Real-world use of ACEI/ARB in diabetic hypertensive patients before the initial diagnosis of obstructive coronary artery disease: patient characteristics and long-term follow-up outcome

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Abstract

Background: Current guidelines recommend angiotensin-converting-enzyme inhibitors (ACEI)/angiotensin receptor blockers (ARB) as a first-line therapy in diabetic hypertensive patients and for secondary prevention in patients with obstructive coronary artery disease (OCAD). However, the effects of using ACEI/ARB before the initial diagnosis of OCAD on major adverse cardiac and cerebral event (MACCE) in diabetic hypertensive patients remain unclear. This study investigated whether using ACEI/ARB before the initial diagnosis of OCAD could be associated with improved clinical outcomes in diabetic hypertensive patients.

Methods: A total of 2501 patients with hypertension and diabetes, who were first diagnosed with OCAD by coronary angiography, were included in the analysis. Of the 2501 patients, 1300 did not use ACEI/ARB before the initial diagnosis of OCAD [the ACEI/ARB(-) group]; 1201 did [the ACEI/ARB(+) group]. Propensity score matching at 1:1 was performed to select 1050 patients from each group. Incidence of acute myocardial infarction (AMI), infarct size in patients with AMI, heart function, and subsequent MACCE during a median of 25.4-month follow-up were determined and compared between the 2 groups.

Results: Compared with the ACEI/ARB(-) group, the ACEI/ARB(+) group had significantly lower incidence of AMI (22.5% vs. 28.4%, p < 0.05), smaller infarct size in patients with AMI (pTNI: 5.7 vs. 6.8 ng/ml, p < 0.05; pCKMB: 21.7 vs. 28.7 ng/ml, p < 0.05), better heart function (LVEF: 60.0 vs. 58.5%, p < 0.05), and lower incidences of non-fatal stroke (2.4% vs. 4.6%, p < 0.05) and composite MACCE (23.1% vs. 29.7%, p < 0.05). No prior ACEI/ARB therapy was significantly and independently associated with non-fatal stroke and composite MACCE.

Conclusions: In diabetic hypertensive patients, treatment with ACEI/ARB before the initial diagnosis of OCAD was associated with decreased incidence of AMI, smaller infarct size, improved heart function, and lower incidences of non-fatal stroke and composite MACCE.
Background
Cardiovascular disease is the leading cause of death worldwide [1–3], especially obstructive coronary artery disease (OCAD). Hypertension and diabetes are strong independent risk factors for OCAD and associated with most of the cardiovascular death globally [4, 5]. Individuals with both hypertension and diabetes are at a higher risk of OCAD than those with either of the two conditions [6].

It has been well known that the renin-angiotensin-aldosterone system (RAAS) plays an important role in regulating cardiovascular and renal function [7, 8]. Randomized clinical trials have confirmed that suppression of RAAS activity by angiotensin-converting-enzyme inhibitors/angiotensin receptor blockers (ACEI/ARB) can protect cardio-renal function and reduce mortality [9–13]. Thus, the current guidelines recommend ACEI/ARB as a first-line therapy for diabetic hypertensive patients [14–16] and for secondary prevention in patients with OCAD [17, 18].

It has been generally accepted that diabetic hypertensive patients can benefit from ACEI/ARB; however, previous studies have found the ACEI/ARB is underutilized in these patients [19–21]. The real-world use of ACEI/ARB in diabetic hypertensive patients in China remains unclear. Diabetic hypertensive patients are prone to develop OCAD. Most previous studies have emphasized the secondary preventive effects of ACEI/ARB on OCAD. Whether starting ACEI/ARB therapy before the initial diagnosis of OCAD could improve patient outcomes is still unknown. The current study aimed to fill this knowledge gap. We used the Cardiovascular Center Beijing Friendship Hospital Database Bank to evaluate the effectiveness of ACEI/ARB therapy on improving major adverse cardiac and cerebral event (MACCE) outcomes in diabetic hypertensive patients.

Methods
Study population
Patients’ records in the Cardiovascular Center of Beijing Friendship Hospital Database Bank were screened. As shown in Fig. 1, the records of 10,098 patients undergoing coronary angiography from December 2012 to February 2019 in our center were screened. Of them, 8385 patients were diagnosed with OCAD. Of the 8385 patients, 5884 were excluded according to the exclusion criteria, which were (1) with prior diagnosis of OCAD, (2) with severe valvulopathy or cardiomyopathy and without hypertension and/or diabetes, (3) with acute infections disease, rheumatic disease, hematological disease, or neoplastic disease, (4) lacking clinical or follow-up data, and (5) with estimated glomerular filtration rate (eGFR) < 30 ml/min/1.73 m². Finally, 2501 patients were included in this analysis. Of the 2501 patients, 1300 were not treated with ACEI/ARB before the initial diagnosis of OCAD; 1201 were confirmed to receive ACEI/ARB treatment before the diagnosis. All patients were followed up to May 31, 2019 with a median follow up of 25.4 months (IQR: 12.3, 48.6 months).

Data collections and definitions
The data collection process was approved by the Institutional Review Board of Beijing Friendship Hospital affiliated to Capital Medical University and was in accordance with the Declaration of Helsinki.

Patients’ demographics, medical and medication history, laboratory test results, echocardiographic and angiographic evaluation results, and clinical outcomes during the hospitalization after the initial diagnosis of OCAD were collected and verified using an electronic medical recording system. The outcomes from MACCE were collected and recorded during clinical follow-up visits.

MACCE included all-cause death, non-fatal myocardial infarction (MI), non-fatal stroke, revascularization, and cardiac rehospitalization (admission because of angina or heart failure). All-cause death was defined as the incidence of cardiovascular death or non-cardiovascular death. Cardiovascular death was defined as fatal stroke and MI, sudden death, and other cardiovascular death. Any coronary revascularization was defined as a revascularization of the target vessel or non-target vessels. Non-fatal MI was defined as chest pain with new ST-segment changes and elevation of myocardial necrosis markers to at least twice of the upper limit of the normal range. Non-fatal stroke, including ischemic and hemorrhagic stroke, was defined as cerebral dysfunction caused by cerebral vascular obstruction or sudden rupture and was diagnosed based on signs of neurological dysfunction or evidence of brain imaging. Cardiac rehospitalization refers to rehospitalization for angina pectoris or heart failure.

Statistical analyses
Continuous variables are presented as mean ± standard deviation (SD) or median (IQR). Comparisons between
the two study groups were analyzed by Student’s t test or Mann–Whitney U-test. Categorical variables are expressed as number and percentage and compared using the Pearson Chi square test or Fisher’s exact test. To control confounding factors, we performed propensity score matching. The cumulative incidence of MACCE was estimated by Kaplan–Meier survival curves. A multivariable Cox regression analysis was performed to identify independent predictors for composite MACCE. Baseline variables that were significantly correlated with outcomes by univariate analysis and clinically relevant were used in the multivariate model. For the COX regression, the outcome event is at least 15–20 times the number of variables. Thus, the included variables for the COX regression

5884 patients were excluded, which were
1. with prior diagnosis of OCAD, n: 4095
2. with severe valvulopathy or cardiomyopathy, without hypertension and/or diabetes, n: 1680
3. with infectious disease, rheumatic disease, hematological disease or neoplastic disease, n: 15
4. lacking clinical or follow-up data, n: 56
5. with eGFR<30ml/min/1.73m², n: 38

Patients with hypertension and diabetes who were first diagnosed of OCAD by CAG, n: 2501

The ACEI/ARB(-) group, n: 1300
The ACEI/ARB(+) group, n: 1201

Fig. 1 Patient flow chart. CBD Cardiovascular Center of Beijing Friendship Hospital Database, CAG coronary angiography, OCAD obstructive coronary artery disease, eGFR estimated glomerular filtration rate, ACEI/ARB angiotensin-converting enzyme inhibitors/angiotensin receptor blockers
were carefully chosen, given the number of events available, to ensure parsimony of the final model. All analyses were two-tailed and P value < 0.05 was considered statistically significant.

**Propensity score matching**

Propensity score matching was used to reduce selection bias in this study. The matching process was conducted with a minimum-distance scoring method and a 1-to-1 match between the ACEI/ARB(-) group and the ACEI/ARB(+) group. In this study, propensity scores were calculated through a binary logistic regression model, including covariates of age, sex, body mass index (BMI), fasting blood glucose (FBG), hemoglobin (HGB), eGFR, low-density lipoprotein cholesterol (LDL-C), history of smoking and dyslipidemia, previous medication history including antiplatelet agent, beta-blocker, and statins. Ultimately, 1050 ACEI/ARB(+) patients were individually 1:1 matched to 1050 ACEI/ARB(-) controls using nearest available score matching. The statistical analysis software SPSS version 24.0 was used for the matching.

**Results**

**Patient characteristics**

As shown in Fig. 1, of the 2501 eligible patients, 1201 patients (48.0%) used ACEI/ARB before the hospital admission; 1300 (52.0%) did not. Comparing with the ACEI/ARB (+) group, the ACEI/ARB (-) group showed significantly higher percent of male, lower BMI, higher heart rate, lower percent of dyslipidemia, and significantly less likely to receive antiplatelet therapy, beta-blocker or statins before the hospital admission for OCAD. In-hospital medical and interventional treatments were similar between the 2 groups except that significantly fewer patients treated with ACEI/ARB in the ACEI/ARB (-) group than in the ACEI/ARB (+) group (52.1% vs. 83.9%, p < 0.001) during hospitalization. Subjects in the ACEI/ARB (-) group had a significant longer average hospital stay (Table 1).

As presented in Table 2, the ACEI/ARB(-) group had significantly higher white cell count, neutrophil count and higher levels of sensitivity C reactive protein (hsCRP), HGB, FBG, random blood glucose (RBG) at admission, eGFR, and LDL-C than the ACEI/ARB(+) group. Echo evaluation showed that the ACEI/ARB(-) group had significantly larger left ventricular end-systolic diameter (LVESD) and left ventricular end-systolic volume (LVESV) and lower left ventricular ejection fraction (LVEF), left ventricular fraction shortening (LVFS) and stroke volume (SV) than the ACEI/ARB(+) group. Angiographically, there was no significant difference between the 2 groups.

Correlation analysis of ACEI/ARB therapy and baseline variables revealed that patients with BMI ≥ 25 kg/m², previous use of antiplatelet agent, beta-blocker and statins were more likely to receive ACEI/ARB therapy before the hospital admission. However, patients with AMI at admission and a previous history of using calcium channel blocker (CCB) were less likely to receive ACEI/ARB therapy before the hospital admission (Fig. 2).

**Propensity score matching**

Propensity scores of 1050 ACEI/ARB users were 1:1 matched to 1050 patients without using ACEI/ARB before the initial diagnosis of OCAD. There were no significant differences in baseline clinical characteristics and medical history between the propensity score matched (PSM) ACEI/ARB(-) and ACEI/ARB(+) groups except that the PSM ACEI/ARB(-) group had significantly fewer patients treated with ACEI/ARB therapy during the hospitalization (51.0% vs. 83.9%, p < 0.001, Table 1).

The ACEI/ARB(-) group had significantly higher hsCRP levels than the ACEI/ARB(+) group. Echo evaluation showed that the ACEI/ARB(-) group had significantly larger LVESD, lower LVEF and LVFS than the ACEI/ARB(+) group (Table 2).

The ACEI/ARB(-) group had a significantly higher incidence of acute myocardial infarction (AMI) at the admission than the ACEI/ARB(+) group (28.4% vs. 22.5%, p < 0.05, Fig. 3). The peak levels of serum myoglobin (Myo), creatinine kinase MB (CKMB), and cardiac troponin I (cTnI) were used to estimate infarct size. We found no difference in pMyo between the 2 groups. The peak levels of serum CKMB and cTnI were significantly higher in the ACEI/ARB(-) group (p-CKMB: 28.7 vs. 21.7 ng/mL, p < 0.05; p-cTnI: 6.8 vs. 5.7 ng/mL, p < 0.05, Table 3).

**In-hospital clinical outcomes**

The ACEI/ARB(-) group had significantly higher incidence of non-fatal stroke than the ACEI/ARB(+) group (Before propensity score matching: 1.6% vs. 0.7%, p < 0.05; After propensity score matching: 1.7% vs. 0.8%, p < 0.05). There was no statistical difference in the other MACCE between the 2 groups.

**Subsequent MACCE and mortality**

During a median of 25.4 months (IQR: 12.3, 48.6 months) follow-up, composite MACCE occurred in 28.7% of patients in the ACEI/ARB (-) group and 23.1% in the ACEI/ARB (+) group (HR = 1.23, 95%CI 1.06, 1.44, p < 0.05, Table 4). Non-fatal stroke occurred in 4.0% of the patients in the ACEI/ARB (-) group and 2.4% in the ACEI/ARB (+) group (HR = 1.62, 95%CI 1.03, 2.56, p < 0.05). The incidences of all cause death, cardiovascular
death, non-fatal MI, revascularization, and cardiac rehospitalization were not statistically different between the 2 groups.

After propensity-score matching, composite MACCE occurred in 29.7% of the patients in the PSM ACEI/ARB(−) group and 23.1% in the PSM ACEI/ARB(+) group (HR = 1.21, 95%CI 1.02, 1.43, p < 0.05, Table 4); non-fatal stroke occurred in 4.6% of the PSM ACEI/ARB(−) group and 2.4% of the PSM ACEI/ARB(+) group (HR = 1.82, 95%CI 1.13, 2.96, p < 0.05). The incidences of all cause death, cardiovascular death, non-fatal MI, revascularization, and cardiac rehospitalization were not statistically different between the 2 groups. The Kaplan–Meier curves show that the ACEI/ARB(−) group had significantly higher cumulative rate of non-fatal stroke and composite MACCE than the ACEI/ARB(+) group (Fig. 4).

Independent association between non-fatal stroke and subsequent MACCE

In the multivariate analysis, we included variables that were identified to be significantly associated with non-fatal stroke and composite MACCE in the univariate model. The multivariate analysis revealed that no prior ACEI/ARB therapy, previous history of stroke, increased number of involved vessels, and lower LVEF were independently associated with non-fatal stroke (Table 5); no prior ACEI/ARB therapy, previous history of stroke, increased number of involved vessels, lower eGFR, lower LVEF, and no-antiplatelet therapy in hospital were significantly and independently associated with subsequent composite MACCE (Table 6).
To the best of our knowledge, the current study was the first to investigate whether ACEI/ARB used before the initial diagnosis of OCAD in diabetic hypertensive patients could be associated with improved clinical outcomes. We found that use of ACEI/ARB before the initial diagnosis of OCAD was associated with reduced incidence of AMI, reduced myocardial infarction size, and improved cardiac function, whereas we found no significant correlation between the prior ACEI/ARB therapy and mortality. However, the incidences of non-fatal stroke and composite MACCE were significantly higher in the ACEI/ARB(−) group than in the ACEI/ARB(+) group. No prior ACEI/ARB therapy was an

| Table 2 Laboratory test results and echocardiographic and angiographic characteristics |
|-----------------------------------------------|-----------------|-----------------|-------------------|-----------------|
| **Laboratory values**                        | **Before PS match** | **ACEI/ARB(-)** | **ACEI/ARB(+)** | **P value** |
| WBC, X10⁹/L                                  | 6.8 (5.6, 8.7)    | 6.6 (5.5, 8.1)  | <0.05            |                 |
| Neutrophil, X10⁹/L                           | 4.5 (3.6, 5.9)    | 4.4 (3.5, 5.5)  | <0.05            |                 |
| Monocyte, X10⁹/L                             | 0.29 (0.16, 0.44) | 0.27 (0.16, 0.41)| NS               |                 |
| Hemoglobin, g/L                              | 134.5 ± 16.5      | 133.7 ± 15.7    | <0.05            |                 |
| Hs-CRP, mg/L                                 | 2.7 (1.0, 10.4)   | 2.0 (0.8, 5.4)  | <0.001           | <0.05           |
| FBG, mmol/L                                  | 7.0 (5.8, 9.0)    | 6.8 (5.7, 8.3)  | <0.05            |                 |
| RBG at admission, mmol/L                     | 9.9 (7.5, 13.3)   | 9.7 (7.2, 12.8) | <0.05            |                 |
| Glycated hemoglobin, %                       | 7.5 ± 1.5         | 7.5 ± 1.6       | NS               |                 |
| ALT, U/L                                     | 190 (13.0, 28.0)  | 180 (13.0, 27.0)| NS               | 22.0 (15.0, 39.0)|                 |
| Creatinine, µmol/L                           | 760 (653, 881)    | 778 (662, 906)  | <0.05            | 818 (706, 969)  |
| eGFR, ml/min/1.73 m²                         | 84.9 (71.4, 97.9) | 82.3 (67.9, 95.3)| <0.001           | 79.1 (64.5, 94.5)| 79.2 (64.0, 95.2)|<0.05           |
| TC, mmol/L                                   | 4.3 (3.6, 5.0)    | 4.2 (3.4, 4.9)  | NS               | 4.45 ± 0.97     |
| TG, mmol/L                                   | 1.5 (1.1, 2.2)    | 1.5 (1.1, 2.1)  | NS               | 1.5 (1.1, 2.0)  |
| LDL-C, mmol/L                                | 2.4 (1.9, 2.9)    | 2.3 (1.8, 2.8)  | <0.001           | 2.6 (2.1, 3.1)  |
| HDL-C, mmol/L                                | 1.0 (0.9, 1.2)    | 1.0 (0.9, 1.2)  | NS               | 1.0 (0.9, 1.2)  |
| NT-Pro BNP, pg/ml                            | 1621 (644, 4214)  | 1585 (521, 4548)| NS               | 1855 (657, 4571)| 1585 (487, 4537)|NS             |
| **Echocardiographic values**                 | **After PS match** | **ACEI/ARB(-)** | **ACEI/ARB(+)** | **P value** |
| LVEDD, cm                                    | 5.06 ± 0.49       | 5.04 ± 0.48     | NS               | 5.13 ± 0.51     |
| LVESD, cm                                    | 3.28 ± 0.57       | 3.21 ± 0.51     | <0.05            | 3.53 ± 0.59     |
| LVEF, %                                      | 63.93 ± 8.92      | 65.66 ± 7.51    | <0.001           | 585.9 ± 9.4     |
| LVFS, %                                      | 35.33 ± 6.22      | 36.57 ± 5.92    | <0.001           | 31.4 ± 6.2      |
| LVEDV, ml                                    | 122.9 ± 28.3      | 122.2 ± 27.9    | NS               | 127.5 ± 30.0    |
| LVESV, ml                                    | 41.0 (32.2, 51.2) | 38.2 (32.2, 48.4)| <0.05            | 47.4 (38.8, 65.9)| 47.4 (36.7, 62.0)|<0.05           |
| LA, cm                                       | 3.71 ± 0.43       | 3.74 ± 0.41     | <0.05            | 3.76 ± 0.45     |
| E/A                                          | 0.80 (0.68, 1.00) | 0.79 (0.68, 1.00)| NS               | 0.84 (0.69, 1.16)| 0.81 (0.67, 1.16)|NS             |
| SV, ml                                       | 76.7 (66.5, 87.9) | 792 (687, 88.9) | <0.05            | 70.8 (63.4, 83.7)| 75.4 (65.4, 87.1)|NS             |
| **Angiography values**                      | **Involved vessel** | **ACEI/ARB(-)** | **ACEI/ARB(+)** | **P value** |
| Single vessel                                 | 180 (13.8)        | 158 (13.2)      | NS               | 157 (15.0)      |
| Multi-vessel/LM                              | 1120 (86.2)       | 1043 (86.8)     | NS               | 893 (85.0)      |
| CTO                                          | 122 (9.4)         | 108 (9.0)       | NS               | 92 (8.8)        |
| Proximal LAD                                  | 697 (53.6)        | 626 (52.1)      | NS               | 557 (53.0)      |

_data are presented as mean ± SD, IQR or n (%)

WBC white blood cell count, Hs-CRP high sensitivity C reactive protein, FBG fasting blood glucose, ALT alanine transaminase, eGFR estimated glomerular filtration rate, TC total cholesterol, TG triglycerides; LDL-C low-density lipoprotein cholesterol, HDL-C high-density lipoprotein cholesterol, NT-Pro BNP N-terminal pro-brain natriuretic peptide, LVEDD left ventricular end-diastolic diameter, LVESD left ventricular end-systolic diameter, LVEF left ventricular ejection fraction, LVFES left ventricular fraction shortening, LVEDV left ventricular end-diastolic volume, LVESV left ventricular end-systolic volume, LA left atrium, E/A ratio of early to late ventricular filling velocities, SV stroke volume, LM left main coronary artery, CTO chronic total occlusions, LA left anterior descending, NS non-significant
independent predictor of non-fatal stroke and composite MACCE.

ACEI promotes vasodilation by inhibiting angiotensin II formation and bradykinin decomposition, and ARB can trigger vasodilation and natriuresis. Therefore, ACEI/ARB are considered as antihypertensive drugs. In addition to the antihypertensive effects, ACEI/ARB has other pleiotropic clinical beneficial effects, such as inhibiting ventricular remodeling, decreasing sympathetic activity, improving insulin resistance, inhibiting atherosclerosis process, inhibiting thrombosis and platelet aggregation, and improving endothelium function and plaque stabilization [22–25]. Previous studies [11, 12, 26] have supported that ACEI/ARB exert clinical beneficial effects beyond blood pressure reduction and can reduce the incidence of major adverse cardiac events.

The current guidelines recommend ACEI/ARB as a first-line drug for diabetic hypertensive patients [14–16]; however, the reported effects of ACEI/ARB on cardiovascular risk in these patients are controversial. Previous studies have found that patients
treated with ACEI/ARB showed lower incidences of AMI [27–29] and stroke [28, 30, 31] than the control group. In addition, the Captopril Prevention Project (CAPPP) [29] has shown that compared with the diuretic/beta-blocker therapy group, the captopril group had lower incidences of cardiovascular mortality and all-cause mortality. A meta-analysis has demonstrated that ACEI/ARB was associated with a 17% reduction in cardiovascular mortality in diabetic hypertensive patients; however, ACEI/ARB was not associated with MI, stroke and all-cause mortality [32]. On the contrary, Bosch et al. [33] have shown that ACEI was not beneficial in the prevention of stroke. In addition, the Candesartan Antihypertensive Survival Evaluation Table 3 The estimated infarction size in patients with AMI

| The peak value of myocardial enzyme | Before PS match | ACEI/ARB(-) | ACEI/ARB(+) | P value | After PS match | ACEI/ARB(-) | ACEI/ARB(+) | P value |
|------------------------------------|----------------|-------------|-------------|---------|----------------|-------------|-------------|---------|
| pMYO,ng/ml                         | 50.1 (260.1, 150.3) | 46.0 (17.4, 146.5) | NS         | 50.4 (28.3, 173.8) | 46.1 (17.8, 150.0) | NS         |
| pCK-MB,ng/ml                       | 28.4 (8.0, 116.0)   | 21.3 (5.2, 89.2)   | <0.05      | 28.7 (8.2, 119.3) | 21.7 (5.2, 90.9) | <0.05      |
| pTNI,ng/ml                         | 7.7 (2.3, 27.0)     | 5.4 (1.0, 22.5)    | <0.05      | 6.8 (2.2, 22.9)  | 5.7 (1.0, 24.3)  | <0.05      |

Data are presented as IQR

AMI acute myocardial infarction, ACEI/ARB angiotensin-converting enzyme inhibitor/angiotensin receptor blocker, pMYO the peak value of myoglobin, pCK-MB the peak value of creatine kinase MB, pTNI the peak value of troponin I, NS non-significant

Table 4 Clinical events during long-term follow-up

| Overall population | ACEI/ARB(-) | ACEI/ARB(+) | HR(95%CI) | P value |
|--------------------|-------------|-------------|-----------|---------|
| Number             | 1300        | 1201        |           |         |
| Composite MACCE    | 373 (28.7)  | 277 (23.1)  | 1.23 (1.06,1.44) | <0.05   |
| All cause death    | 59 (4.5)    | 57 (4.7)    | 0.92 (0.64,1.32) | NS      |
| CV death           | 49 (3.8)    | 37 (3.1)    | 1.18 (0.77,1.81) | NS      |
| Non-fatal MI       | 57 (4.4)    | 37 (3.1)    | 1.38 (0.91,2.08) | NS      |
| Non-fatal stroke   | 52 (4.0)    | 29 (2.4)    | 1.62 (1.03,2.56) | <0.05   |
| Revascularization  | 102 (7.8)   | 92 (7.7)    | 0.98 (0.74,1.29) | NS      |
| Cardiac rehospitalization | 290 (22.3) | 233 (19.4) | 1.12 (0.94,1.33) | NS      |

| Matched population | ACEI/ARB(-) | ACEI/ARB(+) | HR(95%CI) | P value |
|--------------------|-------------|-------------|-----------|---------|
| Number             | 1050        | 1050        |           |         |
| Composite MACCE    | 312 (29.7)  | 242 (23.1)  | 1.21 (1.02,1.43) | <0.05   |
| All cause death    | 53 (5.0)    | 51 (4.9)    | 0.95 (0.65,1.40) | NS      |
| CV death           | 44 (4.2)    | 33 (3.1)    | 1.24 (0.79,1.94) | NS      |
| Non-fatal MI       | 48 (4.6)    | 31 (3.0)    | 1.45 (0.92,2.28) | NS      |
| Non-fatal stroke   | 48 (4.6)    | 25 (2.4)    | 1.82 (1.13,2.96) | <0.05   |
| Revascularization  | 76 (7.2)    | 81 (7.7)    | 0.85 (0.62,1.16) | NS      |
| Cardiac rehospitalization | 241 (23.0) | 202 (19.2) | 1.09 (0.91,1.32) | NS      |

ACEI/ARB angiotensin-converting enzyme inhibitor/angiotensin receptor blocker, MACCE major adverse cardiac and cerebral event, CV cardiovascular, MI myocardial infarction, HR hazard ratio, CI confidence interval, NS non-significant

(See figure on next page.)

Fig. 4 Kaplan-Meier curve for all cause death (a) CV death (b) non-fatal stroke (c) and composite MACCE (d) of the ACEI/ARB(-) group (red line) versus the ACEI/ARB(+) group (green line). a There was no significant difference in the cumulative rate of all cause death between the 2 groups. b There was no significant difference in the cumulative rate of CV death between the 2 groups. c The cumulative rate of non-fatal stroke in the ACEI/ARB(+) group was significantly higher than that in the ACEI/ARB(-) group (p < 0.05). d The cumulative rate of composite MACCE in the ACEI/ARB(-) group was significantly higher than that in the ACEI/ARB(+) group (p < 0.05). ACEI/ARB angiotensin-converting enzyme inhibitor/Angiotensin receptor blocker, CV Cardiovascular, MACCE major adverse cardiac and cerebral event
Japan (CASE-J) trial, which recruited 2018 patients with T2DM, failed to find a reduction in cardiovascular morbidity in patients using ARB [34]. A recent meta-analysis [12] has shown that ACEI significantly reduced the risk of AMI, cardiovascular mortality, and all-cause mortality, whereas treatment with ARBs did not show these benefits. In addition, neither ACEI nor ARB therapy decreased the incidence of stroke in patients with diabetes. Strauss et al. [35] believed that ARB could not reduce the risk of AMI, cardiovascular mortality, or all-cause Mortality. In the current study, we found that compared with diabetic hypertensive patients who did not use ACEI/ARB before the initial diagnosis of OCAD, those who did had significantly

### Table 5 Multivariate COX regression analysis of non-fatal stroke

| Univariate | Multivariate |
|------------|-------------|
|            | HR (95%CI) P | Adjusted HR (95%CI) P |
| Age, y     | 1.02 (0.99, 1.05) NS | 0.99 (0.97, 1.02) NS |
| ACEI/ARB(-) | 1.82 (1.13, 2.96) < 0.05 | 1.72 (1.05, 2.84) < 0.05 |
| Beta-blocker before admission | 0.54 (0.29, 1.01) NS | 0.56 (0.30, 1.06) NS |
| Previous stroke | 4.00 (2.52, 6.35) < 0.001 | 3.70 (2.29, 5.96) < 0.001 |
| Hemoglobin, g/L | 0.98 (0.96, 0.99) < 0.05 | 0.99 (0.97, 1.01) NS |
| AMI at admission | 2.10 (1.32, 3.35) < 0.05 | 1.07 (0.61, 1.86) NS |
| LVEF, % | 0.95 (0.93, 0.97) < 0.001 | 0.96 (0.94, 0.99) < 0.05 |
| Involved vessel | 1.79 (1.28, 2.49) < 0.05 | 1.52 (1.08, 2.14) < 0.05 |
| In-hospital treatment | | |
| ACEI/ARB | 0.72 (0.46, 1.14) NS | 0.85 (0.52, 1.38) NS |
| Antiplatelet agents | 0.34 (0.16, 0.75) < 0.05 | 0.49 (0.19, 1.25) NS |
| Statins | 0.43 (0.24, 0.76) < 0.05 | 0.58 (0.30, 1.11) NS |

ACEI/ARB(-) no Angiotensin-converting enzyme inhibitor/Angiotensin receptor blocker therapy before admission, AMI acute myocardial infarction, LVEF left ventricular ejection fraction; NS Non-significant

### Table 6 Multivariate COX regression analysis of composite MACCE

| Univariate | Multivariate |
|------------|-------------|
|            | HR (95%CI) P | Adjusted HR (95%CI) P |
| Age, y | 1.01 (0.99, 1.02) NS | 1.00 (0.99, 1.01) NS |
| Male, % | 1.05 (0.89, 1.25) NS | 1.05 (0.82, 1.33) NS |
| ACEI/ARB(-) | 1.21 (1.02, 1.43) < 0.05 | 1.24 (1.04, 1.48) < 0.05 |
| Previous stroke | 1.57 (1.29, 1.90) < 0.001 | 1.49 (1.22, 1.82) < 0.001 |
| Smoking | 1.09 (0.93, 1.29) NS | 1.05 (0.85, 1.31) NS |
| Hemoglobin, g/L | 1.00 (0.99, 1.00) NS | 0.99 (0.98, 1.00) NS |
| Glycated hemoglobin, % | 1.08 (1.02, 1.14) < 0.05 | 1.05 (0.99, 1.11) NS |
| LDL-C, mmol/L | 1.15 (1.03, 1.28) < 0.05 | 1.07 (0.95, 1.20) NS |
| eGFR ml/min/1.73 m² | 0.98 (0.97, 0.99) < 0.001 | 0.98 (0.97, 0.99) < 0.05 |
| AMI at admission | 1.62 (1.36, 1.93) < 0.001 | 1.18 (0.96, 1.45) NS |
| LVEF, % | 0.97 (0.96, 0.98) < 0.001 | 0.98 (0.97, 0.99) < 0.001 |
| Involved vessel | 1.45 (1.29, 1.62) < 0.001 | 1.34 (1.19, 1.51) < 0.001 |
| CTO | 1.49 (1.23, 1.81) < 0.001 | 1.07 (0.87, 1.33) NS |
| In-hospital treatment | | |
| Antiplatelet agents | 0.45 (0.33, 0.63) < 0.001 | 0.45 (0.31, 0.65) < 0.001 |
| ACEI/ARB | 0.92 (0.78, 1.09) NS | 1.00 (0.84, 1.19) NS |
| Beta-blocker | 0.93 (0.78, 1.12) NS | 0.87 (0.72, 1.06) NS |
| Statins | 0.75 (0.58, 0.95) < 0.05 | 0.94 (0.72, 1.23) NS |

ACEI/ARB(-) no angiotensin-converting enzyme inhibitor/angiotensin receptor blocker therapy before admission, LDL-C low-density lipoprotein cholesterol, eGFR estimated glomerular filtration rate, AMI acute myocardial infarction, LVEF left ventricular ejection fraction, CTO chronic total occlusions; NS non-significant.
lower incidence of OCAD-associated AMI at the hospital admission and lower incidences of non-fatal stroke and composite MACCE during follow-up. Multivariate analyses revealed that no prior ACEI/ARB therapy was an independent predictor of non-fatal stroke and composite MACCE. These findings have not been reported previously. There were 249 patients (23.7%) using ACEI and 801 patients (76.3%) using ARB before the initial diagnosis of OCAD in the PSM ACEI/ARB(+) group in this study. The incidence of AMI in the ACEI users and ARB users was 24.1% (60/249) and 22.0% (176/801), respectively, and the incidence of AMI in the PSM ACEI/ARB(-) group was 28.4% (298/1050). Therefore, we believed that both ACEI and ARB can reduce AMI development.

Notably, we found that the ACEI/ARB(-) group had significantly higher levels of hs-CRP, pTNI, and pCK-MB and lower LVEF than the ACEI/ARB(+) group, indicating that the patients who did not use ACEI/ARB before the initial diagnosis of OCAD appeared to have higher levels of inflammation, larger myocardial infarction area, and poorer cardiac function. Consistently, Gong et al. [36] also found that previous treatment with ACEI/ARB/β-blocker was associated with better heart function and smaller infarct size. To the best of our knowledge, this is the first study focusing on the effect of ACEI/ARB on the severity of the AMI in diabetic hypertensive patients.

The current study found that 52.0% (1300/2501) of patients with diabetic hypertension and diagnosed with OCAD for the first time did not use ACEI/ARB therapy. Notably, the proportion of patients treated with ACEI/ARB during hospitalization and long-term follow up in the ACEI/ARB(-) group was 51.0% and 41.3%, respectively, which were substantially lower than the proportions in the ACEI/ARB(+) group (83.9% and 67.0%, respectively). A study from the United States (from six states, 57,1483 participants) has shown that about 52.5% of patients with diabetic hypertension were non-adherent to ACEI/ARB therapy, which was related to an increased risk for diabetes-related rehospitalizations [19]. All these findings suggest that the real-world use of ACEI/ARB is seriously insufficient worldwide and the underutilization of ACEI/ARB may lead to poor clinical outcomes. We analyzed the factors associated with the use of ACEI/ARB and found that patients who have previously used CCB were less likely to receive ACEI/ARB therapy before the hospital admission, suggesting that CCB might affect ACEI/ARB’ first-line status in diabetic hypertensive patients. Based on the recommendations of the current guidelines [14–16], we believe the first-line treatment status of ACEI/ARB in patients with diabetic hypertension still need to be emphasized.

Limitations
First, this is a single-center study although including a large sample size; thus, generalization of the findings should be cautious. Second, this is a retrospective observational study. The information on the dosage and duration of ACEI/ARB was limited. Prospective cohort studies are required to confirm our findings.

Conclusions
Use of ACEI/ARB therapy for diabetic hypertensive patients before the initial diagnosis of OCAD was significantly associated with lower incidence of AMI, improved heart function, smaller infarct size, and lower incidences of non-fatal stroke and composite MACCE. No prior ACEI/ARB therapy was significantly and independently associated with non-fatal stroke and composite MACCE. ACEI/ARB therapy was largely underutilized in diabetic hypertensive patients.

Abbreviations
ACEI/ARB: Angiotensin-converting-enzyme inhibitors/angiotensin receptor blockers; OCAD: Obstructive coronary artery disease; MACCE: Major adverse cardiac and cerebral event; AMI: Acute myocardial infarction; Myo: Myoglobin; CKMB: Creatine kinase MB; cTnI: Cardiac troponin I; RAAS: Renin-angiotensin–aldosterone system; BMI: Body mass index; FBG: Fasting blood glucose; RBG: Random blood glucose; HGB: Hemoglobin; eGFR: Estimated glomerular filtration rate; LDL-C: Low-density lipoprotein cholesterol; hsCRP: High sensitivity C reactive protein; LVESV: Left ventricular end-systolic volume; LVEDV: Left ventricular end-diastolic volume; LVEF: Left ventricular ejection fraction; LVFS: Left ventricular fraction shortening; SV: Stroke volume; CCB: Calcium channel blocker; T2DM: Type 2 diabetes mellitus.

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Authors’ contributions
YZ performed study, statistical analysis and wrote manuscript. XSD, BH and QBL participated in study data collection. HC contributed discussion and edited manuscript. XQZ designed study and revised manuscript. WPL designed study, performed statistical analysis and edited manuscript. HWL provided funding support, designed study and reviewed manuscript. All authors read and approved the final manuscript.

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

Ethics approval and consent to participate
The study data collections were approved by the Institutional Review Board of Beijing Friendship Hospital affiliated to Capital Medical University, and written informed consent was obtained from all patients.

Consent for publication
Not applicable.
Competing interests
The authors declare that they have no competing interests.

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References
1. Clark H. NCDs: a challenge to sustainable human development. Lancet. 2013;381(9866):510–1.
2. Roth GA, Johnson C, Abajobir A, et al. Global, regional, and national burden of cardiovascular diseases for 10 causes, 1990 to 2015. J Am Coll Cardiol. 2017;70(1):1–25.
3. Mozaffarian D, Benjamin EJ, Go AS, et al. Heart disease and stroke statistics -2016 update: a report from the American Heart Association. Circulation. 2016;133(4):e38–360.
4. Einarson TR, Acs A, Ludwig C, Panton UH. Prevalence of cardiovascular disease in type 2 diabetes: a systematic literature review of scientific evidence from across the world in 2007–2017. Cardiovasc Diabetol. 2018;17(1):83.
5. Jargannathan R, Patel SA, Ali MK, Narayan KMV. Global updates on cardiovascular disease mortality trends and attribution of traditional risk factors. Curr Diab Rep. 2019;19(7):44.
6. Yamagishi S. Cardiovascular disease in recent onset diabetes mellitus. J Cardiol. 2011;57(3):257–62.
7. Ferrario CM, Strawn WB. Role of the renin-angiotensin-aldosterone system and proinflammatory mediators in cardiovascular disease. Am J Cardiol. 2006;98(12):1–8.
8. Kobori H, Nangaku M, Navar LG, Nishiyama A. The intrarenal renin-angiotensin system and kidney disease. Pharmacol Rev. 2007;59(3):251–87.
9. Darge HJ. Effect of carvedilol on outcome after myocardial infarction in patients with left-ventricular dysfunction: the CAPRICORN randomized trial. Lancet. 2001;357(9266):1385–90.
10. Pfeffer MA, McMurray JJ, Velazquez EJ, et al. Valsartan, captopril, or both in patients with left-ventricular dysfunction or both: the CAPRICORN substudy. Lancet. 2000;355(9200):253–9.
11. Heart Outcomes Prevention Evaluation Study Investigators. Effect of ramipril on cardiovascular and microvascular outcomes in people with diabetes mellitus: results of the HOPE study and MICRO-HOPE substudy. Lancet. 2000;355(9200):253–9.
12. Cheng J, Zhang W, Zhang X, et al. Effect of angiotensin converting enzyme inhibitors and angiotensin II receptor blockers on all-cause mortality, cardiovascular deaths, and cardiovascular events in patients with diabetes mellitus: a meta-analysis. JAMA Intern Med. 2014;174(5):773–86.
13. Lindholm LH, Ibsen H, Dahløf B, et al. Cardiovascular morbidity and mortality in patients with diabetes in the Losartan Intervention For Endpoint reduction in hypertension study (LIFE): a randomised trial against atenolol. Lancet. 2002;359(9311):995–1003.
14. American Diabetes Association. Standards of medical care in diab-etes-2013. Diab Care. 2013;36(Suppl 1):1–66.
15. Williams B, Mancia G, Spiering W, et al. 2018 ESC/ESH guidelines for the management of arterial hypertension. Eur Heart J. 2018;39(33):3021–104.
16. James PA, Oparil S, Carter BL, et al. 2014 evidence-based guideline for the management of high blood pressure in adults: report from the panel members appointed to the Eighth Joint National Committee (JNC 8). JAMA. 2014;311(5):507–20.
17. Amsterdam EA, Wenger NK, Brindis RG, et al. 2014 AHA/ACC guideline for the management of patients with non-ST-elevation acute coronary syndromes: a report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines. J Am Coll Cardiol. 2014;64(24):e139–228.
18. Roff M, Patrocco C, Collet JP, et al. 2015 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation: task force for the management of acute coronary syndromes in patients presenting without persistent ST- Segment Elevation of the European Society of Cardiology (ESC). Eur Heart J. 2016;37(3):267–315.
19. Yang Y, Thumula V, Pace PF, Banahan BF, Wilkin NE, Lobb WB. Nonadhe-rence to angiotensin-converting enzyme inhibitors and/or angiotensin II receptor blockers among high-risk patients with diabetes in medic‑care part D programs. J Am Pharm Assoc. 2010;50(4):527–31.
20. Aparasu RR, Aparasu A. Hypertension management in outpatient visits by diabetic patients. Res Soc Adm Pharm. 2008;4(3):284–91.
21. Johnson ML, Singh H. Patterns of antihypertensive therapy among patients with diabetes. J Gen Intern Med. 2005;20(9):842–6.
22. Schuh JR, Blehm DJ, Friedrich GE, McMahon EG, Blaine EH. Differential effects of renin-angiotensin system blockade on atherogenesis in cholesterol-fed rabbits. J Clin Invest. 1993;91(4):1453–8.
23. Schiffrin EL, Park JB, Intengan HD, Touyz RM. Correction of arterial structure and endothelial dysfunction in human essential hyperten-sion by the angiotensin receptor antagonist losartan. Circulation. 2000;101(14):1653–9.
24. Ido A, Hasebe N, Takeuchi T, Kikuchi K. Effects of temocapril and olmesartan on myocardial sympathetic nervous activity and fatty acid metabolism in rats with chronic beta-adrenergic stimulation. J Cardiovasc Pharmacol. 2003;41(Suppl 1):S133–7.
25. Yamaguchi K, Ura N, Murakami H, et al. Olmesartan ameliorates insulin sensitivity by modulating tumor necrosis factor-alpha and cyclic AMP in skeletal muscle. Hypertens Res. 2005;28(9):773–8.
26. Dahlof B, Devereux RB, Kjeldsen SE, et al. Cardiovascular morbidity and mortality in the Losartan Intervention For Endpoint reduction in hypertension study (LIFE): a randomised trial against atenolol. Lancet. 2002;359(9311):995–1003.
27. Estacio RO, Jeffers BW, Hiatt WR, Biggerstaff SL, Gifford N, Schrier RW. The effect of nisoldipine as compared with enalapril on cardiovascular outcomes in patients with non-insulin dependent diabetes and hyper‑tension. N Engl J Med. 1998;338(10):645–52.
28. Tatti P, Pahor M, Byington RP, et al. Outcome results of the fosinopril versus amiodipine cardiovascular events randomized trial (FACET) in patients with hypertension and NIDDM. Diab Care. 1998;21(4):597–603.
29. Niskanen L, Hedner T, Hansson L, Lanke J, Niklason A, CAPPP Study Group. Reduced cardiovascular morbidity and mortality in hyper‑tensive diabetic patients on first-line therapy with an ACE inhibi‑tor compared with a diuretic/beta-blocker-based treatment regimen: a subanalysis of the Captopril Prevention Project. Diabetes Care. 2001;24(12):2091–6.
30. Mohchuzki S, Dahlof B, Shimizu M, et al. Valsartan in a Japanese popula-tion with hypertension and other cardiovascular disease (Jikei Heart Study): a randomised, open-label, blinded endpoint morbidity-mortal‑ity study. Lancet. 2007;369(9571):1431–9.
31. Sawada T, Yamada H, Dahlof B, Matsubara H, KYOTO HEART Study. Effect of losartan on morbidity and mortality in uncontrolled hyper‑tensive patients with high cardiovascular risks: KYOTO HEART Study. Eur Heart J. 2009;30(20):2461–73.
32. Hao G, Wang ZW, Guo R, et al. Effects of ACEI/ARB in hypertensive patients with type 2 diabetes mellitus: a meta-analysis of randomized controlled studies. BMC Cardiovasc Disord. 2014;14:148.
33. Bosch J, Lonn E, Pogue J, Arnold JM, Dagenais GR, Yusuf S, HOPE/HEPE‑TOO Study Investigators. Long-term effects of ramipril on cardiovascular events and on diabetes: results of the HOPE study extension. Circulation. 2005;112(9):1339–46.
34. Oghara T, Nakao K, Fukui T, Candesartan Antihypertensive Survival Evaluation in Japan Trial Group, et al. Effects of candesartan compared with amiodipine in hypertensive patients with high cardiovascular risks: candesartan antihypertensive survival evaluation in Japan trial. Hypertension. 2008;51(2):393–402.
35. Strauss MH, Hall AS. Angiotensin receptor blockers do not reduce risk of myocardial infarction, cardiovascular death, or total mor-tality: further evidence for the ARB-MI Paradox. Circulation. 2017;135(22):2088–90.
36. Gong XH, Ding XS, Chen H, Li H. Real-world use of angiotensin-converting enzyme inhibitors/angiotensin receptor blockers/β-blocks in Chinese patients before acute myocardial infarction occurs: patient characteristics and hospital follow-up. J Transl Med. 2018;16(1):346.

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