Association Between Postoperative Thrombocytopenia and Outcomes After Coronary Artery Bypass Grafting Surgery

Jinghang Li1†, Dongmin Yu1†, Yuanyuan Song1†, Iokfai Cheang2 and Xiaowei Wang1*

1 Department of Cardiovascular Surgery, The First Affiliated Hospital of Nanjing Medical University, Nanjing, China, 2 Department of Cardiology, The First Affiliated Hospital of Nanjing Medical University, Nanjing, China

Objectives: The effect of postoperative thrombocytopenia on adverse events among coronary artery bypass graft (CABG) patients remains unclear. This study aims to investigate the association between postoperative thrombocytopenia and perioperative outcomes of CABG.

Methods: This is a retrospective study with MIMIC-III (Medical Information Mart for Intensive Care III) database. Adult patients who underwent CABG were included to analyze the impact of thrombocytopenia in patients' outcomes. Postoperative thrombocytopenia was defined as a platelet count <100 × 10^9/L on the first day after CABG surgery. A multivariable logistic regression analysis was utilized to adjust the effect of thrombocytopenia on outcomes for baseline and covariates, and to determine the association with outcomes.

Results: A total of 4,915 patients were included, and postoperative thrombocytopenia occurred in 696 (14.2%) patients. Postoperative thrombocytopenia was not associated with increased 28-day mortality (OR 0.75; 95% CI 0.33–1.72; P = 0.496) or in-hospital mortality (OR 0.75; 95% CI 0.34–1.63; P = 0.463) after adjusting for confounders. Regarding the secondary outcomes, it was associated with a higher risk of a prolonged stay in the intensive care unit (OR 1.53; 95% CI 1.18–1.97; P = 0.001), prolonged hospital stays (OR 1.58; 95% CI 1.21–2.06; P = 0.001), prolonged mechanical ventilation time (OR 1.67; 95% CI 1.14–2.44; P = 0.009), and a trend toward increased occurrence of massive bleeding (OR 1.41; 95% CI 1.00–2.01; P = 0.054). There was no significant association between an increased risk of prolonged vasopressor use and the continuous renal replacement therapy rate.

Conclusions: Postoperative thrombocytopenia was associated with prolonged ICU and hospital stays but not with increased perioperative mortality among CABG patients.

Keywords: coronary artery bypass grafting, thrombocytopenia, mortality, MIMIC-III database, perioperation
INTRODUCTION

The causes of thrombocytopenia in cardiac surgery are multifactorial and complicated. Studies suggested inflammatory response induced by cardiopulmonary bypass (CPB) as a major factor. The activation of inflammatory cytokines (such as TNF-α, IL-8, IL-10, IL-6, and IL-1β) (1, 2) could induce systemic autoimmune platelet clearance (3, 4). The other risk factors included hemodilution, platelet consumption and destruction, heparin-induced thrombocytopenia (HITP), drug-induced thrombocytopenia (DITP), etc. (5).

A low platelet count is one common indicator of organ dysfunction in intensive care unit (ICU), which often relates to the severity and mortality of critical diseases (5). Previous studies showed that preoperative thrombocytopenia was associated with both worse bleeding and adverse outcomes in both cardiac and non-cardiac surgery (6, 7). A recent study of coronary artery bypass grafting (CABG) patients with preoperative thrombocytopenia showed a significant association with severe bleeding and increased 30-day and 1-year mortality (8).

Furthermore, CABG patients receiving antiplatelet therapy exposed to an increased risk of perioperative bleeding and reduced platelet count. As platelet count could reflect a systemic organ function, thrombocytopenia in CABG patients not only relates to bleeding events but also predicts other postoperative adverse events, including acute kidney injury, stroke, and acute myocardial infarction (9–11). Therefore, comprehensively evaluating the impact of perioperative thrombocytopenia in CABG patients is of great importance.

In this study, we characterized the incidence of postoperative thrombocytopenia on the first day after CABG surgery in the MIMIC-III database. Our study aims to explore whether thrombocytopenia in CABG patients has a relationship with an increased risk for postoperative mortality and other adverse events.

METHODS

Sources of Data

The current large single-center retrospective cohort study used publicly available data from the Medical Information Mart for Intensive Care (MIMIC) III database conducted by Beth Israel Deaconess Medical Center (BIDMC, Boston, Massachusetts). Study was in accordance with the MIMIC-III guidelines which approved by the institutional review boards. Further details on MIMIC-III ethics are available from its original publication (12). Database contains data from 46,520 ICU patients admitted to the BIDMC from 2001 to 2012, including diagnosis, demographics, procedures, vital signs, laboratory examination results, input and output information, medications, and other clinical variables.

Data Collection and Definitions

All patients undergoing isolated CABG were enrolled in this analysis. Exclusion criteria were age <18 years old; lack of platelet counts record within 24 h ICU admission, and >5% missing variables. The patients’ baseline characteristics, comorbidities, vital signs, mean arterial pressure (MAP), laboratory results, use of vasopressors, chest drainage volume, continuous renal replacement therapy (CRRT), mechanical ventilation time, etc. were collected. The sequential organ failure assessment (SOFA) score at admission to the ICU was calculated. Postoperative thrombocytopenia was defined as a platelet count <100 × 10⁹/L on the first day after CABG surgery (13).

The primary endpoints were 28-day mortality from the date of ICU admission and in-hospital mortality. Patient mortality information for discharged patients was gathered from the US Social Security Death Index. The secondary outcomes were length of ICU stay, length of hospital stay, mechanical ventilation time, vasopressor use time, chest drainage volume, and CRRT rate.

Statistical Methods

Continuous variables with non-normal distributions in the present study are presented as the median ± interquartile range. Categorical variables are presented as numbers and percentages. Comparison analyses were conducted by using the Mann-Whitney U-test for continuous variables and the χ² test or Fisher exact test for categorical variables.

Univariable logistic regression analysis was conducted to estimate the relationships between postoperative thrombocytopenia and all outcomes. Then, multivariable logistic regression analysis was performed to adjust for the following covariates: comorbidities (hypertension, congestive heart failure, diabetes, chronic pulmonary, renal failure), age, sex, body mass index (BMI), elective surgery, laboratory tests [white blood cell count (WBC), hemoglobin, lactate, activated partial thromboplastin time (APTT), prothrombin time (PT), creatinine], and vital signs [heart rate, mean arterial pressure, respiratory rate, temperature]. To facilitate logistic regression analysis, all continuous outcome variables were transformed into categorical variables according to the 75% interquartile range or clinical definition.

Statistical analysis was performed using STATA version 15 (STATA Corp LLC, College Station, TX, USA) and SPSS version 26.0 (IBM, Armonk, NY). P < 0.05 was considered statistically significant.

RESULTS

Clinical Characteristics of the Study Population

A total of 4,915 CABG patients [median age 68.5 (60.5–76.3) year-old] were included. Median platelet count was 146 [115–184] × 10⁹/L. The data selection procedure is presented in Figure 1.

There were 696 (14.2%) patients had developed postoperative thrombocytopenia. The platelet count was significantly lower in the postoperative thrombocytopenia group (84...
The procedure of data selection and exclusion.

| Variables | Thrombocytopenia | Non-thrombocytopenia | P |
|-----------|------------------|----------------------|----|
| Age, years | 73.6 (65.9, 80.0) | 67.6 (59.6, 75.4) | <0.001 |
| Gender, male, n (%) | 477 (68.5%) | 3,180 (75.4%) | <0.001 |
| BMI | 26.5 (23.6, 29.9) | 28.2 (25.1, 31.9) | <0.001 |

**Comorbidities**
- Hypertension, n (%) | 515 (74.0%) | 3,174 (75.2%) | 0.479 |
- Congestive heart failure, n (%) | 223 (32.0%) | 1,045 (24.8%) | <0.001 |
- Diabetes, n (%) | 220 (31.6%) | 1,667 (39.5%) | <0.001 |
- Chronic pulmonary disease, n (%) | 88 (12.6%) | 625 (14.8%) | 0.146 |
- Renal failure, n (%) | 75 (10.8%) | 419 (9.9%) | 0.496 |

**Laboratory tests**
- WBC, 10^9/L | 12.1 (9.4, 15.0) | 14.0 (11.1, 17.5) | <0.001 |
- Platelet, 10^9/L | 84 (71, 92) | 155 (128, 191) | <0.001 |
- Hemoglobin, g/L | 76 (67, 87) | 86 (75, 97) | <0.001 |
- Lactate, mmol/L | 3.2 (2.2, 4.4) | 2.4 (1.9, 3.2) | <0.001 |
- Creatinine, mg/dL | 1.0 (0.8, 1.2) | 0.9 (0.8, 1.2) | 0.012 |
- APTT, seconds | 47.1 (37.0, 65.2) | 37.0 (31.6, 48.2) | <0.001 |
- PT, seconds | 16.7 (15.5, 18.4) | 15.2 (14.3, 16.3) | <0.001 |

**Vital signs**
- Heart rate, bpm | 84.9 (79.2, 90.0) | 84.7 (78.9, 90.9) | 0.966 |
- MAP, mmHg | 74.3 (70.6, 78.2) | 74.2 (70.6, 78.4) | 0.633 |
- Respiratory rate, bpm | 16.1 (14.6, 18.1) | 16.9 (15.3, 18.8) | <0.001 |
- Temperature, °C | 37.7 (37.3, 38.1) | 37.8 (37.2, 38.1) | 0.04 |
- Elective surgery, n (%) | 263 (37.8%) | 1,509 (35.8%) | 0.307 |
- SOFA score | 6 (5.8) | 4 (3.8) | <0.001 |
- SOFA score>2, n (%) | 685 (98.4%) | 3,298 (78.2%) | <0.001 |

Values are expressed as median (interquartile range), or n (%).

BMI: body mass index; WBC, white blood cell; APTT, activated partial thromboplastin time; PT, prothrombin time; MAP, mean arterial pressure; SOFA, sequential organ failure assessment.

In bold are statistically significant values.

[71–92] vs. 155 [128–191] ×10^9/L, P < 0.001. The baseline characteristics between groups with and without postoperative thrombocytopenia are presented in Table 1. The postoperative thrombocytopenia group was older (median age 73.6 vs. 67.6 years old); had fewer males (68.5 vs. 75.4%); had a lower BMI, WBC, hemoglobin level, respiratory rate, and diabetes rate; and had a higher congestive heart failure rate, lactate level, APTT, PT and SOFA score. There were no significant differences in heart rate, MAP, comorbidity of hypertension, chronic pulmonary disease, renal failure, and elective surgery between groups.

**Related Risk Factors for Postoperative Thrombocytopenia**

The multivariable logistic regression analysis showed that age (OR 1.042; 95% CI 1.028–1.055), lactate (OR 1.221; 95% CI 1.143–1.306), APTT (OR 1.006; 95% CI 1.001–1.011), and
PT (OR 1.083; 95% CI 1.041–1.126) were associated with a significantly higher risk of postoperative thrombocytopenia; BMI (OR 0.963; 95% CI 0.939–0.988), WBC (OR 0.892; 95% CI 0.866–0.918), and hemoglobin (OR 0.831; 95% CI 0.759–0.910) were associated with a significantly lower risk of postoperative thrombocytopenia (Table 2).

**TABLE 2 | The risk factors of postoperative thrombocytopenia in multivariable logistic regression analysis.**

| Variables   | OR       | 95%CI      | P-value |
|-------------|----------|------------|---------|
| Age         | 1.042    | 1.028, 1.055 | <0.001  |
| BMI         | 0.963    | 0.939, 0.988 | 0.004   |
| WBC         | 0.892    | 0.866, 0.918 | <0.001  |
| Hemoglobin  | 0.831    | 0.759, 0.910 | <0.001  |
| Lactate     | 1.221    | 1.143, 1.306 | <0.001  |
| APTT        | 1.006    | 1.001, 1.011 | 0.011   |
| PT          | 1.083    | 1.041, 1.126 | <0.001  |

*Estimates are odds ratios with 95% confidence interval.*

*In bold are statistically significant values.*

Impact of Postoperative Thrombocytopenia on Primary and Secondary Outcomes

Univariate logistic regression analysis showed both the primary outcomes, 28-day mortality (3.0 vs. 1.2%, P = 0.001) and in-hospital mortality (3.4 vs. 1.4%, P = 0.001), were significantly increased in postoperative thrombocytopenia patients. While the secondary outcomes, length of ICU stay (2.8 [1.3, 4.6] vs. 2.2 [1.2, 3.9], P < 0.001), length of hospital stay (8.3 [6.2, 12.7] vs. 7.6 [5.4, 11.0], P < 0.001), length of ventilation time (12.0 [4.7, 21.7] vs. 5.9 [3.5, 15.6], P < 0.001), length of vasopressor usage (14.2 [3.0, 38.4] vs. 12.0 [1.3, 29.9], P < 0.001), and chest drainage volume (1,062.5 [762.5, 1,648.8] vs. 800 [535, 1,150], P < 0.001), were also significantly increased in postoperative thrombocytopenia patients (Table 3).

Further multivariable logistic regression analysis showed that 28-day mortality (OR 0.75; 95% CI 0.33–1.72; P = 0.495) and in-hospital mortality (OR 0.75; 95% CI 0.34–1.63; P = 0.463) did not show significance increased risk in patients with postoperative thrombocytopenia after adjusting with the confounders. For the secondary outcomes, postoperative thrombocytopenia was associated with a higher risk of an ICU stay ≥3 days (OR 1.53; 95% CI 1.18–1.97; P = 0.001), hospital stay ≥10 days (OR 1.58; 95% CI 1.21–2.06; P = 0.001), mechanical ventilation time ≥48 h (OR 1.67; 95% CI 1.14–2.44; P = 0.009), and a trend toward an increasing risk with chest drainage volume ≥1,000 ml (OR 1.41; 95% CI 1.00–2.01; P = 0.054) (Table 4).

**DISCUSSION**

In this retrospective study, postoperative thrombocytopenia (14.2%) often occurred in the first day after CAGB surgery among of 4,915 patients. The primary outcome −28-day mortality and in-hospital mortality were significantly higher in the patients with postoperative thrombocytopenia. In further multivariable logistic regression analysis, although postoperative thrombocytopenia had a significantly increased risk of prolonged ICU stay, prolonged hospital stays, prolonged mechanical ventilation time, increased risk of massive bleeding, results did not find significant difference after adjusting with cofounders in the primary outcome.

Platelets play an important role in the pathogenesis of multiple organ dysfunction, including acute respiratory distress syndrome (ARDS), acute myocardial infarction (AMI), stroke, massive bleeding, and acute kidney injury (AKI), major bleeding, and could act as a powerful prognostic indicator in critical patients (8, 10, 14–16). Therefore, hypothesis suggested that decreased platelet counts might also associated with an inflammatory end-organ injury and coagulation disorders in cardiac surgery.

However, the role of thrombocytopenia especially in postoperative CAGB patients remained controversial. A retrospective study of 4,217 adult CAGB surgery patients showed that postoperative thrombocytopenia (platelet count ≤74 × 10^9/L) was independently associated with an increased risk of 30-day mortality (10). Study of 7,189 CAGB surgery patients, preoperative thrombocytopenia (platelet count ≤150 × 10^9/L) was also associated with an increased risk of both in-hospital/30-day mortality and 1-year mortality (8). On the contrary, the ASCERT study, which included 348,341 CABG patients, did not find platelet count as a predictor of both short-term and long-term mortality after CAGB surgery (17). Our results are similar to ASCERT study, thrombocytopenia after CABG surgery did not show significance in the increased risk of short-term mortality.

Platelet as the key component in the process of hemostasis and thrombosis, thrombocytopenia is an important risk factor for postoperative bleeding in most studies of cardiac and non-cardiac surgery (18). Study reported that reduced platelet count was associated with increased postoperative bleeding after cardiac surgery (19); while prospective study of 323 cardiac surgery patients also demonstrated an association of thrombocytopenia with excessive bleeding, and that preoperative platelet count was an independent risk factor for excessive bleeding after cardiac surgery (20). Therefore, the thrombocytopenia cohort had a higher periocardial/chest drainage volume and a lower hemoglobin level, which consisted in our multivariate model. However, platelet count only reflects partial function of the platelet itself and the coagulation function (21). While thrombocytopenia does associate with the risk of adverse outcomes, proper perioperative management might significantly decrease the impact in mortality, especially for CAGB patients which usually need to receive anti-platelet management (22).

In the inflammatory aspects, cardiac surgery has a higher risk of acute lung injury (ALI) and/or acute respiratory distress syndrome (ARDS) than non-cardiac surgery and often needs longer ventilation support. The inflammatory response associated with cardiac surgery, blood transfusion and microthrombogenesis may contribute to ALI or ARDS after cardiac surgery. Previous studies have revealed that platelets also
TABLE 3 | Clinical outcomes between study cohorts.

| Outcomes                                      | Thrombocytopenia | Non-thrombocytopenia | P-value |
|-----------------------------------------------|------------------|----------------------|---------|
| **Primary outcomes**                          |                  |                      |         |
| 28-day mortality, n (%)                       | 21 (3.0%)        | 50 (1.2%)            | 0.001   |
| In-hospital mortality, n (%)                  | 24 (3.4%)        | 61 (1.4%)            | 0.001   |
| **Secondary outcomes**                        |                  |                      |         |
| Length of ICU stay, days                      | 2.8 (1.3, 4.6)   | 2.2 (1.2, 3.9)       | <0.001  |
| Length of hospital stay, days                 | 8.3 (6.2, 12.7)  | 7.6 (5.4, 11.0)      | <0.001  |
| Length of ventilation time, hours             | 12.0 (4.7, 21.7) | 5.9 (3.5, 15.6)      | <0.001  |
| Length of vasopressor time, hours             | 14.2 (3.0, 38.4) | 12.0 (1.3, 29.9)     | <0.001  |
| Pericardial and chest drainage, ml            | 1,062 (762, 1,648)| 800 (535, 1,150)     | <0.001  |
| CRRT first day                                | 11 (1.6%)        | 62 (1.5%)            | 0.738   |

Values are expressed as median (interquartile range), or n (%).
CRRT, continuous renal replacement therapy.
In bold are statistically significant values.

TABLE 4 | The associations between postoperative thrombocytopenia and clinical outcomes.

|                                      | Univariate analysis | Multivariate analysis |
|--------------------------------------|---------------------|-----------------------|
|                                      | OR (95%CI)          | P                     |
|                                      |                     |                       |
| First outcomes                       |                     |                       |
| 28-day mortality                     | 2.59 (1.55, 4.35)   | <0.001                |
| In-hospital mortality                | 2.43 (1.51, 3.93)   | <0.001                |
|                                      | 0.75 (0.33, 1.72)   | 0.496                 |
|                                      | 0.75 (0.34, 1.63)   | 0.463                 |
| Secondary outcomes                   |                     |                       |
| Length of ICU stay ≥3 days           | 1.50 (1.27, 1.76)   | <0.001                |
| Length of hospital stay ≥10 days     | 1.35 (1.15, 1.60)   | <0.001                |
| Length of ventilation time ≥48 h     | 1.99 (1.57, 2.52)   | <0.001                |
| Length of vasopressor time ≥24 h     | 1.23 (1.04, 1.48)   | 0.018                 |
| Pericardial and chest drainage ≥1,000 ml | 2.39 (1.75, 3.26) | <0.001                |
| CRRT first day                       | 1.08 (0.56, 2.06)   | 0.823                 |
|                                      | 1.13 (0.23, 3.82)   | 0.846                 |

Estimates are odds ratios with 95% confidence interval.
CRRT, continuous renal replacement therapy.
In bold are statistically significant values.

Contribute to the pathophysiological process of ALI (23). Platelets play a critical role in protecting the integrity of the basal alveolar-capillary barrier and regulating alveolar capillary permeability. When platelet counts are severely decreased, the function of the endothelial barrier is often impaired, leading to increased microvessel permeability, alveolar neutrophil infiltration, and lung injury (24). In the present study, the CABG patients with thrombocytopenia had a significantly prolonged mechanical ventilation time (12 vs. 5.9 h) and a 1.67-fold risk of mechanical ventilation time ≥48 h in the multivariate analysis.

Since postoperative thrombocytopenia count may be caused by inflammation associated with CPB, hemodilution, platelet consumption and destruction, the decrease in postoperative platelet contributes to a systemic adverse mechanism and may exert negative effects by increasing the risk of bleeding, blood transfusion, ALI, poor organ perfusion, and systemic inflammatory response. Although the direct mechanism of postoperative thrombocytopenia on these adverse events has not been well-studied. Our results provided additional evidence regarding platelet in the prognosis of CABG patients, and also established a relationship between thrombocytopenia and other major adverse events. The findings of our study add to the accumulating evidence that postoperative thrombocytopenia is an important risk factor in CABG patients.

LIMITATIONS

This study has several limitations. First, this was a retrospective study based on a public database of the MIMIC III, several potential risk factors and covariates were not available for analysis. For example, the perioperative antiplatelet drug use, types of CABG procedures (off-pump/on-pump), perioperative blood transfusion, etc. are lack in our cohort, which might generate certain bias in our results. Second, this is a single-center study, and the quality of perioperative management and surgical practices may differ from other institutions. Finally, we
only studied the impact of the platelet count on CABG surgery, and the influence of platelet function was not considered in this study. Platelet function abnormalities may be a more powerful influencing factor for advent events. Studies with multi-center large cohort, integral covariates analysis, and platelet function are needed to further explore the effects and mechanisms.

CONCLUSION

The results of this retrospective study showed that CABG patients with postoperative thrombocytopenia had a significantly increased risk of a prolonged ICU stay, prolonged hospital stay, prolonged mechanical ventilation time and massive bleeding, but the risk of 28-day and in-hospital death were not significantly increased. Further studies using data from other medical centers will be needed to provide further validation of these findings, and the potential mechanism of postoperative thrombocytopenia and its influence on adverse outcomes should be investigated.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author/s.

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ETHICS STATEMENT

Ethical review and approval was not required for the study on human participants in accordance with the local legislation and institutional requirements. Written informed consent for participation was not required for this study in accordance with the national legislation and the institutional requirements.

AUTHOR CONTRIBUTIONS

JL designed the study and coordinated the study. XW provided administrative support. DY and YS gave provision of study materials or patients. JL and IC collected and assembled the data. IC analyzed and interpreted the data. All authors were participated in the manuscript writing and read and approved the final manuscript.

ACKNOWLEDGMENTS

We appreciate the assistance of Dr. Yanxiu Li and Dr. Zhongwen Zhang in supporting the statistical analysis. We would also like to thank the initial administration support of Dr. Xiangrong Zuo and Prof. Quan Cao.
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