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► **To cite this version:**

C. Villain, M. Metzger, S. Liabeuf, A. Hamroun, S. Laville, et al.. Effectiveness and Tolerance of Renin-Angiotensin System Inhibitors With Aging in Chronic Kidney Disease. *Journal of the American Medical Directors Association*, 2021, S1525-8610 (21), pp.00939-7. 10.1016/j.jamda.2021.10.019 . inserm-03587209

HAL Id: inserm-03587209

<https://inserm.hal.science/inserm-03587209>

Submitted on 22 Jul 2024

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Effectiveness and Tolerance of Renin-Angiotensin System Inhibitors with Aging in Chronic Kidney Disease

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Running Head: RAS inhibitors by age in CKD

Key words: renin angiotensin system inhibitors, chronic kidney disease, propensity score analysis

Funding sources

CKD-REIN is supported by the Agence Nationale de la Recherche through the 2010 *Cohortes-Investissements d’Avenir* program and the 2010 national *Programme Hospitalier de Recherche Clinique* program. CKD-REIN is also supported by a public-private partnership that has included Amgen, Fresenius Medical Care and GlaxoSmithKline (GSK) since 2012,

Lilly France since 2013, Otsuka Pharmaceutical since 2015, Baxter and Merck Sharp & Dohme-Chibret (MSD France) from 2012 to 2017, Sanofi-Genzyme from 2012 to 2015, and Vifor Fresenius since 2018. Inserm Transfert set up the collaboration in 2011 and has managed it since then.

Word count – abstract: 262 (max. 300 words)

Word count – main text: 2962 (excluding Abstract, max. 3000 words)

Number of Figures/Tables: 5 (max. 5)

Number of references: 42 (max. 50)

Brief summary:

In a cohort of CKD patients, RASi prescription was significantly associated with a lower combined risk of end-stage kidney disease and death, without excess risk of adverse effects. Age did not modify these relations.

Acknowledgments

We thank the CKD-REIN study coordination staff for their efforts in setting up the CKD-REIN cohort: Marie Metzger, Elodie Speyer, Céline Lange, Sophie Renault, Reine Ketchemin, and all the clinical research associates. We also thank Jo Ann Cahn and David Fraser (Biotech Communication SARL, Ploudalmézeau, France) for the English revision of our manuscript.

Sponsor's role: None.

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Conflicts of interest

CV, MM, SL, AH, NM, CC, DF, LF, CJ, ML, SB, CA, RPF, and TH have no disclosures relative to this study.

BS has received grants for CKD REIN from the French National Research Agency (ANR-IA-COH-2012/3731), PHRC national 2010, Amgen, Encastra, Fresenius Medical Care, GlaxoSmithKline, Otsuka, Vifor Fresenius, and Merck Sharp and Dohme-Chibret.

ZM has received grants for CKD REIN and other research projects from Amgen, Sanofi-Genzyme, GlaxoSmithKline, Merck Sharp and Dohme-Chibret, Lilly, Fresenius Medical Care, Baxter, Otsuka, Daichi, Astellas, and the French government.

1 **Abstract**

2 **Objectives**

3 Renin-angiotensin system inhibitors (RASi) are recommended for slowing chronic kidney
4 disease (CKD) progression to kidney failure. Their effectiveness and tolerance as patients age
5 remain uncertain because older patients have often been excluded from clinical trials.

6 **Design**

7 CKD-REIN cohort study.

8 **Setting and Participants**

9 We studied 2762 patients with CKD stages 3–4 and a clinical indication for RASi enrolled
10 between 2013 and 2016 in 40 nephrology clinics nationally representative in France.

11 **Methods**

12 The primary outcome was the occurrence of kidney failure or death. The secondary outcomes
13 were the occurrence of cardiovascular events and hospitalizations with acute kidney injury
14 (AKI) or hyperkalemia. A propensity score analysis was performed. We used Cox models to
15 estimate hazard ratios (HRs) for each outcome associated with RASi prescription and tested
16 interactions with age.

17 **Results**

18 Patients' mean age was 67 years, including 841 (30%) aged 75 years and older; 2178 (79%)
19 were prescribed RASi. During a median follow-up of 4.6 years, 33% of patients reached
20 kidney failure or died. RASi prescription was associated with a lower risk of kidney failure or
21 death (HR, 0.79; 95%CI, 0.66-0.95), an association not modified by age (P for
22 interaction=0.72). It was not significantly associated with cardiovascular events. During the
23 first 3 years of follow-up, 14% of patients were hospitalized with AKI or hyperkalemia, but

24 risk was not higher among those prescribed RASi (HR, 0.75; 95%CI, 0.55-1.02) and age did
25 not modify its effect (P for interaction=0.28).

26 **Conclusions and Implications**

27 This study shows that aging does not appear to modify either RASi's beneficial effects on
28 major CKD outcomes or their potential adverse effects.

29

30 **Introduction**

31 Renin-angiotensin system inhibitors (RASi, including angiotensin-converting enzyme [ACE]
32 inhibitors and angiotensin II receptor blockers [ARBs]) are the main drugs recommended for
33 reduction of blood pressure and albuminuria in patients with chronic kidney disease (CKD).¹
34 Since the early 1990s, several randomized trials in patients with CKD have shown that RAS
35 blockade can slow the progression to kidney failure, mainly through their effect on blood
36 pressure and albuminuria.²⁻¹¹ Reduction in mortality and cardiovascular risk is still
37 controversial in patients with CKD.^{2,4,8,9,12,13}

38 Chronic kidney disease is frequent in older people and affects 28% of patients aged 65 years
39 or more in France.¹⁴ In the United States, nearly half of people aged 80 years or older have
40 CKD stages 3-5.¹⁵ Age has been associated with the underuse of RASi in CKD but data on
41 their effectiveness and tolerance are scarce in older patients.^{16,17} In those randomized
42 controlled trials that did not explicitly limit age for study inclusion, the mean age of included
43 patients has not exceeded 67 years.^{2,4-10,13} As most older patients with CKD have low
44 albuminuria and a low risk of progression to kidney failure, but a higher risk of death, the
45 benefit of RASi in this population are questionable.^{18,19} In a randomized trial assessing the
46 nephroprotective efficacy of ARBs in patients with diabetes and nephropathy, those aged 65
47 to 74 years had the same advantages as those younger than 65 years.²⁰ Observational studies,
48 however, report conflicting findings about RASi's benefits in delaying kidney failure and
49 mortality in older patients with CKD.²¹⁻²⁴ This population is also particularly exposed to
50 drug-related adverse effects.²⁵ Randomized controlled trials have not consistently reported the
51 adverse effects of RASi and their frequency. The trial on ARBs mentioned above found
52 similar incidence rates of acute kidney injury (AKI) and hyperkalemia in patients aged 65 to

53 74 years and in those younger than 65 years,²⁰ but none of the observational studies assessed
54 the impact of age on RASi safety in CKD.^{21–24}

55 Accordingly, current evidence about the benefits and risks of RASi use in older patients with
56 CKD is limited, although these may well differ from those in younger patients. We used a
57 propensity score analysis to investigate the impact of age on the effectiveness and tolerance of
58 RASi in patients with CKD enrolled in the CKD-REIN cohort study.

59

60 **Methods**

61 *Population*

62 This study used data from the CKD-REIN cohort (ClinicalTrials.gov identifier:
63 NCT03381950), the design of which has previously been described.²⁶ Briefly, 3033 patients
64 with CKD were recruited from 40 nephrology clinics in France between July 2013 and April
65 2016. During a census phase, we screened all adults with a proven CKD diagnosis and two
66 measures of estimated glomerular filtration rate (eGFR) between 15 and 60 mL/min/1.73m² at
67 least 1 month apart, with no prior chronic dialysis or kidney transplantation. Eligible patients
68 were recruited within the next few weeks during a routine nephrology visit, excluding those
69 who had started kidney replacement therapy in the meantime. Patients under the age of 18 or
70 unable to give informed consent were also excluded. The study protocol was approved by the
71 INSERM institutional review board (reference: IRB00003888), and all participants provided
72 informed consent.

73 For this analysis, we excluded 121 patients with baseline eGFR <15 ml/min/1.73m² and 32
74 treated by renin inhibitors, which also act on the RAS but have a different benefit-risk ratio
75 than ACE inhibitors and ARBs.²⁷ We also excluded 118 patients who had no indication for
76 RASi at baseline, because they did not have hypertension, heart failure, coronary artery
77 disease, severely increased albuminuria, or moderately increased albuminuria associated with
78 diabetes. Finally, this analysis included 2762 patients (Figure 1).

79 *Data Collection*

80 Data were collected from medical records and patient interviews by trained clinical research
81 associates. RASi included ACE inhibitors and ARBs. We calculated eGFR using the Chronic
82 Kidney Disease–Epidemiology Collaboration (CKD-EPI) creatinine equation with correction
83 for ethnicity. Albuminuria or proteinuria were collected either from 24-h or spot urine

84 samples and expressed according to the Kidney Disease: Improving Global Outcomes
85 (KDIGO) albuminuria categories (A1, normal; A2, moderately increased; A3, severely
86 increased).¹ When data on the albumin-to-creatinine ratio were missing, they were imputed
87 from the results of the 24-hour urine collection or the protein-to-creatinine ratio, as described
88 elsewhere.²⁸ Participants were classified as having diabetes if it was reported in their medical
89 records, or they were prescribed a glucose-lowering medication, or they had an HbA1c level
90 $\geq 7\%$. Hypertension was defined as a history of hypertension or antihypertensive medication
91 prescription without a concurrent indication. The Charlson Comorbidity Index was
92 computed.²⁹ Data were available for five activities of daily living (ADL) from the Katz Index
93 (bathing, dressing, going to the toilet, transfer, and feeding) and four instrumental ADL
94 (IADL: ability to use the telephone, handle finances, take medication, and travel).^{30,31}
95 Educational level was classified in two categories: ≤ 12 versus > 12 years. Medication
96 compliance was evaluated with a self-administered questionnaire.³² The method used to
97 collect and assess any adverse drug reactions occurring during follow-up in the cohort has
98 been described previously.³³ Exhaustive data on adverse drug reactions were available for the
99 first three years of follow-up.

100 *Outcomes*

101 The primary outcome was defined as death or kidney failure, defined as initiation of kidney
102 replacement therapy (i.e., kidney transplantation or maintenance dialysis), during follow-up.
103 Secondary outcomes included: i) kidney failure; ii) death before kidney failure occurrence; iii)
104 major cardiovascular events, defined as cardiovascular death, stroke, myocardial infarction, or
105 hospitalization or an emergency department visit for heart failure; iv) AKI or hyperkalemia
106 requiring or occurring during hospitalization. Major cardiovascular events, AKI or
107 hyperkalemia events were identified from medical records and hospital discharge reports.

108 AKI events were further adjudicated by three nephrologists (AH, LF, and ML) using the
109 current KDIGO definition for AKI.³⁴ Data on death and kidney failure were available until
110 February 1, 2020. Data on major cardiovascular events and AKI or hyperkalemia requiring
111 hospitalizations were available for the first three years of follow-up.

112 *Statistical Analysis*

113 We first described patient baseline characteristics according to RASi prescription. Cumulative
114 incidence curves were then computed and compared according to age and RASi prescription
115 for each outcome. To evaluate the impact of age on the effectiveness and tolerance of RASi in
116 patients with CKD and to minimize indication bias due to the observational design of the
117 study, we performed a propensity score analysis using the inverse probability of treatment
118 weighting (IPTW) method.³⁵ A specific propensity score was performed for each outcome as
119 detailed in Appendix 1.³⁶ Hazard ratios (HR) for each outcome were then estimated by using
120 Cox models. Confidence intervals (CI) were obtained by bootstrapping. Interaction between
121 age as a continuous variable and RASi prescription was tested. Hazard ratios were also
122 computed for 4 age groups (<60, 60-69, 70-79, and ≥ 80 years) and 2 groups according to
123 eGFR level (<30 and ≥ 30 ml/min/1.73m²).

124 We conducted secondary analyses to test the robustness of the associations by subgroups
125 according to sex, diabetic status, albuminuria category, and the presence of a cardiovascular
126 disease (CVD) for which RASi are indicated, i.e., heart failure and coronary artery disease.
127 We also tested the interaction by introducing age as a binary variable (< or ≥ 75 years). We
128 also assessed the effect of ACE inhibitors and ARBs separately. For the analysis of secondary
129 outcomes, Fine and Gray competing risk regression models were used with the propensity
130 scores. The events considered for the competing risk analyses were (i) kidney failure for the
131 analysis of death before kidney failure, (ii) death not related to kidney disease for the analysis

132 of kidney failure, (ii) death not related to cardiovascular diseases for the analysis of major
133 cardiovascular events, and (iv) death and kidney failure for the analysis of AKI or
134 hyperkalemia. Lastly, we used time-varying Cox models weighted for the propensity score.
135 We calculated a propensity score for each patient at each time point. Given that the variables
136 other than age, eGFR, and RASi prescription were not available for all the time points, the
137 propensity score was mainly computed with baseline values. Missing data were considered by
138 using multiple imputations. All statistical analyses were performed with R software (R
139 Foundation, Vienna, Austria, version 3.5.1).³⁷ A *P*-value <0.05 was considered statistically
140 significant.

141 **Results**

142 *Baseline Patient Characteristics*

143 Among the 2762 studied patients, 2178 (79%) were prescribed RASi at baseline (Table 1 and
144 Table A1). Their mean age was 67.0 ± 12.8 years; 841 (30%) were aged 75 years or older.

145 Compared to patients to whom RASi were prescribed, those whose physicians did not
146 prescribe it for them were significantly older, more often women and dependent for ADL and
147 IADL, had visited a cardiologist in the past year more often, and had more AKI, heart failure,
148 dysrhythmia, stroke history, and lower levels of both eGFR and blood potassium.

149 *Crude Risk of Kidney Failure, Death, or Major Cardiovascular Events*

150 Over a median follow-up of 4.6 years [interquartile range (IQR) 3.6-5.2], 539 (19.5%)
151 patients reached kidney failure and 377 (13.6%) patients died before kidney failure.

152 Prescription of RASi was significantly associated with a lower risk of the combined outcome
153 of kidney failure or death, and of death before kidney failure, but not of kidney failure alone
154 (Figures A1 to A3). During a median follow-up of 3.0 years (IQR 2.9-3.0), 306 (11.1%)
155 patients experienced a major cardiovascular event, including 29 (9.5%) cardiovascular deaths,
156 49 (16.0%) strokes, 51 (16.7%) myocardial infarctions, and 177 (57.8%) hospitalizations or
157 emergency department visits for heart failure. In the crude analysis, RASi prescription was
158 significantly associated with a lower risk of major cardiovascular events (Figure A4).

159 *Adverse drug reactions*

160 Of the 2178 patients who were prescribed RASi at baseline, 164 experienced at least one
161 RASi-related adverse drug reaction during follow-up; these were mainly serum creatinine
162 elevation, AKI, hypotension, cough, and high serum potassium. Adverse drug reactions
163 occurred in 8.2%, 8.0%, 7.6%, and 5.2% of RASi users in the <60, 60-69, 70-79, and ≥ 80 age
164 classes, respectively ($p=0.47$). In 73% of cases, the drug was discontinued (at least

165 transiently) after the occurrence of the adverse drug reaction, and dose adjustment was
166 necessary in 10% of cases. Over a median follow-up period of 3.0 years, a total of 380
167 (13.8%) patients were hospitalized with AKI (n=322; 11.7%) and/or hyperkalemia (n=104;
168 3.8%). RASi prescription was significantly associated with a lower risk of hospitalization for
169 AKI or hyperkalemia (Figure A5). Patients aged 75 years or older had a higher risk of both
170 events than younger patients.

171 *Propensity Score Analyses*

172 Propensity scores were computed for each outcome. The standardized differences of <10%
173 for each covariate in each score indicate that the treated and untreated patients were well
174 balanced for all variables after weighting (Tables 1 and A2). The propensity score analyses
175 showed a statistically significant association between the prescription of RASi and the risk of
176 kidney failure or death (Table 2). Age was not a significant effect modifier in this association.
177 However, the prescription of RASi was not significantly associated with any of the secondary
178 outcomes and age did not modify these associations. In subgroups of age, RASi prescription
179 was associated with a decreased risk of death or kidney failure, and a decreased risk of kidney
180 failure in patients aged 60-69 years, but all the others studied association were not statistically
181 significant (Table 3). Considering age as a binary variable (< or \geq 75 years) did not change
182 our findings, and all the p values for the interaction between age and RASi prescription were
183 >0.15 for all outcomes studied. The HRs estimated for time-varying Cox models were
184 consistent with those found in the main analysis, for all outcomes studied (Table A3).

185 *Subgroups Analyses*

186 According to eGFR level, RASi prescription was associated with a decreased risk of death or
187 kidney failure in patients with an eGFR <30 ml/min/1.73m², but the others studied association
188 were not statistically significant (Table 4). Age did not modify the association between the

189 outcomes and RASi prescription in the subgroups of eGFR. In patients with and without
190 diabetes, and with and without CVD for which RASi was indicated, as well as with and
191 without moderate or severe albuminuria, this prescription was not significantly associated
192 with the primary outcome but HRs were similar between both groups, and age did not modify
193 the associations (Figure A6). The prescription of RASi was significantly associated with a
194 lower risk of the primary outcome in men, but not in women; nonetheless, these HRs were of
195 the same order of magnitude in both subgroups. Finally, a decrease in the risk of kidney
196 failure or death appeared more strongly associated with prescription of ARBs than ACE
197 inhibitors, but the difference in the risk of this primary outcome between these two classes of
198 RASi was not statistically significant.

199 **Discussion**

200 Using propensity score analyses on data from the CKD-REIN cohort, we found that RASi
201 prescription at baseline was significantly associated with a lower risk of the combined
202 outcome of kidney failure or death in patients with CKD and that age did not modify the
203 effect of the prescription on the outcome. This cohort of patients with CKD receiving
204 nephrology care had no higher risk of hospitalization with AKI or hyperkalemia than those
205 not prescribed RASi. The originality of this study was to assess the impact of age on the
206 effectiveness and tolerance of RASi in this population; a key finding was the lack of an
207 association between older age and a higher risk of RASi-related adverse effects.

208 In our study, prescription of RASi was not associated with a significant decrease in the risk of
209 kidney failure alone. However, in studies evaluating RASi, CKD progression was often
210 defined by multiple endpoints also including a percentage decline of eGFR or a doubling in
211 serum creatinine. Limiting our analysis to kidney failure may have underestimated the
212 association between RASi and CKD progression and reduced statistical power. Moreover,
213 RASi has shown stronger benefits against CKD progression in patients with high levels of
214 albuminuria.^{38,39} In our study, the HRs for the association between the primary outcome and
215 RASi prescription did not differ by albuminuria level. This result should nonetheless be
216 interpreted with caution in these patients whose history of albuminuria before they began
217 RASi treatment is unknown.

218 In older patients with CKD, the risk of death is often higher than the risk of progression to
219 kidney failure, as we observed in our cohort.⁴⁰ Moreover, as conservative management is
220 increasingly proposed to older CKD patients reaching kidney failure, the initiation of kidney
221 replacement therapy may not be an appropriate outcome in this specific population. For this
222 reason, we considered that a combined endpoint (i.e., death or kidney failure) was the most

223 relevant primary outcome. In our study, the prescription of RASi was not significantly
224 associated with a lower risk of death alone, but rather with the risk of the combined primary
225 outcome, and age did not change this association. These results are at odds with a previous
226 study using the same combined endpoint to evaluate RASi prescription in a cohort of
227 veterans.²¹ Like the randomized controlled trials including patients with CKD and
228 intermediate cardiovascular risk, our study did not find a lower risk of cardiovascular events
229 associated with RASi prescription.^{4,8}

230 In our study, prescription of RASi was not associated with a higher risk of hospitalization for
231 AKI or hyperkalemia. This unexpected result may be explained by a selection bias in patients
232 receiving RASi for a long time before inclusion in the study. This bias could be suspected
233 from the observation of a higher prevalence of AKI history in patients not prescribed RASi at
234 baseline, compared to their counterparts with prescriptions. This variable was, however, taken
235 into account in the propensity score and was well balanced after weighting. Another major
236 reason for stopping RASi is hyperkalemia, which usually occurs early after starting the
237 treatment, but the patients' history of hyperkalemia was not available.^{41,42} The fact that
238 patients were receiving nephrology care may also explain this result. However, the main focus
239 of this study was on the potential effect modification by age of the association we studied. Its
240 lack of significance indicates that older patients did not experience more adverse effects
241 related to RASi than younger ones.

242 This study has several strengths. The analysis was based on a large cohort of CKD patients
243 including a high proportion of patients aged 75 years or older. The use of propensity scoring
244 including numerous variables well-balanced between treated and untreated patients minimized
245 indication bias linked to the observational design of the study. Confounding by indication,
246 however, cannot be totally excluded, which is the first limitation of our analysis.

247 Nevertheless, this study population is closer to that of nephrology outpatients than that of
248 controlled randomized trials. The second limitation is that patients receiving RASi at baseline
249 may have been treated for a long time before the study began, and those not receiving them
250 may have used them previously and stopped them due to adverse effects like hyperkalemia
251 and AKI. This may have biased our results, even though AKI history and duration of kidney
252 disease were included in the propensity scores. However, this should not have biased our key
253 finding, which is the lack of interaction with age in these associations. Thirdly, all the
254 variables were not available at each follow-up visit, which might have biased the analyses
255 based on time-varying Cox models. Finally, patient recruitment from nephrology
256 consultations restricts the generalizability of our results.

257 **Conclusions and Implications**

258 To conclude, in CKD patients receiving nephrologist-led care, RASi prescription, compared
259 with its nonprescription, was associated with a lower risk of the combined outcome of death
260 or kidney failure with no apparent increase in the risk of AKI or hyperkalemia requiring
261 hospitalization. RASi are recommended in CKD to control blood pressure and albuminuria,
262 but are often underused in older patients.^{1,16,17} The finding that age did not modify the
263 associations we studied indicates that prescription of RASi appears to be equally beneficial
264 and safe in young and old patients with CKD.

References

1. Kidney Disease Improving Global Outcomes (KDIGO) guidelines. Summary of Recommendation Statements. *Kidney Int Suppl.* 2013;3(1):5-14. doi:10.1038/kisup.2012.77
2. Jafar TH, Schmid CH, Landa M, et al. Angiotensin-converting enzyme inhibitors and progression of nondiabetic renal disease. A meta-analysis of patient-level data. *Ann Intern Med.* 2001;135(2):73-87.
3. Lewis EJ, Hunsicker LG, Bain RP, Rohde RD. The effect of angiotensin-converting-enzyme inhibition on diabetic nephropathy. The Collaborative Study Group. *N Engl J Med.* 1993;329(20):1456-1462. doi:10.1056/NEJM199311113292004
4. Lewis EJ, Hunsicker LG, Clarke WR, et al. Renoprotective effect of the angiotensin-receptor antagonist irbesartan in patients with nephropathy due to type 2 diabetes. *N Engl J Med.* 2001;345(12):851-860. doi:10.1056/NEJMoa011303
5. Hou FF, Zhang X, Zhang GH, et al. Efficacy and Safety of Benazepril for Advanced Chronic Renal Insufficiency. *N Engl J Med.* 2006;354(2):131-140. doi:10.1056/NEJMoa053107
6. Maschio G, Alberti D, Janin G, et al. Effect of the Angiotensin-Converting-Enzyme Inhibitor Benazepril on the Progression of Chronic Renal Insufficiency. *N Engl J Med.* 1996;334(15):939-945. doi:10.1056/NEJM199604113341502
7. Ruggenti P, Perna A, Gherardi G, et al. Renoprotective properties of ACE-inhibition in non-diabetic nephropathies with non-nephrotic proteinuria. *Lancet Lond Engl.* 1999;354(9176):359-364. doi:10.1016/S0140-6736(98)10363-X

8. Brenner BM, Cooper ME, de Zeeuw D, et al. Effects of losartan on renal and cardiovascular outcomes in patients with type 2 diabetes and nephropathy. *N Engl J Med.* 2001;345(12):861-869. doi:10.1056/NEJMoa011161
9. Randomised placebo-controlled trial of effect of ramipril on decline in glomerular filtration rate and risk of terminal renal failure in proteinuric, non-diabetic nephropathy. The GISEN Group (Gruppo Italiano di Studi Epidemiologici in Nefrologia). *Lancet Lond Engl.* 1997;349(9069):1857-1863.
10. Parving HH, Lehnert H, Bröchner-Mortensen J, et al. The effect of irbesartan on the development of diabetic nephropathy in patients with type 2 diabetes. *N Engl J Med.* 2001;345(12):870-878. doi:10.1056/NEJMoa011489
11. Bidani AK, Polichnowski AJ, Loutzenhiser R, Griffin KA. Renal microvascular dysfunction, hypertension and CKD progression. *Curr Opin Nephrol Hypertens.* 2013;22(1):1-9. doi:10.1097/MNH.0b013e32835b36c1
12. Solomon SD, Rice MM, A Jablonski K, et al. Renal function and effectiveness of angiotensin-converting enzyme inhibitor therapy in patients with chronic stable coronary disease in the Prevention of Events with ACE inhibition (PEACE) trial. *Circulation.* 2006;114(1):26-31. doi:10.1161/CIRCULATIONAHA.105.592733
13. Mann JF, Gerstein HC, Pogue J, Bosch J, Yusuf S. Renal insufficiency as a predictor of cardiovascular outcomes and the impact of ramipril: the HOPE randomized trial. *Ann Intern Med.* 2001;134(8):629-636.
14. Stengel B, Metzger M, Froissart M, et al. Epidemiology and prognostic significance of chronic kidney disease in the elderly--the Three-City prospective cohort study. *Nephrol*

- Dial Transplant Off Publ Eur Dial Transpl Assoc - Eur Ren Assoc.* 2011;26(10):3286-3295. doi:10.1093/ndt/gfr323
15. Bowling CB, Muntner P. Epidemiology of chronic kidney disease among older adults: a focus on the oldest old. *J Gerontol A Biol Sci Med Sci.* 2012;67(12):1379-1386. doi:10.1093/gerona/gls173
 16. Pecoits-Filho R, Fliser D, Tu C, et al. Prescription of renin-angiotensin-aldosterone system inhibitors (RAASi) and its determinants in patients with advanced CKD under nephrologist care. *J Clin Hypertens Greenwich Conn.* 2019;21(7):991-1001. doi:10.1111/jch.13563
 17. Villain C, Liabeuf S, Metzger M, et al. Impact of age on cardiovascular drug use in patients with chronic kidney disease. *Clin Kidney J.* 2020;13(2):199-207. doi:10.1093/ckj/sfz063
 18. O'Hare AM, Kaufman JS, Covinsky KE, Landefeld CS, McFarland LV, Larson EB. Current guidelines for using angiotensin-converting enzyme inhibitors and angiotensin II-receptor antagonists in chronic kidney disease: is the evidence base relevant to older adults? *Ann Intern Med.* 2009;150(10):717-724.
 19. O'Hare AM, Hotchkiss JR, Tamura MK, et al. Interpreting treatment effects from clinical trials in the context of real-world risk information: the example of end-stage renal disease prevention in older adults. *JAMA Intern Med.* 2014;174(3):391-397. doi:10.1001/jamainternmed.2013.13328

20. Winkelmayr WC, Zhang Z, Shahinfar S, Cooper ME, Avorn J, Brenner BM. Efficacy and safety of angiotensin II receptor blockade in elderly patients with diabetes. *Diabetes Care*. 2006;29(10):2210-2217. doi:10.2337/dc06-0570
21. Arora P, Golzy M, Patel N, Quigg R, Carter RL, Lohr JW. Renin-Angiotensin-Aldosterone System Blockers in Elderly Adults with Chronic Kidney Disease without Diabetes Mellitus or Proteinuria. *J Am Geriatr Soc*. 2015;63(12):2478-2484. doi:10.1111/jgs.13842
22. Molnar MZ, Kalantar-Zadeh K, Lott EH, et al. ACE Inhibitor and Angiotensin Receptor Blocker Use and Mortality in Patients with Chronic Kidney Disease. *J Am Coll Cardiol*. 2014;63(7):650-658. doi:10.1016/j.jacc.2013.10.050
23. Oh YJ, Kim SM, Shin BC, et al. The Impact of Renin-Angiotensin System Blockade on Renal Outcomes and Mortality in Pre-Dialysis Patients with Advanced Chronic Kidney Disease. *PloS One*. 2017;12(1):e0170874. doi:10.1371/journal.pone.0170874
24. Hsu T-W, Liu J-S, Hung S-C, et al. Renoprotective effect of renin-angiotensin-aldosterone system blockade in patients with predialysis advanced chronic kidney disease, hypertension, and anemia. *JAMA Intern Med*. 2014;174(3):347-354. doi:10.1001/jamainternmed.2013.12700
25. Onder G, van der Cammen TJM, Petrovic M, Somers A, Rajkumar C. Strategies to reduce the risk of iatrogenic illness in complex older adults. *Age Ageing*. 2013;42(3):284-291. doi:10.1093/ageing/aft038
26. Stengel B, Combe C, Jacquelinet C, et al. The French Chronic Kidney Disease-Renal Epidemiology and Information Network (CKD-REIN) cohort study. *Nephrol Dial*

- Transplant Off Publ Eur Dial Transpl Assoc - Eur Ren Assoc.* 2014;29(8):1500-1507.
doi:10.1093/ndt/gft388
27. Musini VM, Lawrence KA, Fortin PM, Bassett K, Wright JM. Blood pressure lowering efficacy of renin inhibitors for primary hypertension. *Cochrane Database Syst Rev.* 2017;4:CD007066. doi:10.1002/14651858.CD007066.pub3
 28. Sumida K, Nadkarni GN, Grams ME, et al. Conversion of Urine Protein-Creatinine Ratio or Urine Dipstick Protein to Urine Albumin-Creatinine Ratio for Use in Chronic Kidney Disease Screening and Prognosis: An Individual Participant-Based Meta-analysis. *Ann Intern Med.* 2020;173(6):426-435. doi:10.7326/M20-0529
 29. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *J Chronic Dis.* 1987;40(5):373-383. doi:10.1016/0021-9681(87)90171-8
 30. Katz S, Ford AB, Moskowitz RW, Jackson BA, Jaffe MW. Studies of Illness in the Aged: The Index of ADL: A Standardized Measure of Biological and Psychosocial Function. *JAMA.* 1963;185(12):914-919. doi:10.1001/jama.1963.03060120024016
 31. Lawton MP, Brody EM. Assessment of older people: self-maintaining and instrumental activities of daily living. *The Gerontologist.* 1969;9(3):179-186.
 32. Questionnaire d'évaluation de l'observance - www.ameli.fr. Accessed November 25, 2019. https://www.ameli.fr/sites/default/files/Documents/5396/document/questionnaire-evaluation-observance_assurance-maladie.pdf

33. Laville SM, Gras-Champel V, Moragny J, et al. Adverse Drug Reactions in Patients with CKD. *Clin J Am Soc Nephrol CJASN*. 2020;15(8):1090-1102. doi:10.2215/CJN.01030120
34. Kellum JA, Lameire N. Diagnosis, evaluation, and management of acute kidney injury: a KDIGO summary (Part 1). *Crit Care*. 2013;17(1):204. doi:10.1186/cc11454
35. Fu EL, Groenwold RHH, Zoccali C, Jager KJ, van Diepen M, Dekker FW. Merits and caveats of propensity scores to adjust for confounding. *Nephrol Dial Transplant Off Publ Eur Dial Transpl Assoc - Eur Ren Assoc*. 2019;34(10):1629-1635. doi:10.1093/ndt/gfy283
36. Austin PC, Stuart EA. Moving towards best practice when using inverse probability of treatment weighting (IPTW) using the propensity score to estimate causal treatment effects in observational studies. *Stat Med*. 2015;34(28):3661-3679. doi:10.1002/sim.6607
37. R Core Team (2013). R: A language and environment for statistical computing. *R Foundation for Statistical Computing, Vienna, Austria*. URL <http://www.R-project.org/>
38. Esnault VLM, Brown EA, Apetrei E, et al. The effects of amlodipine and enalapril on renal function in adults with hypertension and nondiabetic nephropathies: a 3-year, randomized, multicenter, double-blind, placebo-controlled study. *Clin Ther*. 2008;30(3):482-498. doi:10.1016/j.clinthera.2008.03.006
39. Rahman M, Pressel S, Davis BR, et al. Renal outcomes in high-risk hypertensive patients treated with an angiotensin-converting enzyme inhibitor or a calcium channel blocker vs a diuretic: a report from the Antihypertensive and Lipid-Lowering Treatment

- to Prevent Heart Attack Trial (ALLHAT). *Arch Intern Med.* 2005;165(8):936-946.
doi:10.1001/archinte.165.8.936
40. O'Hare AM, Choi AI, Bertenthal D, et al. Age affects outcomes in chronic kidney disease. *J Am Soc Nephrol JASN.* 2007;18(10):2758-2765.
doi:10.1681/ASN.2007040422
41. Kovesdy CP. Management of hyperkalaemia in chronic kidney disease. *Nat Rev Nephrol.* 2014;10(11):653-662. doi:10.1038/nrneph.2014.168
42. Ruiz-Hurtado G, Sarafidis P, Fernández-Alfonso MS, Waeber B, Ruilope LM. Global cardiovascular protection in chronic kidney disease. *Nat Rev Cardiol.* 2016;13(10):603-608. doi:10.1038/nrcardio.2016.48

Figure legend

Figure 1 – Flow chart of the study population. eGFR, estimated glomerular filtration rate according to the CKD EPI equation; RASi, renin-angiotensin system inhibitors.

Tables

Table 1 – Baseline characteristics of the study population

Variable	RASi prescription*	No RASi prescription*	P [†]	Standardized differences after weighting*	Age <60 years	Age 60-69 years	Age 70-79 years	Age ≥80 years
Sample size (%)	2178 (78.9)	584 (21.1)			640 (23.2)	818 (29.6)	887 (32.1)	417 (15.1)
ACE users	999 (45.9)	0 (0.0)			295 (46.1)	308 (37.7)	281 (31.7)	115 (27.6)
ARB users	1310 (60.1)	0 (0.0)			294 (45.9)	397 (48.5)	435 (49.0)	184 (44.1)
ACE or ARB users	2178 (100.0)	0 (0.0)			537 (83.9)	664 (81.2)	686 (77.3)	291 (69.8)
Sociodemographic characteristics and functional status								
Age (years)	66.3 ± 12.7	69.4 ± 12.9	<.001	0.7	48.3 ± 9.1	65.3 ± 2.7	74.4 ± 2.9	83.1 ± 2.8
Men	1474 (67.7)	343 (58.7)	<.001	0.7	373 (58.3)	552 (67.5)	623 (70.2)	269 (64.5)
Sub-Saharan African origin	64 (2.9)	10 (1.7)	<.001	0.7	39 (6.1)	20 (2.4)	13 (1.5)	2 (0.5)
Dependence for ≥1 ADL	175 (8.0)	82 (14.0)	.01	2.8	38 (6.0)	80 (9.8)	88 (9.9)	50 (12.1)
Dependence for ≥1 IADL	522 (24.0)	206 (35.2)	<.001	2.5	105 (16.4)	173 (21.1)	278 (31.4)	171 (41.1)
CKD characteristics								
Baseline eGFR (ml/min/1.73m ²)	34.0 ± 11.6	31.9 ± 11.3		3.6	34.9 ± 12.6	34.9 ± 12.0	32.9 ± 11.0	30.1 ± 9.5
Albuminuria								
A1	600 (27.6)	163 (28.0)	.21	0.4	124 (19.4)	222 (27.1)	288 (32.4)	129 (31.0)
A2	662 (30.4)	199 (34.0)		0.2	166 (25.8)	236 (28.9)	297 (33.5)	162 (38.8)
A3	916 (42.0)	222 (38.0)		0.6	350 (54.7)	360 (44.0)	302 (34.0)	126 (30.2)
Value (mg/g of urine creatinine)	464 ± 1011	384 ± 771	0.08		594 ± 1066	504 ± 1215	353 ± 692	310 ± 669
Causal nephropathy								
Diabetes	493 (22.7)	106 (18.2)	<.001	0.3	91 (14.2)	205 (25.1)	225 (25.4)	78 (18.8)
Glomerulopathy	432 (19.8)	81 (13.8)		0.6	202 (31.6)	155 (19.0)	120 (13.5)	36 (8.5)
Nephroangiosclerosis	504 (23.1)	138 (23.6)		1.1	63 (9.8)	161 (19.6)	251 (28.3)	167 (40.1)
Renal artery stenosis	165 (7.6)	57 (9.7)		0.5	19 (2.9)	54 (6.6)	83 (9.3)	66 (15.9)
Tubulointerstitial nephritis	231 (10.6)	104 (17.8)		0.1	105 (16.4)	102 (12.4)	98 (11.1)	31 (7.4)
Polycystic kidney disease	137 (6.3)	24 (4.1)		0.0	84 (13.1)	52 (6.4)	18 (2.1)	6 (1.4)
Other or unknown	216 (9.9)	74 (12.7)		1.3	77 (12.0)	88 (10.8)	92 (10.4)	33 (8.0)
Duration of kidney disease (years)	8.4 ± 8.9	6.5 ± 7.6	<.001	2.7	10.0 ± 9.4	7.5 ± 8.7	7.5 ± 8.1	6.8 ± 8.0
History of acute kidney injury	432 (19.8)	218 (37.3)	<.001	3.2	156 (24.4)	174 (21.2)	227 (25.6)	93 (22.4)
Serum kalemia (mmol/l)	4.6 ± 0.5	4.4 ± 0.5	<.001		4.6 ± 0.5	4.5 ± 0.5	4.5 ± 0.5	4.6 ± 0.5
Cardiovascular diseases and risk factors								
Hypertension	1966 (90.3)	505 (86.5)	.01	1.8	527 (82.3)	742 (90.7)	815 (91.9)	388 (93.0)
Systolic blood pressure (mmHg)	142 ± 20	144 ± 20	.08		136 ± 19	143 ± 20	145 ± 21	147 ± 20
Diastolic blood pressure (mmHg)	78 ± 12	78 ± 12	.73		81 ± 12	79 ± 12	76 ± 12	76 ± 12
Diabetes	993 (45.6)	240 (41.1)	.05	0.7	167 (26.1)	399 (48.8)	476 (53.7)	190 (45.6)
Body-mass index (kg/m ²)	29.0 ± 5.8	28.2 ± 5.9	.003	1.2	27.4 ± 6.0	29.8 ± 6.4	29.4 ± 5.3	28.0 ± 4.9
Nonsmoker	872 (40.0)	252 (43.1)	.07	2.4	290 (45.3)	281 (34.3)	348 (39.3)	204 (49.0)
Former smoker	1029 (47.3)	278 (47.5)		1.5	206 (32.2)	424 (51.9)	478 (53.9)	198 (47.5)
Current smoker	277 (12.7)	55 (9.4)		0.9	144 (22.5)	113 (13.8)	60 (6.8)	15 (3.5)
Heart failure	252 (11.6)	121 (20.6)	<.001	3.4	46 (7.2)	100 (12.2)	138 (15.6)	88 (21.1)
Coronary artery disease	544 (25.0)	162 (27.7)	.19	1.4	57 (8.9)	205 (25.0)	301 (33.9)	143 (34.3)
Atrial fibrillation	217 (10.0)	107 (18.4)	<.001	1.2	15 (2.4)	76 (9.3)	135 (15.2)	98 (23.5)
Peripheral arteritis	283 (13.0)	87 (14.9)	.23	1.6	35 (5.4)	116 (14.1)	153 (17.3)	66 (15.8)
Ischemic stroke or transient ischemic attack	203 (9.3)	75 (12.8)	.01	0.3	30 (4.6)	73 (9.0)	117 (13.2)	58 (14.0)
Cardiology consultation in the past year	1494 (68.6)	434 (74.3)	.01	3.8	302 (47.2)	574 (70.2)	707 (79.8)	344 (82.5)
Other medical characteristics								
COPD	191 (8.8)	69 (11.8)	.03	1.7	30 (4.7)	88 (10.8)	97 (11.0)	44 (10.6)
History of cancer	436 (20.0)	139 (23.8)	.05	0.6	60 (9.3)	175 (21.3)	224 (25.2)	117 (28.0)
Serum hemoglobin (g/dl)	12.9 ± 2.1	12.9 ± 2.0	.85	3.6	12.9 ± 2.2	13.0 ± 2.3	12.9 ± 2.0	12.7 ± 1.4

Serum albumin (g/l)	40.2 ± 4.2	40.1 ± 4.4	.90	0.8	40.5 ± 4.8	40.2 ± 4.3	40.0 ± 3.9	39.8 ± 4.0
Charlson Comorbidity Index	6.6 ± 2.7	7.3 ± 2.7	<.001		4.0 ± 2.1	6.6 ± 2.2	7.9 ± 2.0	8.9 ± 2.1
Drug prescriptions								
Antiplatelet agents	937 (43.0)	232 (39.7)	.17	1.5	121 (18.9)	353 (43.2)	473 (53.3)	222 (53.2)
Oral anticoagulant	286 (13.1)	126 (21.6)	<.001	1.0	36 (5.6)	99 (12.1)	171 (19.3)	106 (25.4)
Beta blockers	891 (40.9)	282 (48.3)	.002	3.8	210 (32.8)	369 (45.1)	398 (44.9)	196 (47.0)
Statins or ezetimibe	1461 (67.1)	323 (55.3)	<.001	3.3	307 (48.0)	576 (70.4)	628 (70.8)	273 (65.5)
Calcium channel blockers	1061 (48.7)	276 (47.3)	.56	2.4	244 (38.1)	420 (51.3)	451 (50.8)	222 (53.2)
Thiazide diuretics	509 (23.4)	25 (4.3)	<.001	6.0	130 (20.3)	176 (21.5)	176 (19.8)	52 (12.5)
Loop diuretics	769 (35.3)	254 (43.4)	<.001	3.2	123 (19.2)	289 (35.3)	381 (43.0)	230 (55.2)
Potassium-sparing diuretics	95 (4.4)	40 (6.8)	.02	0.6	33 (5.2)	38 (4.6)	48 (5.4)	16 (3.8)
Erythropoietin-stimulating agent	133 (6.1)	58 (9.9)	.002	1.0	43 (6.7)	51 (6.2)	59 (6.7)	38 (9.1)
Total number of drugs used	8.2 ± 3.8	7.8 ± 4.1	.01	0.3	6.4 ± 3.8	8.3 ± 4.0	8.9 ± 3.7	9.0 ± 3.4
Good drug adherence	796 (36.6)	246 (42.1)	.03	1.0	215 (33.5)	294 (35.9)	352 (39.6)	182 (43.6)
Moderate drug adherence	1218 (55.9)	295 (50.5)		1.7	339 (52.9)	478 (58.4)	484 (54.6)	213 (51.1)
Poor drug adherence	163 (7.5)	43 (7.4)		0.7	87 (13.6)	47 (5.7)	51 (5.7)	22 (5.3)

* expressed as sample sizes (percentages) for categorical variables and means ± standard

deviations for continuous variables; † comparison of patients who were prescribed renin-angiotensin system inhibitors (RASi) and those who were not, by logistic regression models for categorical variables and linear regression models for continuous variables; ‡ standardized differences (percentages) between patients who were prescribed RASi and those who were not after weighting with a propensity score for the variables included in the analysis of the primary outcome (mean standardized differences from 20 imputations); ACE, angiotensin-converting enzyme inhibitors; ARB, angiotensin ii receptor blockers; ADL, activities of daily living; IADL, instrumental activities of daily living; eGFR, estimated glomerular filtration rate, with the CKD EPI equation; albuminuria expressed as the KDIGO grade (A1: normal; A2: moderately increased; A3: severely increased); LDL, low density lipoprotein; HDL, high density lipoprotein; COPD, chronic obstructive pulmonary disease; MMSE, mini-mental state examination.

Table 2 – Hazard ratios (HRs) for primary and secondary outcomes assessing effectiveness and tolerance associated with renin-angiotensin system inhibitors (RASi) prescription in patients with CKD and impact of age on these associations. Propensity score analyses.

	Cox models with propensity score	Age-RASi interaction	Fine and Gray models with propensity score	Age-RASi interaction
	HR (95% CI)*	P	sHR (95% CI)*	P
<i>Primary outcome</i>				
Death or kidney failure	0.79 (0.66-0.95)	.72		
<i>Secondary outcomes</i>				
Death before kidney failure	0.84 (0.65 to 1.08)	.92	0.82 (0.63 to 1.06)	.71
Kidney failure	0.80 (0.61 to 1.04)	.25	0.82 (0.63 to 1.06)	.31
Major cardiovascular events	1.06 (0.79-1.43)	.29	1.08 (0.79-1.48)	.30
AKI or hyperkalemia	0.75 (0.55 to 1.02)	.28	0.80 (0.59 to 1.10)	.37

* For prescription of RASi; hazard ratios (95% confidence intervals) (HR [95%CI]) were computed by using models weighted for propensity scores according to the Inverse Probability of Treatment Weighting method and bootstrapping; major cardiovascular events included cardiovascular death, stroke, myocardial infarction, and hospitalization for heart failure; sHR, subdistribution HR; AKI, acute kidney injury; CKD, chronic kidney disease.

Table 3 – Hazard ratios (HRs) for primary and secondary outcomes assessing effectiveness and tolerance associated with renin-angiotensin system inhibitors (RASi) prescription in patients with CKD according to age groups.

	Age category	Event in the RASi prescription group	Event in the RASi nonprescription group	HR (95% CI)*
Death or kidney failure	<60 years	182/537 (33.9)	37/103 (35.9)	0.91 (0.50-1.66)
	60-69 years	172/664 (25.9)	59/154 (38.3)	0.66 (0.45-0.97)
	70-79 years	214/686 (31.2)	76/201 (37.8)	0.88 (0.64-1.21)
	≥80 years	117/291 (40.2)	59/126 (46.8)	0.83 (0.56-1.24)
Death before kidney failure	<60 years	11/537 (2.0)	8/103 (7.8)	0.27 (0.06-1.22)
	60-69 years	65/664 (9.8)	15/154 (9.7)	0.81 (0.34-1.91)
	70-79 years	107/686 (15.6)	46/201 (22.9)	0.78 (0.50-1.19)
	≥80 years	78/291 (26.8)	47/126 (37.3)	0.81 (0.51-1.27)
Kidney failure	<60 years	171/537 (31.8)	26/103 (28.2)	1.27 (0.72-2.25)
	60-69 years	107/664 (16.1)	44/154 (28.6)	0.58 (0.37-0.90)
	70-79 years	107/686 (15.6)	30/201 (14.9)	1.04 (0.65-1.66)
	≥80 years	39/291 (13.4)	12/126 (9.5)	1.24 (0.55-2.81)
Major cardiovascular events	<60 years	22/537 (4.1)	6/103 (5.8)	0.96 (0.10-9.37)
	60-69 years	70/664 (10.5)	13/154 (8.4)	1.56 (0.72-3.37)
	70-79 years	86/686 (12.5)	35/201 (17.4)	0.95 (0.60-1.49)
	≥80 years	45/291 (15.5)	29/126 (23.0)	0.72 (0.40-1.29)
AKI or hyperkalemia	<60 years	52/537 (9.7)	11/103 (10.7)	0.86 (0.36-2.10)
	60-69 years	90/664 (13.6)	23/154 (14.9)	0.73 (0.39-1.35)
	70-79 years	103/686 (15.0)	41/201 (20.4)	0.81 (0.50-1.31)
	≥80 years	38/291 (13.1)	22/126 (17.5)	0.81 (0.42-1.53)

* For prescription of RASi; hazard ratios (95% confidence intervals) (HR [95%CI]) were computed by using models weighted for propensity scores according to the Inverse Probability of Treatment Weighting method and bootstrapping; major cardiovascular events included cardiovascular death, stroke, myocardial infarction, and hospitalization for heart failure; AKI, acute kidney injury; CKD, chronic kidney disease.

Table 4 – Hazard ratios (HRs) for primary and secondary outcomes assessing effectiveness and tolerance associated with renin-angiotensin system inhibitors (RASi) prescription in patients with CKD according to estimated glomerular filtration (eGFR) groups.

	eGFR category	Event in the RASi prescription group	Event in the RASi nonprescription group	HR (95% CI)*	P for age-RASi interaction
Death or kidney failure	≥30 ml/min/1.73m²	217/1264 (17.2)	60/300 (20.0)	0.77 (0.53-1.13)	0.59
	<30 ml/min/1.73m²	468/914 (51.2)	171/284 (60.2)	0.79 (0.64-0.98)	0.67
Death before kidney failure	≥30 ml/min/1.73m²	124/1264 (9.8)	44/300 (14.7)	0.84 (0.53-1.31)	0.54
	<30 ml/min/1.73m²	137/914 (14.9)	72/284 (25.4)	0.71 (0.50-1.01)	0.52
Kidney failure	≥30 ml/min/1.73m²	93/1264 (7.4)	16/300 (5.3)	0.68 (0.33-1.41)	0.07
	<30 ml/min/1.73m²	331/914 (36.2)	99/284 (34.9)	0.92 (0.70-1.22)	0.56
Major cardiovascular events	≥30 ml/min/1.73m²	102/1264 (8.1)	26/300 (6.7)	1.19 (0.70-2.01)	0.21
	<30 ml/min/1.73m²	121/914 (13.2)	57/284 (20.1)	1.00 (0.67-1.48)	0.66
AKI or hyperkalemia	≥30 ml/min/1.73m²	124/1264 (9.8)	35/300 (11.7)	0.68 (0.41-1.13)	0.89
	<30 ml/min/1.73m²	159/914 (17.4)	62/284 (21.8)	0.89 (0.62-1.28)	0.38

* For prescription of RASi; hazard ratios (95% confidence intervals) (HR [95%CI]) were computed by using models weighted for propensity scores according to the Inverse Probability of Treatment Weighting method and bootstrapping; major cardiovascular events included cardiovascular death, stroke, myocardial infarction, and hospitalization for heart failure; AKI, acute kidney injury; CKD, chronic kidney disease; eGFR, estimated glomerular filtration rate, with the CKD EPI equation.

Figure 1

