



The 65th ASH Annual Meeting Abstracts

POSTER ABSTRACTS

114. SICKLE CELL DISEASE, SICKLE CELL TRAIT AND OTHER HEMOGLOBINOPATHIES, EXCLUDING THALASSEMIA: CLINICAL AND EPIDEMIOLOGICAL**Comparing Racial Disparities of Cardiovascular Manifestations in Sickle Cell Trait**

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Introduction

Sickle cell trait (SCT) is a genetic condition that affects millions of people globally, particularly those of African and Mediterranean descent. Various mechanisms have been suggested to contribute to cardiovascular pathology in SCT patients, including chronic anemia, arteriopathy, and cardiac remodeling. However, whether SCT carriers, particularly African Americans (AA), are at an increased risk for cardiovascular disease remains controversial. This study aimed to investigate potential differences in cardiovascular outcomes between AA and non-AA SCT carriers.

Methods

We screened the National Inpatient Sample (NIS) database 2016-2018 for cardiovascular outcomes during hospital admission in African American SCT carriers and compared them to SCT carriers of other races. The primary outcomes were major adverse cardiovascular events, including acute myocardial infarction, cardiac arrest, stroke, acute heart failure, and in-hospital mortality. Secondary outcomes included World Health Organization (WHO) group 5 pulmonary hypertension, acute pulmonary edema, valvular disease, pericarditis, tachyarrhythmia, conduction disorders, acute deep venous thrombosis and/or pulmonary embolism, avascular necrosis, and rhabdomyolysis. Univariate analysis was conducted, with a P value <0.05 considered statistically significant.

Results

A total of 29,060 SCT admissions were analyzed, with 24,659 attributed to African Americans and the remaining to other races. The study found that majority of the patients admitted were female (86% of African American patients and 84% of patients from other races). The average age of patients was 37 years for African Americans and 40 years for other races. African American patients had higher rates of comorbidities such as hypertension, diabetes, chronic kidney disease, and oral anticoagulant use compared to patients of other races.

The study also found that African American SCT patients had a significantly higher risk of acute heart failure (5.7% vs. 3.5%, $p < 0.01$), tachyarrhythmia (4.9% vs. 3.6%, $p < 0.01$), and conduction disorders (1.6% vs. 1.1%, $p = 0.02$) compared to patients of other races. Acute myocardial infarction (1.1% vs. 0.9%), valvular disease (2.7% vs. 2.3%), acute deep vein thrombosis/pulmonary embolism (2.2% vs. 2.0%), and rhabdomyolysis (0.6% vs. 0.5%) occurred at a higher rate in African American patients but did not reach statistical significance. Similarly, in-hospital mortality (0.7% vs. 0.9%), cardiac arrest (0.4% vs. 0.5%), and pericarditis (0.1% vs. 0.2%), although occurring at lower rates in the AA population, did not reach statistical significance. Acute pulmonary edema (0.2% vs. 0.2%), pulmonary hypertension (group 5) (0.1% vs. 0.1%), stroke (1.1% vs. 1.1%) and avascular necrosis (0.1% vs. 0.1%) had similar rates in both AA and other races but did not reach statistical significance.

Conclusion/Discussion

African American patients with sickle cell trait compared to other races had higher rates of acute heart failure, tachyarrhythmias, and other conduction disorders. Aggressive screening for cardiac conditions and management of risk factors including smoking, obesity, high blood pressure, hyperlipidemia, etc., are warranted in this population to improve cardiovascular out-

comes. Although the NIS database provides this study with large patient data, this study may be limited by difference in the number of patients in AA and other race cohorts at baseline, coding error, underreporting, misclassification, retrospective nature, and lack of matching. Further studies analyzing the cardiovascular morbidity and mortality in sickle cell trait carriers, including benefit of use of medications for primary/secondary prevention of cardiovascular conditions are warranted.

Disclosures Ogu: Vertex Pharmaceuticals: Consultancy; Bluebird Bio: Consultancy; Emmaus: Speakers Bureau; Global Blood Therapeutics/Pfizer: Speakers Bureau.

Table 1a - Cardiovascular outcomes in African American SCT patients compared to other races

Outcome	AA (N=24659) *(n, %)	Other races (N=4401) *(n, %)	P value
Acute MI	266 (1.1)	38 (0.9)	0.2
Cardiac arrest	91 (0.4)	20 (0.5)	0.4
Acute Pulmonary edema	49 (0.2)	8 (0.2)	0.82
Valvular disease	662 (2.7)	100 (2.3)	0.12
Pericarditis	18 (0.1)	7 (0.2)	0.07
Acute heart failure	1403 (5.7)	152 (3.5)	<0.01
Pulmonary Hypertension type 5	17 (0.1)	6 (0.1)	0.14
Tachyarrhythmia	1219 (4.9)	159 (3.6)	<0.01
Conduction disorders	329 (1.6)	49 (1.1)	0.02
Stroke (Including TIA)	276 (1.1)	50 (1.1)	0.92
Acute DVT/PE	548 (2.2)	90 (2.0)	0.46
In-Hospital Mortality	177 (0.7)	38 (0.9)	0.3
Osteonecrosis/AVN	16 (0.1)	3 (0.1)	1
Rhabdomyolysis	153 (0.6)	20 (0.5)	0.19

AA, African American; Other races: White, Hispanic, Asian/pacific islander, Native American and others; MI, Myocardial infarction; TIA, Transient Ischemic Attack; DVT, Deep venous thrombosis; PE, Pulmonary embolism; AVN, Avascular necrosis CABG, Coronary artery bypass graft; OSA, Obstructive Sleep apnea; CIED, Cardiac implantable electronic devices; DM, Diabetes Mellitus; HTN, Hypertension; HF, Heart Failure; COPD, Chronic obstructive pulmonary disease; CKD, Chronic kidney disease; DLD, Dyslipidemia; ESRD, End stage renal disease; PVD, Peripheral vascular disease.

Table 1b - Baseline characteristics of SCT patients

Baseline characteristics	Others (4401)	AA (24659)
Female	3779 (85.9)	20592 (83.5)
Age	36.52 (15.31)	39.61 (16.38)
Prior CABG	26 (0.6)	203 (0.8)
OSA	137 (3.1)	1409 (5.7)
CIED	41 (0.9)	371 (1.5)
Oral Anticoagulation	137 (3.1)	1074 (4.4)
DM	578 (13.1)	4718 (19.1)
Antiplatelet	35 (0.8)	272 (1.1)
Aspirin	198 (4.5)	1685 (6.8)
HTN	1076 (24.4)	9376 (38.0)
Cardiomyopathy	86 (2.0)	882 (3.6)
Chronic HF	196 (4.5)	2022 (8.2)
COPD	141 (3.2)	1546 (6.3)
CKD	157 (3.6)	1461 (5.9)
DLD	352 (8.0)	2733 (11.1)
ESRD	118 (2.7)	1275 (5.2)
Carotid Artery Disease	8 (0.2)	84 (0.3)
PVD	18 (0.4)	196 (0.8)

Figure 1

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