Clinical Study

Reduction in Door-to-Needle Time after Transfer of Thrombolysis Site from CCU to Emergency Department

Osama Mohammed, Firjith C. Paramba, Naushad V. Aboobaker, Riyadh A. Mohammed, Nishan K. Purayil, Haitham M. Jassim, Mohammad K. Shariff, Saud M. Aslam, Farook F. Muhsen, Khalid H. Al Noor, and Hani H. Al Kilani

Department of Emergency Medicine, Al Khor Hospital, Hamad Medical Corporation, P.O. Box 21086, Doha, Qatar

Correspondence should be addressed to Osama Mohammed; osamahmohd@gmail.com

Received 25 January 2013; Revised 31 July 2013; Accepted 16 August 2013

Academic Editor: Robert W. Derlet

Copyright © 2013 Osama Mohammed et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Objective. Early restoration of coronary perfusion by thrombolysis or percutaneous coronary intervention is the main modality of treatment to salvage the ischemic myocardium. The earlier the procedure is completed, the greater the benefit is in saving myocardium and restoring its functions. The aim of the study is to compare the door-to-needle time (DNT) in acute ST elevation myocardial infarction (STEMI) in the period prior to December 2008 when the site of thrombolysis was in coronary care unit (CCU) and the period after that when the site was shifted to emergency department (ED).

Methods. A retrospective, descriptive study was conducted at Al Khor Hospital, Qatar, in patients with acute STEMI who underwent thrombolysis at CCU and ED from April 2005 until December 2011, to compare the DNT, duration of hospitalization, and mortality.

Results. A total of 211 patients with acute STEMI were eligible for thrombolysis; 58 patients were thrombolysed in the CCU and 153 in ED. The median DNT was reduced from 33.5 minutes in the CCU to 17 minutes in the ED representing a reduction of more than 50% with a P value of < 0.0001. Conclusion. The transfer of the thrombolysis site from CCU to the ED was associated with a dramatic and significant reduction in median door-to-needle time by more than half.

1. Introduction

Acute reperfusion therapy performed either with thrombolytic therapy or percutaneous coronary intervention (PCI) is the mainstay of treatment for patients with acute ST-segment elevation myocardial infarction (STEMI). The benefit of the perfusion is restoring coronary flow which is time-dependent; the earlier the reperfusion is established, the greater the benefit is in saving the myocardium [1–3]. Randomized clinical trials have shown that early reperfusion therapy reduces the overall 30-day mortality by 17–25%, with increasing benefit as the time from onset of pain to the initiation of thrombolytic therapy is reduced [4–6]. Since symptoms-to-door time (SDT) is beyond the control of the medical team in the hospital, the focus is stressed on decreasing the time from the first medical contact to reperfusion therapy in acute myocardial infarction. Hence, the importance of door-to-needle time (DNT) for thrombolytic therapy and door-to-balloon time for PCI has emerged. These interventions occur within the hospital and can be controlled with proper training of the medical and nursing staff and by applying international practice guidelines. DNT is the time taken from patient’s arrival to a medical facility to the time when thrombolytic therapy is administered. As a result of the importance of the timing of the thrombolysis, DNT time has emerged as an important hospital performance measure for the quality of care of patients with STEMI in the United States and Europe [6–8].

The American College of Cardiology/the American Heart Association (ACC/AHA) and the European Society of Cardiology (ESC) guidelines for STEMI recommend that the DNT for thrombolysis should be within 30 minutes of first medical system contact [6, 7]. Hospitals fail to achieve this goal because of the fact that thrombolytic therapy is often not initiated in the ED [9–13]. In some hospitals therapy may be initiated in the ED, but only a minority of patients are
Emergency Medicine International

2 Emergency Medicine International

Patient presented to ED with chest pain

12-lead ECG

Evaluation by ED physician

Decision to thrombolys

Thrombolysed in ED

ED group of patients

CCU group of patients

Patient presented to ED with chest pain

12-lead ECG

Evaluation by ED physician

Cardiologist informed

Evaluation by cardiologist

Decision to thrombolys

Patient transfer to CCU

Thrombolysed in CCU

Figure 1: Flow chart showing the process of thrombolysis in ED and CCU group.

thrombolysed within this time frame because of varying ED protocols [4, 5, 14, 15]. Hence we decided to conduct a study comparing the DNT of thrombolysis in CCU versus ED in patients presenting with acute STEMI.

2. Patients and Methods

This study was carried out at the ED, Al-Khor Hospital, Hamad Medical Corporation, Qatar. Prior to December 2008, patients presenting with acute STEMI to ED were thrombolysed in the coronary care unit (CCU) after evaluated by the cardiologist. A well-trained team comprising of ED physicians and nurses was formed in the year January 2009 for thrombolysing acute STEMI patients in ED. Patients who were diagnosed of having acute STEMI were evaluated by ED physician and subsequently thrombolysed in the ED (Figure 1). Door-to-needle time for thrombolysis in this group were compared with those of patients who were thrombolysed in CCU prior to 2009. A retrospective data collection was made from the medical records and computerized departmental data base. This included demographic features, comorbid conditions, time of onset of symptoms, time of arrival to hospital, indication for thrombolysis, door-to-needle time, symptom-to-door time, and course in the hospital which were all noted. Patients with delayed presentation and incomplete medical records and who presented with an initial nondiagnostic ECG were excluded. The study was approved by the Ethics Committee of the medical research department (approval number #10170/10).

3. Statistical Analysis

Categorical and continuous values are expressed as frequency, percentage, mean ± SD, median, and range. Descriptive statistics were used to summarise all demographic and other clinical characteristics of the patients. Quantitative variables means for the two thrombolysis sites (independent groups) were analyzed using the unpaired t-test. For nonnormal data (skewed), the corresponding nonparametric Mann-Whitney U test was applied to assess significant difference in DNT between the two thrombolysis sites (CCU and ED). Associations between two or more qualitative or categorical variables were assessed using Chi-square tests. Pictorial representations of the key results were made using appropriate statistical graphs, including box plot and bar diagrams. A two-sided P value < 0.05 was considered as statistically significant. All statistical analyses were conducted using IBM SPSS statistical package V19 (SPSS Inc., Chicago, IL).

4. Results

A total of 302 acute STEMI patients were included in the analysis out of which 211 patients with acute STEMI were eligible for thrombolysis: 153 in the ED and 58 in the CCU. Ninety-one patients were excluded from the study. Details are shown in Figure 2. The base line characteristics and site of infarctions of both the groups are shown in Table 1. There were no significant differences between the two groups. The majority of the patients were men, and most commonly comorbid conditions were hypertension, diabetes, and dyslipidemia.

There was a significant reduction in the DNT of patients thrombolysed in ED. The mean DNT was 17 minutes in the ED group compared to 33.5 minutes in the CCU group, representing more than 50% reduction in DNT with a P value of < 0.0001 (Table 2, Figure 3). The SDT was 120 minutes in both the groups. Mean duration of hospitalization (days) did not differ significantly between the CCU and ED groups (5.3 ± 1.8 and 5.7 ± 2.1; P value = 0.155). One patient from the CCU group and 2 from the ED group died during treatment. Two patients from the ED group were thrombolysed according to the ED STEMI protocol, but in both patients, cardiac enzymes remained normal and there was no regional wall abnormality on the Echocardiography. The symptoms were probably due to coronary artery spasm.

5. Discussion

Recent advancements in the medical technologies have revolutionized the treatment of acute myocardial infarction. Even
Emergency Medicine International

3

302 patients in the study with STEMI

211 patients were eligible for thrombolysis

91 patients excluded from the study

58 patients were thrombolysed in CCU

133 patients were thrombolysed in ED

3 patients presented to ED with normal ECG

42 patients excluded because of incomplete or missing medical

46 patients not thrombolysed because of CI

24 in ED 18 in CCU 35 in ED 11 in CCU

Figure 2: Flow chart of patient allocation.

Figure 3: Box plot graph comparing DNT in both the ED and the CCU patient groups. CCU: coronary care unit and ED: emergency department.

Table 1: Demographic and clinical data in patients with acute STEMI.

| Variable       | CCU group N = 58 | ED group N = 153 |
|----------------|------------------|------------------|
| Age            |                  |                  |
| Mean ± SD      | 46.05 ± 7.47     | 47.31 ± 7.99     |
| Median (range) | 46 (27–73)       | 47 (25–63)       |
| Gender         |                  |                  |
| Male = 56 (96.55%) | Male = 150 (98.04%) |
| Female = 2 (3.45%) | Female = 3 (1.96%) |
| Hypertension   | 13 (27%)         | 31 (28%)         |
| Diabetes       | 18 (31%)         | 43 (30%)         |
| High LDL > 3 mmol/L* | 25 (46%)         | 59 (44%)         |
| Smoking        | 32 (68%)         | 68 (61%)         |
| Type of MI     |                  |                  |
| Anterior MI    | 29 (50%)         | 66 (43%)         |
| Nonanterior MI | 29 (50%)         | 87 (57%)         |

* According to the European Society of Cardiology guidelines 2012.

Figure 4

Comparison of Median DTN time between ED and ICU and the international benchmark

Comparison though primary PTCA has been the mainstay of treatment in the present era, it is limited to a few tertiary care centers and is not affordable for all the patients especially in the developing world. Hence thrombolysis is still the treatment of choice for reperfusion in most parts of the world [6, 16]. Earlier trials have demonstrated the benefit of giving thrombolysis as early as possible in acute STEMI patients [17–21].

In our study, shifting thrombolysis to ED was associated with a major reduction in the DNT (from 33.5 to 17 minutes), which is well within the international guidelines (Figure 4).

This result was much better than the results reported in previous studies [4–6, 8, 10, 11, 13–15]. Although the DNT reduction was highly significant, there was no significant reduction in hospitalization days or mortality among either of the groups which is similar to what was found in some
Acknowledgment

The authors thank Prem Chandra from the Medical Research Center, Hamad Medical Corporation, for his advice and statistical data review. They give special thanks to Ann Christine Mercado, the Head Nurse, Emergency department, and all the nursing staff for their cooperation and proper documentation which helped tremendously in retrieving data beside the doctor’s documentation. Special thanks to the staff of medical records department for their support.

References

[1] R. L. McNamara, J. Herrin, Y. Wang et al., “Impact of delay in door-to-needle time on mortality in patients with ST-segment elevation myocardial infarction,” The American Journal of Cardiology, vol. 100, no. 8, pp. 1227–1232, 2007.

[2] J. G. Jollis, R. H. Mehta, M. L. Roettig, P. B. Berger, J. D. Babb, and C. B. Granger, “Reperfusion of acute myocardial infarction in North Carolina emergency departments (RACE): study design,” American Heart Journal, vol. 152, no. 5, pp. 851.e1–851.e11, 2006.

[3] Y. T. Taylali, “Door-to-needle times in acute myocardial infarction,” Asian Cardiovascular and Thoracic Annals, vol. 18, no. 2, pp. 122–126, 2010.

[4] C. T. Hourigan, D. Mountain, P. E. Langton et al., “Changing the site of delivery of thrombolytic treatment for acute myocardial infarction from the coronary care unit to the emergency department greatly reduces door to needle time,” Heart, vol. 84, no. 2, pp. 157–163, 2000.

[5] M. J. Schull, M. Vermeulen, G. Slaughter, L. Morrison, and P. Daly, “Emergency department crowding and thrombolyis delays in acute myocardial infarction,” Annals of Emergency Medicine, vol. 44, no. 6, pp. 577–585, 2004.

[6] F. van de Werf, J. Bax, A. Betriu et al., “Management of acute myocardial infarction in patients presenting with persistent ST-segment elevation,” European Heart Journal, vol. 29, no. 23, pp. 2909–2945, 2008.

[7] F. A. Masoudi, R. O. Bonow, R. G. Brindis et al., “ACC/AHA 2008 statement on performance measurement and reperfusion therapy: a report of the ACC/AHA Task Force on Performance Measures (Work group to address the challenges of performance measurement and reperfusion therapy),” Circulation, vol. 118, no. 24, pp. 2649–2661, 2008.

[8] R. L. McNamara, J. Herrin, E. H. Bradley et al., “Hospital improvement in time to reperfusion in patients with acute myocardial infarction, 1999 to 2002,” Journal of the American College of Cardiology, vol. 47, no. 1, pp. 45–51, 2006.

[9] P. Wilmshurst, A. Purchase, C. Webb, C. Jowett, and T. Quinn, “Improving door to needle times with nurse initiated thrombolyis,” Heart, vol. 84, no. 3, pp. 262–266, 2000.

[10] R. C. Maharaj, H. Geduld, and L. A. Wallis, “Door-to-needle time for administration of fibrinolytics in acute myocardial infarction in Cape Town,” South African Medical Journal, vol. 102, no. 4, pp. 241–244, 2012.

[11] P. Panduranga, I. Al-Zakwani, K. Sulaiman et al., “Clinical profile and mortality of ST-segment elevation myocardial infarction patients receiving thrombolytic therapy in the Middle East,” Heart Views, vol. 13, no. 2, pp. 35–41, 2012.

[12] I. Al-Zakwani, M. Zubaid, A. Al-Riyami et al., “Primary coronary intervention versus thrombolytic therapy in myocardial
infarction patients in the Middle East,” *International Journal of Clinical Pharmacy*, vol. 34, no. 3, pp. 445–451, 2012.

[13] A. A. Abba, B. Wani A., R. A. Rahmatullah, M. Z. Khalil, A. M. Kumo, and M. A. Ghonaim, “Door to needle time in administering thrombolytic therapy for acute myocardial infarction,” *Saudi Medical Journal*, vol. 24, no. 4, pp. 361–364, 2003.

[14] A. R. Corfield, C. A. Graham, J. N. Adams, I. Booth, and A. C. McGuffie, “Emergency department thrombolysis improves door to needle times,” *Emergency Medicine Journal*, vol. 21, no. 6, pp. 676–680, 2004.

[15] P. J. Zed, R. B. Abu-Laban, T. M. Cadieu, R. A. Pursell, and L. Filiatrault, “Fibrinolytic administration for acute myocardial infarction in a tertiary ED: factors associated with an increased door-to-needle time,” *American Journal of Emergency Medicine*, vol. 22, no. 3, pp. 192–196, 2004.

[16] E. M. Antman, D. T. Anbe, P. W. Armstrong et al., “ACC/AHA guidelines for the management of patients with ST-elevation myocardial infarction-executive summary: a report of the American College of Cardiology/American Heart Association Task force on practice guidelines,” *Circulation*, vol. 110, no. 5, pp. 588–636, 2004.

[17] P. Appleby, C. Baigent, R. Collins et al., “Indications for fibrinolytic therapy in suspected acute myocardial infarction: collaborative overview of early mortality and major morbidity results from all randomised trials of more than 1000 patients,” *The Lancet*, vol. 343, no. 8893, pp. 311–322, 1994.

[18] E. Raik, “Thrombolytic therapy for myocardial infarction: standards versus clinical indicators (letter),” *Medical Journal of Australia*, vol. 169, no. 1, p. 59, 1998.

[19] J. M. Rawles, “Quantification of the benefit of earlier thrombolytic therapy: five-year results of the Grampian Region Early Anistreplase Trial (GREAT),” *Journal of the American College of Cardiology*, vol. 30, no. 5, pp. 1181–1186, 1997.

[20] S. Coccolini, G. Berti, S. Bosi, M. Pretolani, and G. Tumiotto, “Prehospital thrombolysis in rural emergency room and subsequent transport to a coronary care unit: ravenna myocardial infarction (RaMI) trial,” *International Journal of Cardiology*, vol. 49, pp. S47–S58, 1995.

[21] E. Boersma, A. C. P. Maas, J. W. Deckers, and M. L. Simoons, “Early thrombolytic treatment in acute myocardial infarction: reappraisal of the golden hour,” *The Lancet*, vol. 348, no. 9030, pp. 771–775, 1996.

[22] W. D. Weaver, M. Cerqueira, A. P. Hallstrom et al., “Prehospital-initiated vs hospital-initiated thrombolytic therapy: the myocardial infarction triage and intervention trial,” *Journal of the American Medical Association*, vol. 270, no. 10, pp. 1211–1216, 1993.

[23] Y. Fu, S. Goodman, W.-C. Chang, F. van de Werf, C. B. Granger, and P. W. Armstrong, “Time to treatment influences the impact of ST-segment resolution on one-year prognosis: insights from the Assessment of the Safety and Efficacy of a New Thrombolytic (ASSENT-2) trial,” *Circulation*, vol. 104, no. 22, pp. 2653–2659, 2001.