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# Racial and Ethnic Disparities in the Usage and Outcomes of Ischemic Stroke Treatment in the United States

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# Abstract

**Objectives**—This study explores racial and ethnic differences in 1) receiving tissue plasminogen activator (tPA) and endovascular thrombectomy (EVT) as treatment for ischemic stroke and 2) outcomes and quality of care after use of tPA or EVT in the US.

**Materials and Methods**—An observational analysis of 89,035 ischemic stroke patients from the 2019 National Inpatient Sample was conducted. We performed weighted logistic regressions between race and ethnicity and 1) tPA and EVT utilization and 2) in-hospital mortality. We also performed a weighted Poisson regression between race and ethnicity and length of stay (LOS) after tPA or EVT.

**Results**—Black patients had significantly lower odds of receiving tPA and EVT than White patients and minority populations (including but not limited to Black, Hispanic, Pacific Islander, Native American, and Asian) had significantly longer hospital LOS after treatment with tPA or EVT. We failed to find a significant difference between race/ethnicity and in-hospital mortality post-tPA or EVT.

**Conclusions**—Black ischemic stroke patients were less likely to receive tPA and EVT than White patients, and among patients who received tPA or EVT, minority patients had significantly longer hospital LOSs than White patients. While we failed to find a difference in in-hospital mortality, racial and ethnic disparities are still evident in the decreased usage of tPA and EVT and longer LOSs for minority patients. This study calls for interventions to expand the utilization of tPA and EVT and advance quality of care post-tPA or EVT in order to improve stroke care for minority patients.

### Keywords

Ischemic stroke; Tissue Plasminogen Activator; Endovascular Thrombectomy; Racial and Ethnic Disparities; Ischemic Stroke Treatment

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None.

# Introduction

Every year, more than 795,000 people in the United States have a stroke as reported by the American Heart Association (AHA) [1]. Strokes are the fifth leading cause of adult death in the United States and are a leading cause of long-term disability [1]. Strokes can occur in any population, but studies have shown that there are racial and ethnic discrepancies in stroke incidence and post-stroke outcomes nationally. Black and Hispanic patients experience a higher incidence of stroke and have strokes that occur earlier in life [2]. The AHA reports that Black patients have nearly a two-fold higher risk of first-time stroke when compared to Whites [1], and the Centers for Disease Control and Prevention reports that Black patients have the highest mortality rate due to strokes [3]. Black patients are also less likely to receive tissue plasminogen activator (tPA) [4], have higher door-to-imaging times [5], and are less likely to receive endovascular thrombectomy (EVT) than White patients [6]; all of which may contribute to the higher mortality rate post-stroke in Black patients.

TPA and EVT have been shown to improve functional outcomes after ischemic stroke [7,8,9] and are both widely accepted in guidelines for treating ischemic stroke [10,11]. While the national use of tPA and EVT have increased over the years, this increase in usage is not proportionate between races as tPA and EVT use is lower in Black populations [12]. In this study, we aim to continue the exploration of racial disparities in the usage of tPA and EVT nationally as well as to investigate outcomes in ischemic stroke patients who received either tPA or EVT. We hypothesize that tPA and EVT usage will be lower in Black patients as compared to White patients. Additionally, we hypothesize that Black patients will have higher in-hospital mortality and longer hospital length of stay (LOS) after being treated with tPA or EVT than White patients.

# Methods

#### Data and sample

We used data from the 2019 National Inpatient Sample (NIS) collected by the Healthcare Cost Utilization Project (HCUP) of the Agency for Healthcare Research and Quality (AHRQ) for this observational analysis. The NIS data were de-identified, and a data user agreement was signed before we analyzed the data. The HCUP is Health Insurance Portability and Accountability Act-compliant, and therefore review by an institutional review board is not required. The 2019 NIS is the most recent dataset available for analyses. The NIS is the largest hospital discharge sample available and includes discharge records from all HCUP-participating hospitals covering more than 97% of the U.S. population. This NIS sample includes HCUP-participating community hospitals while excluding patients in prison hospitals, inpatient rehabilitation hospitals, and long-term acute care facilities. Sample weights were provided to generate nationally representative estimates.

The current study's sample includes adults aged 18 or older with a primary diagnosis of ischemic stroke who received tPA or EVT. The *International Classification of Diseases, 10th Revision, Clinical Modification* (ICD-10-CM) diagnosis and procedure codes were used

for classification. Patients discharged with ischemic stroke were identified using principal diagnosis ICD-10 codes: 163.x [13].

#### Variable selection

Our initial analysis examined the usage of tPA and EVT in various racial and ethnic groups. Utilization of TPA and EVT served as the outcome variables and race and ethnicity was the primary predictor variable. Race and ethnicity were defined by NIS as Non-Hispanic (NH) White, NH Black, Hispanic, and Other which included but was not limited to Asian, Pacific Islanders, and Native Americans.

The procedural code ICD-10-PCS 3E03317 was used to define patients who received tPA [14]. The following procedural codes were used to define patients who underwent EVT: ICD-10-PCS 03CG3Z7, 03CG3Z2, 03CG4Z6, 03CG4ZZ, 03CH3Z6, 03CH3Z7, 03CH3Z7, 03CH3Z6, 03CH4Z6, 03CH4Z2, 03CH4Z2, 03CJ3Z6, 03CJ3Z7, 03CJ3Z2, 03CJ4Z6, 03CL4ZZ, 03CK3Z7, 03CK3Z7, 03CK4Z6, 03CK4ZZ, 03CP4Z6, 03CL4ZZ, 03CQ3Z2, 03CQ4Z6, 03CQ4ZZ [12].

For our secondary analysis, we observed the outcomes after ischemic stroke in the patient subsample who received either tPA or EVT. We compared the differences in clinical outcomes, in-hospital mortality and length of stay (LOS), by using race and ethnicity as the primary predictor variable. In-hospital mortality is a binary variable with patients who died coded as 1 and patients who did not die coded as 0. LOS is a continuous variable that was defined by the NIS by subtracting the admission date from the discharge date.

Confounding variables included age, sex, neighborhood median household income [15], payers [16], calendar quarter of discharge [17], number of comorbidities [18], hospital's location (rural or urban) [19], hospital teaching status [20.21], bed size [20,21, 22] and weekend discharge [23]. Age groups were divided into 18-44, 45-64, 65-74, and 75 and older. Neighborhood median household income quartiles were defined by the 2019 NIS income ranges: \$1-24,999, \$25,000-34,999, \$35,000-44,999, \$45,000 or more. NIS categorizes "payers" as Medicare, Medicaid, private insurance, self-pay, and no charge. The discharge quarters were categorized into four quarters: January-March, April-June, July-September, and October-December. We used the Charlson Comorbidity Index to categorize the number of comorbidities into two groups: 1 comorbidity or 2 or more comorbidities. Hospital location is a binary variable categorized as rural or urban. Teaching status is a binary variable categorized as teaching and non-teaching hospital. NIS categorizes hospital size as small, medium, or large based on number of beds depending on hospital location, region, and teaching status [24]. Weekend discharge describes whether the patient was discharged on a weekday (Monday-Friday) or a weekend (Saturday or Sunday). Existing literature that establishes differences in in-hospital mortality based on patient, neighborhood, and hospital characteristics guided our choice of these confounding variables [25].

#### Statistical Analysis

We first assessed the percentage of missing values in the study sample. There was a total of 3,807 (4.1%) observations that contained missing values. Previous studies have suggested

that when an analytic sample has less than 5% missing values, performing imputations does not significantly reduce biases [26]. Therefore, with 4.1% missing values in our sample, we decided not to perform imputation but conduct the analysis in the complete sample.

In our first analysis, we examined whether there was a statistically significant difference in the rate of utilization of tPA and EVT in our sample (age 18 and primary diagnosis of ischemic stroke) for NH White, NH Black, Hispanic, and Other racial and ethnic groups (Asian, Pacific Islanders, Native Americans, and others), using Pearson Chi-Square (x2) test. We then performed weighted logistic regression to test the association between race and ethnicity and use of tPA and EVT, while controlling for sex, neighborhood median household income, payers, calendar quarter of discharge, number of comorbidities, hospital's location, hospital teaching status, rural-urban status, bed size, and weekend discharge. The odds ratios (OR) and 95% confidence intervals (95% CI) are presented below.

For the second analysis, we examined the differences in-hospital mortality within our patient subsample who received tPA or EVT across racial and ethnic groups and tested their statistical significance using the x2 test. We then performed a weighted logistic regression to test the association between race and ethnicity and in-hospital mortality, while controlling for sex, neighborhood median household income, payers, calendar quarter of discharge, number of comorbidities, hospital's location, hospital teaching status, rural-urban status, bed size, and weekend discharge. The OR and 95% CI are presented below.

For our third analysis, as LOS is a count measure, we performed a weighted Poisson regression to test the association between race and ethnicity and LOS, controlling for all the aforementioned confounding variables. The incidence rate ratios and 95% CI are presented below.

All analyses were performed using sampling weights, and Stata SE 17 was used for all statistical analyses (StataCorp, College Station, TX). One author (D.Z.) had full access to all the data in the study and takes responsibility for its integrity and the data analysis.

# Results

The full sample included 89,035 ischemic stroke patients. Of ischemic stroke patients in our sample, 68.32% were NH White, 17.35% were NH Black, 8.11% were Hispanic, and 6.21% were Asian, Pacific Islanders, Native Americans, or others. Within our ischemic stroke sample, 10.23% received tPA, 6.57% received EVT, and 4.08% died in the hospital. Additional sample demographics including age, sex, number of comorbidities, household income quartiles, insurance type, time of discharge, location, teaching status, and bed size of hospital are shown in Table 1.

The proportion of ischemic stroke patients who received tPA and EVT as categorized by race and ethnicity is shown in Figure 1 and Figure 2, respectively. NH Black patients had the lowest proportion of tPA usage at 9.22% (P < 0.001) signifying that of those who received tPA, NH Black patients made up the lowest percentage. Similarly, NH Black patients also had the lowest proportion of EVT usage at 5.76% (P < 0.001) as seen in Figure 2.

The findings from the logistic regression assessing the association between race and ethnicity and the usage of tPA and EVT are described in Table 2. In the unadjusted analysis, NH Black patients were found to have significantly lower odds of receiving tPA than NH White patients (OR=0.88, 95% CI=0.83-0.94, P=<0.001) and that Hispanic patients had significantly higher odds of receiving tPA than NH White patients (OR=1.11, 95% CI=1.03-1.20, P=0.006). When adjusting for confounding variables, NH Black patients continued to have significantly lower odds of receiving tPA than NH White patients (AOR=0.85, 95%) *CI: 0.80–0.91, P=<0.001)*, but tPA usage in Hispanic patients was no longer significantly different than NH White patients. In the unadjusted analysis, NH Black patients were found to have significantly lower odds of receiving EVT than NH White patients (AOR=0.86, 95% CI=0.80-0.93, P=<0.001) and that Asian, Pacific Islander, and Native American patients had significantly higher odds of receiving EVT than NH Whites (AOR=1.24, 95%) CI=1.12-1.37, P=<0.001). When adjusting for confounding variables, NH Black patients continued to have significantly lower odds of receiving EVT than NH White patients (AOR=0.75, 95% CI: 0.70-0.82, P=<0.001), and EVT usage in Asian, Pacific Islander, Native American patients was still significantly higher than NH White patients, though the strength of significance lessened (AOR=1.11, 95% CI=1.00-1.24, P=0.048). Overall, Table 2 demonstrates that NH Black patients were less likely to receive tPA and EVT than NH White patients after ischemic stroke.

The results on the association between race and ethnicity and in-hospital mortality are shown in Table 3. We did not find a significant racial or ethnic difference for in-hospital mortality after receiving tPA or EVT in either the unadjusted or the adjusted analyses. This demonstrates that in-hospital mortality post-ischemic stroke did not significantly differ between racial and ethnic groups among those who received tPA or EVT.

The results of the Poisson regression assessing the association between race and ethnicity and hospital LOS in patients who received tPA or EVT are shown in Table 4. Unadjusted and adjusted analyses showed that minority patients (NH Black, Hispanic, Asian, Native American, and Pacific Islander) who received tPA all had significantly longer LOSs than White patients (*Black patients: Adjusted Incidence Rate Ratio* [*AIRR*]=1.31, 95% CI: 1.21-1.41, P < 0.001, *Hispanic patients: AIRR*=1.15, 95% CI: 1.04-1.28, P=0.008, Other minority patients: AIRR=1.16, 95% CI: 1.07-1.27, P=0.001). Similarly, all minority patient groups who received EVT had significantly longer hospital LOSs than White patients within the unadjusted and adjusted analyses (*Black patients: AIRR*=1.30, 95% CI: 1.19-1.44, P < 0.001, *Hispanic patients: AIRR*=1.23, 95% CI: 1.10-1.38, P < 0.001, Other minority patients: AIRR=1.14, 95% CI: 1.14-1.44, P < 0.001). These findings demonstrate that racial and ethnic minority patients who received either tPA or EVT stayed in the hospital significantly longer than White patients.

#### Discussion

The current study compares the usage of tPA and EVT between races and ethnicities and furthers this analysis by assessing the racial and ethnic differences for in-hospital mortality and hospital length of stay after receiving tPA or EVT. In accordance with previous literature, our study demonstrates that NH Black ischemic stroke patients are less likely

to receive tPA and EVT than NH White patients. Both tPA and EVT have been shown to be effective in improving outcomes after ischemic stroke<sup>7,9</sup>, and the use of tPA and EVT has increased over the years [12,27]. However, even with the increase in usage of tPA and EVT, Black patients still receive these treatments at a lower proportion than White patients [6]. Additional research is needed to explore why the rate of use of EVT and tPA is increasing throughout America, yet the usage of EVT and tPA in Black patients is not equally increasing.

Of the patients who received tPA or EVT, we failed to find a racial or ethnic difference in inhospital mortality. Despite our insignificant finding, we did find that patients from minority populations (NH Black, Hispanic, Asian, Pacific Islander, and Native American) all had a significantly longer hospital LOSs than NH White patients. This signifies that racial and ethnic minority patients who received tPA or EVT stayed in the hospital longer than White patients. This finding is consistent with previous research which demonstrated a longer LOS for minority ischemic stroke patients during 2011–2012 [28]. This increased LOS for minority patients may indicate suboptimal quality of care or poorer clinical and functional outcomes post tPA or EVT. LOS may be used as metric of quality of care as previous studies have demonstrated that shorter length of stay for stroke patients is associated with higher quality of care [29]. The current study reveals two important disparities in minority patient ischemic stroke care: 1. less utilization of tPA and EVT in NH Black patients and 2. longer length of stay for minority patients who received tPA and EVT when compared to NH White patients.

Delays in care could be a reason why Black patients are less likely to receive tPA and EVT than White patients. TPA is most effective when given within 4.5 hours of stroke onset and EVT is most effective when given within 6 hours of stroke onset [10]. The use of emergency medical services (EMS) for transportation to the hospital is associated with shorter transport time, however Black patients are less likely to use EMS at the onset of stroke thus often delaying their care [30, 31]. Black patients who use EMS experience prolonged time from calling 911 to arriving at the hospital as compared to White patients [32]. Once Black patients arrive to the hospital, they face longer wait times<sup>33</sup> and experience longer door-to-imaging time which further delays care [5,34]. Furthermore, even for patients who meet the recommended time period, Black patients are still less likely to receive tPA than White patients [35]. Further research on why these delays occur could guide the development of interventions to improve the efficiency of care for Black ischemic stroke patients.

Limited access to hospitals that provide EVT and tPA may be another reason why Black patients are less likely to receive these treatments. There is significantly lower access to hospitals that provide tPA in rural areas [36], and we see a similar trend with EVT. EVT is performed primarily at large stroke centers that tend to be urban teaching hospitals [37]. Only 37% of stroke centers in the nation are able to perform EVT, and only 20% of Americans can be transported within 15 minutes to a hospital that provides EVT [38]. Black patients are less likely to arrive at hospitals that perform EVT procedures [37] which demonstrates limited access. Transfer to a hospital that provides EVT services can further delay care which may lessen the likelihood of Black patients receiving this treatment.

Expansion of tPA and EVT use to more regional areas may improve access to stroke care for minority patients.

Lastly, implicit biases cannot go unrecognized when considering the racial and ethnic differences in usage of tPA and EVT and LOS after these procedures. Many healthcare providers have implicit or unconscious racial biases that can impact the relationship between the patient and physician [39]. One study showed that physician implicit bias against Black patients were correlated with a decreased likelihood of recommending thrombolysis treatment for myocardial infarction to Black patients and an increased likelihood of recommending thrombolysis treatment to White patients [40]. Studies have cited that medical provider racial biases may play a role in the racial disparities present in stroke care [30]. A study from the AHA Get With The Guidelines Stroke Program demonstrated that Black stroke patients are less likely to receive patient-centered evidence-based care than White stroke patients [4]. Evidence based care can contribute to both quality of care and length of stay. The AHA also suggests that a lack of minority physicians may play a role in the racial and ethnic disparities present in stroke care [30]. When discussing the disparities in the usage and outcomes of tPA and EVT in ischemic stroke patients, we must not overlook medical providers' implicit racial and ethnic biases that impact stroke care in minority patients.

In addition to disparities in the utilization of tPA and EVT between non-Hispanic Black and non-Hispanic White patients, we did not observe significant differences in the receipt of these standard-of-care treatments for ischemic stroke among other racial and ethnic groups when compared to non-Hispanic White patients. Notably, the Other racial group that includes Asian, Pacific Islander, and Native American patients, demonstrated a higher likelihood of receiving EVT treatment, even after adjusting for potential confounding variables. Several factors may contribute to this phenomenon: First, the Other racial group is highly diverse, potentially encompassing individuals who maintain healthier lifestyles and better cardiovascular health (e.g., certain Asian groups) when compared to White patients [41]. However, it's important to acknowledge that this categorization may obscure disparities faced by specific subgroups within the Other racial group, such as Native Americans, who may experience delayed access to health care, but they are less represented in our current grouping methodology [42,43]. Second, the Other racial group has a much smaller sample size compared to the White group, which could introduce potential biases in our estimates.

#### Limitations

Our study has several key limitations. First, we did not consider stroke severity in our analysis. While coding of the National Institutes of Health Stroke Scale (NIHSS) in the NIS dataset has been increasing over the past few years, the NIHSS is not yet coded for each patient<sup>12</sup>. Including only patients with coded NIHSS would have limited our sample size. However, we are hopeful that in future studies we can control for NIHSS as the coding in the NIS dataset increases. Additionally, the use of tPA and EVT is a patient specific medical decision. Within the NIS dataset, we cannot control for all reasons as to why a patient did or did not receive tPA or EVT such as time of arrival to hospital since onset of stroke, medical

contraindications for treatment, or patient medical history. Also, a small number of patients received both tPA and EVT, and this was not controlled for in our analysis. Additionally, while we did not find a racial or ethnic difference in in-hospital mortality after EVT or tPA, this may be due to a lack of statistical power as only a small subset of our population died in the hospital. Because the dataset comes from medical records, there is a possibility of data coding errors regarding demographics, tPA and EVT use, mortality, and discharge status. Lastly, we cannot account for all comorbidities and confounding variables that may be underreported.

# Conclusion

The current study demonstrates that Black ischemic stroke patients are less likely to receive tPA and EVT than White patients. Additionally, when assessing post-tPA or EVT outcomes, we failed to find a racial or ethnic difference for in-hospital mortality, but we did find that minority patients (including but not limited to Black, Hispanic, Asian, Pacific Islander, and Native American) had significantly longer LOS in the hospital than White patients. Even with an increasing trend in the usage of tPA and EVT in the US, we still see disparities when caring for minority patients with ischemic stroke. Black patients are less likely to receive both tPA and EVT, and minority populations have longer hospital stays than White patients which may be an indicator of suboptimal quality of care or poor clinical outcome. We hope that with additional research, we can gain a greater understanding of the racial and ethnic disparities in usage of the life-saving treatments of tPA and EVT in ischemic stroke patients to further encourage interventions to improve stroke care for minority patients.

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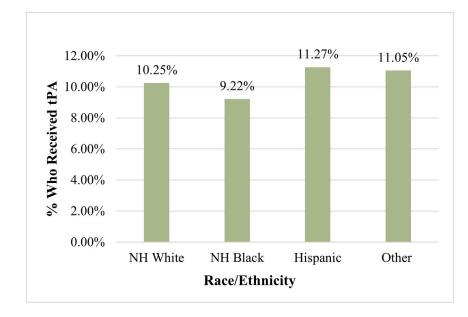
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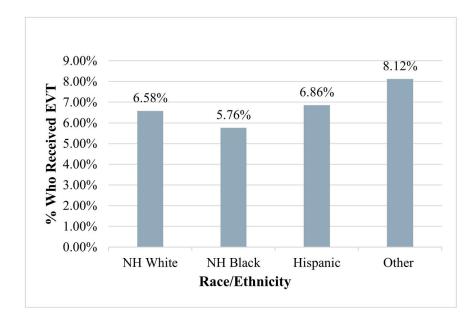
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# Figure 1.

Proportion of ischemic stroke patients who received tPA (N=89,035) \*tPA: Tissue Plasminogen Activator

<sup>†</sup> The "Other" category of race/ethnicity includes Asian, Pacific Islanders, Native Americans, and "others" as classified by the NIS



# Figure 2.

Proportion of Ischemic Stroke Patients Who Received EVT (N=89,035) \*EVT: Endovascular Thrombectomy

<sup>†</sup> The "Other" category of race/ethnicity includes Asian, Pacific Islanders, Native Americans, and "others" as classified by the NIS

#### Table 1.

Sample characteristics by tPA and EVT for patients with ischemic stroke, National Inpatient Sample 2019 (n=89,035; N=445,174).

Variable	tPA	P-value	EVT	P-valu
Race/Ethnicity		< 0.001		< 0.001
NH White	6242 (10.25%)		4007 (6.58%)	
NH Black	1425 (9.22%)		890 (5.76%)	
Hispanic	811 (11.27%)		494 (6.86%)	
Other	608 (11.05%)		447 (8.12%)	
Age		< 0.001		< 0.001
18–44	587 (14.77%)		342 (8.61%)	
45–64	2821 (11.04%)		1719 (6.73%)	
65–74	2095 (9.58%)		1377 (6.3%)	
75+	3583 (9.52%)		2400 (6.37%)	
Sex		0.8457		0.029
Male	4571 (10.19%)		2862 (6.38%)	
Female	4515 (10.22%)		2976 (6.74%)	
Household income quartiles		< 0.001		< 0.001
\$1-24,999	2461 (8.88%)		1688 (6.09%)	
\$25,000-34,999	2206 (9.73%)		1483 (6.54%)	
\$35,000-44,999	2396 (11.08%)		1475 (6.82%)	
\$45,000+	2023 (11.87%)		1192 (6.99%)	
Payers		< 0.001		< 0.001
Medicare	5571 (9.59%)		3602 (6.2%)	
Medicaid	813 (9.86%)		572 (6.94%)	
Private insurance	2054 (12.27%)		1292 (7.72%)	
Other	648 (10.88%)		372 (6.25%)	
Discharge Quarter		0.6387		0.0956
Jan-Mar	2213 (10.06%)		1385 (6.3%)	
Apr-Jun	2261 (10.09%)		1437 (6.41%)	
July-Sep	2280 (10.29%)		1484 (6.7%)	
Oct-Dec	2332 (10.37%)		1532 (6.82%)	
Discharge Day		< 0.001		0.3832
Weekday Discharge	6601 (10%)		4299 (6.51%)	
Weekend Discharge	2485 (10.79%)		1539 (6.68%)	
Location		< 0.001		< 0.001
Urban	8811 (10.61%)		5815 (7%)	
Rural	275 (4.57%)		23 (0.38%)	
Teaching Status		0.2986		< 0.001
No	7591 (10.25%)		5471 (7.39%)	
Yes	1495 (9.97%)		367 (2.45%)	
Hospital bed size		< 0.001		< 0.001

Variable	tPA	P-value	EVT	P-value
Small	1375 (8.54%)		392 (2.44%)	
Medium	2746 (10.54%)		1274 (4.89%)	
Large	4965 (10.59%)		4172 (8.9%)	
Comorbidities		< 0.001		< 0.001
1	937 (7.51%)		356 (2.85%)	
2+	8149 (10.64%)		5482 (7.16%)	

\* The "Other" category of race/ethnicity includes but is not limited to Asian, Pacific Islanders, and Native Americans

#### Table 2.

Logit Regression Assessing the Association Between the Use of tPA and EVT and Race/Ethnicity in Ischemic Stroke Patients, National Inpatient Sample 2019 (n=89,035; N= 445,174).

Variable	Unadjusted Odds Ratio	Unadjusted 95% Confidence Interval	Adjusted Odds Ratio	Adjusted 95% Confidence Interval
Race/Ethnicity				
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	0.88 ***	[0.83–0.94]	0.85 ***	[0.80–0.91]
Hispanic	1.11***	[1.03–1.20]	0.99	[0.92–1.08]
Other	1.08	[0.99-1.18]	0.95	[0.87-1.04]

Ischemic stroke patients who received EVT

Variable	Unadjusted Odds Ratio	Unadjusted 95% Confidence Interval	Adjusted Odds Ratio	Adjusted 95% Confidence Interval
Race/Ethnicity				
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	0.86***	[0.80-0.93]	0.75 ***	[0.70-0.82]
Hispanic	1.06	[0.96–1.17]	0.93	[0.84–1.03]
Other	1.24 ***	[1.12–1.37]	1.11*	[1.00–1.24]

\* \* P < .05;

\*\* P<.01;

\*\*\* P < .001.

 $\vec{t}$  Models adjusted for patients' age, sex, neighborhood median household income, payers, discharge quarter, number of comorbidities, hospital location, hospital teaching status, hospital bed size, and weekend discharge. Marginal probabilities were estimated from the logit regression model and converted from odds ratios. Note: P-values were calculated using Chi-square tests.

 $^{\$}$ All statistics were adjusted using sampling weights

<sup>//</sup>The "Other" category of race/ethnicity includes but is not limited to Asian, Pacific Islanders, and Native Americans

#### Table 3.

Logit Regression Assessing the Association Between In-hospital Mortality and Race/Ethnicity in Ischemic Stroke Patients who Received tPA or EVT, National Inpatient Sample 2019.

Variable	Unadjusted Odds Ratio	Unadjusted 95% Confidence Interval	Adjusted Odds Ratio	Adjusted 95% Confidence Interval
Race/Ethnicit	У			
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	0.94	[0.70–1.26]	1.19	[0.86–1.64]
Hispanic	0.98	[0.68–1.42]	1.05	[0.70–1.58]
Other	1.37	[0.95–1.98]	1.38	[0.94–2.02]
In-hospital m	ortality for EVT patients (n=8	9,035 ; N=29,185)		
Variable	Unadjusted Odds Ratio	Unadjusted 95% Confidence Interval	Adjusted Odds Ratio	Adjusted 95% Confidence Interval
Race/Ethnicit	у			
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	0.77	[0.60-0.98]	0.85	[0.65-1.10]
Hispanic	1.07	[0.81–1.41]	1.08	[0.81–1.46]

\*\* P < .05;

\*\* P < .01;

\*\*\* P<.001.

 $\dot{\tau}$  Results were presented as weighted N (%).

<sup>*i*</sup>Models adjusted for patients' age, sex, neighborhood median household income, payers, discharge quarter, number of comorbidities, hospital location, hospital teaching status, hospital bed size, and weekend discharge. Marginal probabilities were estimated from the logit regression model and converted from odds ratios. Note: P-values were calculated using Chi-square tests.

 $^{\$}$ All statistics were adjusted using sampling weights

<sup>//</sup>The "Other" category of race/ethnicity includes but is not limited to Asian, Pacific Islanders, and Native Americans

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#### Table 4.

Poisson Regression Assessing the Association Between Hospital Length of Stay and Race/Ethnicity in Ischemic Stroke Patients Who Received tPA or EVT, National Inpatient Sample 2019.

Variable	Unadjusted Incidence Rate Ratio	Unadjusted 5% Confidence Interval	Adjusted Incidence Rate Ratio	Adjusted 95% Confidence Interval
Race/Ethnicity				
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	1.37 ***	[1.281.48]	1.31 ***	[1.21–1.41]
Hispanic	1.18**	[1.06–1.31]	1.15 **	[1.04–1.28]
Other	1.19***	[1.09–1.30]	1.16***	[1.07–1.27]

Variable	Unadjusted Incidence Rate Ratio	Unadjusted 95% Confidence Interval	Adjusted Incidence Rate Ratio	Adjusted 95% Confidence Interval
Race/Ethnicity				
NH White	1.00	[1.00-1.00]	1.00	[1.00-1.00]
NH Black	1.45 ***	[1.33–1.58]	1.30 ***	[1.19–1.43]
Hispanic	1.33***	[1.17–1.50]	1.23 ***	[1.10–1.38]
Other	1.32***	[1.17–1.49]	1.28 ***	[1.14–1.44]

\*\* P < .05;

\*\* P < .01;

\*\*\* P < .001.

 ${}^{\dagger}$ Results were presented as weighted N (%).

<sup>*t*</sup>Models adjusted for patients' age, sex, neighborhood median household income, payers, discharge quarter, number of comorbidities, hospital location, hospital teaching status, hospital bed size, and weekend discharge. Marginal probabilities were estimated from the logit regression model and converted from odds ratios. Note: P-values were calculated using Chi-square tests.

 $^{\$}$ All statistics were adjusted using sampling weights

// The "Other" category of race/ethnicity includes but is not limited to Asian, Pacific Islanders, and Native Americans