



Trends in Thrombolytic Use for Ischemic Stroke in the United States

Citation

Fang, Margaret C., David M. Cutler, and Allison B. Rosen. 2010. Trends in thrombolytic use for ischemic stroke in the United States. Journal of Hospital Medicine 5(7): 406-9.

Published Version

doi:10.1002/jhm.689

Permanent link

http://nrs.harvard.edu/urn-3:HUL.InstRepos:5343031

Terms of Use

This article was downloaded from Harvard University's DASH repository, and is made available under the terms and conditions applicable to Open Access Policy Articles, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#OAP

Share Your Story

The Harvard community has made this article openly available. Please share how this access benefits you. <u>Submit a story</u>.

Accessibility

Trends in Thrombolytic Use for Ischemic Stroke in the United States

Margaret C. Fang, MD, MPH¹

David M. Cutler, PhD^{2,3}

Allison B. Rosen, MD, MPH, ScD⁴

Corresponding Author:

Margaret C. Fang, MD, MPH

University of California, San Francisco

Division of Hospital Medicine

533 Parnassus Ave., Box 0131

San Francisco, CA 94143

Phone: (415) 502-7100

Fax: (415) 514-2094

Email: mfang@medicine.ucsf.edu

¹ University of California, San Francisco Department of Medicine (San Francisco, CA)

² Harvard University Department of Economics and Kennedy School of Government (Cambridge, MA)

³ National Bureau of Economic Research (Cambridge, MA)

⁴ University of Michigan Department of Medicine (Ann Arbor, MI)

Acknowledgments

We would like to thank Mr. Loren Yglecias for his assistance with manuscript text and references.

Funding

This work was supported by a National Institute on Aging grant P01 AG17625. Dr. Fang was supported by a National Institute on Aging Paul B. Beeson Career Development Award AG028978.

Conflicts of Interest/Disclosures

None of the authors have significant conflicts of interest to disclose

Title: Trends in Thrombolytic Use for Ischemic Stroke in the United States

Cover title: Trends in thrombolytic use for stroke

Text word count: 1560

Tables: 2

Figures: 1

.emic stroke, Outcomes Measator, Thrombolysis Index terms: Acute stroke, Ischemic stroke, Outcomes Measurement, Quality Improvement,

Tissue plasminogen activator, Thrombolysis

ABSTRACT

Background

Although recombinant tissue plasminogen activator (tPA) improves outcomes from ischemic stroke, prior studies have found low rates of administration. Recent guidelines and regulatory agencies have advocated for increased tPA administration in appropriate patients, but it is unclear how many patients actually receive tPA.

Objective

To determine whether national rates of tPA use for ischemic stroke have increased over time.

Methods

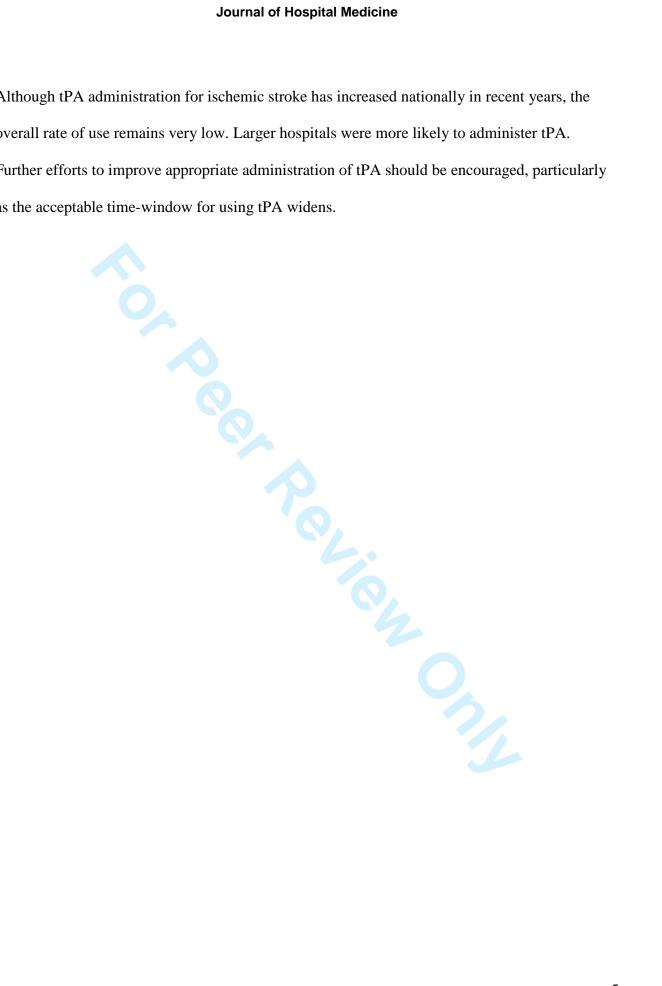
We identified all patients with a primary diagnosis of ischemic stroke from years 2001-2006 in the National Hospital Discharge Survey, a nationally representative sample of inpatient hospitalizations, and searched for procedure codes for intravenous thrombolytic administration. Clinical and demographic factors were obtained from the survey and multivariable logistic regression used to identify independent predictors associated with thrombolytic use.

Results

Among the 22,842 patients hospitalized with ischemic stroke, tPA administration rates increased from 0.87% in 2001 to 2.40% in 2006 (p<0.001 for trend). Older patients were less likely to receive tPA (adjusted odds ratio and 95% confidence interval (OR) = 0.4 [0.3-0.6] for patients \geq 80 years vs. < 60 years), as were African-American patients (OR=0.4 [0.3-0.7]). Larger hospitals were more likely to administer tPA (OR=3.3 [2.0-5.6] in hospitals with at least 300 beds compared to those with 6-99 beds).

Conclusions

Although tPA administration for ischemic stroke has increased nationally in recent years, the overall rate of use remains very low. Larger hospitals were more likely to administer tPA. Further efforts to improve appropriate administration of tPA should be encouraged, particularly as the acceptable time-window for using tPA widens.



Background

Recombinant tissue plasminogen activator (tPA), approved for use in the United States for the treatment for acute ischemic stroke since 1996, improves overall outcomes from ischemic stroke when administered to selected patients¹⁻⁴. Several prominent guidelines, including the Brain Attack Coalition and the American Stroke Association, have recommended increasing the use of tPA for acute ischemic stroke⁵⁻⁷. In addition, in 2003 the Joint Commission on Accreditation of Healthcare Organizations developed a disease-specific certification program to designate certain institutions "Primary Stroke Centers", with one of the performance measures being the availability of thrombolysis⁸.

Despite guidelines and regulatory agencies promoting the use of thrombolysis for ischemic stroke, previous studies have shown disappointingly low rates of use⁹⁻¹². The goals of this study were to assess whether national trends in the use of thrombolysis for acute ischemic stroke have increased in light of increased regulatory activity as well as to identify patient characteristics associated with thrombolytic administration.

Materials and Methods

Data for this study were obtained from the 2001 through 2006 National Hospital Discharge Survey (NHDS), a nationally representative sample of inpatient hospitalizations conducted annually by the National Center for Health Statistics¹³. The NHDS collects data from approximately 300,000 patients from about 500 short-stay nonfederal hospitals in the United States and uses a 3 stage sampling strategy that allows for extrapolation to national level estimates. Response rates typically exceed 90% from participating hospitals. The survey collects demographic data, including age, sex, race, hospital geographic region, hospital bedsize and

patient insurance status. In addition, up to 7 diagnostic and 4 procedural codes from the hospitalization are available, as is hospitalization length of stay and patient discharge disposition. No information on timing of symptoms, degree of neurologic compromise, or results of imaging tests were available in the NHDS.

We searched for all patients aged 18 years or older with a primary diagnostic code of ischemic stroke using the *International Classification of Diseases*, 9th Edition, Clinical Modification (ICD-9-CM) codes 433, 434, 436, 437.0, and 437.1, excluding codes with a fifth digit of 0 (which indicated arterial occlusion without mention of cerebral infarction). We then searched for the presence of an ICD-9-CM procedure code for injection or infusion of thrombolytic agent (code 99.10). Specific comorbid conditions associated with ischemic stroke were identified by searching for specific ICD-9-CM codes, including for heart failure, coronary artery disease, hypertension, diabetes mellitus, and atrial fibrillation. To provide a general assessment of the severity of illness of the patients, we calculated an adapted Charlson comorbidity score for each patient using available secondary discharge diagnosis codes¹⁴. We also searched for codes corresponding to intracranial hemorrhage, a complication associated with tPA administration.

Statistical Analysis

We defined thrombolytic utilization rates as the number of patients hospitalized with a primary diagnosis of ischemic stroke who had a procedure code for thrombolysis divided by the total number of patients hospitalized with ischemic stroke. To calculate nationally representative prevalence rates, we used the sample weights provided by the NHDS to account for the complex sampling design of the survey. Differences in thrombolytic administration rates by patient and

hospital characteristics were tested using chi-squared tests for categorical variables and t-tests for continuous variables. Variables underwent a backwards selection process with a significance level of 0.05 to develop the final multivariable model of predictors of thrombolytic administration. Length of stay and hospital discharge status were not included in the variable selection process, as the focus was on predictors of initial administration of thrombolytics. All analyses were conducted using SAS Version 9.1 (SAS Institute Inc., Cary, NC).

Results

From years 2001 through 2006, we identified 22,842 patients with a primary diagnosis of ischemic stroke. Of these, 313 (1.37%, 95% confidence interval [CI]: 1.22%-1.53%) had a procedure code for injection or infusion of thrombolysis. Using NHDS sample weights, these numbers corresponded to an estimated 2.55 million hospitalizations for ischemic stroke in the United States during the time period and to 35,082 patients receiving intravenous thrombolytics. Although the overall rate of thrombolysis administration was quite low overall, the administration rate increased over time, from 0.87% [95%CI: 0.61%-1.22%] of stroke patients in year 2001 to 2.40% [95%CI: 1.95-2.93%] in year 2006 and with a particular increase especially noted after year 2003 (p<0.001 for trend, **Figure 1**).

On bivariate analysis, a lower proportion of African-American patients received tPA compared to white patients (0.8% vs. 1.5%, p=0.003), while a higher proportion of patients with atrial fibrillation received tPA (2.3% vs. 1.2%, p<0.001). Older patients were less likely than younger patients to receive tPA (**Table 1**). The rate of intracranial hemorrhage was significantly higher in patients who received tPA (5.4% vs. 0.6%, p<0.001) and the overall inpatient mortality in patients who received tPA was 9.0%. Mortality in patients receiving tPA continued to be

higher than in patients who did not receive tPA even when patients with intracranial hemorrhage were excluded (8.1% vs. 5.3%, p<0.001). Larger hospitals were more likely to administer tPA to patients with ischemic stroke, with a 1.79% administration rate in hospitals with \geq 300 beds compared to 0.90% in hospitals with 100-199 beds and 0.52% in hospitals with 6-99 beds (p<0.001).

After adjusting for patient and hospital characteristics, the absolute rate of thrombolysis administration increased by an average of 0.19% per year [95% confidence interval 0.12%-0.26%]. Factors that were significantly associated with administration of thrombolytics included being hospitalized in a larger hospital, having a history of atrial fibrillation, and a higher Charlson comorbidity index (**Table 2**). Patients aged 80 years or older, African-American patients, and those with diabetes mellitus were significantly less likely to receive thrombolysis.

Discussion

Despite strong recommendations from guidelines and regulatory agencies, national rates of intravenous thrombolysis for ischemic stroke continue to be quite low overall. However, tPA administration appears to have increased from previous years and particularly increased in years after the Joint Commission began to accredit institutions as "Primary Stroke Centers". The oldest patients and African-Americans were less likely to receive thrombolytics, while patients with atrial fibrillation were more likely to receive thrombolysis, potentially related to atrial fibrillation causing more severe strokes¹⁵. Five point four percent of patients who received tPA were diagnosed with intracranial hemorrhage, and the inpatient mortality rate of patients with tPA was 9.0%.

The exact optimal rate of thrombolysis administration for the patients in our study is unknown, as the NHDS database lacked detailed information on factors that would preclude tPA administration such as late timing of presentation and mild stroke symptoms³. Studies conducted in stroke registries and regional settings have found that only approximately 15%-32% of patients presenting with ischemic stroke arrive within 3 hours of symptom onset, and of these, only about 40%-50% are eligible for tPA clinically^{9, 10, 16-19}. However, even among presumed eligible patients, tPA administration rates only range between 25% and 43% ^{17, 19, 20}, and the ideal rate is likely to be higher than the very low rates we observed in our study. Newer evidence that extending the time window where tPA may be given safely may increase the number of eligible patients²¹.

Patients who received thrombolysis had higher mortality rates than patients who did not. Although we were unable to determine a causal association, prior observational studies of tPA administration for acute stroke have found that patients with more severe neurologic deficits were more likely to receive thrombolysis ^{17,18}. The 9.0% inpatient case-fatality rate observed in our study compares favorably to the 13.4% mortality rate after tPA reported in a post-approval meta-analysis of safety outcomes ²² and the rate of intracranial hemorrhage in our analysis was similar to those observed in other settings ^{9, 22-25}. We were unable to determine whether intracranial hemorrhages in our study were as a result of tPA administration or whether patients who received tPA were more likely to have intracranial hemorrhages detected, such as may be due to increased frequency of head imaging.

Larger hospitals were more likely to administer tPA. This may reflect regionalization of stroke care, particularly in those designated as stroke "centers of excellence". As well, there is some evidence that there is a "learning curve" with thrombolysis administration, where

guideline-recommended practice and use of tPA increases with additional experience with the drug^{9, 26}. Promoting systems that allow for rapid triage and diagnosis of acute stroke should be encouraged and hospital leaders should develop strategies that allow for early recognition of potential tPA candidates.

There are several limitations to our analysis. The NHDS does not collect detailed data on clinical or presenting features of stroke, and so we lacked information on stroke severity and eligibility for administration of thrombolysis. Our study may have underestimated the overall rates of thrombolysis, as it was dependent on diagnostic codes. A previous study of 34 patients who received tPA found that although the 99.10 code was 100% specific, the code identified only 17 patients who actually received tPA (sensitivity of 50%)²⁰. Another study comparing Medicare administrative claims data to actual pharmacy billing charges for tPA found that administrative data underestimated the rate of tPA administration by about 25%-30% ¹². If a diagnostic code sensitivity of 50% was assumed, rates of tPA administration in our study may have been as high as 4.8% [95% CI: 4.1%-5.5%] by year 2006.

In conclusion, the use of intravenous thrombolysis in patients admitted with acute ischemic stroke in the United States has risen over time, but overall use remains very low. Further efforts to improve appropriate administration rates should be encouraged, particularly as the acceptable time-window for using tPA widens.

Figure 1. Proportion of patients admitted with a primary diagnosis of ischemic stroke who received thrombolysis



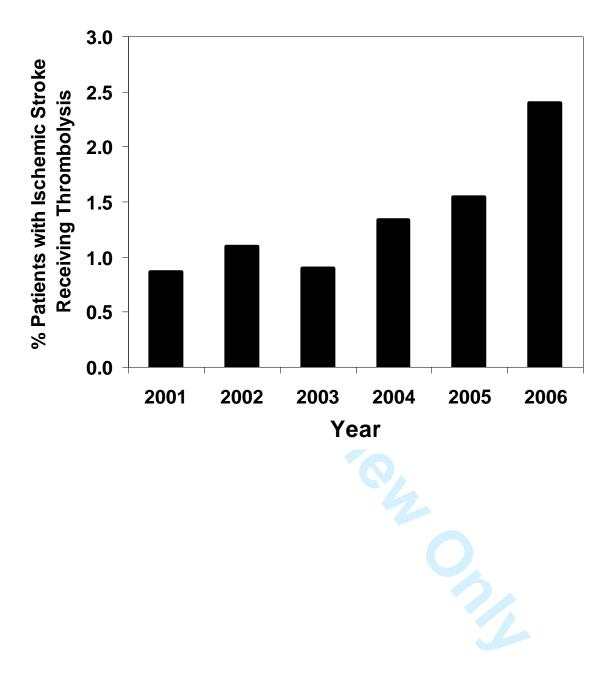


Table 1. Clinical characteristics of 22,842 patients with a primary diagnosis of ischemic stroke

	Thrombolysis	No thrombolysis	P value
	N (%)	N (%)	
Age			0.001
< 60	76 (24.3)	4478 (19.9)	
60-69	73 (23.3)	3942 (17.5)	
70-79	82 (26.2)	6265 (27.8)	
80+	82 (26.2)	7844 (34.8)	
Female	155 (49.5)	12625 (56.0)	0.02
Race			0.003
White	173 (55.3)	11542 (51.2)	
African-American	29 (9.3)	3774 (16.8)	
Other	16 (5.1)	814 (3.6)	
Not stated	95 (30.4)	6399 (28.4)	
Region			0.05
Northeast	78 (24.9)	4570 (20.3)	
Midwest	84 (26.8)	6924 (30.7)	
South	104 (33.2)	8284 (36.8)	
West	47 (15.0)	2751 (12.2)	
Type of admission			< 0.001
Emergent	247 (78.9)	14233 (63.2)	
Urgent	33 (10.5)	3703 (16.4)	
Elective	5 (1.6)	1346 (6.0)	
Unknown	28 (9.0)	3247 (14.4)	
Length of stay, days [95% CI]	7.2 [6.6-7.8]	6.0 [5.9-6.1]	< 0.001
Hospital bedsize			< 0.001
6-99	16 (5.1)	3065 (13.6)	
100-199	48 (15.3)	5289 (23.5)	
200-299	86 (27.5)	5212 (23.1)	
300+	163 (52.1)	8963 (39.8)	

Payment type			< 0.001
Medicare	176 (56.2)	15197 (67.5)	
Medicaid	21 (6.7)	1245 (5.5)	
Private	45 (14.4)	2483 (11.0)	
HMO/PPO	39 (12.5)	2224 (9.9)	
Other/unknown	32 (10.2)	1380 (6.1)	
Discharge status			< 0.001
Home	98 (31.3)	9507 (42.2)	
Short term care facility	25 (8.0)	1271 (5.6)	
Long term care facility	62 (19.8)	5400 (24.0)	
Alive, status unknown	89 (28.4)	4514 (20.0)	
Death	28 (9.0)	1218 (5.4)	
Unknown	11 (3.5)	619 (2.8)	
Comorbid conditions			
Congestive heart failure	48 (15.3)	2769 (12.3)	0.10
Coronary artery disease	49 (15.7)	4082 (18.1)	0.26
Hypertension	164 (52.4)	12480 (55.4)	0.29
Diabetes mellitus	42 (13.4)	4965 (22.0)	< 0.001
Atrial fibrillation	96 (30.7)	4096 (18.2)	< 0.001
Intracranial hemorrhage	17 (5.4)	139 (0.6)	< 0.001
Charlson score (mean)	2.48 [2.32-2.64]	2.38 [2.36-2.40]	0.23

Table 2. Independent predictors of receiving thrombolysis in 22,842 hospitalized patients with a primary diagnosis of ischemic stroke

Characteristic	Adjusted odds ratio and 95% confidence
	intervals
Year (per year, from 2001 to 2006)	1.2 [1.1-1.3]
Age	
<60	Referent
60-69	1.0 [0.7-1.4]
70-79	0.6 [0.5-0.9]
80+	0.4 [0.3-0.6]
Race	
Not African-American	Referent
African-American	0.4 [0.3-0.7]
Unknown	1.0 [0.8-1.2]
Hospital bedsize	
6-99	Referent
100-199	1.7 [1.0-3.1]
200-299	3.2 [1.8-5.4]
300+	3.3 [2.0-5.6]
Diabetes mellitus	0.5 [0.3-0.6]
Atrial fibrillation	2.2 [1.7-2.9]
Charlson comorbidity score ¹⁴ (per point increase)	1.1 [1.1-1.2]

REFERENCES

- The National Institute of Neurological Disorders and Stroke rt-PA Stroke Study Group.
 Tissue plasminogen activator for acute ischemic stroke. N Engl J Med. 1995;333:1581-1587
- 2. Kwiatkowski TG, Libman RB, Frankel M, Tilley BC, Morgenstern LB, Lu M, Broderick JP, Lewandowski CA, Marler JR, Levine SR, Brott T. Effects of tissue plasminogen activator for acute ischemic stroke at one year. National Institute of Neurological Disorders and Stroke Recombinant Tissue Plasminogen Activator Stroke Study Group. N Engl J Med. 1999;340:1781-1787
- 3. Hacke W, Donnan G, Fieschi C, Kaste M, von Kummer R, Broderick JP, Brott T, Frankel M, Grotta JC, Haley EC, Jr., Kwiatkowski T, Levine SR, Lewandowski C, Lu M, Lyden P, Marler JR, Patel S, Tilley BC, Albers G, Bluhmki E, Wilhelm M, Hamilton S. Association of outcome with early stroke treatment: Pooled analysis of ATLANTIS, ECASS, and NINDS rT-PA stroke trials. *Lancet*. 2004;363:768-774
- 4. Wahlgren N, Ahmed N, Davalos A, Ford GA, Grond M, Hacke W, Hennerici MG, Kaste M, Kuelkens S, Larrue V, Lees KR, Roine RO, Soinne L, Toni D, Vanhooren G. Thrombolysis with alteplase for acute ischaemic stroke in the safe implementation of thrombolysis in stroke-monitoring study (SITS-MOST): An observational study. *Lancet*. 2007;369:275-282
- 5. Alberts MJ, Hademenos G, Latchaw RE, Jagoda A, Marler JR, Mayberg MR, Starke RD, Todd HW, Viste KM, Girgus M, Shephard T, Emr M, Shwayder P, Walker MD, for the Brain Attack Coalition. Recommendations for the establishment of primary stroke centers. *JAMA*. 2000;283:3102-3109

- Klijn CJ, Hankey GJ. Management of acute ischaemic stroke: New guidelines from the American Stroke Association and European Stroke Initiative. *Lancet Neurol*. 2003;2:698-701
- 7. Adams HP, Jr., del Zoppo G, Alberts MJ, Bhatt DL, Brass L, Furlan A, Grubb RL, Higashida RT, Jauch EC, Kidwell C, Lyden PD, Morgenstern LB, Qureshi AI, Rosenwasser RH, Scott PA, Wijdicks EF. Guidelines for the early management of adults with ischemic stroke: A guideline from the American Heart Association/American Stroke Association Stroke Council, Clinical Cardiology Council, Cardiovascular Radiology and Intervention council, and the Atherosclerotic Peripheral Vascular Disease and Quality of Care Outcomes in Research Interdisciplinary Working groups: The American Academy of Neurology affirms the value of this guideline as an educational tool for neurologists. *Circulation*. 2007;115:e478-534
- 8. The Joint Commission Primary Stroke Center Certification.

 http://www.jointcommission.org/CertificationPrograms/PrimaryStrokeCenters. Last accessed October 1, 2009.
- 9. Katzan IL, Furlan AJ, Lloyd LE, Frank JI, Harper DL, Hinchey JA, Hammel JP, Qu A, Sila CA. Use of tissue-type plasminogen activator for acute ischemic stroke: The Cleveland area experience. *JAMA*. 2000;283:1151-1158
- California Acute Stroke Pilot Registry (CASPR) Investigators. Prioritizing interventions to improve rates of thrombolysis for ischemic stroke. *Neurology*. 2005;64:654-659
- 11. Qureshi AI, Suri MF, Nasar A, He W, Kirmani JF, Divani AA, Prestigiacomo CJ, Low RB. Thrombolysis for ischemic stroke in the united states: Data from National Hospital Discharge Survey 1999-2001. *Neurosurgery*. 2005;57:647-654; discussion 647-654

- 12. Kleindorfer D, Lindsell CJ, Brass L, Koroshetz W, Broderick JP. National US estimates of recombinant tissue plasminogen activator use: ICD-9 codes substantially underestimate. *Stroke*. 2008;39:924-928
- 13. US Department of Health and Human Services Public Health Service and National Center for Health Statistics. National Hospital Discharge Durvey 1991-2006. Multi-year public-use data file documentation.
- 14. Deyo R, Cherkin D, Ciol M. Adapting a clinical comorbidity index for use with ICD-9-CM administrative databases. *J Clin Epidemiol*. 1992;45:613-619
- Lin HJ, Wolf PA, Kelly-Hayes M, Beiser AS, Kase CS, Benjamin EJ, D. Agostino R.
 Stroke severity in atrial fibrillation. The Framingham study. *Stroke*. 1996;27:1760-1764
- 16. Barber P, Zhang J, Demchuk A, Hill M, Buchan A. Why are stroke patients excluded from tPA therapy? An analysis of patient eligibility. *Neurology*. 2001;57:1739-1740
- Katzan IL, Hammer MD, Hixson ED, Furlan AJ, Abou-Chebl A, Nadzam DM.
 Utilization of intravenous tissue plasminogen activator for acute ischemic stroke. *Arch Neurol*. 2004;61:346-350
- 18. Kleindorfer D, Kissela B, Schneider A, Woo D, Khoury J, Miller R, Alwell K, Gebel J, Szaflarski J, Pancioli A, Jauch E, Moomaw C, Shukla R, Broderick JP, for The Neuroscience I. Eligibility for recombinant tissue plasminogen activator in acute ischemic stroke: A population-based study. *Stroke* 2004;35:27-29
- 19. Nadeau JO, Shi S, Fang J, Kapral MK, Richards JA, Silver FL, Hill MD. Tpa use for stroke in the registry of the Canadian stroke network. *Can J Neurol Sci.* 2005;32:433-439

- 20. Johnston SC, Fung LH, Gillum LA, Smith WS, Brass LM, Lichtman JH, Brown AN. Utilization of intravenous tissue-type plasminogen activator for ischemic stroke at academic medical centers: The influence of ethnicity. *Stroke*. 2001;32:1061-1068
- 21. Saver JL, Gornbein J, Grotta J, Liebeskind D, Lutsep H, Schwamm L, Scott P, Starkman S. Number needed to treat to benefit and to harm for intravenous tissue plasminogen activator therapy in the 3- to 4.5-hour window: Joint outcome table analysis of the ECASS 3 trial. *Stroke* 2009;40:2433-2437
- Graham GD. Tissue plasminogen activator for acute ischemic stroke in clinical practice:
 A meta-analysis of safety data. *Stroke*. 2003;34:2847-2850
- 23. Grond M, Stenzel C, Schmulling S, Rudolf J, Neveling M, Lechleuthner A, Schneweis S, Heiss WD. Early intravenous thrombolysis for acute ischemic stroke in a community-based approach. *Stroke*. 1998;29:1544-1549
- 24. Bateman BT, Schumacher HC, Boden-Albala B, Berman MF, Mohr JP, Sacco RL, Pile-Spellman J. Factors associated with in-hospital mortality after administration of thrombolysis in acute ischemic stroke patients: An analysis of the Nationwide Inpatient Sample 1999 to 2002. *Stroke*. 2006;37:440-446
- 25. Zweifler RM, Brody ML, Graves GC, U TT, Drinkard R, Cunningham S, Rothrock JF.
 Intravenous t-PA for acute ischemic stroke: Therapeutic yield of a stroke code system.
 Neurology. 1998;50:501-503
- 26. Grotta JC, Burgin WS, El-Mitwalli A, Long M, Campbell M, Morgenstern LB, Malkoff M, Alexandrov AV. Intravenous tissue-type plasminogen activator therapy for ischemic stroke: Houston experience 1996 to 2000. *Arch Neurol*. 2001;58:2009-2013