The role of late reperfusion in ST-segment elevation myocardial infarction: a real-world retrospective cohort study

Qixin Guo 1, Jinyu Huang 1*, Yong Shen 2, Guoxin Tong 3, Hong Li 3 and Shasha Meng 3

Abstract

Background: Early reperfusion of the coronary artery has become the first choice for patients with ST-segment elevation myocardial infarction (STEMI). How to deal with patients who miss the time window for early reperfusion is still controversial. Based on real-world data, this study was conducted to explore whether percutaneous coronary intervention (PCI) has an advantage over standard drug therapy in patients who miss the optimal treatment window.

Methods: Consecutive patients who were diagnosed with STEMI and met the inclusion criteria between 2009 and 2018 in our center were retrospectively included in this cohort study. The primary endpoint events were major adverse cardiac events (MACEs), including heart failure, sudden cardiac death, malignant arrhythmia, thrombi and bleeding events during the period of admission. Secondary endpoint events were components of MACEs. At the same time, we also evaluated angina pectoris at admission and discharge through Canadian Cardiovascular Society (CCS) grading.

Results: This study enrolled 417 STEMI patients and divided them into four groups (PCI < 3 days, 14.87%; 3 days < PCI < 7 days, 21.104%; PCI > 7 days, 34.29%; MED, 29.74%). During the period of admission, MACEs occurred in 52 cases. The incidence of MACEs was 11.29, 7.95, 4.20 and 25.81% in the four respective groups (p < 0.0001). The MED group had higher rates of MACEs (OR = 3.074; 95% CI 0.1.116–8.469, p = 0.03) and cardiac death (OR = 3.027; 95% CI 1.121–8.169, p = 0.029) compared to the PCI group. Although both treatments were effective in improving CCS grade at discharge, the PCI group improved more significantly (p < 0.0001).

Conclusions: In the real world, delayed PCI can be more effective in patients with angina symptoms at discharge and reduce the incidence of MACEs and cardiac death during hospitalization. The timing of intervention was independent of the occurrence of MACEs during hospitalization and of improvement in symptoms.

Keywords: ST-segment elevation myocardial infarction, Reperfusion therapy, Percutaneous Transluminal coronary intervention
Background
STEMI is on the rise in China, and the turning point in cardiovascular events has not occurred yet. Optimal treatment for STEMI includes early reperfusion or thrombolytic therapy. This is the cornerstone of contemporary treatment of STEMI, preventing myocardial necrosis and its consequences. Considering the current national conditions in China, it is difficult to carry out thrombolysis in primary hospitals \[1\–3\]. After being transferred to the chest pain center \[4\], many patients miss the optimal PCI time or even refuse PCI. In contrast to developed countries \[5, 6\], approximately one-third of eligible patients receive primary PCI \[7\]. The rest are treated with either delayed PCI or conservative medication.

There is no definitive treatment strategy for STEMI patients who miss the optimal PCI window. Previous studies have shown that delaying PCI may be effective in maintaining cardiac function and improving cardiac remodeling in patients and can effectively alleviate electrophysiological disorders \[8–11\]. There have been some large randomized controlled trials (RCTs) that have challenged these ideas, and even though they have had similar experimental designs, they have had completely opposite results \[12–16\]. From the perspective of clinical analysis, their conclusions are very meaningful. The root cause is that RCT screening is so rigorous that the results apply only to a subset of the population. Therefore, these conclusions cannot be widely extended to clinical practice. Due to the sparse data from the real-world setting, the optimal management strategy for delayed patients with STEMI remains controversial. The aim of this study was to explore whether PCI has an advantage over standard drug therapy in a cohort of STEMI patients who miss the optimal treatment window based on real-world data from Hangzhou First People’s Hospital.

Methods
Study population
We retrospectively included all consecutive patients referred to Hangzhou First People's Hospital for further treatment of myocardial infarction between 2009 and 2018. Treatment decisions were made by the physician and the patient in consultation, and the procedure and location of the stent placement was entirely up to the surgeon. Patients who met the inclusion criteria but did not meet the exclusion criteria were enrolled. The enrolled patients were divided into four subgroups based on predesigned criteria. In total, we included a total of 417 patients treated with PCI or conservative medications. A flow chart illustrating the patient selection process is presented in Fig. 1.

Criteria for inclusion and exclusion

Inclusion criteria
The patient was diagnosed with STEMI according to the STEMI guidelines in China \[17\]
The onset of chest pain was greater than 12 h earlier

Exclusion criteria
Previous PCI or coronary artery bypass graft (CABG) or thrombolytic therapy
Myocardial reinfarction
Severe myocardial infarction complications before admission
Coronary angiography followed by no stent implantation

Treatment
All patients were treated with optimal medications, including dual antiplatelet drugs (aspirin and clopidogrel), anticoagulants, angiotensin-converting enzyme inhibitors, β receptor blockers, and lipid-lowering therapy, for as long as the heart rate and blood pressure were not adversely affected, unless the use of these medications was clearly contraindicated. The clinician determined whether to give a dose of antiplatelet drugs by predicting whether the drug would reach an effective blood concentration at the time of surgery. Low-molecular-weight heparin (LMWH) was discontinued on the morning of PCI, common heparin was used intraoperatively to maintain the patient’s activated coagulation time (ACT) level, and LMWH was continued for 3–4 days after PCI. The use of additional instruments and drugs required during the procedure was entirely up to cardiovascular interventionists, such as thrombus aspiration catheters and β2/α3 receptor blockers. When myocardial infarction was complicated with multiple lesions, the culprit diseased vessels were treated preferentially, and non-culprit lesions were treated after 1 month.

Data extraction
All blood biochemical results were obtained from the first blood samples taken within 24 h of admission. The evaluation of the coronary angiography results was performed entirely in the catheterization room by the surgeon and the first assistant. All CCS scores were assessed by the attending physician and recorded in the course of the disease. The extraction of the coronary angiography results and the CCS grades on admission and discharge were performed separately by the two authors (Qixin Guo and Yong Shen).
**Endpoint**

The primary endpoint events were major adverse cardiac events (MACEs), including heart failure, sudden cardiac death, malignant arrhythmia, thrombus and bleeding events during the period of admission. Secondary endpoint events were components of major adverse cardiac events. We also evaluated angina pectoris at admission and discharge through CCS grading.

**Statistical analysis**

Before the statistical operation, all the data were drawn into scatter graphs and tested for normality and homogeneity of variance. The main baseline characteristics of
patients are described as frequencies for categorical variables and as mean ± standard deviation (SD) for continuous variables (normally distributed) or median with interquartile range (not normally distributed). The means of different groups were compared by one-way ANOVA (normality and independence) or Kruskal-Wallis H rank sum test (independence but not normality). The comparison of multiple rates was performed using the common chi-square test. Pairwise comparisons between the groups were performed using either the chi-square test (ordered result variable) or the Mann-Whitney U test (disordered result variable) after correcting the P value.

Before we proceeded to multifactor logistic regression, we drew directed acyclic graphs (DAGs) [18, 19] to exclude possible mediating variables (Fig. 2). Then, we screened covariates through the effect change method and imported all possible covariates into the regression equation by using the

**Table 1** Effect change method

| variable | First round | second round | third round | fourth round | fifth round | sixth round | seventh round | eighth round | ninth round | tenth round |
|----------|-------------|--------------|-------------|--------------|-------------|-------------|---------------|--------------|-------------|-------------|
| EF       | 3.328       | 3.379        | 3.402       | 3.398        | 3.379       | 3.27        | 3.169         | 3.218        | 3.256       | 3.581       |
| LDL      | 2.876       | 2.871        | 2.871       |              |             |             |               |              |             |             |
| HDL      | 3.175       | 3.17         | 3.218       | 3.134        | 3.127       | 3.014       | 3.123         | 2.902        |             |             |
| UA       | 3.373       | 3.374        | 3.412       | 3.351        | 3.351       | 3.367       | 4.281         | 3.903        | 3.77        | 4.298       |
| Scr      | 3.09        | 3.095        | 3.088       | 3.067        | 3.056       | 2.964       |               |              |             |             |
| Sex      | 2.846       | 2.849        | 2.851       | 2.846        | 2.804       |             |               |              |             |             |
| Hypertension | 2.8       | 2.796        | 2.807       | 2.775        | 2.763       | 2.614       | 2.76          |               |             |             |
| Diabetes | 2.895       | 2.897        |             |              |             |             |               |              |             |             |
| Smoking  | 2.859       | 2.854        | 2.856       | 2.863        |             |             |               |              |             |             |
| Alcohol  | 2.895       |              |             |              |             |             |               |              |             |             |
| Age      | 3.144       | 3.142        | 3.155       | 3.141        | 3.141       | 2.997       | 3.154         | 2.933        | 3.074       |             |

**EF** ejection fraction, **LDL** low density lipoprotein, **HDL** high-density lipoprotein, **SCR** serum creatinine, **UA** uric acid

The OR value in the range of 2.6037 to 3.1823 indicates that the change of OR value is less than 10%. After 10 rounds of screening, EF and UA are the variables that must be included in the regression model.
enter method. The OR value of the drug treatment compared with PCI was recorded. Each variable was removed one by one, and a regression model was constructed to obtain the OR values of different treatment methods. We removed the variable that had the least effect on the OR value, and the OR value did not change by more than 10%. One by one, other variables were eliminated in the same way until all irrelevant variables were eliminated (Table 1). Finally, the selected covariates and control variables were combined to construct the regression model [20].
All statistical analyses were conducted using SPSS software (version 23.0). A two-tailed $P$-value $< 0.05$ was considered statistically significant.

**Results**

**Baseline characteristics**

Table 2 summarizes baseline patient characteristics according to treatment modality and timing of intervention. The mean age was 71 years (58–79), and 68.3% of the patients were male. In the study, 58% had a medical history of hypertension, 2.6% had hyperlipidemia, and 20.9% had diabetes. Most of the patients in the MED group were elderly (compared with the other three groups, $P < 0.0001$). Significant differences between pairs of groups were also observed in CCS rank, UA, sex, and SCR.

**Primary endpoint**

During hospitalization, 52 patients (12.47%) experienced a MACE: 26 (6.23%) had heart failure, 21 (5.04%) had cardiovascular death, 2 (0.48%) had myocardial infarction, 7 (1.68%) had malignant arrhythmia, and 6 (1.44%) had bleeding or thrombotic events (Table 3). The incidence of MACEs was 11.29, 7.95, 4.20 and 25.81% in the four respective groups ($p < 0.0001$). However, in the pairwise comparison results, no significant differences were found between the three PCI subgroups (Table 4), even after group merging. The MED group had higher rates of MACEs (OR = 3.074; 95% CI 0.1.116–8.469, $p = 0.03$) and cardiac death (OR = 3.027; 95% CI 1.121–8.169, $p = 0.029$) compared to the PCI group. Ejection fraction (EF), different treatment modalities and Uric acid (UA) were independent risk factors for MACEs in hospitals, while in-hospital deaths were only correlated with age and treatment modality. No statistically significant differences were found in the exploration results for other secondary end points (details can be found in Additional file 1).

**CCS classification score**

There were significant differences in CCS classification at discharge and admission. When the differences between the groups were further studied, we found that there was no significant difference in CCS grade between groups 2 and 3 at admission, and the overall value was group 1 > group 4 > group 2 ≈ group 3 (Table 4). The same method was used to evaluate the CCS grade at discharge, and there was no statistically significant difference in CCS grade among the three PCI groups, with $P$ values of 0.901, 0.468 and 0.491, respectively. The overall ranking was group 4 > group 1 = group 2 = group 3 (Table 4). Both conservative drug therapy and interventional therapy showed a significant decrease in CCS grading. In contrast, the interventional therapy was better alleviating patients’ subjective symptoms, but

### Table 3 Primary and secondary outcomes

| Event                              | PCI < 3 days ($n = 62$) | 3 days < PCI < 7 days ($n = 88$) | PCI > 7 days ($n = 143$) | MED ($n = 124$) | $P$     |
|------------------------------------|-------------------------|----------------------------------|-------------------------|-----------------|---------|
| MACE                               | 7 (11.29)               | 7 (7.95)                         | 6 (4.20)                | 32 (25.81)      | < 0.0001|
| Heart failure                      | 3                       | 4                                | 2                       | 17              | < 0.0001|
| Cardiac death                      | 2                       | 3                                | 2                       | 14              | 0.003   |
| Recurrent MI                        | 0                       | 0                                | 1                       | 1               | 0.328   |
| Malignant arrhythmia               | 2                       | 0                                | 0                       | 5               | 0.009   |
| Bleeding or thrombotic events      | 1                       | 0                                | 1                       | 4               | 0.192   |

MACE major adverse cardiovascular events, MI myocardial infarction

The reason why the accumulative sum of secondary endpoint events is greater than that of primary endpoint events is that patients have accumulated a variety of adverse events during hospitalization.

### Table 4 Pairwise comparison of $P$ values

| CCS1 | 2   | 3   | 4   | $P$   |
|------|-----|-----|-----|-------|
| 1    | < 0.0001 | < 0.0001 | 0.018 |
| 2    | 0.072 |       |     | 0.001 |
| 3    |       | < 0.0001 |     |       |

| CCS2 | 1   | 2   | 3   | $P$   |
|------|-----|-----|-----|-------|
| 1    | 0.901 | 0.468 | < 0.0001 |
| 2    | 0.491 |       | < 0.0001 |
| 3    |       | < 0.0001 |     |       |

| MACE  | 1   | 2   | 3   | $P$   |
|------|-----|-----|-----|-------|
| 1    | 0.491 | 0.056 | 0.022 |
| 2    | 0.23  |       | 0.001 |
| 3    |       | < 0.0001 |     |       |

| Heart failure | 1   | 2   | 3   | $P$   |
|---------------|-----|-----|-----|-------|
| 1             | 0.933 | 0.143 | 0.066 |
| 2             | 0.145 |       | 0.028 |
| 3             |       | < 0.0001 |     |       |

| Cardiac death | 1   | 2   | 3   | $P$   |
|---------------|-----|-----|-----|-------|
| 1             | 0.951 | 0.386 | 0.065 |
| 2             | 0.309 |       | 0.038 |
| 3             |       | < 0.0001 |     |       |

$1$ for PCI < 3 days group; $2$ for 3 days < PCI < 7 days; $3$ on behalf of the PCI > 7 days

CCS1: CCS score at admission
CCS2 CCS score at discharge, MACE major adverse cardiovascular events

The adjusted $P$ value was 0.008333. The results were statistically significant only if the $P$ value was less than 0.008333.
symptom alleviation had little relationship with the timing of intervention (Table 4, Fig. 3).

**Discussion**

The principal findings of this real-world study of patients with STEMI who exceeded the optimal reperfusion window were as follows. 1. PCI treatment significantly reduced the incidence of MACEs and deaths in the hospital compared to optimal drug therapy. 2. PCI can significantly improve the patient’s symptoms after treatment and increase the satisfaction with the outcome of hospitalization.

This study focuses on areas that currently are not specifically recommended in the guidelines. Based on the conclusions of previous RCTs, this study can be a good supplement. The study group missed the optimal reperfusion window, and most of the patients still had significant angina symptoms when admitted to the hospital. The disease may deteriorate at any time. For these patients, the use of reperfusion therapy as soon as possible may be able to successfully protect the patient through the crisis period. The clinician’s focus on when reperfusion offers the greatest long-term benefit may actually increase the patient’s risk. It is theoretically and ethically impossible to include this population in RCTs. The results from observational studies have high intrinsic validity and can provide important references for clinical practice.

The current guidelines recommend direct PCI for STEMI patients with chest pain up to 12 h, with the indication extending up to 48 h in some patients [21, 22]. Residual anterograde coronary artery blood flow and reverse collateral circulation after myocardial infarction might have significant clinical value.
The detailed division of the definition of myocardial infarction contributed to the accuracy of the study. Recent research on non-ST-segment elevation acute coronary syndrome showed that the delayed group did not have an increased adverse prognosis compared to the early intervention group [22], but it was significantly better than conservative treatment [23]. Depending on the pathophysiology of different myocardial infarction types, the changes in nosocomial conditions and long-term prognosis are totally different [24]. Our study found that although PCI significantly improved the incidence of endpoint events, there was no difference between the time groups of different interventions, which is inconsistent with previous research conclusions [25, 26]. The combination of the PCI < 3 days group and the 3 days < PCI < 7 days group was compared with the PCI > 7 days group, and no significant results were obtained. This may be due to the limited number of cases and lack of long-term outcomes. In summary, based on the current evidence-based medical evidence, as an important influencing factor, PCI was performed within 3 days for patients whose condition might change in a short period of time and after 7 days for patients whose condition was relatively stable, which strongly correlated with the improvements of in-hospital events.

**Limitation**

The present study has the following limitations: 1. this study is a single-center retrospective cohort study, the sample size is not large enough, and the exact results need to be supported by large-database studies. In addition, the retrospective nature of our study ensures that the results cannot be conclusive. 2. Many patients in our center were referred by local hospitals, which may result in deviations and partial data loss. For example, many patients in the group with PCI > 7 days were treated locally and transferred to our hospital for interventional surgery after stabilization. 3. During treatment, some patients were transferred to other hospitals for treatment or discharged automatically with unknown results. 4. Finally, we lost the follow-up records for our patients when we moved. Further research should focus on two areas. The first is to continue to follow up the patients and fill in the missing data. Survival analysis may yield more accurate results. Second, as cardiac intervention instruments and drugs are updated, large multicentric RCTs are also needed to guide the current treatment strategies.

**Conclusions**

In the real world, our data suggested that delayed PCI can be more effective in patients with angina symptoms at discharge and reduce the incidence and MACEs and cardiac death during hospitalization. The timing of intervention was independent of the occurrence of MACEs during hospitalization and of the improvement in symptoms. We recommend further clinical trials to confirm this conclusion.

**Supplementary information**

**Supplementary information** accompanies this paper at [https://doi.org/10.1186/s12872-020-01479-0](https://doi.org/10.1186/s12872-020-01479-0).

**Additional file 1.**

**Abbreviations**

STEMI: ST-segment elevation myocardial infarction; PCI: Percutaneous coronary intervention; MACEs: Major adverse cardiac events; CCS: Canadian cardiovascular society; RCTs: Randomized controlled trials; CABG: Coronary artery bypass graft; LMWH: Low-molecular-weight heparin; ACT: Activated coagulation time; SD: Standard deviation; DAG: Directed acyclic graphs; EF: Ejection fraction; UA: Uric acid

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**Authors’ contributions**

JYH provided the general direction of the thesis, while QXG mainly completed data extraction, statistical analysis and article writing. YS, GXT, HL and SSM provide their own Suggestions for revision when writing is completed. All authors read and approved the final version of the manuscript.

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**Availability of data and materials**

All data that support the findings of this study are included in this published article (and its supplementary information files). The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

**Ethics approval and consent to participate**

Not applicable.

**Consent for publication**

Not applicable.
Competing interests
We declare that we do not have any commercial or associative-interest that represents a conflict of interest in connection with the work submitted.

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