

# Association of Inpatient Use of Angiotensin Converting Enzyme Inhibitors and Angiotensin II Receptor Blockers with Mortality Among Patients With Hypertension Hospitalized With COVID-19

Peng Zhang\*<sup>1,4</sup>; Lihua Zhu\*<sup>1,3</sup>; Jingjing Cai\*<sup>5</sup>; Fang Lei\*<sup>3</sup>; Juan-Juan Qin\*<sup>1,3</sup>; Jing Xie<sup>1</sup>; Ye-Mao Liu<sup>1,3</sup>; Yan-Ci Zhao<sup>1,3</sup>; Xuwei Huang<sup>1,3</sup>; Lijin Lin<sup>3,4</sup>; Meng Xia<sup>3</sup>; Ming-Ming Chen<sup>1,3</sup>; Xu Cheng<sup>1,3</sup>; Xiao Zhang<sup>6</sup>; Deliang Guo<sup>7</sup>; Yuanyuan Peng<sup>8</sup>; Yan-Xiao Ji<sup>2,3</sup>; Jing Chen<sup>3</sup>; Zhi-Gang She<sup>1,3</sup>; Yibin Wang<sup>8</sup>; Qingbo Xu<sup>9</sup>; Renfu Tan<sup>10</sup>; Haitao Wang<sup>7</sup>; Jun Lin<sup>12</sup>; Pengcheng Luo<sup>13</sup>; Shouzhi Fu<sup>14</sup>; Hongbin Cai<sup>15</sup>; Ping Ye<sup>16</sup>; Bing Xiao<sup>17</sup>; Weiming Mao<sup>18</sup>; Liming Liu<sup>19</sup>; Youqin Yan<sup>20</sup>; Mingyu Liu<sup>15</sup>; Manhua Chen<sup>16</sup>; Xiao-Jing Zhang<sup>1,3</sup>; Xinghuan Wang<sup>21</sup>; Rhian M. Touyz<sup>22</sup>; Jiahong Xia<sup>23</sup>; Bing-Hong Zhang<sup>24</sup>; Xiaodong Huang<sup>25</sup>; Yufeng Yuan<sup>7</sup>; Loomba Rohit<sup>26</sup>; Peter P. Liu<sup>27</sup>; Hongliang Li<sup>1-4</sup>

<sup>1</sup>Cardiology, Renmin Hospital of Wuhan University; <sup>2</sup>Medical Science Research Center, Zhongnan Hospital of Wuhan University; <sup>3</sup>Institute of Model Animal of Wuhan University; <sup>4</sup>Basic Medical School, Wuhan University; <sup>5</sup>Cardiology, The Third Xiangya Hospital, Central South University, Changsha, China; <sup>6</sup>Eye Center, Renmin Hospital of Wuhan University; <sup>7</sup>Hepatobiliary and Pancreatic Surgery, Zhongnan Hospital of Wuhan University; <sup>8</sup>Cardiology, Zhongnan Hospital, Wuhan University; <sup>9</sup>Anesthesiology, Cardiovascular Research Laboratories, David Geffen School of Medicine, University of California, Los Angeles, USA; <sup>10</sup>Centre for Clinic Pharmacology, The William Harvey Research Institute, Queen Mary University of London, UK; <sup>11</sup>Wuhan Kanghuashuai Technology Company, Wuhan; <sup>12</sup>Gastroenterology, Zhongnan Hospital of Wuhan University, Wuhan; <sup>13</sup>Urology, Wuhan Third Hospital & Tongren Hospital of Wuhan University; <sup>14</sup>Intensive Care Unit, Wuhan Third Hospital & Tongren Hospital of Wuhan University; <sup>15</sup>Wuhan Ninth Hospital; <sup>16</sup>Cardiology, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology; <sup>17</sup>Stomatology, Xiantao First People's Hospital; <sup>18</sup>General Surgery, Huanggang Central Hospital, Wuhan, China; <sup>19</sup>General Surgery, Ezhou Central Hospital; <sup>20</sup>Infections Department, Wuhan Seventh Hospital; <sup>21</sup>Center for Evidence-Based and Translational Medicine, Zhongnan Hospital of Wuhan University, Wuhan, China; <sup>22</sup>Urology, Zhongnan Hospital of Wuhan University; <sup>23</sup>Institute of Cardiovascular and Medical Sciences, BHF Glasgow Cardiovascular Research Centre, University of Glasgow, Glasgow, UK; <sup>24</sup>Cardiovascular Surgery, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology; <sup>25</sup>Neonatology, Renmin Hospital of Wuhan University; <sup>26</sup>Gastroenterology, Wuhan Third Hospital & Tongren Hospital of

Wuhan University; <sup>27</sup>NAFLD Research Center, Division of Gastroenterology and Epidemiology, University of California San Diego, San Diego, CA, USA, and; <sup>28</sup>Division of Cardiology, Department of Medicine, University of Ottawa Heart Institute, Ottawa, Ontario, Canada.

\*These authors contributed equally.

H.L., P.P.L., L.R., and Y.Y. are co-corresponding authors.

**Running title:** ACEI/ARB on COVID-19 in patients with hypertension



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**Subject Terms:**

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**Address correspondence to:**

Dr. Hongliang Li

Cardiology, Renmin Hospital of Wuhan University

99 Zhangzhidong Road

Wuhan 430060

lihl@whu.edu.cn

## ABSTRACT

**Rationale:** Use of angiotensin-converting enzyme inhibitors (ACEIs) and angiotensin II receptor blockers (ARBs) is a major concern for clinicians treating coronavirus disease 2019 (COVID-19) in patients with hypertension.

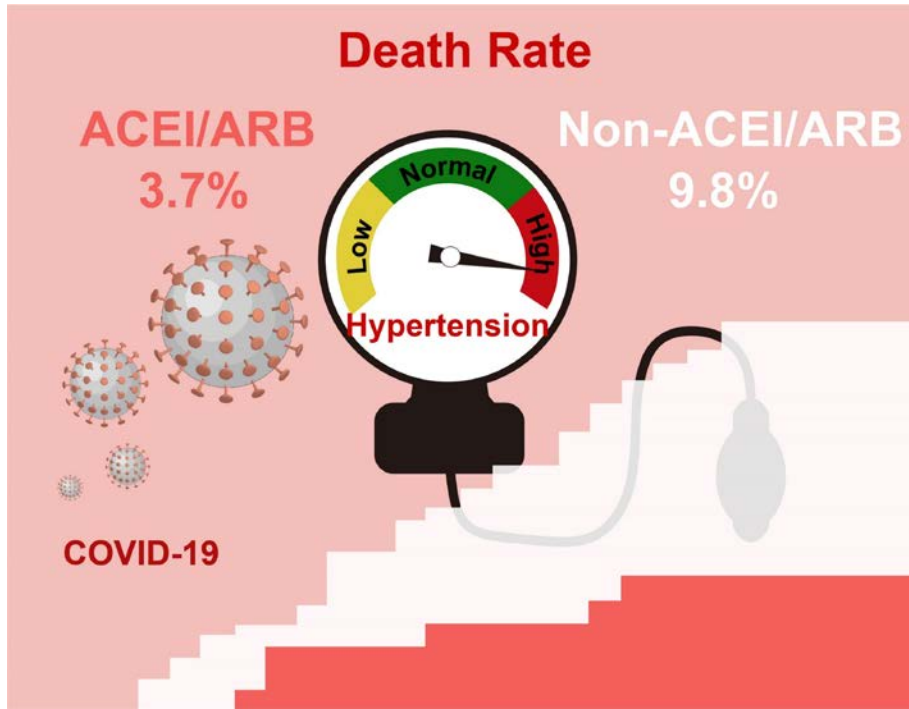
**Objective:** To determine the association between in-hospital use of ACEI/ARB and all-cause mortality in COVID-19 patients with hypertension.

**Methods and Results:** This retrospective, multi-center study included 1128 adult patients with hypertension diagnosed with COVID-19, including 188 taking ACEI/ARB (ACEI/ARB group; median age 64 [IQR 55-68] years; 53.2% men) and 940 without using ACEI/ARB (non-ACEI/ARB group; median age 64 [IQR 57-69]; 53.5% men), who were admitted to nine hospitals in Hubei Province, China from December 31, 2019 to February 20, 2020. Unadjusted mortality rate was lower in the ACEI/ARB group versus the non-ACEI/ARB group (3.7% vs. 9.8%;  $P = 0.01$ ). In mixed-effect Cox model treating site as a random effect, after adjusting for age, gender, comorbidities, and in-hospital medications, the detected risk for all-cause mortality was lower in the ACEI/ARB group versus the non-ACEI/ARB group (adjusted HR, 0.42; 95% CI, 0.19-0.92;  $P = 0.03$ ). In a propensity score-matched analysis followed by adjusting imbalanced variables in mixed-effect Cox model, the results consistently demonstrated lower risk of COVID-19 mortality in patients who received ACEI/ARB versus those who did not receive ACEI/ARB (adjusted HR, 0.37; 95% CI, 0.15-0.89;  $P = 0.03$ ). Further subgroup propensity score-matched analysis indicated that, compared to use of other antihypertensive drugs, ACEI/ARB was also associated with decreased mortality (adjusted HR, 0.30; 95% CI, 0.12-0.70;  $P = 0.01$ ) in COVID-19 patients with hypertension.

**Conclusions:** Among hospitalized COVID-19 patients with hypertension, inpatient use of ACEI/ARB was associated with lower risk of all-cause mortality compared with ACEI/ARB non-users. While study interpretation needs to consider the potential for residual confounders, it is unlikely that in-hospital use of ACEI/ARB was associated with an increased mortality risk.

### Key Words:

COVID-19, hypertension, ACEI/ARB, SARS-COV-2, pneumonia, infectious disease, lung.



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## Nonstandard Abbreviations and Acronyms:

ACC	American College of Cardiology
ACE	Angiotensin-converting enzyme
ACEI	Angiotensin-converting enzyme inhibitor
AHA	American Heart Association
AIDS	Acquired immune deficiency syndrome
ALT	Alanine transaminase
ARB	Angiotensin II receptor blocker
ARDS	Acute respiratory distress syndrome
AST	Aspartate transaminase
BNP	B-type natriuretic peptide
CCB	Calcium channel blocker
COPD	Chronic obstructive pulmonary disease
COVID-19	Coronavirus disease 2019
CRP	C-reactive protein
CT	Chest computerized tomography
cTNI	Cardiac troponin I
cTNT	Cardiac troponin T
DBP	Diastolic blood pressure
DIC	Disseminated intravascular coagulation
HFSA	Heart Failure Society of America
HR	Hazard ratio
hs-cTNI	High sensitivity cardiac troponin I
IQR	Interquartile range
IRDs	Incidence rate differences
ISH	International Society of Hypertension
ISTH	International Society on Thrombosis and Hemostasis
LDL-c	Low density lipoprotein cholesterol
MERS	Middle East Respiratory Syndrome
NT-proBNP	N-Terminal pro-brain natriuretic peptide
OR	Odds ratio
RAS	Renin-angiotensin-aldosterone system
RT-PCR	Reverse transcription-polymerase chain reaction
SaO <sub>2</sub>	Oxyhemoglobin saturation
SARS	Severe acute respiratory syndrome
SARS-COV	Severe acute respiratory syndrome coronavirus
SBP	Systolic blood pressure
SD	Standardized difference
ULN	Upper limit of normal
WHO	World Health Organization



## INTRODUCTION

The coronavirus disease 2019 (COVID-19) epidemic is caused by an infection with a novel coronavirus, officially named severe acute respiratory syndrome coronavirus 2 (SARS-COV-2).<sup>1</sup> Among patients with COVID-19 admitted to a hospital, emerging data suggest that hypertension may be associated with an increased risk of mortality due to COVID-19.<sup>2-4</sup>

Angiotensin converting enzyme inhibitor (ACEIs) and angiotensin II receptor blockers (ARBs) are part of renin-angiotensin-aldosterone system (RAS) inhibiting agents and considered as one of the first-line medications for the management of a large proportion of patients with hypertension.<sup>5,6</sup> However, continued use of ACEI/ARB has become controversial in the setting of COVID-19. The reason for this controversy stems from the fact that ACEIs and ARBs use may increase the expression of ACE2 receptor in animal-based studies,<sup>7, 8</sup> which is the known cellular receptor and a necessary entry point for SARS-COV-2 infection<sup>9</sup>. Conversely, it has been indicated that ACE2 expression is downregulated following SARS infection, resulting in excessive activation of RAS and exacerbated pneumonia progression.<sup>10</sup> Therefore, administration of ACEI/ARB may in turn be beneficial by blocking ACE2 downregulation-induced hyperactivation of RAS and thereby preventing acute lung injury and risk of adult respiratory distress syndrome. However, due to lack of sufficient clinical data supporting either the beneficial or harmful effects of ACEI/ARB use in patients with COVID-19, the optimal strategy for the management of hypertension in COVID-19 is uncertain and remains to be elucidated. The aim of this retrospective cohort study was to determine the association between in-hospital use of ACEI/ARB and all-cause mortality in COVID-19 among patients with hypertension.

## METHODS

Materials and data that support the findings of this study are available from the corresponding authors upon reasonable request.

### *Study design and participants.*

This retrospective, multi-center study included 1128 patients with hypertension and COVID-19 infection admitted to nine hospitals in Hubei, China, including Renmin Hospital of Wuhan University, Zhongnan Hospital of Wuhan University, Wuhan First Hospital, Wuhan Third Hospital, Wuhan Seventh Hospital, Wuhan Ninth Hospital, Thunder Mountain Hospital, Huanggang Central Hospital, and the Central Hospital of Enshi Tujia and Miao Autonomous Prefecture. The study protocols were approved by central ethics committee, and all collaborating hospitals either approved study protocol by local ethics committees or accepted the central ethics approval. Patient informed consent was waived by each ethics committee. Patients were admitted between December 31, 2019 and February 20th, 2020. The final date of follow up was March 7, 2020.



COVID-19 was diagnosed by meeting one or both criteria of chest computerized tomography (CT) manifestations and reverse transcription-polymerase chain reaction (RT-PCR) according to the New Coronavirus Pneumonia Prevention and Control Program (5th edition) published by the National Health Commission of China and WHO interim guidance (**Online Table I**).<sup>11, 12</sup> We used the following inclusion and exclusion criteria to determine the study cohort. The inclusion criteria included patients with COVID-19, aged from 18 to 74 years, who were admitted to the above-mentioned hospitals in Hubei, China from December 31st, 2019 to February 20th, 2020. The exclusion criteria included incomplete medical records (*e.g.*, transfer to any other hospital), pregnancy, acute lethal organ injury (*e.g.*, acute myocardial infarction, acute coronary syndrome, acute pulmonary embolism, or acute stroke), decompensated or end stage of chronic organ dysfunction (*e.g.*, decompensated cirrhosis, decompensated chronic renal insufficiency, or severe congestive heart failure), acquired immune deficiency syndrome (AIDS), or leukemia or malignancy. Patients with hypertension were classified based upon clearly documented medical history with systolic blood pressure (SBP)  $\geq 140$  mmHg or diastolic blood pressure (DBP)  $\geq 90$  mmHg.<sup>5</sup>

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### ***Data collection.***

Following data were collected including patient demographic information, medical history, clinical characteristics, laboratory data, radiologic report data, history of comorbidities, therapeutic interventions during the hospitalization, and clinical outcomes. The patient demographic information (age and gender), clinical characteristics (fever, cough, fatigue, dyspnea, heart rate, respiratory rate, and blood pressure), and durations from symptom onset to admission were collected from electronic medical system. The radiologic report data (chest computed tomography [CT]-diagnosed unilateral and bilateral lesions) were obtained from picture archiving and communication system. Laboratory data (blood cell count, C-reactive protein [CRP], procalcitonin, D-dimer, organ function markers, K<sup>+</sup>, LDL-c [low density lipoprotein cholesterol], SaO<sub>2</sub>, and blood glucose) were collected from laboratory information system. Comorbidities (hypertension, diabetes, coronary heart disease, chronic renal diseases, cerebrovascular diseases, chronic liver disease, and chronic obstructive pulmonary disease) were extracted from medical history. The in-hospital medications and interventions were collected from doctor advices. Personal health identifying information (*e.g.*, name and ID) was anonymized and each participant was given a study ID using an electronic coding system before data extraction to preserve patient privacy. Data were carefully reviewed and confirmed by experienced physicians and were double-checked to guarantee the accuracy of the data extraction procedures.

### ***Definition.***

The onset of COVID-19 was defined as the time point when the symptoms were first noticed. Patients with hypertension who received ACEI/ARB during hospitalization were classified as ACEI/ARB group. Patients with hypertension who did not receive ACEI/ARB during hospitalization were classified as non-ACEI/ARB group. In the subgroup propensity score-matched cohort analysis among patients taking antihypertensive medications, participants taking ACEI/ARB and other antihypertensive drugs (non-ACEI/ARB) during in-hospital stay were included. Patients discontinued treatment of hypertension due to inability to take medications (*e.g.*, hypotension, mechanical ventilation without nasal feeding and unable to orally taking medicines, or increase of creatinine level) during hospitalization were still included from the cohort. The primary endpoint was defined as 28-day all-cause death. ARDS and septic shock were defined according to the WHO interim guideline “Clinical management of severe acute respiratory infection when novel coronavirus (2019-nCoV) infection is suspected”<sup>13</sup> (**Online Table I**). Acute kidney injury was defined as



an elevation in serum creatinine level equal or above 26.5  $\mu\text{mol/L}$  within 48 hours.<sup>14</sup> Cardiac injury was defined as the serum level of cardiac troponin I (cTNI), cardiac troponin T (cTNT), or high sensitivity cardiac troponin I (hs-cTNI) above the upper limit of normal (ULN). Disseminated intravascular coagulation (DIC) was defined according to the criteria defined by the International Society on Thrombosis and Hemostasis (ISTH).<sup>15</sup> The increase in blood cell count and biochemical indexes were defined as over their ULN according to the criteria by the laboratory standards in each hospital (ULNs for CRP, procalcitonin, creatinine, D-dimer and platelets were shown in **Online Table II**).

### ***Propensity score-matched analysis.***

Propensity score-matched cohorts were created based on variables which were expected to be potential confounders associated with exposure to ACEI/ARB, including age, gender, fever, cough, dyspnea, comorbidities (diabetes, coronary heart disease, and chronic renal disease), CT-diagnosed bilateral lung lesions, and incidence of increased CRP and creatinine. We adjusted for imbalanced variables (D-dimer, procalcitonin, and unilateral lesion) and in-hospital medications (antiviral drug and lipid lowering drug) between ACEI/ARB versus non-ACEI/ARB groups in following mixed-effect Cox model. We used non-parametric missing value imputation, based on the missForest procedure in the R, to account for missing data on the laboratory variable of increased creatinine, CRP levels, procalcitonin, D-dimer, and unilateral lesion.<sup>16</sup> A random forest model using the rest of the variables in the data set was performed to predict the missing values for these variables. The internally cross-validated errors were also estimated. ACEI/ARB and non-ACEI/ARB users were paired according to the propensity scores using exact matching with a caliper size of 0.05. The balance of covariates was evaluated by estimating standardized differences (SD) before and after matching, and small absolute value less than 0.1 was considered successful balancing between the two groups. For cohort analysis in all patients with hypertension, and in those treated with antihypertensive therapies, ACEI/ARB and non-ACEI/ARB ratios were paired at 1:2 and 1:1, respectively.

### ***Statistical analysis.***

Continuous variables were expressed as median and interquartile range (IQR), and categorical variables were expressed as number and percentage (%). Statistical differences between two groups were analyzed using the Mann-Whitney U test for continuous variables, while categorical variables were compared using Fisher's exact test or  $\chi^2$  test. The risk of composite endpoints and corresponding hazard ratio (HR) were

calculated using the Cox proportional hazard model comparing ACEI/ARB group vs. non-ACEI/ARB group. Multi-variable adjusted including age, gender, comorbidities (diabetes, coronary heart disease, cerebrovascular disease, and chronic renal disease), and in-hospital medications (antiviral drug and lipid lowering drug) were performed. We modeled site as a random effect in the mixed-effect Cox model. The proportional hazard assumptions were verified using correlation testing based on Schoenfeld residuals. Incidence rate differences (IRDs) were calculated to provide incidence difference on absolute change. The cumulative rates of death were compared using the Kaplan-Meier method. A two-side  $\alpha$  less than 0.05 was considered statistically different. Because of the potential for type 1 error due to multiple comparisons, findings for analyses of secondary endpoints should be interpreted as exploratory. Data were analyzed in R-3.6.3 (R Foundation for Statistical Computing, Vienna, Austria) and SPSS Statistics (version 23.0, IBM, Armonk, NY, USA).

### ***Sensitivity analysis.***

E-value analysis was conducted to assess the robustness of the association between ACEI/ARB use and all-cause mortality in the mixed-effect Cox model to address unmeasured confounding using the methodology of VanderWeele and Ding.<sup>17-19</sup> The E-value is an alternative approach to sensitivity analyses for unmeasured confounding in our studies that avoids making assumptions that, in turn, require subjective assignment of inputs for some formulas. If the strength of unmeasured confounding is weaker than indicated by the E-value, then the main study result could not be overturned to one of the unmeasured confounder. E-values can therefore help to assess the robustness of the main study result by considering whether unmeasured confounding of this magnitude is plausible. We performed two sensitivity analyses to evaluate the robustness of propensity score-matched cohort analyses, among all patients with hypertension, using pairs of 1:2. In the first sensitivity analysis, comorbid diabetes was not included in matching, while the second sensitivity analysis was conducted adding cerebrovascular disease as a matching variable. We conducted a subset analysis restricted to patients who were taking an anti-hypertensive medication, applying matching variables as above with the pairing ratio at 1:1.



## RESULTS

### *Participants.*

This study cohort included 3611 patients with COVID-19 who were admitted to these nine hospitals in Hubei, China. After excluding 181 participants following our exclusion criteria, 3430 participants comprising 1128 hypertensive and 2302 normotensive cases were included in subsequent analysis (Figure 1; Online Table III). Among the 1128 participants with hypertension and COVID-19, 188 were classified as ACEI/ARB group (median age 64 [IQR 55-68] years; 53.2% men) and the remaining 940 were classified as non-ACEI/ARB group (median age 64 [IQR 57-69]; 53.5% men). The characteristics of the ACEI/ARB group versus the non-ACEI/ARB group on admission were provided in Table 1. Compared to the ACEI/ARB group, the non-ACEI/ARB group had higher prevalence of fever, dyspnea, and bilateral lung lesion at presentation. The dynamic changes in blood pressure during a 28-day follow-up period after presentation were similar between the ACEI/ARB and non-ACEI/ARB groups (Online Figure I). In terms of in-hospital treatment, the ACEI/ARB group had a higher percentage of patients receiving antiviral (88.8% vs. 81.7%;  $P = 0.02$ ) and lipid-lowering therapies (22.9% vs. 10.0%;  $P = 1.51E-6$ ) than patients in the non-ACEI/ARB group (Table 2).

### *Primary outcomes.*

During a 28-day follow-up duration, 99 deaths died out of the 1128 patients with hypertension and COVID-19. The risk of 28-day all-cause mortality was significantly lower in ACEI/ARB group versus non-ACEI/ARB group (3.7% [7/188] vs. 9.8% [92/940];  $P = 0.01$ ). In the mixed-effect Cox model using site as a random effect, after adjusting for age, gender, comorbidities, and in-hospital medications (antiviral and lipid lowering drugs), use of ACEI/ARB was associated with lower all-cause mortality (adjusted HR, 0.42; 95% CI, 0.19-0.92;  $P = 0.03$ ) versus use of non-ACEI/ARB (**Figure 2A** and **Table 3**). The E value for the point estimate of primary endpoint was 4.22 with upper limit of CI at 1.41. In our study, the adjusted hazard ratios (HR) for association of known variables for all-cause mortality due to COVID-19 were 1.08 (95% CI, 1.04-1.11;  $P = 3.30E-6$ ) for age, 2.23 (95% CI, 1.45-3.43;  $P = 2.78E-4$ ) for gender, 1.47 (95% CI, 0.86-2.52;  $P = 0.16$ ) for coronary heart disease, and 1.35 (95% CI, 0.58-3.16;  $P = 0.49$ ) for cerebrovascular disease.

We further conducted a propensity score-matched analysis to account for confounding that may have resulted in a protective association between ACEI/ARB use and all-cause mortality. We successfully matched 174 patients with hypertension in the ACEI/ARB group to non-ACEI/ARB group at a ratio of 1:2, and 522 individuals were included in the propensity score-matched samples. Imbalanced in-hospital medications (antiviral drug and lipid lowering drug), increase of D-dimer, and unilateral lung lesion after matching were adjusted in mixed-effect Cox model using site as a random effect. The results remained consistent and statistically significant, demonstrating lower risk of all-cause mortality in patients who received ACEI/ARB (adjusted HR, 0.37; 95%CI, 0.15-0.89;  $P = 0.03$ ) versus those who did not receive ACEI/ARB using this propensity score-matched analysis (**Figure 2B** and **Table 3**). To further assess the robustness of the association between ACEI/ARB use and mortality, we performed additional sensitivity analyses by using different matching variables. The results remained consistent and statistically significant in these sensitivity analyses with HRs of 0.34 (95%CI, 0.14-0.82;  $P = 0.02$ ) in the first sensitivity analysis and of 0.33 (95% CI, 0.13-0.80;  $P = 0.01$ ) in the second analysis.



Since there were 34.0% patients with hypertension who did not receive antihypertensive drugs during hospitalization, we performed subgroup propensity score-matching analysis on the remaining 745 patients who received at least one anti-hypertensive medication during hospitalization to further minimize the potential bias from non-users. One hundred eighty-one patients using ACEI/ARB versus those using other anti-hypertensive drugs were paired at 1:1. Characteristics of matched and unmatched participants in this analysis were listed in **Online Table IV**. The results demonstrated that the in-hospital use of ACEI/ARB was associated with lower risk of all-cause mortality (adjusted HR, 0.29; 95%CI, 0.12-0.69;  $P = 0.005$ ) due to COVID-19 (**Online Figure II** and **Online Table V**). This association was further supported by sensitivity analyses with adjusted HR of 0.29 (95% CI, 0.12-0.70;  $P = 0.01$ ) in the first sensitivity analysis not including diabetes as a matching variable and of 0.30 (95%CI, 0.13-0.71;  $P = 0.01$ ) in the second sensitivity analysis adding cerebrovascular disease as a matching variable.

#### *Secondary outcomes.*

The incidence of septic shock (3.2% in ACEI/ARB vs. 8.0% in non-ACEI/ARB [ $P = 0.03$ ]; IRD, -0.19 [95%CI, -0.36 – -0.01]) and DIC (0.0% vs. 2.3%,  $P = 0.04$ ; IRD, -0.23 [95%CI, -0.52 – -0.07]) were lower in the ACEI/ARB group than the non-ACEI/ARB group (**Table 3**). In the mixed-effect Cox model,

use of ACEI/ARB was associated with lower risk of septic shock (adjusted HR, 0.36; 95% CI, 0.16-0.84;  $P = 0.01$ ) compared to non-ACEI/ARB group (**Table 3**).

In propensity score-matched cohort analysis, the risk of septic shock was lower in ACEI/ARB group (adjusted HR, 0.32 [95% CI, 0.13-0.80;  $P = 0.01$ ]; IRD, -0.20[95% CI, -0.39 – -0.01]) than non-ACEI/ARB group among all individuals with hypertension (**Table 3**). In a sub-group of patients, who received at least one anti-hypertensive medication during hospitalization, the findings remained consistent (adjusted HR, 0.24 [95% CI, 0.10-0.63;  $P = 0.003$ ]; IRD, -0.31 [95% CI, -0.54 – -0.09]) (**Online Table V**).

#### *Comparison of patients with hypertension vs non-hypertension.*

Compared to patients in the normotensive group, the hypertensive group had a higher percentage of dyspnea, comorbidities, and abnormal laboratory markers on admission, and they received more intensive in-hospital interventions (**Online Table III** and **Online Table VI**). During the 28-day follow-up period, co-existing hypertension was significantly associated with the higher risk of all-cause mortality (adjusted HR, 1.41; 95% CI, 1.03-1.94;  $P = 0.03$ ) and the occurrence of multi-organ injury (**Online Figure III** and **Online Table VII**).

## DISCUSSION

In this multicenter retrospective study, in-hospital use of ACEI/ARB was associated with lower risk of all-cause mortality due to COVID-19 compared with either nonuse of ACEI/ARB or use of a different class of anti-hypertensive agent among patients with hypertension. Although it is plausible that unmeasured confounding may have contributed to the observed protective association, these data suggest that in-hospital use of ACEI/ARB was not associated with increased mortality in COVID-19. These findings provide clinical evidence in support of recently published guidance statements by several international societies to continue ACEI/ARB in patients with COVID-19<sup>20, 21</sup>. Given the retrospective nature of this study, these data need further validation in geographically diverse, prospective, cohort studies. Randomized controlled trials are needed to examine the efficacy of ACEI/ARB use in patients with hypertension and COVID-19.

Previous clinical studies have shown that hypertension is a risk factor for higher mortality in patients suffering from SARS and Middle East Respiratory Syndrome (MERS).<sup>22-24</sup> Similar to the association between hypertension and other previous outbreaks from novel coronavirus infections, recent studies in patients with COVID-19 have also reported that hypertension is associated with higher mortality in COVID-19 compared to normotensions.<sup>3,4</sup> The underlying pathogenic mechanism linking hypertension and severity of COVID-19 infection remains to be elucidated. It has been hypothesized that excessive activation of RAS system might contribute to progression of COVID-19 related lung injury by promoting inflammatory response and cytokine storm,<sup>25</sup> stimulating the NADH/NADPH oxidase system,<sup>26</sup> and triggering cell contraction and vasoconstriction<sup>27</sup>. Further studies are needed to better understand the underlying biological mechanisms involved in the association between hypertension and adverse outcomes in COVID-19.

In the management of COVID-19 patients with hypertension, use of ACEI and ARB has been a contentious issue. These therapeutic effects involve ACE2, a known cellular receptor of SARS-COV-2 which is required for viral entry and propagation in host cells. ACEI and ARB treatment can increase ACE2 expression in animal-based studies.<sup>7, 8</sup> Potentially, ACEI/ARB might increase ACE2 expression, thus promoting SARS-COV-2 susceptibility and disease severity of COVID-19. Conversely, ACE2 negatively regulates RAS and serves as a counterbalance to ACE function. Its expression is significantly downregulated after SARS-COV infection, contributing to hyper-activated RAS cascades.<sup>28</sup> As a consequence, loss of ACE2 in mice confers resistance to SARS-COV infection, but also results in exacerbated vascular permeability, lung edema, neutrophil accumulation, and pulmonary dysfunction.<sup>10, 29</sup> Recombinant ACE2 protein protects against acute lung injury in mouse models of ARDS and SARS. A retrospective review of 539 consecutive hospitalized patients with viral pneumonia indicates that continuing in-hospital use of ACEI or ARB may reduce the risk of pneumonia and death (ACEI, OR, 0.64 for risk of pneumonia; OR, 0.25 for in-hospital death; ARB, OR, 0.48 for risk of pneumonia; OR, 0.75 for in-hospital death).<sup>30</sup> A more recent study by Liu *et al.* shows that plasma AngII concentration is significantly elevated after SARS-COV-2 infection.<sup>31</sup> The influence of ACEI/ARB on COVID-19 related outcomes has not been fully investigated. Furthermore, among patients with COPD, it has been suggested that ARB might be more effective than ACEI to reduce the severity and mortality due to COPD.<sup>32</sup> This study was not powered to assess the differential beneficial effects of ACEI versus ARB versus combination of ACEI and ARB in

improving outcomes in COVID-19. The differential efficacy of ACEI and ARB use in improving COVID related outcomes also needs to be examined in further studies.

A statement jointly published by American Heart Association (AHA), the Heart Failure Society of America (HFSA) and the American College of Cardiology (ACC),<sup>20</sup> and another statement from the International Society of Hypertension (ISH) on COVID-19<sup>21</sup> strongly recommended continuation of ACEI or ARB among patients with co-existing hypertension and COVID-19. The main argument in favor of continuation of ACEI/ARB was that there are no clinical data to show whether ACEI and ARB would either improve or worsen COVID19. Therefore, it is not recommended to modify or change antihypertensive therapy before fully evaluating the possible influence of ACEI/ARB among patients with COVID-19. Our findings in this paper provide evidence supporting continuous use of ACEI/ARB for patients with hypertension infected with SARS-COV-2. Comorbidities and in-hospital medications might lead confounding to the association between ACEI/ARB application with COVID-19 mortality. Previous studies have suggested that hypokalemia may be a marker of unopposed Ang II,<sup>33, 34</sup> and therefore, hypokalemia may modify the association between anti-hypertensive therapy and outcomes in COVID-19 infection. Further studies are needed to examine the role of hypokalemia and hypertension and ACEI/ARB use and its influence on COVID-19 severity. The protection effect of ACEI/ARB also may result from the favorable effect on microvascular complications, thus reduces cardiovascular and renal morbidity.<sup>35, 36</sup>

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Overall, these findings suggested potential beneficial effects observed with continued use of ACEI/ARB therapy and the possible contribution of RAS activation in the pathogenesis of severity of COVID-19 in patients with hypertension. Considering the impact of unmeasured potential confounders, we conducted E-value analysis and found that E value was substantially greater than accepted risk factors for COVID-19 mortality. Therefore, it is not likely that an unmeasured confounder exists to modify the conclusion that ACEI/ARB use was not associated with increased COVID-19 mortality as observed in this study.

### **Limitations.**

This study has several limitations. First, the study population originated from nine hospitals in Hubei Province and may reflect natural history of hospitalized patients with COVID-19. It is plausible that the

beneficial effects of ACEI/ARB may be different among patients who are managed in the outpatient setting or in ethnically or geographically diverse populations. Second, the study sample-size was modest and included 188 patients who received ACEI/ARB. It did not have the power to detect if there was a differential effect between ACEI and ARB. Third, since the retrospective nature of this study, some parameters were not available in all patients, and in-hospital medications might be not fully recorded. For instance, some patients with hypertension may have failed to report antihypertensive medication though self-administration (e.g., Traditional Chinese Medicines) or might stop antihypertensive medication due to well-controlled blood pressure. Fourth, application of antihypertensive drugs was not matched or adjusted when comparing ACEI/ARB and non-ACEI/ARB groups since it is a key comparative factor for outcomes. The differences in proportions of patients using beta-blocker and diuretics between ACEI/ARB and non-ACEI/ARB groups might induce unappreciated confounding to conclusion. Fifth, this study was not able to retrieve pre-hospital self-medications from in-hospital electronic record systems in the urgent circumstance of the COVID-19 pandemic, and thus was unable to account for their influence on the severity of presentation and outcomes. Therefore, large-scale prospective cohort studies and randomized controlled trials are needed in ethnically and geographically diverse cohorts to better understand the association between ACEI/ARB and survival in COVID-19.

### **Conclusions.**

Among patients with hypertension hospitalized with COVID-19, inpatient treatment with ACEI/ARB was associated with lower risk of all-cause mortality compared with ACEI/ARB non-users. While study interpretation needs to consider the potential for residual confounders, it is unlikely that inpatient ACEI/ARB would be associated with an increased risk of mortality.

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PZ, LZ, FL and JC designed study, collected and analyzed data, and wrote manuscript. JJQ, YCZ, XH, LL, MX, MMC, YXJ, RT, HW, JL, PL, SF, HC, PY, BX, WM, LL, YY, ML, MC, BHZ, XW, JX, BHZ, and XH collected and reviewed clinical, laboratory, and radiological data. YML, XC and JC performed statistical analysis. JX, XZ, DG and YP reviewed, interpreted, and checked clinical data. XJZ, ZGS, YW,



QX and R.M.T. wrote manuscript and provided valuable suggestions for study design and data analysis. YY, L.R., P.L. and H.L. contributed equally, designed the project, edited manuscript, and supervised the study. All authors have approved the final version of this paper.

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## DISCLOSURES

None.



## SUPPLEMENTAL MATERIALS

Online Tables I - VII

Online Figures I - III

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## REFERENCES

1. Coronaviridae Study Group of the International Committee on Taxonomy of V. The species Severe acute respiratory syndrome-related coronavirus: classifying 2019-nCoV and naming it SARS-CoV-2. *Nat Microbiol.* 2020;doi:10.1038/s41564-020-0695-z
2. Guan WJ, Ni ZY, Hu Y, Liang WH, Ou CQ, He JX, Liu L, Shan H, Lei CL, Hui DSC, et al. Clinical Characteristics of Coronavirus Disease 2019 in China. *N Engl J Med.* 2020;doi:10.1056/NEJMoa2002032
3. Wang D, Hu B, Hu C, Zhu F, Liu X, Zhang J, Wang B, Xiang H, Cheng Z, Xiong Y, et al. Clinical Characteristics of 138 Hospitalized Patients With 2019 Novel Coronavirus-Infected Pneumonia in Wuhan, China. *Jama.* 2020;doi:10.1001/jama.2020.1585
4. Zhou F, Yu T, Du R, Fan G, Liu Y, Liu Z, Xiang J, Wang Y, Song B, Gu X, et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study.

*Lancet (London, England)*. 2020;doi:10.1016/S0140-6736(20)30566-3

5. Flack JM and Adekola B. Blood pressure and the new ACC/AHA hypertension guidelines. *Trends Cardiovasc Med*. 2020;30:160-164. doi:10.1016/j.tcm.2019.05.003
6. Kovell LC, Ahmed HM, Misra S, Whelton SP, Prokopowicz GP, Blumenthal RS and McEvoy JW. US Hypertension Management Guidelines: A Review of the Recent Past and Recommendations for the Future. *J Am Heart Assoc*. 2015;4doi:10.1161/JAHA.115.002315
7. Igase M, Kohara K, Nagai T, Miki T and Ferrario CM. Increased expression of angiotensin converting enzyme 2 in conjunction with reduction of neointima by angiotensin II type 1 receptor blockade. *Hypertens Res*. 2008;31:553-9. doi:10.1291/hypres.31.553
8. Ferrario CM, Jessup J, Chappell MC, Averill DB, Brosnihan KB, Tallant EA, Diz DI and Gallagher PE. Effect of angiotensin-converting enzyme inhibition and angiotensin II receptor blockers on cardiac angiotensin-converting enzyme 2. *Circulation*. 2005;111:2605-10. doi:10.1161/CIRCULATIONAHA.104.510461
9. Wu F, Zhao S, Yu B, Chen Y-M, Wang W, Song Z-G, Hu Y, Tao Z-W, Tian J-H, Pei Y-Y, et al. A new coronavirus associated with human respiratory disease in China. *Nature*. 2020;doi:10.1038/s41586-020-2008-3
10. Kuba K, Imai Y, Rao S, Gao H, Guo F, Guan B, Huan Y, Yang P, Zhang Y, Deng W, et al. A crucial role of angiotensin converting enzyme 2 (ACE2) in SARS coronavirus-induced lung injury. *Nat Med*. 2005;11:875-9. doi:10.1038/nm1267
11. National Health Commission of China. New Coronavirus Pneumonia Prevention and Control Program. 2020.
12. World Health Organization. Laboratory testing for 2019 novel coronavirus (2019-nCoV) in suspected human cases Interim guidance. 2020. <https://www.who.int/publications-detail/laboratory-testing-for-2019-novel-coronavirus-in-suspected-human-cases-20200117>
13. World Health Organization. Clinical management of severe acute respiratory infection when novel coronavirus (2019-nCoV) infection is suspected. 2020. [https://www.who.int/publications-detail/clinical-management-of-severe-acute-respiratory-infection-when-novel-coronavirus-\(ncov\)-infection-is-suspected](https://www.who.int/publications-detail/clinical-management-of-severe-acute-respiratory-infection-when-novel-coronavirus-(ncov)-infection-is-suspected)
14. Kellum JA, Lameire N, Aspelin P, Barsoum RS, Burdmann EA, Goldstein SL, Herzog CA, Joannidis M, Kribben A, Levey AS, et al. Kidney disease: Improving global outcomes (KDIGO) acute kidney injury work group. KDIGO clinical practice guideline for acute kidney injury. *Kidney International Supplements*.



Circulation  
Research

2012;2:1-138. doi:10.1038/kisup.2012.1

15. Gando S, Wada H, Thachil J, Scientific, Standardization Committee on DICotISOt and Haemostasis. Differentiating disseminated intravascular coagulation (DIC) with the fibrinolytic phenotype from coagulopathy of trauma and acute coagulopathy of trauma-shock (COT/ACOTS). *J Thromb Haemost.* 2013;11:826-35. doi:10.1111/jth.12190

16. Waljee AK, Mukherjee A, Singal AG, Zhang Y, Warren J, Balis U, Marrero J, Zhu J and Higgins PD. Comparison of imputation methods for missing laboratory data in medicine. *BMJ Open.* 2013;3doi:10.1136/bmjopen-2013-002847

17. VanderWeele TJ and Ding P. Sensitivity Analysis in Observational Research: Introducing the E-Value. *Ann Intern Med.* 2017;167:268-274. doi:10.7326/M16-2607

18. Mathur MB, Ding P, Riddell CA and VanderWeele TJ. Web Site and R Package for Computing E-values. *Epidemiology.* 2018;29:e45-e47. doi:10.1097/EDE.0000000000000864

19. Haneuse S, VanderWeele TJ and Arterburn D. Using the E-Value to Assess the Potential Effect of Unmeasured Confounding in Observational Studies. *Jama.* 2019;321:602-603. doi:10.1001/jama.2018.21554

20. American Heart Association (AHA), Heart Failure Society of America (HFSA) and American College of Cardiology (ACC). Patients taking ACE-i and ARBs who contract COVID-19 should continue treatment, unless otherwise advised by their physician. 2020.

21. International Society of Hypertension. A statement from the International Society of Hypertension on COVID-19. 2020.

22. Morra ME, Van Thanh L, Kamel MG, Ghazy AA, Altibi AMA, Dat LM, Thy TNX, Vuong NL, Mostafa MR, Ahmed SI, et al. Clinical outcomes of current medical approaches for Middle East respiratory syndrome: A systematic review and meta-analysis. *Rev Med Virol.* 2018;28:e1977. doi:10.1002/rmv.1977

23. Matsuyama R, Nishiura H, Kutsuna S, Hayakawa K and Ohmagari N. Clinical determinants of the severity of Middle East respiratory syndrome (MERS): a systematic review and meta-analysis. *BMC Public Health.* 2016;16:1203. doi:10.1186/s12889-016-3881-4

24. Chen M, Yang ZD, Smith KM, Carter JD and Nadler JL. Activation of 12-lipoxygenase in proinflammatory cytokine-mediated beta cell toxicity. *Diabetologia.* 2005;48:486-95. doi:10.1007/s00125-005-1673-y

25. Rodriguez-Iturbe B, Pons H and Johnson RJ. Role of the Immune System in Hypertension. *Physiol*

Rev. 2017;97:1127-1164. doi:10.1152/physrev.00031.2016

26. Zhang Y, Murugesan P, Huang K and Cai H. NADPH oxidases and oxidase crosstalk in cardiovascular diseases: novel therapeutic targets. *Nat Rev Cardiol.* 2020;17:170-194. doi:10.1038/s41569-019-0260-8

27. Dhaun N and Webb DJ. Endothelins in cardiovascular biology and therapeutics. *Nat Rev Cardiol.* 2019;16:491-502. doi:10.1038/s41569-019-0176-3

28. Kuba K, Imai Y and Penninger JM. Angiotensin-converting enzyme 2 in lung diseases. *Curr Opin Pharmacol.* 2006;6:271-6. doi:10.1016/j.coph.2006.03.001

29. Imai Y, Kuba K, Rao S, Huan Y, Guo F, Guan B, Yang P, Sarao R, Wada T, Leong-Poi H, et al. Angiotensin-converting enzyme 2 protects from severe acute lung failure. *Nature.* 2005;436:112-6. doi:10.1038/nature03712

30. Henry C, Zaizafoun M, Stock E, Ghamande S, Arroliga AC and White HD. Impact of angiotensin-converting enzyme inhibitors and statins on viral pneumonia. *Proc (Bayl Univ Med Cent).* 2018;31:419-423. doi:10.1080/08998280.2018.1499293

31. Liu Y, Yang Y, Zhang C, Huang F, Wang F, Yuan J, Wang Z, Li J, Li J, Feng C, et al. Clinical and biochemical indexes from 2019-nCoV infected patients linked to viral loads and lung injury. *Sci China Life Sci.* 2020;doi:10.1007/s11427-020-1643-8

32. Lai CC, Wang YH, Wang CY, Wang HC, Yu CJ and Chen L. Comparative effects of angiotensin-converting enzyme inhibitors and angiotensin II receptor blockers on the risk of pneumonia and severe exacerbations in patients with COPD. *Int J Chron Obstruct Pulmon Dis.* 2018;13:867-874. doi:10.2147/COPD.S158634

33. Te Riet L, van Esch JH, Roks AJ, van den Meiracker AH and Danser AH. Hypertension: renin-angiotensin-aldosterone system alterations. *Circ Res.* 2015;116:960-75. doi:10.1161/CIRCRESAHA.116.303587

34. Kozhuharov N, Goudev A, Flores D, Maeder MT, Walter J, Shrestha S, Gualandro DM, de Oliveira Junior MT, Sabti Z, Muller B, et al. Effect of a Strategy of Comprehensive Vasodilation vs Usual Care on Mortality and Heart Failure Rehospitalization Among Patients With Acute Heart Failure: The GALACTIC Randomized Clinical Trial. *Jama.* 2019;322:2292-2302. doi:10.1001/jama.2019.18598

35. Paulus WJ and Tschope C. A novel paradigm for heart failure with preserved ejection fraction: comorbidities drive myocardial dysfunction and remodeling through coronary microvascular endothelial inflammation. *J Am Coll Cardiol.* 2013;62:263-71. doi:10.1016/j.jacc.2013.02.092



Circulation  
Research

36. Wang TZ, Kodiyankal RPL and Calfee DP. Antimicrobial resistance in nephrology. *Nat Rev Nephrol.* 2019;15:463-481. doi:10.1038/s41581-019-0150-7



# Circulation Research

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## NOVELTY AND SIGNIFICANCE

### *What Is Known?*

- Hypertension is the most common comorbidity of the coronavirus disease 2019 (COVID-19) and had been suggested to be associated with increased mortality.
- Angiotensin converting enzyme inhibitor (ACEIs) and angiotensin II receptor blockers (ARBs) are first-line medications for a large proportion of patients with hypertension.
- Use of ACEI/ARB is a major concern for clinicians in treating COVID-19 patients with hypertension because of the potential effect of ACEI/ARB on increasing the expression of ACE2, the binding receptor and entry point of the coronavirus.



### *What New Information Does This Article Contribute?*

- The incidence of the 28-day all-cause death among patients who had inpatient treatment with ACEI/ARB is significant lower compared with ACEI/ARB non-users, based on the analysis of 1128 hospitalized COVID-19 patients with hypertension.
- After matching and adjusting variables may interfering the effect of ACEI/ARB, in-hospital use of ACEI/ARB still exhibits remarkable association with reduced all-cause mortality of COVID-19 patients with hypertension.
- These findings clearly support recently published recommendations regarding continuation of ACEI or ARB among patients with co-existing hypertension and COVID-19.

Application of ACEI/ARB is a one of major concerns in treating COVID-19 among patients with hypertension, for the possible increased risk by ACEI/ARB on increased ACE2, a direct receptor and entry point of SARS-COV-2. However, the association between ACEI/ARB and COVID-19 mortality in patients with hypertension is largely unknown. Here, the multicentered retrospective cohort study indicated that in-hospital use of ACEI/ARB is associated reduced 28-day all-cause mortality of COVID-19 compared to non-ACEI/ARB group in patients with hypertension. Findings in this study provide direct clinical data to show the potential benefit of ACEI/ARB in treating patients with hypertension and hospitalized with COVID-19 and support the continuous use of ACEI/ARB in those patients as recently recommend by societies.

## FIGURE LEGENDS

**Figure 1.** The flowchart showing the strategy of participant enrollment.

- a, 1128 participants with a history of hypertension enrolled in the hypertension cohort.
- b, 2302 participants without a history of hypertension enrolled in the non-hypertension cohort.
- c, 188 patients with hypertension who taking ACEI and/or ARB during hospitalization were enrolled in the ACEI/ARB cohort. Patients discontinued treatment of hypertension due to inability to take medications or hypotension were not excluded from the cohort.
- d, 940 patients with hypertension who never taking ACEI and ARB during hospitalization were enrolled in the non-ACEI/ARB cohort.
- e, Propensity score-matched age, gender, cough, dyspnea, comorbidities (diabetes, coronary heart disease and chronic renal disease), CT-diagnosed lung lesions, and incidence of increased CRP and creatine. Hospital site as a random effect in the mixed-effect Cox model.
- f, 557 patients with antihypertension drug who never taking ACEI and ARB during hospitalization were enrolled in the secondary non-ACEI/ARB cohort.



**Figure 2.** Kaplan-Meier Curves for cumulative probability of COVID-19 mortality during 28-day follow-up duration in ACEI/ARB or non-ACEI/ARB cohort among 1128 patients with hypertension in unmatched model. The median (IQR) observation time was 28(20-28) in ACEI/ARB cohort and 28(19-28) in non-ACEI/ARB cohort (A). Propensity-score matched model and the median (IQR) observation time was 28(20.5-28) in ACEI/ARB cohort and 28(18-28) in non-ACEI/ARB cohort (B). The blips indicate censoring.

**Table 1** Characteristics of patients with hypertension in ACEI/ARB and non-ACEI/ARB groups before and after propensity score matching.

Parameters	Unmatched			Matched (1:2)		
	ACEI/ARB <sup>†</sup> (n = 188)	Non-ACEI/ARB <sup>‡</sup> (n = 940)	SD <sup>§</sup>	ACEI/ARB <sup>†</sup> (n = 174)	Non-ACEI/ARB <sup>‡</sup> (n = 348)	SD <sup>§</sup>
<b>Clinical characteristics on admission</b>						
Age, median(IQR)	64 (55-68)	64 (57-69)	-0.035	64(56-68)	64(56-69)	-0.046
Male gender, n (%)	100(53.2)	503(53.5)	-0.006	94(54.0)	197(56.6)	-0.052
Female gender, n (%)	88(46.8)	437(46.5)	0.006	80(46.0)	151(43.4)	0.052
Heart rate, median(IQR)	82.0(76.0-95.3)	84.0(78.0-96.0)	-0.062	82(76-94)	82(77-94)	-0.004
Respiratory rate, median(IQR)	20.0(19.0-21.0)	20.0(19.0-22.0)	-0.049	20(19-21)	20(18-21)	0.013
SBP, median(IQR)	132.5(123.0-145.8)	132.0(120.0-144.0)	0.064	134(124-148)	132(121-145)	0.027
DBP, median(IQR)	80.0(72.0-87.8)	80.0(72.0-88.0)	-0.020	80(74-88)	80(73-87)	0.072
Symptom onset to admission, median(IQR), day	10.0(7.0-15.0)	10.0(7.0-15.0)	-0.055	10(7-15)	10(7-15)	-0.111
Fever, n(%)	126(67.0)	700(74.5)	-0.164	120(69.0)	235(67.5)	0.031
Cough, n(%)	122(64.9)	653(69.5)	-0.098	111(63.8)	223(64.1)	-0.006
Fatigue, n(%)	65(34.6)	363(38.6)	-	60(34.5)	131(37.6)	-0.066
Dyspnea, n(%)	38(20.2)	260(27.7)	-0.175	34(19.5)	66(19.0)	0.015
<b>Comorbidities on admission</b>						
Diabetes, n(%)	44(23.4)	196(20.9)	-	40(23.0)	86(24.7)	-0.040
Coronary heart disease, n(%)	29(15.4)	102(10.9)	0.136	24(13.8)	46(13.2)	0.017
Chronic renal diseases, n(%)	7(3.7)	28(3.0)	0.041	7(4.0)	11(3.2)	0.046
Cerebrovascular diseases, n(%)	5(2.7)	36(3.8)	-0.066	4(2.3)	8(2.3)	0.000
Chronic liver disease, n(%)	4(2.1)	17(1.8)	0.023	4(2.3)	5(1.4)	0.064
Chronic obstructive pulmonary disease, n(%)	1(0.5)	5(0.5)	0.000	1(0.6)	1(0.3)	0.044
<b>Chest CT on admission</b>						
Unilateral lesion, n/N(%)	16/173(9.3)	42/895(4.7)	0.180	13/159(8.2)	18/329(5.5)	0.107
Bilateral lesions, n/N (%)	146/173(84.4)	795/895(88.8)	-0.130	137/159(86.2)	284/329(86.3)	-0.005
<b>Laboratory examination on admission</b>						
Leukocyte count > 9.5, 10 <sup>9</sup> /L, n/N (%)	22/183(12.0)	101/883(11.4)	0.018	20/170(11.8)	32/325(9.9)	0.062



Neutrophil count > 6.3, 9.5 10 <sup>9</sup> /L ,n/N (%)	32/183(17.5)	179/882(20.3)	-0.072	28/170(16.5)	64/324(19.8)	-0.085
Lymphocyte count < 1.1, 9.5 10 <sup>9</sup> /L, n/N (%)	82/183(44.8)	426/883(48.2)	-0.069	77/170(45.3)	157/325(48.3)	-0.060
	<b>Unmatched</b>			<b>Matched (1:2)</b>		
<b>Parameters</b>	<b>ACEI/ARB<sup>†</sup> (n = 188)</b>	<b>Non-ACEI/ARB<sup>‡</sup> (n = 940)</b>	<b>SD<sup>§</sup></b>	<b>ACEI/ARB<sup>†</sup> (n = 174)</b>	<b>Non-ACEI/ARB<sup>‡</sup> (n = 348)</b>	<b>SD<sup>§</sup></b>
C-reactive protein increase>ULN*, n/N (%)	78/133(58.7)	443/641(69.1)	-0.219	74/124(59.7)	136/221(61.5)	-0.038
Procalcitonin level increase> ULN*, n/N (%)	40/144(27.8)	237/727(32.6)	-0.105	38/134(28.4)	83/266(31.2)	-0.062
ALT increase> 40 U/L, n/N (%)	33/179(18.4)	170/877(19.4)	-0.024	30/166(18.1)	52/323(16.1)	0.052
AST increase> 40 U/L, n/N (%)	40/179(22.4)	247/877(28.2)	-0.134	37/166(22.3)	79/323(24.5)	-0.051
Creatinine>ULN*, n/N (%)	14/166(8.4)	92/845(10.9)	-0.083	11/153(7.2)	25/308(8.1)	-0.035
D-dimer> ULN*, n/N (%)	67/149(45.0)	392/743(52.8)	-0.156	63/137(46.0)	140/273(51.3)	0.106
K <sup>+</sup> < 3.5 mmol/L, n/N (%)	51/179(28.5)	233/873(26.7)	0.040	47/166(28.3)	88/323(27.2)	0.024
Platelets < ULN*, n/N (%)	22/183(12.0)	106/883(12.0)	0.000	21/170(12.35)	44/325(13.54)	-0.733
LDL-c, mmol/L, median (IQR)	2.2(1.8-2.8) [n = 151]	2.3(1.8-2.8) [n = 721]	-0.010	2.2(1.8-2.8) [n = 140]	2.2(1.8-2.7) [n = 257]	0.042
SaO <sub>2</sub> , <95%, n/N (%)	30/138(21.7)	146/692(21.1)	0.016	28/125(22.4)	53/254(20.9)	0.037
Blood glucose, median (IQR), mmol/L	6.0(5.1-8.3) [n = 163]	6.2(5.2-8.4) [n = 803]	-0.011	6(5.1-8.3) [n = 151]	6.2(5.3-8.7) [n = 298]	-0.047

Abbreviations: ACEI, angiotensin converting enzyme inhibitor; ARB, angiotensin II receptor blocker; SBP, Systolic blood pressure; DBP, Diastolic blood pressure; COPD, Chronic obstructive pulmonary disease; ALT, alanine transaminase; AST, Aspartate transaminase; IQR, Interquartile range; SD, Standardized difference.

\*Upper limit of normal (ULN) was defined according to criteria in each hospital and normal ranges of tests in each hospital were provided in Online Table II. Increase or decrease in a biomarker indicates the number of patients have increased or decreased levels of biomarker at baseline.

<sup>†</sup>Patients with hypertension who taking ACEI and/or ARB during hospitalization were enrolled in the ACEI/ARB cohort. Patients discontinued treatment of hypertension due to inability to take medications or hypotension were not excluded from the cohort.

<sup>‡</sup>Patients with hypertension who never taking ACEI and ARB during hospitalization were enrolled in the non-ACEI/ARB cohort.

<sup>§</sup>SD, Standardized differences be used to compare the mean of baseline covariate between ACEI/ARB and non-ACEI/ARB group.

**Table 2** In-hospital management of ACEI/ARB and non-ACEI/ARB groups.

Treatment	ACEI/ARB* (n = 188)	Non-ACEI/ARB† (n = 940)	Absolute difference (95% CI), %	P value‡
Traditional Chinese medicine n (%)	171(91.0)	807(85.9)	5.1(-0.2 to 10.4)	0.08
Antiviral drug, n (%)	167(88.8)	768(81.7)	7.1(1.2 to 13.0)	0.02
Antibiotics drug, n (%)	144(76.6)	718(76.4)	0.2(-6.4 to 6.9)	0.99
Nasal cannula oxygen inhalation§, n (%)	142(75.5)	745(79.3)	-3.7(-10.1 to 2.7)	0.30
Systemic corticosteroids, n (%)	66(35.1)	380(40.4)	-6.2(-13.9 to 1.5)	0.20
Antidiabetic drug, n (%)	55(29.3)	226(24.0)	5.2(-1.6 to 12.0)	0.16
Lipid lowering drug, n (%)	43(22.9)	94(10.0)	12.8(7.8 to 18.0)	1.51E-6
Immunoglobulin, n (%)	51(27.1)	279(29.7)	21.2(16.7 to 25.6)	0.54
Vasoactive drug, n (%)	18(9.6)	106(11.3)	-1.7(-6.6 to 3.2)	0.58
Noninvasive ventilation  , n (%)	15(8.0)	20(11.4)	5.9(3.1 to 8.6)	0.21
Invasive ventilation  , n (%)	9(5.0)	51(5.4)	3.5(1.4 to 5.6)	0.86
Antifungal medications, n (%)	7(3.7)	40(4.3)	-0.5(-3.7 to 2.6)	0.89
Renal replacement therapy, n(%)	1(0.5)	14(1.5)	0.1(-0.9 to 1.2)	0.49
Extracorporeal membrane oxygenation  , n (%)	0(0.0)	4(0.4)	-0.4(-1.4 to 0.5)	1.00
Diuretic, n (%)	58(30.9)	120(12.8)	18.1(12.4 to 23.8)	1.06E-9
CCB, n (%)	103(54.8)	489(52.0)	2.8(-5.1 to 10.6)	0.54
beta-blocker, n (%)	53(28.2)	168(17.9)	10.3(4.1 to 16.5)	0.002
alpha-blocker, n (%)	4(2.1)	21(2.2)	-0.1(-2.4 to 2.2)	1.00
ACEI, n (%)	31(16.5)	0(0.0)	16.5(13.9 to 19.1)	3.34E-35



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ARB, n (%)	157(83.5)	0(0.0)	83.5(78.1 to 88.9)	8.16E-199
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\*Patients with hypertension who taking ACEI and/or ARB during hospitalization were enrolled in the ACEI/ARB cohort. Nine patients in ACEI/ARB group were taken off ACEI/ARB due to hypotension (2 cases), mechanical ventilation without nasal feeding and unable to orally taking medicines (1 case), and increased creatinine level (> 1.5-fold of ULN; 6 cases). These patients were still included in ACEI/ARB group. Patients placed on ACEI/ARB on admission were included in the ACEI/ARB group.

†Patients with hypertension who never taking ACEI and ARB during hospitalization were enrolled in the non-ACEI/ARB cohort.

‡The *P* value was calculated by Fisher’s exact test or  $\chi^2$  test.

§Nasal cannula oxygen inhalation was taken in isolation.

||Noninvasive ventilation, invasive ventilation, and extracorporeal membrane oxygenation are at mutually exclusive.



# Circulation Research

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**Table 3** Hazard ratios and incidence rate ratios for outcomes in ACEI/ARB group versus non-ACEI/ARB group under mixed-effect Cox model and propensity score-matching model

ACEI/ARB vs non-ACEI/ARB	Unmatched				Matched (1:2) <sup>†</sup>			
	IRD (100 Person-Day) (95% CI)	Crude HR(95% CI)	P value <sup>§</sup>	Mixed-effect Model* HR(95% CI)	P value <sup>  </sup>	Mixed-effect Model IRD (100 Person-Day) (95% CI)	HR(95% CI)	P value <sup>  </sup>
All-cause mortality	-0.24(-0.43,-0.05)	0.37(0.17,0.79)	0.01	0.42(0.19,0.92)	0.03	-0.27(-0.48,-0.07)	0.37(0.15-0.89)	0.03
Septic shock	-0.19(-0.36,-0.01)	0.38(0.17,0.87)	0.02	0.36(0.16,0.84)	0.02	-0.20(-0.39,-0.01)	0.32(0.13-0.80)	0.01
ARDS	-0.23(-0.52,0.07)	0.70(0.47,1.02)	0.06	0.69(0.47,1.02)	0.06	-0.32(-0.65,0.01)	0.65(0.41-1.04)	0.07
DIC	-0.09(-0.19,0.00)	0.04(0.0, 4.50)	0.18	‡	‡	-0.10(-0.19,0.00)	‡	‡
Acute kidney injury	0.05(-0.10,0.20)	1.17(0.60,2.25)	0.65	1.03(0.51,2.07)	0.94	-0.04(-0.23,0.16)	0.78(0.37-1.65)	0.52
Acute heart injury	-0.10(-0.29,0.08)	0.91(0.57,1.45)	0.70	0.89(0.55,1.44)	0.64	0.00(-0.27,0.27)	0.76(0.44-1.32)	0.33

Abbreviations: HR, Hazard ratio; CI, Confidence interval; IRD, Incidence rate differences, an absolute difference in the incidence between two groups.

\*The mixed-effect Cox model is a conditional model including a random per-center effect. Unmatched were adjusted for age, gender, comorbidities (diabetes, coronary heart disease, cerebrovascular disease, and chronic renal disease), and in-hospital medications (antiviral drug and lipid lowering drug).

†In the mixed-effect Cox model after propensity score matching, age, gender, cough, dyspnea, comorbidities (diabetes, coronary heart disease and chronic renal disease), CT-diagnosed lung lesions, and incidence of increased CRP and creatinine were matched. Modeled center as a random effect in the multivariate analyses followed by adjustment for antiviral drug, lipid lowering drug, D-dimer, and unilateral lesion.

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‡The incidence of DIC was 0 in the ACEI/ARB cohort. HR in DIC was not calculated.

§P values were calculated based on Cox proportional hazard model.

||P values were calculated based on mixed-effect Cox model.



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