Predictive value of elevated alanine aminotransferase for in-hospital mortality in patients with acute myocardial infarction

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Abstract

Background and aims: Liver enzymes, including alanine aminotransferase (ALT) and aspartate aminotransferase (AST), are markers of hepatic dysfunction and fatty liver disease. Although ALT and AST have been suggested as risk factors for cardiovascular disease, their role as predictors of mortality after acute myocardial infarction (AMI) has not been established. The objective of this study was to investigate the predictive value of ALT and AST for mortality in patients with AMI.

Methods: We analyzed records of 712 patients with AMI and no known liver disease treated at the Department of Cardiovascular Center in the First Hospital of Jilin University. The primary outcome was all-cause in-hospital mortality. Relationships between primary outcome and various risk factors, including serum transaminase levels, were assessed using multivariate logistic regression analysis.

Results: Age (P < 0.001), hypertension (P = 0.034), prior myocardial infarction (P < 0.001), AST (P < 0.001), ALT (P < 0.001), creatinine (P = 0.007), blood urea nitrogen (P = 0.006), and troponin I (P < 0.001) differed significantly between ST-segment elevation myocardial infarction (STEMI) and non-STEMI. The following factors were associated with an increased risk of in-hospital all-cause mortality in patients with AMI: ALT ≥ 2ULN (adjusted odds ratio [AOR] 2.240 [95% confidence interval (CI), 1.331–3.771]; P = 0.002); age ≥ 65 year (AOR 4.320 [95% CI 2.687–6.947]; P < 0.001); increased fasting plasma glucose (FPG) (AOR 2.319 [95% CI 1.564–3.438]; P < 0.001); elevated D-dimer (AOR 2.117 [95% CI 1.407–3.184]; P < 0.001); elevated fibrinogen (AOR 1.601 [95% CI 1.077–2.380]; P = 0.20); and reduced estimated glomerular filtration rate (eGFR) (AOR 2.279 [95% CI 1.519–3.419]; P < 0.001).

Conclusions: Our findings demonstrated that elevated ALT was independently associated with increased in-hospital all-cause mortality in patients with AMI. Other risk factors were increased age, FPG, D-dimer, and fibrinogen and decreased eGFR.

Keywords: Liver enzymes, Alanine transaminase, eGFR, Fasting plasma glucose, Acute myocardial infarction, Mortality

Introduction

Coronary atherosclerotic disease (CAD) is the main source of cardiovascular disease (CVD), causing more than 7 million deaths worldwide each year. In recent decades, there have been substantial declines in cardiovascular disease mortality across much of Europe [1]. The latest guidelines from the European Heart Association, published in 2019, indicate that CAD is a pathologic
process characterized by atherosclerotic plaque accumulation in the epicardial arteries, which is either obstructive or non-obstructive [2]. Acute coronary syndrome (ACS) refers to an acute ischemic syndrome of the heart caused by new thrombosis secondary to rupture or erosion of an unstable coronary artery atherosclerotic plaque. It includes unstable angina and acute myocardial infarction (AMI), either ST-segment elevation myocardial infarction (STEMI) or non-STEMI (NSTEMI).

AMI is associated with high morbidity and mortality. Mortality is influenced by many factors, including advanced age, smoking status, and the presence of hypertension, diabetes mellitus (DM), dyslipidemia, or obesity. Elevated levels of liver transaminases, specifically alanine aminotransferase (ALT) and aspartate aminotransferase (AST), are used as markers of hepatic dysfunction and have attracted attention as emerging risk factors for CVD.

Recent studies have investigated the role of liver transaminases as independent predictors of cardiac-related morbidity and mortality [3, 4]. Several prospective epidemiologic studies have suggested that hepatic dysfunction is common in cardiac disease [5, 6]. When no other causes of liver injury are identified, elevations of liver transaminases are associated with an elevated risk of cardiac-related mortality [7]. The goal of this study was to evaluate the association between elevated liver transaminases and in-hospital all-cause mortality in patients with AMI.

Methods
Study population
A population-based retrospective case–control study was conducted in the First Hospital of Jilin University, Changchun City, Jilin Province. General characteristics and routine laboratory tests of patients diagnosed with AMI were screened to identify patients for inclusion in the study. A total of 712 patients were included in the study (Fig. 1). Baseline demographic data, medical history, laboratory data, and clinical data during hospitalization were retrieved from our departmental heart disease electronic database (Table 1) and all methods were performed in accordance with the relevant guidelines and regulations.

Definitions
The diagnostic criteria for AMI were in accordance with the 2017 guidelines from the European Society of Cardiology (ESC) [8]. Myocardial injury was defined as at least one cardiac troponin value above the 99th percentile upper reference limit. STEMI was defined as persistent chest discomfort or other symptoms suggestive of ischemia and ST-segment elevation in at least two contiguous leads. Ischemic symptoms in the absence of ST-segment elevation at presentation were designated as NSTEMI [8]. Hypoxic liver injury (HLI) was defined as at least one serum transaminase level more than twice the upper limit of normal (ULN). The ULN for ALT was defined as 50 U/L for men and 40 U/L for women, whereas the ULN for AST was defined as 40 U/L for men and 35 U/L for women. The World Health Organization (WHO) defines impaired fasting glucose as a fasting plasma glucose (FPG) value of 6.1 mmol/L or higher, whereas the American Diabetes Association recommends a cutoff value of 5.6 mmol/L [9]. In this study, we used the WHO cutoff value. Estimated glomerular filtration rate (eGFR) was calculated as 175 × plasma creatinine⁻¹.234 × age⁻⁰.⁰¹⁷⁹ × 0.79 if female [10]. Elevated D-dimer level was defined as a serum concentration ≥ 400 ng/mL and increased fibrinogen was defined as a serum concentration ≥ 3.5 g/L.

Exclusion criteria
Patients with incomplete data were excluded from the study. We also excluded patients with elevated transaminases for various non-AMI reasons, including hepatitis and hepatic cirrhosis, chronic schistosomiasis, hepatobiliary obstructive disease, bone disease, pancreatitis; cardiomyopathy, severe heart failure, severe renal insufficiency, and infectious diseases. It is well known that hyperlipidemia and fatty liver disease are risk factors for CVD. Patients diagnosed with fatty liver disease were not excluded from our study, but we did exclude patients with liver dysfunction attributed to fatty liver disease.
Laboratory data
Peripheral blood samples obtained at the time of admission were tested for the following: complete blood count, creatinine, cardiac troponin I (Tn I), creatine kinase MB (CK-MB), myoglobin, potassium, sodium, brain natriuretic peptide (BNP), prothrombin
time (PT), PT activity (PTA), fibrinogen, and D-dimer. Total cholesterol (TC), triglycerides, high-density lipoprotein cholesterol (HDL-C), low-density lipoprotein cholesterol (LDL-C), FPG, ALT, AST, alkaline phosphatase (ALP), γ-glutamyl transpeptidase (GGT), cholinesterase, albumin, total bilirubin (T-Bil), and direct bilirubin (D-Bil) were measured after 12 h of fasting following admission. All blood samples obtained at the time of admission were analyzed in the certified laboratory department of First Hospital of Jilin University.

Statistical analysis
All data analysis was performed using SPSS version 22.0, and P < 0.05 indicated statistical significance. Continuous variables were represented by the median (25th and 75th percentiles) or mean ± standard deviation. Categorical data were described by counts and percentages. Continuous variables were compared using two-tailed independent sample t-tests, and categorical variables were compared using the chi-square test. Multivariate logistic regression models were used to identify risk factors for mortality in patients with AMI. These models included adjustments for potential confounding variables, and adjusted odds ratios (AORs) and 95% confidence intervals (CIs) were calculated. The logistic regression models were calibrated by Hosmer–Lemeshow test.

Results
Demographic and clinical characteristics
Baseline demographic and clinical characteristics of the 714 patients included in the study are shown in Table 1. Of the 474 patients with STEMI, 327 were male, 147 were female, and the median age was 65.0 years. In this group, 270 patients (57.0%) had a smoking history, 133 (28.1%) had a history of drinking alcohol, 247 (52.1%) had hypertension, 135 (28.5%) had DM, 47 (9.9%) had a prior myocardial infarction, and 72 (15.2%) had a prior cerebral infarction. Of the 238 patients with NSTEMI, 153 were male and 85 were female, with a median age of 67.0 years. In this group, 138 patients (58.0%) had a smoking history, 67 (28.2%) had a history of drinking alcohol, 144 (60.5%) had hypertension, 81 (34.0%) had DM, 52 (21.8%) had a previous myocardial infarction, and 27 (11.3%) had a prior cerebral infarction.

Patients with NSTEMI were older and more likely to have DM than those in the STEMI group. AST, ALT, white blood cell count, neutrophil count, Tn I, myoglobin, CK-MB, and in-hospital mortality were also higher in the STEMI group than in the NSTEMI group.

Factors associated with in-hospital mortality in all patients
Multivariate logistic regression analysis was used to identify risk factors for in-hospital mortality in all 712 patients with AMI (Table 2). Age, sex, DM, smoking history, FPG, TC, triglycerides, AST, ALT, GGT, Tn I, D-dimer, fibrinogen, and eGFR were included in the multivariate analysis. Study participants with an ALT ≥ 2 ULN had an AOR of 2.240 [95% CI 1.331–3.771; P = 0.002], when compared with patients with an ALT < 2 ULN. Other factors associated with an increased odds of in-hospital mortality were age ≥ 65 years (AOR 4.320 [95% CI 2.687–6.947]; P < 0.001); FPG (AOR 2.319 [95% CI 1.564–3.438]; P < 0.001); decreased eGFR (AOR 2.279 [95% CI 1.519–3.419]; P < 0.001); elevated fibrinogen (AOR 1.601 [95% CI 1.077–2.380]; P = 0.020); and elevated D-dimer (AOR 2.117 [95% CI 1.407–3.184]; P < 0.001).

Figures 2, 3 and 4 show the differences in mortality rates according to age, ALT, and eGFR subgroups for the entire study population. These figures clearly illustrate the increase in mortality with increasing age and ALT and decreasing eGFR.

Factors associated with in-hospital mortality according to type of myocardial infarction
Multivariate logistic regression analysis was used to identify risk factors for all-cause in-hospital death in 474 patients with STEMI and 238 patients with NSTEMI (Tables 3 and 4). Factors included in the multivariate analysis were the same as those included when analyzing all study participants. Patients with an ALT ≥ 2 ULN had a higher risk of in-hospital mortality in both the STEMI and NSTEMI groups (AOR 1.915 [95% CI 1.028–3.565] and AOR 3.698 [95% CI 1.302–10.500], respectively; P < 0.001 for both). Other factors associated in-hospital mortality in both the STEMI and NSTEMI groups were age ≥ 65 year (AOR 4.668 [95% CI 2.684–8.117] and AOR 2.753 [95% CI 1.053–7.199], respectively); FPG (AOR 2.010 [95% CI 1.236–3.267] and AOR 3.823 [95% CI 1.820–7.808], respectively; P < 0.001 for both); and elevated D-dimer (AOR 1.937 [95% CI 1.175–3.194] and AOR 3.053 [95% CI 1.471–6.334], respectively). Elevated fibrinogen was associated with an increased odds of in-hospital mortality in patients with STEMI (AOR 1.810 [95% CI 1.113–2.944]; P = 0.017), but not in patients with NSTEMI.

Discussion
One of the most important factors associated with an increased risk of in-hospital mortality in patients with an AMI was an elevated ALT. The liver is a vital organ that is very sensitive to hemodynamic changes because of its complex vascular system and high metabolic activity.
Table 2  Multivariate analysis of variables associated with in-hospital mortality in all study patients

| Variable                  | Survivors (n = 520) | Non-survivors (n = 192) | P value | AOR (95% CI) |
|---------------------------|---------------------|-------------------------|---------|--------------|
| Age (y)                   |                     |                         | <0.001  | 4.320 (2.687–6.947) |
| < 65, n (%)               | 283 (54.4)          | 31 (16.1)               |         |              |
| ≥ 65, n (%)               | 237 (45.6)          | 161 (83.9)              |         |              |
| Sex                       |                     |                         | 0.298   | 0.795 (0.517–1.224) |
| Female, n (%)             | 149 (28.7)          | 83 (43.2)               |         |              |
| Male, n (%)               | 371 (71.3)          | 109 (56.8)              |         |              |
| DM                        |                     |                         | 0.229   | 1.317 (0.841–2.063) |
| No, n (%)                 | 387 (74.4)          | 109 (56.8)              |         |              |
| Yes, n (%)                | 133 (25.6)          |                         |         |              |
| History of smoking        |                     |                         | 0.364   | 1.317 (0.798–1.853) |
| No, n (%)                 | 208 (40.0)          | 96 (50.0)               |         |              |
| Yes, n (%)                | 312 (60.0)          | 96 (50.0)               |         |              |
| FPG (mmol/L)              |                     |                         | <0.001  | 2.319 (1.564–3.438) |
| < 6.1, n (%)              | 302 (58.1)          | 65 (33.9)               |         |              |
| ≥ 6.1, n (%)              | 218 (41.9)          | 127 (66.1)              |         |              |
| Triglycerides (mmol/L)    |                     |                         | 0.184   | 0.739 (0.472–1.155) |
| < 1.8, n (%)              | 326 (62.7)          | 142 (74.0)              |         |              |
| ≥ 1.8, n (%)              | 194 (37.3)          | 50 (26.0)               |         |              |
| TC (mmol/L)               |                     |                         | 0.689   | 1.194 (0.502–2.843) |
| < 6.6, n (%)              | 490 (94.2)          | 180 (93.7)              |         |              |
| ≥ 6.6, n (%)              | 30 (5.8)            | 12 (6.3)                |         |              |
| ALT (U/L)                 |                     |                         | 0.764   | 1.075 (0.671–1.722) |
| < 2 ULN, n (%)            | 237 (45.6)          | 79 (41.1)               |         |              |
| ≥ 2 ULN, n (%)            | 283 (54.4)          | 113 (58.9)              |         |              |
| AST (U/L)                 |                     |                         | 0.002   | 2.240 (1.331–3.771) |
| < 2 ULN, n (%)            | 432 (83.1)          | 140 (72.9)              |         |              |
| ≥ 2 ULN, n (%)            | 88 (16.9)           | 52 (27.1)               |         |              |
| GGT (U/L)                 |                     |                         | 0.643   | 1.161 (0.618–2.182) |
| < 2 ULN, n (%)            | 468 (90.0)          | 163 (84.9)              |         |              |
| ≥ 2 ULN, n (%)            | 52 (10.0)           | 29 (15.1)               |         |              |
| D-dimer (ng/mL)           |                     |                         | <0.001  | 2.117 (1.407–3.184) |
| < 400, n (%)              | 374 (71.9)          | 67 (34.9)               |         |              |
| ≥ 400, n (%)              | 146 (28.1)          | 125 (65.1)              |         |              |
| Tn I (ng/mL)              |                     |                         | 0.736   | 0.929 (0.604–1.429) |
| < 2.5, n (%)              | 266 (51.2)          | 87 (45.3)               |         |              |
| ≥ 2.5, n (%)              | 254 (48.8)          | 105 (54.7)              |         |              |
| Fibrinogen (g/L)          |                     |                         | 0.020   | 1.601 (1.077–2.380) |
| < 3.5, n (%)              | 289 (55.6)          | 73 (38.0)               |         |              |
| ≥ 3.5, n (%)              | 231 (44.4)          | 119 (62.0)              |         |              |
| eGFR (mL/min/1.73 m²)     |                     |                         | <0.001  | 2.279 (1.519–3.419) |
| ≥ 90, n (%)               | 373 (71.7)          | 65 (33.9)               |         |              |
| < 90, n (%)               | 147 (28.3)          | 127 (66.1)              |         |              |

Age, sex, DM, smoking history, FPG, TC, triglycerides, AST, ALT, GGT, Tn I, D-dimer, fibrinogen, and eGFR were included in the multivariate logistic regression analysis

ALT, alanine aminotransferase; AOR, adjusted odds ratio; AST, aspartate aminotransferase; CI, confidence interval; DM, diabetes mellitus; eGFR, estimated glomerular filtration rate; FPG, fasting plasma glucose; GGT, γ-glutamyl transpeptidase; NSTEMI, non–ST-segment elevation myocardial infarction; STEMI, ST-segment elevation myocardial infarction; T-Bil, total bilirubin; TC, total cholesterol; Tn I, cardiac troponin I; ULN, upper limit of normal
Every 10-mm Hg decrease in arterial blood pressure decreases hepatic arterial blood flow by approximately 10% [12]; therefore, abnormal serum transaminases are often observed in patients with AMI. A few recent studies reported that elevated serum transaminases were independently associated with poor clinical outcomes in patients with AMI. The extent of liver injury, in particular in patients with preexisting metabolic syndrome, may have a direct impact on cardiac outcomes [13].

Hypoxic liver injury (HLI) is commonly diagnosed in the emergency room in patients with STEMI (in 22% of patients), and those patients with HLI have higher mortality rate and higher frequency of major adverse cardiovascular events (MACE) after percutaneous coronary intervention [14]. Using multivariable regression analysis, Jantti et al. reported that a > 20% increase in ALT was associated with an increased risk of 90-day mortality independent of other known risk factors [15]. However, the association between liver transaminases and CVD remains a subject of debate. Authors of a narrative review based on prospective data concluded that the available evidence did not support a linear relationship between ALT and CVD events [16]. Differences in the effects of specific aminotransferases have not been clearly elucidated and require further research.

A few potential mechanisms have been proposed to explain the association between elevated ALT levels and increased in-hospital all-cause mortality in patients with AMI. For example, the liver has high metabolic activity and perfusion rates, and acute circulatory changes, such as AMI-related hypotension, will directly influence hepatic blood flow and thereby increase ALT and AST. Liver transaminases may also be elevated because of hepatic congestion from acute right ventricular dysfunction. Current research suggests that venous congestion, reduced capability of hepatocytes to extract oxygen, and reperfusion injury are of particular importance [12]. Although the source of elevated serum transaminases might be ischemic myocardial tissue, increased levels of these enzymes often reflect HLI secondary to both impaired forward perfusion and passive backward congestion, which are prevalent during STEMI [6]. In addition, an analysis of asymptomatic individuals found that increased levels of serum ALT (even high-normal levels) were associated with markers of CVD [17]. It is known that incomplete peripheral lipolysis of very low–density lipoprotein (VLDL) particles over-enriched with triglycerides increases the number of small dense LDL (sdLDL) particles, which carry a disproportionately large amount of LDL-C [17–19]. ALT levels are directly related to desmosterol: TC ratio, a marker of cholesterol synthetic activity [20], thereby suggesting that ALT elevations are associated with increased hepatic triglyceride

[11]. Every 10-mm Hg decrease in arterial blood pressure decreases hepatic arterial blood flow by approximately 10% [12]; therefore, abnormal serum transaminases are often observed in patients with AMI. A few recent studies reported that elevated serum transaminases were independently associated with poor clinical outcomes in patients with AMI. The extent of liver injury, in particular in patients with preexisting metabolic syndrome, may have a direct impact on cardiac outcomes [13].

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and cholesterol output in VLDL particles and lead to increased production of sdLDL [17].

In this study, we found that increased patient age was also significantly associated with mortality, which is consistent with previous reports of higher mortality in elderly patients with AMI [21–23]. In the CADIL-LAC trial, both cardiac and noncardiac causes of death were more frequent in older patients [21]. Myocytes are continuously lost as the heart ages, and senescent cells exhibit increased susceptibility to microvasculature reperfusion injury. D-dimer, one of the degradation products of fibrin, is produced when fibrinolytic enzymes degrade fibrin clots produced by thrombin. Choi et al. reported that in STEMI patients undergoing primary percutaneous coronary intervention, high D-dimer levels on admission were associated with a larger myocardial infarction.

**Table 3 Multivariate analysis of variables associated with in-hospital mortality in patients with STEMI**

| Variables | Survivors (n = 35) | Non-survivors (n = 120) | P value | AOR (95% CI) |
|-----------|-------------------|-----------------------|--------|--------------|
| Age ≥ 65 year, n (%) | 151 (42.7) | 97 (80.8) | <0.001 | 4.668 (2.684–8.117) |
| Male, n (%) | 256 (72.3) | 71 (59.2) | 0.542 | 0.848 (0.498–1.441) |
| DM, n (%) | 64 (25.4) | 45 (37.5) | 0.610 | 1.161 (0.654–2.062) |
| Smoking, n (%) | 212 (59.9) | 58 (48.3) | 0.864 | 1.046 (0.627–1.743) |
| FPG ≥ 6.1 mmol/L, n (%) | 154 (43.5) | 76 (63.3) | 0.005 | 2.010 (1.236–3.267) |
| TG ≥ 1.8 mmol/L, n (%) | 134 (37.9) | 31 (25.8) | 0.208 | 0.699 (0.400–1.221) |
| TC ≥ 6.6 mmol/L, n (%) | 19 (5.4) | 5 (4.2) | 0.930 | 0.948 (0.288–3.117) |
| AST ≥ 2 ULN, n (%) | 230 (65.0) | 76 (63.3) | 0.529 | 0.832 (0.470–1.474) |
| ALT ≥ 2 ULN, n (%) | 74 (15.6) | 27 (22.5) | 0.041 | 1.915 (1.028–3.565) |
| GGT ≥ 2 ULN, n (%) | 35 (0.9) | 18 (15.0) | 0.287 | 1.502 (0.710–3.178) |
| D-dimer ≥ 400 ng/mL, n (%) | 103 (29.1) | 75 (62.5) | 0.207 | 1.937 (1.170–3.194) |
| Tn I ≥ 2.5 ng/mL, n (%) | 35 (0.9) | 18 (15.0) | 0.287 | 1.502 (0.710–3.178) |
| Fibrinogen ≥ 3.5 g/L, n (%) | 163 (46.0) | 76 (63.3) | 0.17 | 1.810 (1.123–2.944) |
| eGFR (mL/min/1.73 m²) < 90, n (%) | 95 (26.8) | 70 (58.3) | 0.017 | 1.834 (1.112–3.023) |

**Table 4 Multivariate analysis of variables associated with in-hospital mortality in patients with NSTEMI**

| Variables | Survivors (n = 166) | Non-survivors (n = 72) | P value | AOR (95% CI) |
|-----------|-------------------|-----------------------|--------|--------------|
| Age ≥ 65y, n (%) | 86 (51.8) | 64 (88.9) | 0.039 | 2.753 (1.053–7.199) |
| Male, n (%) | 115 (69.3) | 38 (52.8) | 0.225 | 0.604 (0.267–1.365) |
| DM, n (%) | 43 (25.9) | 38 (52.8) | 0.163 | 1.786 (0.790–4.038) |
| Smoking, n (%) | 100 (60.2) | 38 (52.8) | 0.123 | 1.920 (0.838–4.403) |
| FPG ≥ 6.1 mmol/L, n (%) | 101 (60.2) | 38 (52.8) | 0.123 | 1.920 (0.838–4.403) |
| TG ≥ 1.8 mmol/L, n (%) | 60 (36.1) | 19 (26.4) | 0.886 | 1.065 (0.452–2.506) |
| TC ≥ 6.6 mmol/L, n (%) | 11 (6.6) | 7 (9.7) | 0.617 | 1.431 (0.351–5.835) |
| AST ≥ 2 ULN, n (%) | 53 (31.9) | 37 (51.4) | 0.277 | 1.681 (0.659–4.286) |
| ALT ≥ 2 ULN, n (%) | 13 (7.8) | 19 (26.4) | 0.014 | 3.698 (1.302–10.500) |
| GGT ≥ 2 ULN, n (%) | 17 (10.2) | 11 (15.3) | 0.076 | 0.280 (0.069–1.143) |
| D-dimer ≥ 400 ng/mL, n (%) | 43 (25.9) | 50 (69.4) | 0.003 | 3.053 (0.471–6.334) |
| Tn I ≥ 2.5 ng/mL, n (%) | 53 (31.9) | 28 (38.9) | 0.712 | 0.854 (0.371–1.969) |
| Fibrinogen ≥ 3.5 g/L, n (%) | 68 (41.0) | 43 (59.7) | 0.612 | 1.210 (1.578–2.533) |
| eGFR < 90 mL/min/1.73 m², n (%) | 52 (31.3) | 57 (79.2) | <0.001 | 4.063 (1.890–8.737) |

Age, sex, DM, smoking history, FPG, TC, triglycerides, AST, ALT, GGT, Tn I, D-dimer, fibrinogen, and eGFR were included in the multivariate logistic regression analysis ALT, alanine aminotransferase; AOR, adjusted odds ratio; AST, aspartate aminotransferase; CI, confidence interval; DM, diabetes mellitus; eGFR, estimated glomerular filtration rate; FPG, fasting plasma glucose; GGT, γ-glutamyl transpeptidase; NSTEMI, non-ST-segment elevation myocardial infarction; STEMI, ST-segment elevation myocardial infarction; T-Bil, total bilirubin; TC, total cholesterol; Tn I, cardiac troponin I; ULN, upper limit of normal.
size [24]. In the current study, higher D-dimer levels were associated with higher all-cause in-hospital mortality in patients with AMI; this finding is consistent with the results reported by Kikkert et al. [25].

In the current study, another important predictor of early mortality in patients with AMI was reduced eGFR. A previous study reported that among patients with AMI, any degree of preexisting renal impairment should be considered a potent, independent, and easily identifiable risk factor for cardiovascular complications, with each 10-unit reduction in eGFR being associated with a 10% increased risk for death and nonfatal cardiovascular outcomes [26]. In general, patients with chronic kidney disease have more cardiovascular risk factors and a higher incidence of MACE with AMI and acute kidney injury (AKI) was revealed to be an independent long-term prognostic indicator in patients with STEMI [27]. Even small, subclinical elevations of serum creatinine that do not fulfill the consensus criteria for acute kidney injury have been independently associated with a twofold increased risk of adverse in-hospital outcomes after AMI [28]. Our findings suggested that serum eGFR was an independent predictor of in-hospital all-cause mortality after AMI, with mortality increasing as eGFR decreased. Various factors associated with impaired renal function, such as insulin resistance, oxidative stress, inflammation, endothelial dysfunction, vascular calcifications, and hypercoagulability, may contribute to adverse outcomes in patients with ACS [29].

The 2019 ESC guidelines regarding diabetes, pre-diabetes, and cardiovascular diseases developed in collaboration with the European Association for the Study of Diabetes concluded that patients with DM and established CVD have very high cardiovascular risk [30]. Our findings that elevated FPG, but not DM, was associated with mortality suggest that glycemic control is the critical determinant of the risk of death. Previous studies have shown that severe hyperglycemia on admission was associated with increased in-hospital mortality in AMI patients with or without DM [31, 32]. Hyperglycemia can induce abnormalities in endothelial and vascular smooth muscle cell function, as well as a propensity to thrombosis, thereby contributing to atherosclerosis and its complications [33], as well as hypercoagulability and impaired fibrinolysis [34].

This study had several limitations. As a retrospective study, it relied on the accuracy of data in the medical records. It was also based on data from a single center, whose population consisted primarily of a single ethnic group of individuals located in Northeast China. Because of the climate, diet, and other factors, the incidence of cardiovascular disease in this region is significantly higher than in other regions, and the data may not reflect a general population of patients with AMI. Furthermore, although we excluded patients diagnosed with and treated for various liver diseases (including hepatobiliary malignancies, liver fibrosis and cirrhosis, and viral hepatitis) based on medical records, some factors potentially affecting transaminase levels, such as undiagnosed liver disease or medications affecting liver function, may have remained undetected and therefore not been considered when excluding patients.

Conclusions
This retrospective study focused on non-invasive markers of in-hospital death from any cause after AMI. Increased risk of in-hospital all-cause mortality was significantly associated with increased levels of ALT, older age, FPG, D-dimer, and fibrinogen and decreased eGFR. Our results therefore suggest that elevation of the serum transaminase ALT, which likely reflects HLI, may be a useful marker of increased risk of early mortality in patients with AMI.

Abbreviations
ALP: Alkaline phosphatase; ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; BNP: Brain natriuretic peptide; BUN: Blood urea nitrogen; CK-MB: Creatine kinase MB; CVD: Cardiovascular disease; D-Bil: Direct bilirubin; FPG: Fasting plasma glucose; GGT: γ-Glutamyl transpeptidase; HDL-C: High-density lipoprotein cholesterol; INR: International normalized ratio; LDL-C: Low-density lipoprotein cholesterol; MI: Myocardial infarction; NSTEMI: Non–ST-segment elevation myocardial infarction; PT: Prothrombin time; PTA: Prothrombin time activity; RBC: Red blood cell; STEMI: ST-segment elevation myocardial infarction; T-Bil: Total bilirubin; TC: Total cholesterol; Tn I: Cardiac troponin I; WBC: White blood cell; AOR: Adjusted odds ratio; CI: Confidence interval; eGFR: Estimated glomerular filtration rate; CAD: Coronary atherosclerotic disease; DM: Diabetes mellitus; HLI: Hypoxic liver injury; MACE: Major adverse cardiovascular events; ESC: European Society of Cardiology; WHO: World Health Organization.

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None.

Authors’ contributions
XL and JL made main contribution to design of the study, drafted the manuscript, and performed the statistical analysis and participated. ZZ and MJJ performed data collection and helped to draft the manuscript and make the statistical analysis. HJ and GY participated in the design and coordination of the study and participated in the data collection. All authors have read and approved the final manuscript.

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Availability of data and materials
The datasets used and analyzed during the current study are available from the corresponding author on request.
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