Elevated lipoprotein(a) levels as an independent predictor of long-term recurrent events in patients with acute coronary syndrome: an observational, retrospective cohort study

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Background Whether lipoprotein(a) [Lp(a)] is associated with recurrent cardiovascular events (RCVEs) still remains controversial. The present study aimed to investigate the prognostic value of Lp(a) for long-term RCVEs and each component of it in people with acute coronary syndrome (ACS).

Methods This multicenter, observational and retrospective study enrolled 765 ACS patients at 11 hospitals in Chengdu from January 2014 to June 2019. Patients were assigned to low-Lp(a) group [Lp(a) < 30 mg/dl] and high-Lp(a) group [Lp(a) ≥ 30 mg/dl]. The primary and secondary endpoints were defined as RCVEs and their elements, including all-cause death, nonfatal myocardial infarction (MI), nonfatal stroke and unplanned revascularization.

Results Over a median 17-month follow-up, 113 (14.8%) patients presented with RCVEs were reported, among which we observed 57 (75%) all-cause deaths, 22 (2.9%) cases of nonfatal stroke, 13 (1.7%) cases of nonfatal MI and 33 (4.3%) cases of unplanned revascularization. The incidences of RCVEs and revascularization in the high-Lp(a) group were significantly higher than those in the low-Lp(a) group (P < 0.05), whereas rates of all-cause death, nonfatal stroke and nonfatal MI were not statistically different (P > 0.05). Kaplan–Meier analysis also revealed the same trend. Multivariate Cox proportional hazards analysis showed that 1-SD increase of Lp(a) was independently associated with both the primary endpoint event [hazard ratio (HR), 1.285 per 1-SD; 95% confidence interval (CI), 1.112–1.484; P < 0.001] and revascularization (HR, 1.588 per 1-SD; 95% CI, 1.305–1.932; P < 0.001), but not with the other secondary events.

Conclusion Increased Lp(a) is an independent predictor of RCVEs and unplanned revascularization in patients with ACS. Coron Artery Dis 33: 385–393 Copyright © 2022 The Author(s). Published by Wolters Kluwer Health, Inc.

Introduction Acute coronary syndrome (ACS), including acute myocardial infarction (MI) and unstable angina, is a type of clinical critical syndrome and is a major cause of hospitality and death of coronary heart disease (CHD). More seriously, despite significant advances in the optimal secondary prevention treatment, such as statin therapy, there still remain substantial residual risks in patients who had cardiovascular events (CVEs) before [1,2]. Therefore, besides the existing indicators, it is urgent to find a new one to help predict the recurrent CVEs (RCVEs) improve the prognosis.

Recently, lipoprotein(a) [Lp(a)] has been recognized as a novel independent risk factor for the incidence of CHD [3–7], which is consisted of a low-density lipoprotein-like particle and apolipoprotein B100, with apolipoprotein(a) [apo(a)], the characteristic protein of Lp(a), covalently binding to it via a disulfide binding [8]. Compositionally, Lp(a) shows higher pathogenicity compared with LDL-cholesterol (LDL-C) in CHD due to the presence of apo(a), which is regarded as the major causative factor of atherosclerosis, thrombosis and inflammation [9]. It has been demonstrated the predictive value of Lp(a) on cardiovascular diseases (CVD) in the general people around the world and current guideline suggests Lp(a) should be one-off measured to stratify people who have a substantial lifetime risk of CVD [10]. However, controversy still exists in people who had previous CVEs, especially ACS [11–13]. Aside from the negative results, among studies that reported poor prognosis, there is also evidence to suggest differential predictive values of Lp(a) in individual components of RCVEs [7,12,14–17]. Therefore, we conducted this study to further discuss the
effects of Lp(a) on predicting RCVEs and components of it in patients with ACS.

Methods

Study design and population

This retrospective, multicenter and observational cohort study (registration number for clinical trials:ChiCTR19000025138) consecutively recruited Chinese patients with ACS [defined as ST-segment elevation MI (STEMI), non-ST-segment elevation MI and unstable angina pectoris, and definitions were determined by current guidelines [18]] from 11 tertiary general hospitals located in Chengdu between January 2014 and June 2019 (http://www.medresman.org). Patients were excluded according to the criteria as follows: (a) younger than 18 years of age; (b) severe liver and kidney diseases, severe infectious diseases, decompensated heart failure and malignant tumors; (c) incomplete plasma Lp(a) records on admission; (d) died in the hospital at baseline and (e) lost to follow-up. Finally, 765 patients were included in the analysis. After admission, all enrolled patients received optimal secondary prevention therapy. Demographic and clinical data, medical history and medicine at discharge were collected from hospital records at baseline. Since plasma Lp(a) concentrations ≥30 mg/dl have been reported to have positive relation to increased risk of CVD [6,19], the study population was assigned to high-Lp(a) group [Lp(a) ≥ 30 mg/dl] (n = 203) and low-Lp(a) group [Lp(a) < 30 mg/dl] (n = 562). The study was approved by local ethics review board.

Follow-up and clinical endpoints

Patients were followed up routinely at the data of discharge, 1, 6 and 12 months, and then annually after that. The information about RCVEs was obtained through telephone contacts with patients and their family members, or records from outpatient service and readmission. The baseline and follow-up data were collected by trained cardiovascular professionals. The primary observational endpoint of the present study was RCVEs and was the composite of all-cause death, nonfatal MI, nonfatal stroke and unplanned revascularization. The secondary observational endpoints were each component of the primary composite endpoint. Nonfatal MI was defined as the value of at least one of myocardial biomarkers greater than the upper limit of reference and with at least one of the following: (a) symptoms of myocardial ischemia and (b) electrocardiogram changes: new ST-T segment changes or left bundle branch block or emergence of pathological Q waves. Stroke was defined as acute or focal brain dysfunction with imaging changes due to a variety of vascular (ischemic or hemorrhagic) etiologies. Unplanned revascularization was defined as revascularization of any ischemic vessel by percutaneous coronary intervention (PCI) or coronary artery bypass grafting (CABG), but scheduled revascularization such as second-stage operation was not included.

Statistical analysis

Data were statistically analyzed with the SPSS version 26.0 software. Continuous data of normal or nonnormal distribution were presented as mean ± SD or median (25th–75th percentiles: Q1–Q3), and the differences of two groups were analyzed by using unpaired t-test or Mann–Whitney U test, respectively. Categorical variables were expressed as number (percentages) and examined by Chi-square test or Fisher’s exact test, as appropriate. The cumulative incidences of primary endpoint and secondary endpoints were assessed by Kaplan–Meier survival analysis, and we used the log-rank test to compare between the two groups. In order to estimate whether elevated Lp(a) was a predictor of poor prognosis, based on the univariate regression analysis for Lp(a), three models were established using Cox proportional hazards regression analysis: (a) model 1: sex, age and types of ACS were adjusted; (b) model 2: variables in model 1 and smoking, diabetes and hypertension were adjusted and (c) model 3: adjusted for variables that were included in model 2, LDL-C, serum creatinine and PCI. The association of Lp(a) with each component of RCVEs was also evaluated by model 3. Lp(a) was analyzed in two ways: (a) as a categorical variable and (b) as a continuous variable. A two-tailed P value <0.05 was considered to indicate statistical significance.

Results

Baseline characteristics of the population

A total of 765 patients were enrolled in this study, including 574 males (75%) and 191 females (25%). The average age was 65.67 ± 13.40 years. The median and mean Lp(a) level was 13.41 mg/dl (25th percentile–75th percentile: 7.14–31.05 mg/dl) and 26.04 ± 31.41 mg/dl. Lp(a) levels ranged from 0.1 to 237.0 mg/dl (Fig. 1).

The demographic, socioeconomic, clinical and medical characteristics of the study population at discharge were listed in Table 1. Patients in the high-Lp(a) group had higher prevalence of STEMI and multivessel diseases; moreover, they were presented with lower level of triglyceride. Apart from that, there were no significant differences in age, sex, smoker, pre-existing conditions, clinical symptoms, blood pressure, heart rate, other results of laboratory measurements, PCI, medicine at discharge, days in hospital and costs between the two groups (P > 0.05).

Clinical outcomes and Kaplan–Meier analysis

During the 17 months (10 and 24 months) follow-up period, 113 RCVEs (14.8%) were occurred, including 57 (7.5%) all-cause death, 22 (2.9%) nonfatal stroke, 13 (1.7%) nonfatal MI and 33 (4.3%) coronary revascularization. The
prevalence of RCVEs was higher in the high-Lp(a) group
than in the low-Lp(a) group (20.7% vs. 12.6%; \(P = 0.006\)),
and this was mainly driven by revascularization (8.9% vs.
2.7%; \(P < 0.001\)) (Fig. 2). Additionally, the Kaplan–Meier
survival curves confirmed the same trend. The curves for
primary endpoint and revascularization showed a signif-
icant difference between two groups (Fig. 3a, Log-rank
\(P = 0.002\); Fig. 3e, Log-rank \(P < 0.001\)), whereas subjects
in higher Lp(a) levels failed to show statistical distinction
of all-cause death, nonfatal stroke and MI (Fig. 3b–d, Log-
rank \(P = 0.475\); \(P = 0.776\); \(P = 0.252\), respectively).

**Cox proportional hazard regression analyses to
evaluate the association of lipoprotein(a) with
recurrent cardiovascular events**

Based on the univariate regression analysis for Lp(a),
three models constructed by using Cox proportion haz-
ard analyses (model 1–model 3, as above) were applied
to evaluate the relationship between Lp(a) and RCVEs.
Whether Lp(a) was a nominal variable or continuous
variable, univariate Cox regression analysis (Crude
Model) showed that Lp(a) was significantly correlated
with RCVEs [as a nominal variable: HR (95% CI), 1.819
(1.241–2.666); \(P = 0.002\), and as a continuous variable: HR
(95% CI), 1.255 (1.094–1.439); \(P = 0.001\)]. After adjusting
for sex, age, type of ACS, smoking, diabetes, hyperten-
sion, LDL-C, serum creatinine and PCI in model 3, this
association still existed [as a nominal variable: HR (95%
CI), 2.068 (1.366–3.132); \(P < 0.001\), and as a continuous
variable: HR (95% CI), 1.285 (1.112–1.484); \(P < 0.001\)]
(Table 2).

Furthermore, we used model 3 to evaluate the association
of Lp(a) with each component of primary endpoint. When
the RCVEs were considered separately, we observed that
only the rate of revascularization rose significantly with
elevated Lp(a) levels (\(P < 0.001\)), with a 4.387-fold higher
risk in high-Lp(a) group compared with the reference
group and a 1.588-fold higher risk 1-SD increment in
Lp(a). The results showed that higher Lp(a) levels were
independently associated with both RCVEs and revascu-
larization, but not with nonfatal stroke and MI (Table 3).

**Discussion**

This multicenter, observational study retrospectively
investigated the predictive significance of Lp(a) levels
obtained at admission for RVCEs in patients with ACS.
We demonstrated that elevated Lp(a) mass concentra-
tion was a strong independent predictor of RVCEs and
revascularization (as PCI or CABG), but not significantly
associated with all-cause death, nonfatal stroke and MI
irrespective of appropriate lipid-lowering therapy.

To the best of our knowledge, a large number of studies
have demonstrated that Lp(a) plays an important role in
the risk of CVD as primary prevention in the general peo-
ple with different races, such as White, Black, Asian and
mixed [3–5,20–22]. As for secondary prevention, the asso-
ciation between Lp(a) levels and the risk of subsequent
CVEs has been recognized in the patients with established
coronary artery diseases (CAD), even they were compli-
cated with diabetes [6,7,19,23]. However, among studies
that were targeted at subjects who had ever-experienced
CVEs revealed inconsistent results [11–15,24]. As early as
2012, Zhou et al. [11] have described the positive corre-
lation between elevated Lp(a) levels and the incidence
of consequent major adverse cardiovascular events combined with definite or suspected cardiac death, nonfatal MI, revascularization and fatal or nonfatal stroke within 713 patients. The HR (95% CI) was 1.51 (1.20–1.91) and 1.38 (1.08–1.77) for patients with ACS and those who underwent PCI, respectively [11]. Besides, several studies that enrolled ACS patients who underwent PCI also revealed patients with higher Lp(a) levels (≥20 mg/dl or ≥118 mmol/l) were associated with poor prognosis [12,15]. However, they did not find any significant distinction in death, stroke and MI between the two groups, whereas the occurrence of revascularization was obviously higher when Lp(a) concentrations were elevated. In contrast, the dal-outcomes randomized clinical trial that pooled 969 ACS patients’ and 3170 control patients’ data from 27 countries indicated that there was no association between baseline Lp(a) and ischemic CVEs defined as CAD death, nonfatal coronary event and ischemic stroke [13]. Recently, Xu et al. [24] reported that among 6714 patients who received PCI with an average of 874 days follow-up, plasma Lp(a) was not an independent predictor of long-term cardiovascular outcomes. The present study indeed found higher Lp(a) level was a useful marker for the prognosis of RCVEs. The discrepancy among the existed studies might be probably attributed to multiple confounding factors. For example, the conflicting results between our study and dal-outcomes trial could be explained by Lp(a) and potential LDL-C concentrations. Previous studies have already demonstrated that the risk ratio was continuously elevated with an increasing Lp(a) concentration, and

| Table 1 Baseline clinical characteristics grouped by lipoprotein(a) levels |
| --- |
| Variable | Low-Lp(a) group (<30 mg/dl) | High-Lp(a) group (≥30 mg/dl) | P value |
| --- | --- | --- | --- |
| Demographic | | | |
| Age, years | 65.74 ± 13.24 | 65.48 ± 13.88 | 0.813 |
| Male, n (%) | 418 (74.4) | 156 (76.6) | 0.486 |
| Social benefit | | | |
| Hospital stay, days | 9.00 (7.00, 12.00) | 9.00 (7.00, 11.00) | 0.722 |
| Hospitalized cost, 10 000 yuan | 3.52 (1.29, 4.88) | 3.83 (2.03, 5.28) | 0.058 |
| Medical history | | | |
| Smoking, n (%) | 229 (40.7) | 80 (39.4) | 0.739 |
| Diabetes, n (%) | 159 (28.3) | 46 (22.7) | 0.120 |
| Hypertension, n (%) | 331 (58.9) | 126 (62.1) | 0.430 |
| Dyslipidemia, n (%) | 82 (11.0) | 24 (12.0) | 0.710 |
| Prior CHD, n (%) | 91 (16.2) | 36 (17.7) | 0.613 |
| Prior MI, n (%) | 33 (5.9) | 15 (7.4) | 0.145 |
| Prior PCI, n (%) | 39 (7.0) | 16 (7.9) | 0.665 |
| Prior stroke, n (%) | 26 (4.6) | 9 (4.5) | 0.917 |
| Chronic obstructive pulmonary disease, n (%) | 20 (3.8) | 13 (6.4) | 0.088 |
| Peripheral arterial disease, n (%) | 8 (1.4) | 2 (1.0) | 0.919 |
| Type of ACS, n (%) | | | |
| STEMI | 304 (54.1) | 127 (62.6) | 0.037 |
| NSTEMI | 148 (26.3) | 42 (20.7) | 0.111 |
| UAP | 110 (19.6) | 34 (17.7) | 0.378 |
| Clinical presentation | | | |
| Chest pain/chest tightness | 534 (95.0) | 189 (93.1) | 0.249 |
| Dyspnea | 17 (3.0) | 10 (4.9) | 0.208 |
| Nausea and vomiting | 41 (7.3) | 14 (6.9) | 0.852 |
| Sweat | 124 (22.1) | 43 (21.2) | 0.797 |
| Systolic blood pressure, mmHg | 130.0 (115.0, 148.0) | 130.0 (115.0, 148.0) | 0.701 |
| Heart rate, beats per minute | 78.0 (68.0, 91.0) | 80.0 (69.0, 89.0) | 0.941 |
| Laboratory measurements | | | |
| Lp(a), mg/dl | 9.76 (5.62, 16.23) | 55.40 (40.46, 84.79) | <0.001 |
| Total cholesterol, mmol/l | 4.32 (3.59, 5.17) | 4.23 (3.55, 5.08) | 0.727 |
| LDL-C, mmol/l | 2.60 (1.98, 3.27) | 2.52 (2.05, 3.15) | 0.915 |
| HDL-C, mmol/l | 1.13 (0.94, 1.36) | 1.14 (0.96, 1.40) | 0.324 |
| Triglyceride, mmol/l | 1.44 (1.01, 2.17) | 1.15 (0.84, 1.78) | <0.001 |
| Serum creatinine, μmol/l | 77.3 (64.5, 94.7) | 77.9 (66.0, 97.2) | 0.633 |
| eGFR, ml/(minutes × 1.73 m²) | 85.36 (65.22, 98.74) | 84.98 (65.47, 97.46) | 0.779 |
| Fibrinogen, g/l | 3.18 (2.60, 4.02) | 3.35 (2.69, 4.43) | 0.095 |
| Hemoglobin, g/l | 133.0 (121.0, 145.0) | 131.0 (116.8, 146.3) | 0.377 |
| Multiple coronary artery lesions, n (%) | 159 (36.2) | 78 (45) | 0.047 |
| PCI, n (%) | 401 (71.4) | 196 (76.8) | 0.132 |
| Postdischarge medication | | | |
| Antiplatelet drugs, n (%) | 193 (99.0) | 526 (97.4) | 0.318 |
| Statins, n (%) | 189 (96.9) | 518 (95.9) | 0.533 |
| B-blocker, n (%) | 142 (72.8) | 384 (71.1) | 0.650 |
| ACEI/ARB, n (%) | 99 (50.8) | 270 (50.0) | 0.854 |
| Diuretics, n (%) | 43 (22.1) | 97 (18.0) | 0.213 |

ACEI, angiotensin converting enzyme inhibitor; ACS, acute coronary syndrome; ARB, angiotensin receptor blocker; CHD, coronary heart disease; eGFR, estimated glomerular filtration rate; HDL-C, HDL-cholesterol; LDL-C, LDL-cholesterol; Lp(a), lipoprotein(a); MI, myocardial infarction; NSTEMI, non-ST-segment elevation myocardial infarction; PCI, percutaneous coronary intervention; STEMI, ST-segment elevation myocardial infarction; UAP, unstable angina pectoris.
management of LDL-C was of great importance in CVD prevention [4,11,14,19,23,25,26], although Lp(a) seems to play a more important role in causing CAD [15,27,28]. The median Lp(a) and LDL-C levels of our study (13.41 mg/dl and 2.70 mmol/l) were relatively higher than those in the dal-outcomes trial (12.30 mg/dl and 1.94 mmol/l), which may properly explain the inconsistent effects. Another interference factor could be different characteristics in the enrolled patients. In a Chinese prospective study with a large sample size, patients who received PCI without a history of MI or PCI/CABG were included [24], whereas the present study enrolled subjects with ACS and did not exclude those with prior MI, PCI or CABG, causing more prevalence of adverse events. In addition, the ACS population of our study is characterized by the high use of medication [aspirin, P2Y12 inhibitors (94%) and statin (92.4%)] at discharge, which was unlikely to be the point of the disparity, although we have no information about the medication compliance during the follow-up.

In addition, we found that unplanned revascularization drove the composite outcome, which was to some extent in agreement with the former researchers [12,15]. This could be explained by the pathogenicity of Lp(a), which included atherosclerosis, inflammation and thrombosis [9]. Another possible reason may be that patients with high-Lp(a) levels were more likely to have lesions in multiple coronary arteries, and the lesions usually need to be treated several times. If these lesions were not addressed through planned revascularization, they were probably those that were addressed in unplanned revascularization.

Furthermore, several studies indicated that high-Lp(a) levels were related to the occurrence of MI, CVD death or stroke [14,16,17]. A meta-analysis enrolled CAD patients, though, showed negative findings [7]. It is concordant with what we observed in the present study, even though we found there were still stepwise increments in the HR for all-cause death and nonfatal MI, but not for nonfatal stroke [1.147, 1.096 and 0.898 for continuous Lp(a), respectively]. Notably, elevated Lp(a) was independent of the increased risk of CVD mortality and all-cause death that were reported in the preceding mentioned meta-analysis [29], implying higher Lp(a) probably was not predictive for all-cause death. Allocating fatal-MI/stroke to all-cause death may cause the predictive value of nonfatal MI/stroke to be NS. What is more, subjects with previous cerebrovascular issues had greater RCVEs risk than those with a previous ACS [14]. As the proportion of patients who had ever experienced stroke in the present study was only 4.6%, this low rate may also be part of the reason why our results differed. Because of the unified conclusions about the controversies, further studies should be explored by standardizing the assay methods of Lp(a), extending follow-up time and expanding sample's quantifies.

As is well known, plasma Lp(a) levels were genetically determined (by Lp(a) gene located on chromosome 6q26) and not reduced by any intervention, such as diet, age and exercise. By contrast, having diet in obese individuals whether with diabetes or not led to an increase in Lp(a) levels, despite improvement in LDL-C levels [30]. Likewise, statins that lower lipid-like LDL-C also cause a slight increase in Lp(a) concentrations, which could probably be explained by apo(a) expression increment and a new buffer for oxidized phospholipid in the circulation due to the decrease of LDL-C [31,32]. In order to modify Lp(a) levels, some drugs have been tested to figure out whether they did work. Niacin and proprotein convertase subtilisin/kexin type 9 (PCSK9) were reported to be associated with a significant reduction in Lp(a) levels (about 20–30%), and simultaneously in LDL-C levels, but the clinical effects
Kaplan–Meier curves for the composite endpoint of RCVEs and endpoint events that RCVEs included. (a) Kaplan–Meier curves for primary endpoint (RCVEs); (b) Kaplan–Meier curves for all-cause death; (c) Kaplan–Meier curves for nonfatal stroke; (d) Kaplan–Meier curves for nonfatal MI and (e) Kaplan–Meier curves for revascularization. Lp(a), lipoprotein(a); MI, myocardial infarction; RCVEs, recurrent cardiovascular events.
were not identical [33–38]. Niacin did not reduce the risk of major CVEs and, even worse, increased it [33–35]. Instead, PCSK9, such as evolocumab and alirocumab, made it possible to achieve favorable effects on clinical outcomes [36–38]. Recently, antisense therapy was also developed. Several randomized, double-blind and placebo-controlled trials investigated that antisense DNA oligonucleotides, like ISIS-APO(a) Rx, IONIS-APO(a) Rx and AKCEA-APO(a)-LRx, were indeed a novel, dose-dependent and potent way that benefited Lp(a) lowering without safety concerns [42]. What was interesting was that the proportion of Lp(a) lowering without safety concerns [39–41], and this was achieved by targeting apo(a), which was by binding to the complementary apo(a) mRNA sequence to lead to the reduction of translation of apo(a) [42]. What was interesting was that the proportion of Lp(a) reduction by using AKCEA-APO(a)-LRx in CVD patients was significantly larger than that by using PCSK9 (47% vs. 16%). More importantly, the former one reduced the proinflammatory activation of circulating monocytes as well [43], which may indicate a better treatment for patients with elevated Lp(a) levels. Whether isolated Lp(a)-lowering therapy would contribute to a decreased risk of CVD or not still needs to be explored.

Our findings were limited by several factors. First, the present study used retrospective data with a modest sample size, which might potentially result in selection and recall bias. Due to the limitation of sample size, we did not classify patients into more groups based on different Lp(a) levels, which may provide more information on the relation between Lp(a) concentrations and RCVEs. Second, plasma Lp(a) levels were only measured on admission, but since Lp(a) was inherently determined, it may remain relatively stable in the whole life if no Lp(a)-lowering drugs were taken. Third, we were not able to obtain the information about medication use during the follow-up time, and the adherence was unknown. The lack of aspirin, P2Y12, or statin use might influence our results to a certain degree. Fourth, since multivessel CAD probably could be found in patients with high-Lp(a) levels, unplanned revascularization may be conducted for the lesions that were already present at baseline. The missing data of coronary angiography might also affect our results. In addition, Lp(a) may be differently measured in each hospital, and that would probably bring systematic errors.

**Conclusion**

Our study found that a rise in Lp(a) levels is an independent risk factor of RCVEs in patients with ACS. Among components that made up the RCVEs, interestingly, only

| Table 2 | Different models to evaluate association of lipoprotein(a) with recurrent cardiovascular events |
|----------|------------------------------------------------------------------------------------------|
| **Model** | **LP(a) as a nominal variable** | **LP(a) as a continuous variable** |
|          | HR   | 95% CI   | P value | HR   | 95% CI   | P value |
| Crude model | 1.819 | 1.241–2.666 | 0.002 | 1.255 | 1.094–1.439 | 0.001 |
| Model 1   | 1.850 | 1.262–2.713 | 0.002 | 1.264 | 1.100–1.453 | <0.001 |
| Model 2   | 1.937 | 1.313–2.856 | <0.001 | 1.280 | 1.115–1.470 | <0.001 |
| Model 3   | 2.068 | 1.366–3.132 | <0.001 | 1.285 | 1.112–1.484 | <0.001 |

Model 1: adjusted for age, sex and type of ACS.
Model 2: adjusted for variables in model 1 and smoking, diabetes and hypertension.
Model 3: adjusted for variables in model 2 and LDL-C, serum creatinine and PCI.

ACSC, acute coronary syndrome; CI, confidence interval; HR, hazard ratio; Lp(a), lipoprotein(a); PCI, percutaneous coronary intervention.

*aThe HR was examined regarding low-Lp(a) group as reference.

*bThe HR was examined by per 1-SD increase of Lp(a).

| Table 3 | Association of lipoprotein(a) with recurrent cardiovascular events and components of recurrent cardiovascular events by using model 3 |
|----------|------------------------------------------------------------------------------------------|
| **End point** | **Univariate analysis** | **Multivariate analysis** |
|          | HR | 95% CI | P value | HR | 95% CI | P value |
| Lp(a) as a nominal variable** | | | | | | |
| Primary endpoint | 1.819 | 1.241–2.666 | 0.002 | 2.068 | 1.366–3.132 | <0.001 |
| All-cause death | 1.230 | 0.697–2.170 | 0.475 | 1.370 | 0.737–2.550 | 0.320 |
| Nonfatal stroke | 0.866 | 0.319–2.349 | 0.778 | 0.770 | 0.253–2.346 | 0.646 |
| Nonfatal MI | 1.901 | 0.621–5.813 | 0.260 | 2.239 | 0.699–7.172 | 0.175 |
| Unplanned revascularization | 3.765 | 1.896–7.476 | <0.001 | 4.387 | 2.052–9.382 | <0.001 |
| Lp(a) as a continuous variable** | | | | | | |
| Primary endpoint | 1.255 | 1.094–1.439 | 0.001 | 1.285 | 1.112–1.484 | <0.001 |
| All-cause death | 1.082 | 0.855–1.369 | 0.513 | 1.104 | 0.852–1.431 | 0.453 |
| Nonfatal stroke | 0.976 | 0.631–1.510 | 0.912 | 0.897 | 0.526–1.527 | 0.888 |
| Nonfatal MI | 1.034 | 0.626–1.711 | 0.897 | 1.081 | 0.662–1.799 | 0.733 |
| Unplanned revascularization | 1.539 | 1.279–1.851 | <0.001 | 1.588 | 1.305–1.932 | <0.001 |

CI, confidence interval; HR, hazard ratio; Lp(a), lipoprotein(a); PCI, percutaneous coronary intervention.

*aThe HR was examined regarding low-Lp(a) group as reference.

*bThe HR was examined by per 1-SD increase of Lp(a).

*cThe multivariate analysis was performed by using model 3.
revascularization showed the positive correlation with Lp(a). These findings provided additional information about potentially important role of Lp(a), which is not currently widely screened, but maybe should be done in the general population.

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Conflicts of interest
The authors declare no conflict of interest.

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