

Thai Registry in Acute Coronary Syndrome (TRACS) - An Extension of Thai Acute Coronary Syndrome Registry (TACS) Group: Lower In-Hospital but Still High Mortality at One-Year

Suphot Srimahachota MD*,
Smonporn Boonyaratavej MD*, Rungsrit Kanjanavanit MD**,
Piyamitr Sritara MD***, Rungroj Krittayaphong MD****,
Rapephon Kunjara-Na-Ayudhya MD*****, Pyatat Tatsanavivat MD*****
for the TR ACS group

* Cardiac Center and Division of Cardiovascular Disease, Department of Medicine, King Chulalongkorn Memorial Hospital, Chulalongkorn University, Bangkok, Thailand

** Division of Cardiology, Department of Medicine, Maharaj Nakorn Chiang Mai Hospital, Chiang Mai, Thailand

*** Division of Cardiology, Department of Medicine, Ramathibodi Hospital, Bangkok, Thailand

**** Division of Cardiology, Department of Medicine Siriraj Hospital, Bangkok, Thailand

***** Vichaiyut Hospital, Bangkok, Thailand

***** Division of Cardiology, Department of Medicine, Srinakarin Hospital, Khon Kaen University, Khon Kaen, Thailand

Background: The Thai Registry of Acute Coronary Syndrome (TRACS) registry was conducted five years after the first Thai Acute Coronary Syndrome (ACS) registry.

Objective: To describe demographics, management practices, and in-hospital outcomes of current Thai ACS patients and to seek for any significant changes in this registry from the earlier first Thai ACS registry.

Material and Method: The TRACS is a multi-centers, prospective, nation-wide registration with 39 participating medical centers. Web-based data entry was used and the data were centrally managed and analyzed.

Results: Between October 2007 and December 2008, 2,007 patients were enrolled. Fifty-five percent had ST elevation myocardial infarction (STEMI), 33% had non-ST-elevation myocardial infarction (NSTEMI), and 12% had unstable angina (UA). Overall prevalence of diabetes was 50.7%. The STEMI group was younger, predominantly male, with less diabetes than NSTEMI. At presentation, lower percent of cardiogenic shock (7.9%) and cardiac arrest (2.8%) were noted. Sixty seven percent of the STEMI received reperfusion therapy. Thrombolysis was given in 42.6% and primary percutaneous coronary intervention (PCI) was performed in 24.7% of all STEMI patients. Median door-to-needle and door-to-balloon time were 65 and 127 minutes. The median time-to-treatment was 285 min in the thrombolysis group and 324 min in the primary PCI group. Regarding NSTEMI-ACS, coronary angiography was performed in 38.4% and about one-fourth received revascularization either by PCI or bypass surgery during index admission. In-hospital mortality was 5.3% for STEMI, 5.1% for NSTEMI, and 1.7% for UA. When following the patients up to 12 months, the mortality was 14.1%, 25.0%, and 13.8% respectively.

Conclusion: The TRACS registry showed differences in demographic, management practices and in-hospital outcomes of the Thai ACS patients. Although mortality rate in this registry decreased significantly as compared to the first Thai ACS registry, the results had to be interpreted with caution because of the difference in characteristics and severity of the enrolled patients. At 12-month follow-up, the mortality rate was significantly higher in NSTEMI than STEMI or UA patients. Practice management should be considered particularly for the invasive strategy for these groups of patients.

Keywords: Acute coronary syndrome registry, ST-elevation myocardial infarction, non-ST-elevation myocardial infarction, Unstable angina

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Correspondence to:

Srimahachota S, Division of Cardiovascular Diseases, Department of Medicine King Chulalongkorn Memorial Hospital, Bangkok 10330, Thailand.

Phone & Fax: 0-2256-4291

E-mail: s_srimahachota@yahoo.co.th

The first Thai Acute Coronary Syndrome (TACS) registry, which was conducted between 2002 and 2004⁽¹⁾, demonstrated two times higher in-hospital mortality when compared with the GRACE registry⁽²⁾. However, some baseline characteristics were different particularly the severity of patients and much higher prevalence of diabetes. The door-to-balloon time in the first registry was longer than standard guideline recommendation^(3,4). The delay reperfusion affects the worst outcomes of the patients. National Health Security Office realized this situation and set-up the network fast track for ST-elevation myocardial infarction (STEMI) to minimize the door-to-balloon time and expect to increase the reperfusion rate. This article would like to compare demographics, management practices, and in-hospital outcomes with the first TACS registry.

Material and Method

The TRACS or second Thai ACS registry is a multi-centered, prospective, nation-wide registration with 39 participating medical centers. Web-based data entry was used and the data were centrally managed and analyzed. The present study was approved by the Ethics Committee and informed consents had to be obtained before the enrollment.

Participating hospitals

Thirty-nine hospitals, governmental and private, from every region in Thailand voluntarily participated in the TRACS. Each hospital had to enroll

consecutive 50 to 80 patients. The characteristics of the participating hospitals are shown in Table 1. Distribution of participating hospitals was more generalized than the first TACS registry.

Inclusion and exclusion criteria

The authors used the same inclusion criteria as described in the first TACS registry⁽¹⁾. Consecutive patients were enrolled prospectively. The index ACS symptoms, e.g. chest pain or angina equivalents, had to occur within 14 days before enrollment and accompanied by electrocardiographic ST segment deviations or T wave changes. At discharge, the patients were classified into one of the following categories: ST-segment-elevation myocardial infarction (STEMI), non-STEMI (NSTEMI), or unstable angina (UA). The only criteria diagnosis for diabetes has been changed by including the criteria of HbA1C greater than 6.5%. The authors excluded the patients who had to be re-admitted because of ACS.

Statistical analysis

This is a descriptive study. Categorical data were summarized as frequencies and percentages. Continuous variables were reported as mean \pm SD or median, 25th, and 75th percentiles. The STATA version 8 was used for data analysis.

Results

The participant hospitals in the TRACS had more distribution into regional areas and less to have

Table 1. Characteristics of participating hospitals

	1 st registry (2002-2004)	2 nd registry (2007-2008)
Number of hospitals	17	39
Location		
Metropolitan, n (%)	13 (76.5%)	16 (41.0%)
Government, n (%)	10 (58.8%)	10 (25.6%)
Private, n (%)	3 (17.6%)	6 (15.4%)
Regional government, n (%)	4 (23.5%)	23 (59.0%)
Government, n (%)	3 (75%)	22 (56.4%)
Private, n (%)	1 (25%)	1 (2.6%)
Number of hospital beds, median (IQR)	755 (418-1,110)	602 (100-2,279)
Number of CCU beds, median (IQR)	6 (5-8)	6 (0-22)
Number of ACS admission/year, mean (range)	275 (150-400)	428 (150-1,140)
Cardiac catheterization, n (%)	16 (94.1%)	22 (56.4%)
Emergency on call for primary PCI, n (%)	11 (64.7%)	17 (43.6%)
Open-heart surgery, n (%)	16 (94.1%)	22 (56.4%)

ACS = acute coronary syndrome; PCI = percutaneous coronary intervention; IQR = interquartile range

catheterization or open-heart surgery facility than the previous registry (Table 1). Between October 2007 and December 2008, 2,007 patients were consecutively enrolled. The STEMI patients were enrolled in higher percent than the previous registry (from 40.9% up to 54.9%). Average mean age had slightly decreased and males seemed to be more predominant in second registry (Table 2). Severity at presentation using Killip classification showed that patients in TRACS had less cardiogenic shock particularly in STEMI and less cardiac arrest than the first registry. Diabetes still had high prevalence in this registry (50.8%). NSTEMI had highest prevalence of diabetes (57.5%).

In the STEMI group, percent of patients who received reperfusion treatment was increased from 52.6% in first registry to 67.2% in second registry (Table 3). Of these, 42.6% received thrombolysis therapy and 24.7% received primary percutaneous coronary intervention. Median door-to-needle time has significantly improved by a decrease from 85 minutes in the first registry to 65 minutes in the second registry. However, door-to-balloon time has not changed. Streptokinase was still the thrombolysis of choice. Overall coronary angiography (CAG) was performed in only half of the patients. In non-ST elevation ACS group, CAG was also less performed than the first registry (Table 4). Only 12.9% in UA patients and 16.7% in NSTEMI patients received early invasive strategy according to the guideline. Important medications are shown in Table 5. Adenosine di-phosphate (ADP) inhibitor and statin were given in higher percent than the previous registry. However, beta-blocker and angiotensin converting enzyme inhibitor (ACEI)/angiotensin receptor blocker (ARB) agent were still used in lower percent than the guideline recommendation.

In-hospital mortality in the second registry was significantly reduced in all categories of ACS patients when compared with the first registry (Table 6). Mortality rate in the STEMI group markedly reduced from 17.0% to 5.3%. As well as for the NSTEMI group, the mortality rate decreased from 13.1% to 5.1%. Because of the less severity in the second registry due to less cardiogenic shock and less cardiac arrest at presentation, the authors classified the severity using Killip classification and looked into the in-hospital mortality in each group. The authors found that the in-hospital mortality according to Killip class was also reduced from the first registry in almost all categories. Other outcomes such as congestive heart failure, serious cardiac arrhythmia, stroke or bleeding

complications had a trend in the same direction as in-hospital mortality. Overall length of stay decreased from 10.1 ± 11.8 days to 7.4 ± 8.3 days. Fig. 1 shows the number of patients who received follow-up for six and 12 months. The mortality rate was highest in NSTEMI followed by STEMI and UA respectively (Fig. 2).

Discussion

The second Thai ACS registry (TRACS) was started five years after the first registry. There are some differences in participant hospitals and demographic data. In this registry, the distribution of participant hospitals covered the whole area of Thailand not only

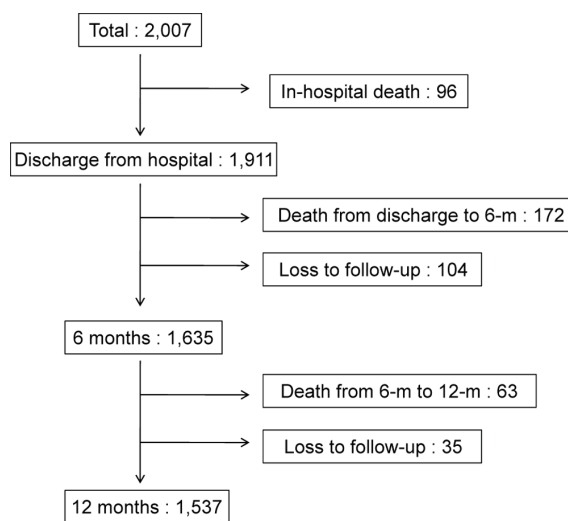


Fig. 1 Demonstrate of patients in the cohort, number of patients enrolled during admission, at discharge, 6-month follow-up and 12-month follow-up

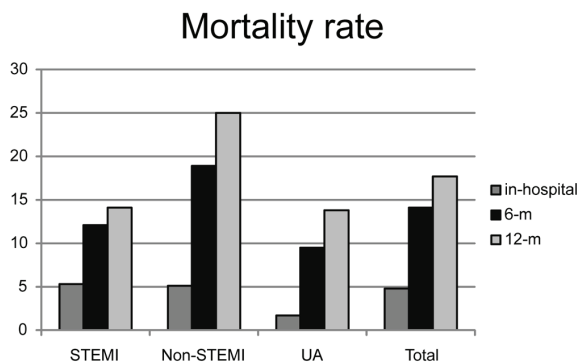


Fig. 2 Mortality rate in-hospital, 6-m and 12-m in patients with STEMI, Non-STEMI and unstable angina

Table 2. Baseline characteristics of the patients according to discharge diagnosis

	STEMI		NSTEMI		UA		Total	
	1 st registry n = 3,836 (40.9%)	2 nd registry n = 1,102 (54.9%)	1 st registry n = 3,548 (37.9%)	2 nd registry n = 664 (33.1%)	1 st registry n = 1,989 (21.2%)	2 nd registry n = 241 (12.0%)	1 st registry n = 9,373	2 nd registry n = 2,007
Mean age (y)	62.2 ± 12.8	60.9 ± 13.0	68.0 ± 11.6	67.1 ± 11.9	65.8 ± 11.0	65.5 ± 11.6	65.2 ± 12.3	63.5 ± 12.8
Male (%)	68.1	75.7	54.9	58.7	52.5	54.4	59.8	67.5
Presenting symptom								
Typical angina (%)	82.0	91.7	71.7	81.6	84.2	85.1	78.6	87.6
Atypical angina (%)	10.0	5.8	15.7	14.2	13.1	13.7	12.8	9.5
Shock (%)	16.3	16.3	6.3	9.5	1.2	5.8	9.3	12.8
Cardiac arrest (%)	7.3	3.81	2.7	1.96	0.9	0.4	4.2	2.79
Killip classification								
Killip I (%)	59.3	75.2	46.9	52.3	73.5	74.7	57.6	67.6
Killip II (%)	15.1	12.3	30.3	29.5	21.9	18.7	21.9	18.7
Killip III (%)	8.32	4.6	5.0	8.3	10.2	4.1	10.2	5.8
Killip IV (%)	17.3	7.9	17.8	9.9	1.2	2.5	10.3	7.9
Risk factors								
Diabetes (%)	37.2	47.6	50.9	57.4	45.5	46.5	44.2	50.7
Hypertension (%)	51.4	49.8	71.7	70.2	73.9	74.7	63.9	59.5
Dyslipidemia (%)	72.5	8.9	76.7	83.4	78.4	88.4	75.4	83.2
Smoking (%)	42.7	41.8	25.3	22.0	23.4	15.9	32.0	32.1
Family Hx of CAD	10.0	10.0	8.1	8.4	10.3	8.7	9.3	9.3
Refer (%)	54.2	54.0	31.4	42.5	25.1	33.2	39.5	49.3

UA = unstable angina; NSTEMI = non-ST-elevation myocardial infarction; STEMI = ST-elevation myocardial infarction; CAD = coronary artery disease

Table 3. Reperfusion treatment in STEMI

	STEMI	
	1 st registry (n = 3,836)	2 nd registry (n = 1,102)
Reperfusion treatment (%)	2,018 (52.6)	741 (67.2)
Thrombolytic (%)	1,165 (30.4)	469 (42.6)
Tissue plasminogen activator (%)	68 (1.8)	4 (0.4)
Streptokinase (%)	1,097 (28.6)	454 (41.2)
Tenecteplase	0	11 (1.0)
Door to needle time (no refer)	(n = 664)	(n = 169)
- median (min)	85.0	65.0
- mean (min) ± SD	114.0 ± 96	86.5 ± 73.4
Time to treatment (no refer)	(n = 973)	(n = 169)
- median (min)	240.0	285.0
- mean (min) ± SD	283.0 ± 190.6	224.4 ± 137.6
Door to needle time within 30 min (%)	6.0	5.9
Primary-PCI (%)	853 (22.2)	272 (24.7)
Door to balloon time (no refer)	(n = 829)	(n = 162)
- median (min)	122.0	127.0
- mean (min) ± SD	171.7 ± 180.7	183.0 ± 172.5
Time to treatment (no refer)	(n = 831)	(n = 162)
- median (min)	359.0	323.5
- mean (min) ± SD	452.0 ± 299.8	409.3 ± 300.0
Door to balloon time within 90 min (%)	34	12.3
Rescued PCI (%)	128 (3.3)	30 (2.7)
Emergency CABG (%)	94 (2.5)	6 (0.5)
Elective PCI (%)	761 (19.8)	117 (10.6)
Elective CABG (%)	136 (3.6)	13 (1.2)
CAG (%)	2,406 (62.7)	552 (50.1)
Abnormal (%)	2,349 (97.6)	535 (97.5)

Reperfusion treatment indicated the patient who presented within 12 hours from onset of chest pain or more than 12 hours but within 24 hours with persistent chest pain
 STEMI = ST-elevation myocardial infarction; PCI = percutaneous coronary intervention; CABG = coronary artery bypass grafting; CAG = coronary angiography

Table 4. The utilization of catheterization and revascularization during hospitalization in patients with NSTEMI and UA

	NSTEMI		UA	
	1 st registry, n (%)	2 nd registry, n (%)	1 st registry, n (%)	2 st registry, n (%)
Number of patients	3,548	664	1,989	241
Coronary angiography	1,556 (43.9)	256 (38.4)	920 (46.3)	82 (34.0)
Abnormal	1,492 (95.9)	248 (96.9)	801 (87.0)	75 (91.5)
Revascularization	938 (26.4)	177 (26.6)	492 (24.7)	55 (22.8)
PCI	701 (19.8)	146 (22.0)	382 (19.2)	43 (17.8)
Early invasive	464 (13.1)	111 (16.7)	224 (11.3)	31 (12.9)
Elective	237 (6.7)	35 (5.3)	158 (7.9)	12 (5.0)
CABG	285 (8.0)	31 (4.7)	111 (5.6)	12 (5.0)
Emergency	60 (1.7)	3 (0.5)	16 (0.8)	1 (0.4)
Elective	225 (6.3)	28 (4.2)	95 (4.8)	11 (4.6)

UA = unstable angina; NSTEMI = non-ST-elevation myocardial infarction; PCI = percutaneous coronary intervention; CABG = coronary artery bypass grafting

Table 5. Pharmacological treatment during hospitalization

Medications	STEMI		NSTEMI		UA	
	1 st registry	2 st registry	1 st registry	2 st registry	1 st registry	2 st registry
Number of patients	3,836	1,102	3,548	664	1,989	241
Aspirin (%)	95.2	99.5	94.6	98.6	94.1	98.8
ADP inhibitor (%)	60.4	97.8	58.5	83.4	53.5	75.9
LMWH (%)	50.7	56.7	72.2	87.1	62.3	75.5
Unfractionated heparin (%)	28.3	39.5	21.3	28.5	16.4	21.6
GP IIb/IIIa (%)	19.5	19.5	5.3	4.5	3.2	2.5
Beta-blocker (%)	58.3	59.2	61.6	62.4	71.9	71.8
ACE inhibitor (%)	59.4	66.6	57.5	58.3	55.3	54.8
ARB (%)	5.3	6.3	8.8	11.6	11.2	12.0
Statin (%)	77.5	96.1	81.5	91.4	81.7	92.5

UA = unstable angina; NSTEMI = non-ST-elevation myocardial infarction; STEMI = ST-elevation myocardial infarction; ADP = adenosine di-phosphate; LMWH = low molecular weight heparin; GP = glycoprotein; ACE = angiotensin converting enzyme; ARB = angiotensin receptor blocker

Table 6. Hospital outcomes, 6-m and 12-m mortality in patients with unstable angina, non-ST elevation myocardial infarction and ST elevation myocardial infarction

Outcomes	STEMI		NSTEMI		UA		Total	
	1 st registry	2 st registry	1 st registry	2 st registry	1 st registry	2 st registry	1 st registry	2 st registry
Number of patients	3,836	1,102	3,548	664	1,989	241	9,373	2,009
CHF (%)	44.1	27.1	56.2	50.3	27.4	28.6	45.1	34.9
CHF after 48 hrs (%)	7.8	2.4	5.5	2.6	2.9	3.3	6.0	2.5
Serious cardiac arrhythmia (%)	29.1	17.1	10.6	7.5	3.2	3.3	16.6	12.2
Heart block	11.5	6.2	3.1	1.7	1.4	1.7	6.2	4.1
Ventricular arrhythmia	19.4	8.8	8.1	1.2	1.8	1.2	11.4	6.8
Both		2.1		0.3		0.4		1.3
Stroke (%)	2.5	1.6	2.1	0.3	0.8	0	2.0	1.2
Ischemic	1.9	0.9	1.7	0.8	0.8	0	1.6	0.7
Hemorrhagic	0.6	0.7	0.1	0	0	0	0.3	0.4
Both		0		0.2		0		0.1
Unknown		0		0.2		0		0.1
Major bleeding	7.9	5.3	6.0	4.1	2.0	1.7	5.9	4.4
Length of stay (days)								
Mean ± SD	9.4 ± 12.3	6.7 ± 7.5	11.8 ± 12.6	8.8 ± 9.7	8.6 ± 8.8	6.4 ± 6.6	10.1 ± 11.8	7.4 ± 8.3
Median	6.0	4.9	8.0	5.6	6.0	4.2	6.8	5.0
Death								
In-hospital	17.0	5.3	13.1	5.1	3.0	1.7	12.6	4.8
Killip I	8.0	2.8	7.2	3.2	1.9	0.6	6.1	2.6
Killip II	13.1	6.7	11.6	3.6	2.5	2.2	10.2	4.5
Killip III	18.8	11.8	17.1	5.5	8.8	10.0	16.8	8.6
Killip IV	50.5	23.0	47.3	19.7	54.2	16.7	49.6	21.4
Cardiac	14.7	4.4	8.6	4.4	2.4	1.7	9.8	4.1
Non-cardiac	2.2	0.8	4.5	0.8	0.6	0	2.8	0.7
6-month		12.1		18.9		9.5		14.1
12-month		14.1		25.0		13.8		17.7

UA = unstable angina; NSTEMI = non-ST-elevation myocardial infarction; STEMI-ST = elevation myocardial infarction; CHF = congestive heart failure

metropolitan area, more general hospitals but also less catheterization laboratory and CABG facility. The enrollment required informed consent for this registry, which may cause selection bias by excluding the seriously conditioned patients out of the present study. Cardiogenic shock and cardiac arrest at presentation were less than the first registry. This may cause the much lower in-hospital mortality in this registry. Nevertheless, the improvement of median door-to-needle time from 85 minutes to 65 minutes can contribute to lower in-hospital mortality. As the authors know that, lower the pain-to-treatment or door-to-treatment time can lower the mortality⁽⁵⁻¹²⁾. The authors would like to give this credit to the Fast Track committee of National Health Security Office to setup network of STEMI management. This committee was developed after the first result TACS registry outcomes; unaccepted door to needle time and two to three time higher mortality than the GRACE registry⁽²⁾. This project potentiates and encourages the physicians to make rapid diagnosis and treatment for STEMI patients in Thailand.

Diabetes is still a problem for our patients in the present registry. Many reports showed poor short- and long-term outcomes⁽¹³⁻²⁴⁾. Because of using new American Diabetes Association criteria diagnosis for diabetes including the HbA1C level more than 6.5%⁽²⁵⁾, the prevalence of diabetes in this registry has increased up to 50%. Nevertheless, a higher percent of diabetes does not affect the in-hospital mortality. Anyway, the authors need to post hoc subgroup analysis in this field. In terms of medication utilization, higher percent of the patients received ADP inhibitor particularly in the STEMI as well as statin was given in more than 90% of each category of ACS. The cost of the clopidogrel is not the issue because the government started the compulsory license policy for this medication. Most of the ACS patients can be assessed under this policy. Nevertheless, beta-blocker and ACEI/ARB are still underused.

Coronary angiography in this registry was performed less than the first registry. The latest guideline^(3,4) recommended primary PCI is the reperfusion therapy of choice and pharmaco-invasive strategy within 24 hours for almost STEMI patients after thrombolysis. For high-risk non-ST elevation ACS patients as in most cases of our registry, early invasive strategy is recommended. As previously mentioned, the participant hospitals have catheterization facility less than in the first TACS registry, this may be the reason to explain the reduction of rate of CAG for these groups of patients.

Many questions came to the present study about the low in-hospital mortality as mentioned previously caused by the participant hospitals itself. The authors had post hoc analysis between old participant centers versus new participant centers. The authors selected the important factors to compare the outcomes between the old and new centers (Table 7). The in-hospital mortality was significantly lower in old centers than new centers even some of poor prognostic factors seemed to be higher in the old centers. The management practices were significantly difference. The old sites have strategy of reperfusion by using primary PCI in the majority of the patients but the new centers used thrombolysis in most cases. CAG was also performed in three fourth of the patients in old centers whereas 14.1% were performed in new centers.

At 6-month and 12-month follow-up, the mortality rate was significantly higher in the NSTEMI than STEMI or UA. The new centers also had higher mortality rate than the old centers during follow-up. Less aggressiveness perform CAG may be the factor contributing to intermediate outcome particularly in NSTEMI.

Limitations

The most consideration for this registry is all patients required informed consent. This can cause selection bias by excluding some very high-risk patients out of the present study. However, when the authors looked into the in-hospital mortality, the authors can minimize this effect by using the severity classification (Killip) to determine the mortality rate instead of overall mortality. Small sample is another issue. This registry contained sample sizes of only one-fifth of the first registry, however, the authors counteracted by enrolling more general hospitals and distributed to the whole country. Because of budget constraint, the authors hypothesized that two thousand patients do not behave differently when compared with ten thousand patients.

Conclusion

The second Thai ACS registry (TRACS) showed differences in demographic, management practices and in-hospital outcomes of the Thai ACS patients. Although mortality rate in this registry decreased significantly as compared to the first TACS registry, the results had to be interpreted with caution because of the difference in characteristics and severity of the enrolled patients. However, at the 12-month

Table 7. In-hospital outcome according to participant hospital

	2 nd ACS registry (TRACS)		p-value
	Old centers (n = 15)	New centers (n = 24)	
Total number of patients	978	1,029	
Mean age (y)	63.6 ± 12.8	63.5 ± 12.8	0.876
Male gender (%)	67.6	67.4	0.946
Killip classification			<0.001
Killip I	62.9	72.0	
Killip II	23.9	13.8	
Killip III	5.4	6.1	
Killip IV	7.8	8.1	
Cardiac arrest (%)	3.6	2.0	0.037
Diabetes (%)	58.7	43.2	<0.001
LV EF (%)	48.6 ± 14.6	49.4 ± 14.7	0.419
Refer (%)	43.1	55.3	<0.001
Reperfusion indicated (%)	76.0	83.1	0.305
Received thrombolysis	41.3	65.2	<0.001
Received 1-PCI (%)	21.7	5.8	<0.001
CAG perform	76.2	14.1	<0.001
Mortality			
In-hospital	4.6	5.0	<0.001
6-month	11.4	16.6	0.001
12-month	14.8	20.5	0.001

LV EF = left ventricular ejection fraction; 1-PCI = primary percutaneous coronary intervention; CAG = coronary angiography

follow-up, the mortality was significantly increased particularly in the NSTEMI patients.

Contributors

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The following individuals are involved in the Registry.

Steering committee

Prasart Laothavorn, Yingnoi Uboldejpracharuk, Nithi Mahanondh, Taworn Sutthichaiyakul, Rapeephon Kunjara Na Ayudhya, Chadsri Prachuabmoh.

Executive committee

Chadsri Prachuabmoh(Chair), Rapeephon Kunjara Na Ayudhya, Pyatat Tatsanavivat, Gampanart Veerakul, Damras Tresukosol, Piyamitr Sritara, Suphot Srimahachota, Sopon Sanguanwong, Rungsrit Kanjanavanit, Worachart Moleerergpoom, Pisit

Hutayanon, Watana Boonsom, Napa Sirivivattanakul.

Writing and publication committee

Tada Yipintsoi, Pyatat Tatsanavivat, Supachai Tanomsup, Piyamitr Sritara, Damras Tresukosol, Rapeephon Kunjara Na Ayudhya.

Data coordinating center

Prapanrat Kivwongngam, Institution and investigators involved in data collection: Bangkok Hospital Bangkok: Prachueb Sirivongrungson; Bangkok Hospital Phuket: Sopon Krisanarungson, Khorawit Sooklim; Bangkok Metropolitan Administration Medical College and Vajira Hospital: Watana Boonsom; Bhumibol Adulyadej Hospital: Krisada Sastravaha; Buddhachinaraj Hospital: Poj Jianmongkol; Central Chest Institute of Thailand: Boonjong Saejueng; Chaiyaphum Hospital: Moragot Pattarajpongson; Chaophraya Yommarat Hospital: Pairoj Pinjeesekikul, Aranya Kanlayanaphotporn; Chumphon Khet Udomsak Hospital: Kanokrat Petsrichan; Chiangrai Prachanukroh Hospital: Wattana Wongtheptien; Kasemrad Hospital: Suchai Kanjanatarayont, Wirote Tantikosoom; King

Chulalongkorn Memorial Hospital: Suphot Srimahachota, Polpat Euswas; Maharaj Nakorn Chiangmai Hospital: Rungsrin Kanjanavanit; Pattalung Hospital: Ketthip Buakaew; Piyavate Hospital: Paisan Bunsiricomchai; Phramongkutklao Hospital: Sopon Sanguanwong; Phranakorn Sri Ayutthaya Hospital: Chatree Jaroenchaiwattana; Police Hospital: Worachat Moreererpoog, Kasem Ratanasumawong; Prapokklao hospital: Piyapong Permlarp, Utai Puntitpong; Prae Hospital: Mongkol Maraprasertsak; Punnga: Aramwong Taweelap, Umawadee Tohmad; Rajavithi Hospital: Napa Siriwiwattanakul; Ramathibodi Hospital: Piyamitr Sritara; Ramkhamhaeng Hospital: Wasan Udayachalerm, Ekarat Sirikarin; Ranong Hospital: Norrathep Assawapatchara; Samutprakarn Hospital: Suwat Nantanawat; Saraburi Hospital: Prajak Suchatsuntorn; Sawanpracharuk Hospital: Nutt Nompannopas; Synphaet Hospital: Chatchavet Sirikharin; Siriraj Hospital: Damras Tresukosol; Songkhla Hospital: Uraivan Parinyasiri; Songklanagarind Hospital: Woravut Jintapakorn; Srinakarin Hospital, Khonkaen University: Songsak Kiatchoosakun; Suratthani Hospital: Kanchanee Janwanitsthaporn; Surin Hospital: Chalongchai Toondee, Mookda Su; Thammasat University Hospital: Pisit Hutayanon; Vajira Phuket Hospital: Nara Kingkaew; Vichaiyut Hospital: Sarayut Wiboonchutikul; Udonthani Hospital: Sumon Tangsuntornwiwat.

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Potential conflicts of interest

None.

References

- Srimahachota S, Kanjanavanit R, Boonyaratavej S, Boonsom W, Veerakul G, Tresukosol D. Demographic, management practices and in-hospital outcomes of Thai Acute Coronary Syndrome Registry (TACSR): the difference from the Western world. *J Med Assoc Thai* 2007; 90(Suppl 1): 1-11.
- Steg PG, Goldberg RJ, Gore JM, Fox KA, Eagle KA, Flather MD, et al. Baseline characteristics, management practices, and in-hospital outcomes of patients hospitalized with acute coronary syndromes in the Global Registry of Acute Coronary Events (GRACE). *Am J Cardiol* 2002; 90: 358-63.
- Van de WF, Bax J, Betriu A, Blomstrom-Lundqvist C, Crea F, Falk V, et al. Management of acute myocardial infarction in patients presenting with persistent ST-segment elevation: the Task Force on the Management of ST-Segment Elevation Acute Myocardial Infarction of the European Society of Cardiology. *Eur Heart J* 2008; 29: 2909-45.
- Kushner FG, Hand M, Smith SC Jr, King SB III, Anderson JL, Antman EM, et al. 2009 focused updates: ACC/AHA guidelines for the management of patients with ST-elevation myocardial infarction (updating the 2004 guideline and 2007 focused update) and ACC/AHA/SCAI guidelines on percutaneous coronary intervention (updating the 2005 guideline and 2007 focused update) a report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. *J Am Coll Cardiol* 2009; 54: 2205-41.
- Newby LK, Rutsch WR, Califf RM, Simoons ML, Aylward PE, Armstrong PW, et al. Time from symptom onset to treatment and outcomes after thrombolytic therapy. GUSTO-1 Investigators. *J Am Coll Cardiol* 1996; 27: 1646-55.
- O'Keefe JH Jr, Rutherford BD, McConahay DR, Ligon RW, Johnson WL Jr, Giorgi LV, et al. Early and late results of coronary angioplasty without antecedent thrombolytic therapy for acute myocardial infarction. *Am J Cardiol* 1989; 64: 1221-30.
- Weaver WD. Time to thrombolytic treatment: factors affecting delay and their influence on outcome. *J Am Coll Cardiol* 1995; 25 (7 Suppl): 3S-9S.
- Cannon CP. Time to treatment: a crucial factor in thrombolysis and primary angioplasty. *J Thromb Thrombolysis* 1996; 3: 249-55.
- Sharkey SW, Bruneete DD, Ruiz E, Hession WT, Wysham DG, Goldenberg IF, et al. An analysis of time delays preceding thrombolysis for acute myocardial infarction. *JAMA* 1989; 262: 3171-4.
- Antoniucci D, Valenti R, Migliorini A, Moschi G, Trapani M, Buonamici P, et al. Relation of time

- to treatment and mortality in patients with acute myocardial infarction undergoing primary coronary angioplasty. *Am J Cardiol* 2002; 89: 1248-52.
11. Brodie BR, Stuckey TD, Wall TC, Kissling G, Hansen CJ, Muncy DB, et al. Importance of time to reperfusion for 30-day and late survival and recovery of left ventricular function after primary angioplasty for acute myocardial infarction. *J Am Coll Cardiol* 1998; 32: 1312-9.
 12. Brodie BR, Stone GW, Morice MC, Cox DA, Garcia E, Mattos LA, et al. Importance of time to reperfusion on outcomes with primary coronary angioplasty for acute myocardial infarction (results from the Stent Primary Angioplasty in Myocardial Infarction Trial). *Am J Cardiol* 2001; 88: 1085-90.
 13. Jaffe AS, Spadaro JJ, Schechtman K, Roberts R, Geltman EM, Sobel BE. Increased congestive heart failure after myocardial infarction of modest extent in patients with diabetes mellitus. *Am Heart J* 1984; 108: 31-7.
 14. Hasdai D, Granger CB, Srivatsa SS, Criger DA, Ellis SG, Califf RM, et al. Diabetes mellitus and outcome after primary coronary angioplasty for acute myocardial infarction: lessons from the GUSTO-IIb Angioplasty Substudy. *Global Use of Strategies to Open Occluded Arteries in Acute Coronary Syndromes. J Am Coll Cardiol* 2000; 35: 1502-12.
 15. Franklin K, Goldberg RJ, Spencer F, Klein W, Budaj A, Brieger D, et al. Implications of diabetes in patients with acute coronary syndromes. The Global Registry of Acute Coronary Events. *Arch Intern Med* 2004; 164: 1457-63.
 16. Zuanetti G, Latini R, Maggioni AP, Santoro L, Franzosi MG. Influence of diabetes on mortality in acute myocardial infarction: data from the GISSI-2 study. *J Am Coll Cardiol* 1993; 22: 1788-94.
 17. Lehto S, Pyorala K, Miettinen H, Ronnema T, Palomaki P, Tuomilehto J, et al. Myocardial infarct size and mortality in patients with non-insulin-dependent diabetes mellitus. *J Intern Med* 1994; 236: 291-7.
 18. Malmberg K, Ryden L. Myocardial infarction in patients with diabetes mellitus. *Eur Heart J* 1988; 9: 259-64.
 19. Timmer JR, van der Horst IC, de Luca G, Ottervanger JP, Hoorntje JC, de Boer MJ, et al. Comparison of myocardial perfusion after successful primary percutaneous coronary intervention in patients with ST-elevation myocardial infarction with versus without diabetes mellitus. *Am J Cardiol* 2005; 95: 1375-7.
 20. Chun BY, Dobson AJ, Heller RF. The impact of diabetes on survival among patients with first myocardial infarction. *Diabetes Care* 1997; 20: 704-8.
 21. Malmberg K, Yusuf S, Gerstein HC, Brown J, Zhao F, Hunt D, et al. Impact of diabetes on long-term prognosis in patients with unstable angina and non-Q-wave myocardial infarction: results of the OASIS (Organization to Assess Strategies for Ischemic Syndromes) Registry. *Circulation* 2000; 102: 1014-9.
 22. Miettinen H, Lehto S, Salomaa V, Mahonen M, Niemela M, Haffner SM, et al. Impact of diabetes on mortality after the first myocardial infarction. The FINMONICA Myocardial Infarction Register Study Group. *Diabetes Care* 1998; 21: 69-75.
 23. Otter W, Kleybrink S, Doering W, Standl E, Schnell O. Hospital outcome of acute myocardial infarction in patients with and without diabetes mellitus. *Diabet Med* 2004; 21: 183-7.
 24. Sala J, Masia R, Gonzalez de Molina FJ, Fernandez-Real JM, Gil M, Bosch D, et al. Short-term mortality of myocardial infarction patients with diabetes or hyperglycaemia during admission. *J Epidemiol Community Health* 2002; 56: 707-12.
 25. American Diabetes Association. Executive summary: standards of medical care in diabetes—2010. *Diabetes Care* 2010; 33 (Suppl 1): S4-10.

ทะเบียนผู้ป่วยกล้ามเนื้อหัวใจขาดเลือดเฉียบพลันของไทย (TRACS) - การศึกษาเพิ่มเติมจากการศึกษาท่อนหน้า (TACS): อัตราเสียชีวิตในโรงพยาบาลลดลงแต่การเสียชีวิตที่ 1 ปี ยังคงสูง

สุพจน์ ศรีมหาโชตะ, สมนพร บุญยะรัตเวช, รังสฤษฎ์ กาญจนะวณิชย์, ปิยมิตร ศรีธรรมา, รุ่งโรจน์ กฤตยพงษ์, รพีพล กฤษกร ณ อยุธยา, ปิยทัศน์ ทศนาวิวัฒน์, สำหรับคณะวิจัย TRACS

ภูมิหลัง: การศึกษาทะเบียนผู้ป่วยกล้ามเนื้อหัวใจขาดเลือดเฉียบพลัน (TRACS) เป็นการศึกษา 5 ปีหลังจากเริ่มเก็บข้อมูลทะเบียนผู้ป่วยกล้ามเนื้อหัวใจขาดเลือดเฉียบพลันครั้งแรก (TACS)

วัตถุประสงค์: เพื่อศึกษาลักษณะข้อมูลพื้นฐาน การรักษา และ อัตราตายของผู้ป่วยกล้ามเนื้อหัวใจขาดเลือดเฉียบพลัน นอกจากนี้ ยังศึกษาความแตกต่างของผู้ป่วยกลุ่มนี้เมื่อเทียบกับการศึกษาทะเบียนผู้ป่วยครั้งแรก

วิธีการศึกษา: TRACS เป็นการศึกษาไปข้างหน้าแบบสหสถาบันทั่วประเทศไทย 39 สถาบัน โดยใช้การลงข้อมูลผ่านทางเว็บไซต์ และประมวลและวิเคราะห์ข้อมูลที่ส่งกลาง การเก็บข้อมูลจะต้องได้รับการยินยอมจากผู้ป่วยหรือญาติ

ผลการศึกษา: การเก็บข้อมูลเริ่มตั้งแต่เดือนตุลาคม พ.ศ. 2550 ถึงเดือนธันวาคม พ.ศ. 2551 โดยมีผู้ป่วยจำนวน 2,007 ราย เข้าร่วมการศึกษา ร้อยละ 55 เป็นผู้ป่วยที่มี STEMI โดยที่ ร้อยละ 33 เป็นผู้ป่วย non-STEMI และร้อยละ 12 เป็น unstable angina พบผู้ป่วยเบาหวานร้อยละ 50.7 ผู้ป่วยกลุ่ม STEMI จะมีอายุน้อยกว่า เป็นผู้ชายมากกว่า แต่มีเบาหวานน้อยกว่า เมื่อเปรียบเทียบกับทะเบียนครั้งแรก พบว่า ผู้ป่วย STEMI มีภาวะช็อก (7.9%) และหัวใจหยุดเต้น (2.8%) น้อยกว่าร้อยละ 67 ของผู้ป่วย STEMI ได้รับการรักษาด้วย reperfusion โดยที่ร้อยละ 42.6 ได้ยาละลายลิ้มเลือด และร้อยละ 24.7 ได้รับการทำบอลลูนขยายหลอดเลือดทันที ค่ามัธยฐานของเวลา door-to-needle และ door-to-balloon เท่ากับ 65 และ 127 นาที ค่ามัธยฐานของเวลา time-to-treatment ในผู้ป่วยได้ยาละลายลิ้มเลือด และผู้ที่ได้รับการทำบอลลูนเท่ากับ 285 และ 324 นาทีตามลำดับ สำหรับผู้ป่วย non-STE-ACS พบว่าได้รับการฉีดสตีดูหลอดเลือดหัวใจเท่ากับร้อยละ 38.4 และประมาณ 1 ใน 4 ได้รับการทำบอลลูน หรือ การทำผ่าตัดต่อหลอดเลือดระหว่างที่อยู่ในโรงพยาบาล อัตราตายในโรงพยาบาลเท่ากับร้อยละ 5.3, 5.1 และ 1.7 สำหรับผู้ป่วย STEMI, non-STEMI และ unstable angina เมื่อติดตามผู้ป่วยเป็นเวลา 12 เดือน พบอัตราตาย ร้อยละ 14.1, 25.0 และ 13.8 ตามลำดับ

สรุป: ทะเบียนผู้ป่วย TRACS แสดงให้เห็นความแตกต่างของข้อมูลพื้นฐาน การรักษาและอัตราตายในโรงพยาบาลเมื่อเทียบกับฐานข้อมูลครั้งแรก ในการศึกษาครั้งนี้ อัตราตายในโรงพยาบาลน้อยแต่การแปลผลต้องใช้ความระมัดระวังเนื่องจากความรุนแรงของผู้ป่วยไม่เหมือนกัน และเมื่อติดตามการรักษาไป 1 ปี พบว่า ผู้ป่วยทั้ง STEMI, non-STEMI และ unstable angina ยังมีอัตราตายที่สูงอยู่ ทำให้ต้องคำนึงถึงการรักษาผู้ป่วยกลุ่มนี้เป็นพิเศษ โดยเฉพาะการพิจารณาทำการฉีดสตีดูหลอดเลือดหัวใจ