Risk factors for 5-year mortality in people with HIV after cancer diagnosis (2000-2017)

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Introduction

- Studies demonstrating higher mortality following a cancer diagnosis among people with versus without HIV compels characterizing the factors driving poorer prognosis
- We estimated 5-year survival and risk factors for 5-year mortality among people with HIV (PWH) diagnosed with any cancer in North America from 2000-2017

Vethods

Study Population:

- PWH, ≥18 years old, participating in the North American A Collaboration on Research and Design (NA-ACCORD)
- Participated in cohorts collecting ICD-O-3 site/histology data
- Diagnosed with any validated cancer (except for non-mela cancer) from 2000-2017

Observation time:

Study entry: date of cancer diagnosis

Study exit, earliest of: death, administrative censoring (12/31) to-follow-up, or 5 years of follow-up after cancer diagnosis

Timescale: time since cancer diagnosis

Statistical Analysis:

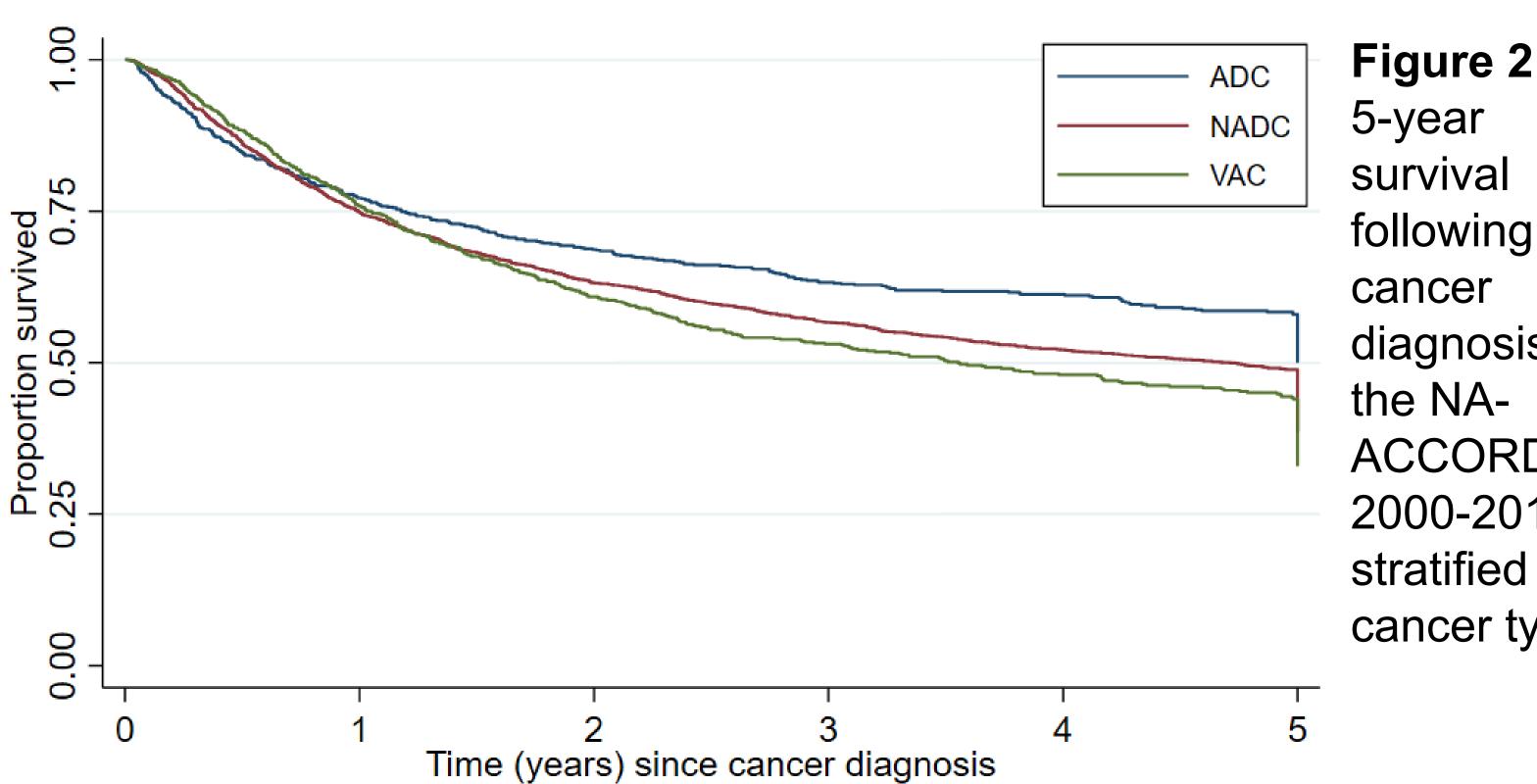
- Estimated 5-year survival by cancer type using Kaplan Meier methods
- Cancer types were categorized as:
 - AIDS-defining cancer (ADC)
 - Virally-associated non-AIDS-defining cancer (VAC): certain oral cavity/pharyngeal cancers, hepatocellular carcinoma, vulvar, penile, vaginal and anal squamous cell carcinoma, and Hodgkin's Lymphoma
 - Non-AIDS-defining cancer (NADC): all other cancer types
- Patients could contribute multiple cancer diagnoses
- Assessed association between demographic/clinical factors and 5-year all-cause mortality using Cox proportional hazards models
- Risk factors included:
 - Age at cancer diagnosis (per 5-year increase)
 - Race/ethnicity (non-Hispanic [NH] white, NH-Black, Hispanic, Other
 - Sex (male or female)
 - Viral suppression at cancer diagnosis (≥200 copies/mL vs. <200 copies/mL)
 - CD4 count at cancer diagnosis (<200 cells/ μ L, 200-350 cells/ μ L, \geq 350 cells/µL)
 - AIDS-defining illness ([ADI] clinical diagnosis or CD4 <200 cells/µL) prior to cancer diagnosis (yes, no))
 - Calendar year of cancer diagnosis (per 1-year increase)

Results

Associations between all-cause mortality and race/ethnicity, CD4 count and viral suppression merit further exploration into social, structural, and clinical/provider driven factors that may explain poor outcomes following a cancer diagnosis in people with HIV

er in North		
		AIDS-defining cance
	Age-	
	Female vs. male	
AIDS Cohort	NH Black vs. NH white -	
	Other vs. NH white-	•
lata	Hispanic vs. NH white	
anoma skin	Calendar year -	•
1/2017), loss-	CD4: 200-<350 vs. <200 cells/uL-	
	CD4: 350+ vs. <200 cells/uL-	_ _
	Virally suppressed (yes vs. no)-	
	History of AIDS-defining illness-	
		0.5 1 Adjusted Hazard Ra

- Mortality following ADCs and VACs was higher in Black vs. white patients; mortality following NADCs was lower for Hispanic vs. white patients
- For ADCs and NADCs, prior ADI was associated with increased mortality risk



There were 4556 cancer diagnoses (835 ADC, 897 VAC, 2824 NADC) in 4103 patients (12185 person-years); 5-year survival: for ADCs, 50% (95% CI 46%, 54%); for VACs, 33% (95% CI 29%, 37%); for NADCs, 39% (95% CI 36, 41%)



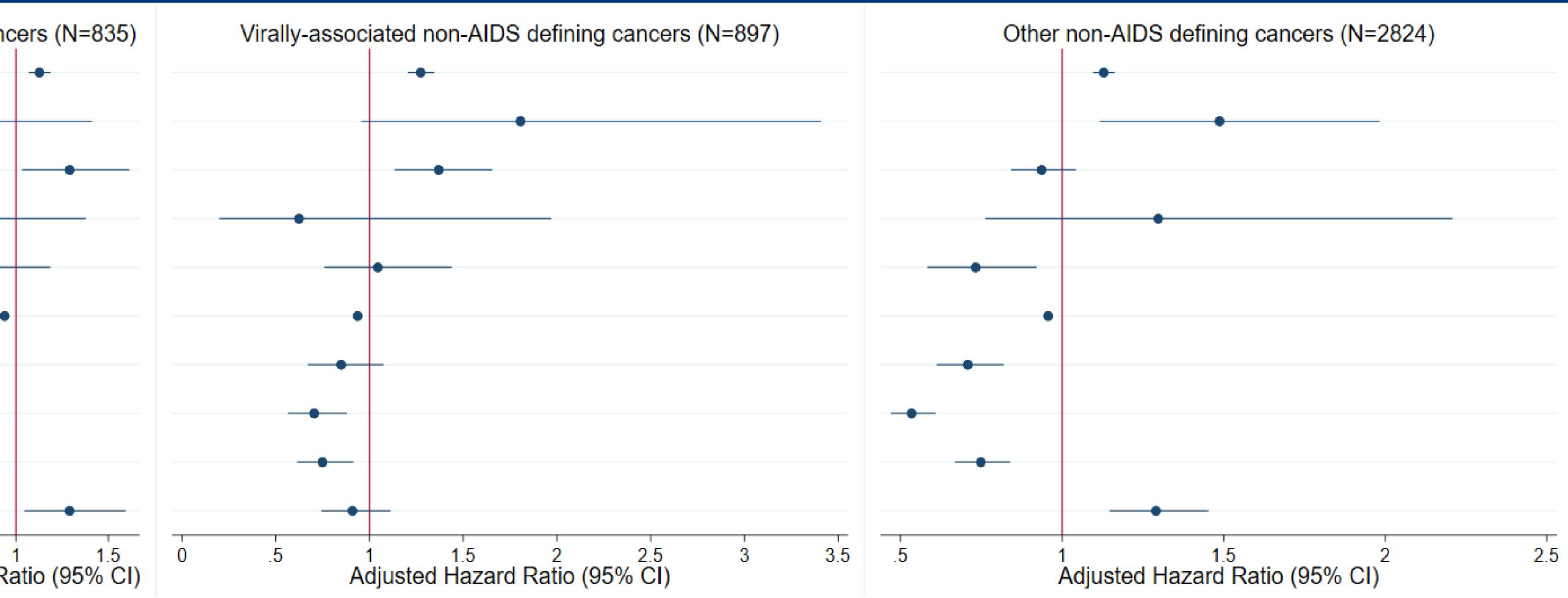


Figure 1. Risk factors for 5-year mortality following a cancer diagnosis in the NA-ACCORD, 2000-2017

survival

cancer

the NA-

ACCORD,

2000-2017

stratified by

cancer type

following a

diagnosis in

- Viral suppression and high CD4 count (≥350 cells/µL) were inversely associated with 5-year mortality for all cancer types
- Age was an independent predictor of mortality for all cancer types

Conclusions

- Lack of viral suppression and low CD4 count at cancer diagnosis increased 5-year mortality risk following cancer diagnosis for all cancer types
- Inconsistent racial disparities in mortality were observed by cancer type
- Limitations include assessment of all-cause, not cause-specific mortality
- Findings underscore importance of successful HIV treatment and exploring potential etiology for NADCs/VACs
- Future work will incorporate cancer treatment, stage, individual cancer site assessment, and longitudinal HIV viremia/ immune status

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ata were collected by cancer registries participating in the National Program of Cancer Registries (NPCR) of the Centers for Disease Control and Prevention (CDC) estitute the first sentence of Acknowledgments with: The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health or the Centers for Disease Control and Prevention

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