82-0.95	1.03 (0.04)	0.95-1.11
Cognition impairment × Black	ı		0.80 (0.05)***	0.72-0.90	ı	ı	ı	1

*P<0.05, **P<0.01, ***P<0.001, CMC: Chronic medical conditions; CES-D: Center for epidemiologic studies depression. CI: Confidence interval, SRH: Self-rated health, SE: Standard error



as stated by Moody-Ayers *et al.* in 2005, "efforts to understand cognitive function may enhance our understanding of Black-White disparities in health outcomes^[11]."

The Black-White variation in the predictive role of baseline cognitive score on mortality may be considered in line with some other studies suggesting that baseline characteristics fail to predict long-term mortality among Blacks. [12,13,31] In two studies in 2015, baseline SRH and depression, which predict mortality of Whites failed to predict the same outcome for Blacks.[12,13] In 2001 Ferraro and Kelley-Moore showed that although SRH as a time-varying covariate predicts mortality among Blacks, baseline SRH is not as good predictor of long-term mortality for Blacks as Whites.[31] Other studies have also shown that SRH may lose its predictive validity for mortality among individuals with lower socio-economic status. [32,33] This is again in support of our findings, given race is a strong proxy of social class and socio-economic status in US.[34]

Regarding the main effect of race on cognitive ability, a number of studies in U.S. have shown that Blacks have lower unadjusted scores on cognitive function than Whites. [11,35] This difference is largely due to lower socio-economic status [35] and also an accelerated decline in health [36] among Blacks compared to Whites. In both races, higher socio-economic status indicators such as education was associated with lower cognitive decline, a finding which is supported by the literature. [37,38] It has been shown that years of schooling is a strong predictor of cognitive reserve. [39]

Research on the association between baseline cognitive impairment, cognitive decline over time, and long-term mortality has provided mixed findings. [6] While the effect of severe cognitive impairment on mortality stays significant above and beyond the effects of demographic, socio-economic, and health status, [40-43] the long-term effect of mild cognitive impairment at baseline as a predictor of mortality may be more debatable, as shown only by some [44,45] but not all [40,46] studies. Similarly, findings are mixed regarding the effect of cognitive decline over time on subsequent mortality, which is shown by some [40] but not all [41] studies.

Among Whites, a higher risk of mortality with cognitive impairment at baseline was independent of demographics, socio-economic status, physical health, and depressive symptoms. This finding is in line with the literature. [3,5,6] The previous

population-based studies have shown that the risk of death is about 50% higher among those with low cognitive test scores at baseline.^[47-50]

Among Blacks, however, baseline cognitive test score failed to predict long-term all-cause mortality similarly in the absence or presence of baseline demographics, socio-economics, and health status. Interestingly, some, [51,52] if not all [7,53] studies, have shown higher survival rates of Blacks with Alzheimer's disease compared to their White counterparts. In a study among individuals with Alzheimer's disease, cognitive impairment had a stronger effect on mortality among Blacks than Whites, which is in contrast to our results. [53]

We showed that Black-White differences in the cognitive function - mortality link are not due to racial differences in baseline socio-economic or health status, as the interaction between race and baseline cognitive test score on mortality remained significant with potential confounders (or mediators) in the model. The exact mechanism by which baseline cognitive ability predicts mortality is still unknown. As poor health status is associated with both cognitive impairment and mortality, it may confound or mediate the cognitive ability - mortality association. Several chronic medical conditions that adversely affect cognitive test scores^[54] also increases the risk of mortality. [53,55] Longitudinal studies have shown that cognitive impairment at baseline predicts incident dementia, [56] which reduces survival. [9,55] According to the population-based studies, [57-74] individuals with Alzheimer's disease are up to 3 times more likely to die than those without the condition.^[53] Thus, studies on the link between cognition and mortality require adjustment for chronic diseases. As explained above, higher socio-economic status is another major determinant of cognitive function inside and across populations.[37-39]

The result of our study has clinical and public health implications. Cognitive impairment is a modifiable risk factor for premature mortality, and some policies and interventions such as education may improve cognitive functioning of the most atrisk populations. [51,53,75-77] For instance, the ACTIVE trial has shown significant benefits for memory, reasoning, and processing speed training at 5 and even 10 year follow-ups in cognitively intact older adults. [76,77] However, some methodological issues of the ACTIVE study should be addressed. [78] Despite this, some promising results have suggested that cognitive function can be enhanced.



[79-84] As cognitive ability plays a different role in predicting mortality of Blacks and Whites, [5,7,8] interventions that enhance cognitive reserve of the communities may differently benefit populations' survival. Design and development of community-based and clinical interventions that use cognitive maintenance/enhancement as a strategy to reduce mortality of populations will be improved by similar research-based information on the magnitude of the relationship between cognitive test score and long-term mortality across population groups. Such information may be used to decide between universal or population-specific programs that wish to promote longevity of the public through enhancement of cognitive reserve. [55]

Our findings are relevant to the complex causes the Black-White economic and health gap in the United States.^[11] We argue that it is not race or risk factors but race and risk factors that cause disparities. In a study in 2005, Moody-Ayers *et al.* measured cognitive function and functional decline among 779 Black and 4892 White community samples of older adults (age 70 or older) and showed that cognitive function might mediate the higher functional decline among Blacks.^[55]

The findings are also in line with the Black-White health paradox, defined as a disproportionately lower prevalence of a wide range of undesired subjective health outcomes, despite higher levels of objective health problems such as chronic disease among Black Americans, in comparison to White Americans. [17,64-68] Based on the Black-White health paradox, Blacks do not show some of the expected associations that are commonly found among Whites.[17,64-68] The results are also in line with the Minorities Diminished Returns theory^[85-93] and also other studies that have shown resources better serve Whites than Blacks,[12-14] and that the predictive role of baseline risk and protective factors on long-term outcomes are larger for Whites than Blacks. These patterns may be due to biases in measurement,[94] or racism.[95-99]

Our study had some limitations that need attention. First, cognitive ability is subject to change; however, this study did not model cognitive decline over time. This is particularly important because both baseline and decline in cognitive ability are shown to predict mortality. [69-72] Another limitation of this study is how we measured cognitive function, which may cause some measurement bias, which may itself depend on culture, ethnicity, or race. [35,73,74] The study did not control for specific chronic neurological (e.g.,

dementia, and Alzheimer's disease) or psychiatric disorders (e.g., major clinical depression) conditions that impair cognitive abilities. Using a nationally representative sample and having a 25 year follow-up period were, however, major strengths of this study.

Conclusions

Baseline cognitive test score predicts long-term all-cause mortality among Whites but not Blacks. Future research should explore factors that may explain why minority ethnic differences exist in the role of cognitive test score on all-cause mortality risk in the United States. It is likely that socio-economics, physical health, and mental health factors differently confound or mediate the links between baseline cognitive test scores and subsequent mortality across race. Culture, access to health care, resource utilization, pattern of comorbid conditions, and health risk behaviors are also other potential explanatory factors behind Black-White differences in the cognitive test score mortality link.

Author Contributions

Shervin Assari designed and analyzed this work and drafted the manuscript and conducted the revisions. Shervin Assari had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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