# Racial Disparity and Awareness of Disease-Risk Genetic Testing: A Nationwide Sampling Survey Analysis



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### Background:

Genetic tests for certain mutations provide opportunities to reduce cancer risk by offering preventive strategies to those with elevated risk to mitigate or even

The disparity in awareness may translate into disparities in the uptake of genetic testing, health behavior change, and disease prevention through precision and personalized medicine

### Objectives:

This study aimed to examine the race/ethnicity disparity in disease-risk genetic testing (DRGT) awareness in the United States and further identify the factors associated with disease-risk genetic testing awareness among race/ethnicity groups

### Methods:

A retrospective cross-sectional analysis was conducted using 2022 Health Information National Trends Survey(HINTS) data. HINTS is a nationally representative survey that has been administered every few years by the National Cancer Institute since 2003. The HINTS target population is all adults aged 18 or older in the civilian non-institutionalized population of the United States and it is designed to collect data on the American public's need for, access to, and use of health-related information and health-related behaviors, perceptions, and

If participants reported that they have heard of disease risk genetic testing, they were considered as having awareness of disease risk genetic testing.

Descriptive analyses were performed among the participants with and without disease-risk genetic testing awareness. Survey-weighted logistic regressions with stepwise model selection were used to identify the factors influencing awareness of disease-risk genetic testing. A significance level of 0.2 s required to allow a variable entry into and stay in the model. Akaike Information Criterion (AIC) were used to assess the model performance. Further, factors related to disease-risk genetic testing among different race/ethnicity groups were identified. Statistical analyses were performed with SAS software, version 9.4 (SAS Institute,

Carv. NC. USA).

- · Among 5968 responders, 58.5% were aware of disease-risk genetic testing . The awareness varies among different racial/ethnic groups. (Figure 1)
- Compared with non-Hispanic white participants, Hispanic, non-Hispanic African American (nH-AA), and Asian American and Pacific Islander (AAPI) participants were less likely to have DRGT awareness (adjusted-OR 0.49, 0.53, and 0.40, respectively,
- Participants with disease-risk genetic testing awareness are more likely to be younger, female, white, to have a higher education and household income, insurance, a regular provider, internet access, and a family history of cancer. Perceived genetic susceptibility, worry about cancer, and perceived progress of cancer treatment and prevention are higher among those with awareness. (All pvalue<0.05) (Table 1)
- Significant factors influencing DRGT awareness included age, race/ethnicity, education, family cancer history, and perceptions of the importance of genetics for
- Factors associated with awareness of DRGT varied among racial/ethnic groups. (Table 2) For non-Hispanic White participants, age, regular provider visits, and family cancer history were significant. (Table 2.b) Among Hispanics, education, fatalistic beliefs, and cancer worry were significant factors. (Table 2.a) nH-AA showed significance in access to the internet and family cancer history. (Table 2.c) No. statistically significant factors were observed for AAPI. (Table 2.d)

- Response rate of 2022 Health Information National Trends Survey was only 28%.
- Awareness of disease risk genetic testing is based on self-reported survey response. It does not reflect the extent of knowledge and accuracy of knowledge of disease
- This study was limited mall sample size among subgroup, especially for AAPI population. Further study focus on each racial/ethnic groups is warranted.

- Disparities in disease risk genetic testing awareness exist, particularly among AAPI and their white counterparts.
- Age, race, education, family cancer history, and perceptions of genetic and cancer prevention significantly impact DRGT awareness.
- Factors associated with awareness of DRGT differ among racial/ethnic groups.
- Addressing these differences is crucial in developing targeted promotion strategies.

Table 1 Descriptive characteristics of individuals who have awareness of disease risk genetic testing

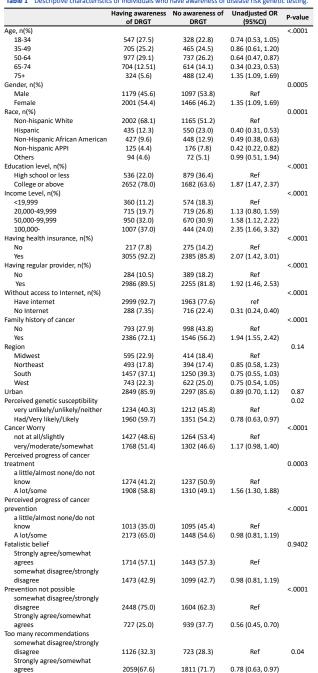


Figure 1. Awareness of disease risk genetic testing by race/ethnicity group

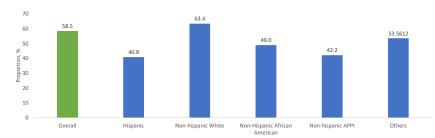


Figure 2. Associated factors of awareness of disease risk genetic testing

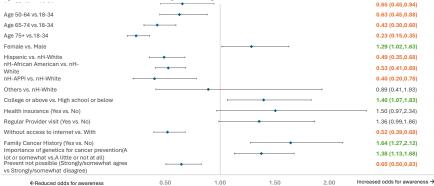


Table 2. Adjusted odds ratios for associated factors of awareness of disease risk genetic testing among race/ethnicity groups, respectively

a. Altiong mispanic participants				b. Among
	95%CI			
	aOR	Lower	Upper	
College or above education (Ref: High school or below)	2.48	1.38	4.45	Age (Ref: 18 35-49 50-64
Insurance (Yes vs. No)	1.91	0.91	4.01	65-74
Having access to internet (No vs. Yes)	0.41	0.22	0.78	75+
Urban Area (Ref: rural)	0.53	0.25	1.14	Female (Ref:
Perception Fatalistic belief (Strongly/somewhat disagree vs Strongly/somewhat agree)	1.99	1.22	3.24	College or ab Regular Prov Having acces Family Cance
Cancer Worry (not at all/slightly vs. very/moderate/somewhat)	1.93	1.02	3.64	Perception Perceived
Prevent not possible (Strongly/somewhat agree vs. Strongly/somewhat disagree)	0.73	0.44	1.21	vs. a lot or so
Prevention not possible (Strongly/somewhat agree vs Strongly/somewhat disagree)	1.45	0.91	2.32	somewhat vs Prevention vs Strongly/s

		95%CI		
	aOR	Lower	Upper	
Age (Ref: 18-34)				
35-49	0.63	0.35	1.11	
50-64	0.56	0.32	0.97	
65-74	0.37	0.22	0.60	
75+	0.17	0.10	0.30	
Female (Ref: Male)	1.24	0.93	1.65	
College or above education (Ref: High school or below)	1.33	0.92	1.94	
Regular Provider visit (Yes vs. No)	1.74	1.05	2.89	
Having access to internet (No vs. Yes)	0.68	0.45	1.02	
Family Cancer History (Yes vs. No)	1.80	1.28	2.54	
Perception				
Perceived genetic susceptible (A little or not at all	0.79	0.59	1.07	
vs. a lot or somewhat)				
Importance of genetics for cancer cure(A lot or	1.23	0.90	1.67	
somewhat vs.A little or not at all)				
Prevention not possible (Strongly/somewhat agree	0.56	0.40	0.80	
vs Strongly/somewhat disagree)				

d Among non-Hispanic AAPI participants

### c Among non-Hispanic African American participants

				a. Among non mapanio Asi i participanto			
		95%CI			95%CI		%CI
	aOR	Lower	Upper		aOR	Lower	Upper
Having access to internet (No vs. Yes)	0.32	0.15	0.70	Having health insurance (Yes vs. No)	2.48	0.48	12.84
Family Cancer History (Yes vs. No)	1.77	1.04	3.02	•			
College or above education (Ref: High school or below)	1.29	0.75	2.21	Having access to internet (No vs. Yes)	0.11	0.00	5.95
Perception				Region (Ref: Midwest)	2.32	0.36	15.18
Importance of genetics for cancer cure (A lot or	1.67	0.96	2.92	Northeast	0.50	0.14	1.87
somewhat vs. A little or not at all)				South	1.22	0.28	5.39
Importance of genetics for cancer prevention (A lot	1.70	0.92	3.13				
or somewhat vs. A little or not at all)				West	0.40	0.15	1.03

Note: The risk factors of awareness of disease risk genetic testing among race/ethnic group, respectively, were selected via survey-weighted logistic regressions with stepwise model relection with same potential variable list. The potential variable list included age, gender, race, education level, income level, having health insurance, having regular provider, access to internet, family history of cancer, region, urban, perceived genetic susceptibility, cancer worry, perceived progress of cancer treatment and prevention., fatalistic belief, prevention not possible, too many recommendation. A significance level of 0.35 is required to allow a variable entry into and stay in the model.

Abbreviation: AAPI, Asian American and Pacific Islander: aOR, adjusted Odds ratio: DRGT, Disease Risk Genetic testing: OR, odds ratio: CI, Confidence Interval: Reference: Loomans-Kropp, H. A., & Umar, A. (2019). Cancer prevention and screening: the next step in the era of precision medicine. NPJ precision oncology, 3(1), 3