Effect of Renin-Angiotensin-Aldosterone System Blockade Therapy on Incidence of Contrast-induced Nephropathy in Patients with Chronic Kidney Disease

Christin Spatz, Lawand Saadulla, Apurva Lapsiwala, Amin Parhizgar, Nasrollah Ghahramani

Introduction. The incidence of contrast-induced nephropathy (CIN) ranges between 10% and 50% among high-risk patients. Whether medications that affect rennin-angiotensin-aldosterone system (RAAS) have any impact on the development of CIN remains uncertain.

Materials and Methods. We performed a retrospective study of patients with CKD stages 3 and 4 who were either on or off RAAS blockade therapy at the time of coronary angiography. Development of CIN was defined by a 25% increase of serum creatinine from baseline or an increase in serum creatinine by 0.5 mg/dL from baseline. Serum creatinine values were recorded before contrast exposure and for 5 days after coronary angiography.

Results. A total of 178 patients with CKD who had coronary angiography during the study period were included, of whom 62 (35%) were on ACE inhibitors, 12 (7%) were on ARBs, and 1 (1%) was on combination of ACE inhibitors and ARBs. The estimated glomerular filtration rate was 44.0 ± 11.5 mL/min. The odds ratio of acute kidney failure on day 5 was 0.73 (95% confidence interval, 0.31 to 1.69) for the ACE inhibitors and 0.46 (95% confidence interval, 0.06 to 3.70) for ARBs. Multivariable analysis revealed the findings to be independent of demographic variables, comorbidities, type of contrast medium, and the prophylactic strategies.

Conclusions. Patients on RAAS blockade therapy before contrast exposure did not have an increased incidence of CIN. There was also no increased incidence of CIN with ACE inhibitors or ARBs in the subgroups at higher risk, such as those with diabetes mellitus.

INTRODUCTION

Contrast-induced nephropathy (CIN) is considered the 3rd leading cause of iatrogenic acute kidney injury in high-risk patients undergoing radiographic procedures. It is associated with substantial in-hospital morbidity and mortality and usually develops within 48 hours of contrast administration. Contrast-induced nephropathy is diagnosed by an elevation of serum creatinine by a proportionate 25% to 50% rise from the baseline and in the absence of other explanations. Other criteria used to define CIN are based on creatinine clearance, glomerular filtration rate (GFR), and a fixed rise in serum creatinine of approximately 0.5 mg/dL to 1 mg/dL above baseline. The precise mechanism by which contrast agents cause kidney injury is unclear; however, several mechanisms have been proposed, including direct renal toxicity, hemodynamic changes, and a potential role for the renin-angiotensin-aldosterone system.
failure is not fully understood. The postulated pathogeneses of CIN include direct tubular injury, tubular obstruction and dysfunction and renal vasomotor constriction, resulting in medullary hypoxemia. The reported incidence of CIN has varied widely in the medical literature. Numerous studies have reported an estimated incidence of CIN, ranging from as low as 2% to as high as 50%. This variation is generally based on the presence or absence of risk factors for development of CIN. Some identified risk factors include preexisting chronic kidney disease (CKD), diabetic nephropathy, true or relative hypovolemia, percutaneous coronary interventions, concomitant nephrotoxic exposure, and osmolality, volume and toxicity of radiocontrast media.

A study by Pakfetrat and colleagues using multiple logistic regression analysis showed diabetes mellitus to be the strongest predictor for at risk of contrast-related acute kidney injury, followed by hypercholesterolemia, and an estimated glomerular filtration rate lower than 90 mL/min/1.73 m².

Angiotensin-converting enzyme (ACE) inhibitors and angiotensin receptor blockers (ARBs) have been associated with a decline in GFR, primarily mediated through the reduced effects of angiotensin II-induced increase in resistance at the efferent (postglomerular) arteriole. By blocking the response of angiotensin II, the efferent arteriole relaxes, which lowers intraglomerular pressure and reduces the GFR. This reduction in GFR can be severe in certain patients such as those with bilateral renal artery stenosis, heart failure, diabetes, vascular disease or CKD. A significant drop in GFR can also occur in those who already have reduced intrarenal perfusion pressure such as acute volume loss. This retrospective study aims to investigate the possible impact of use of the rennin-angiotensin-aldosterone system (RAAS) medications (ACE inhibitors and ARBs) on the incidence of CIN in patients with mild-to-moderate CKD who received coronary angiography.

MATERIALS AND METHODS

Study Protocol

The protocol was reviewed and approved by the Institutional Review Board. For this retrospective cross-sectional study, data was obtained by medical chart review. Inclusion criteria were as follows: age greater than 18, contrast medium exposure during a coronary angiography, CKD stages 3 or 4 (estimated GFR between 15 mL/min to 59 mL/min) as determined by the Modification of Diet in Renal Disease (MDRD) equation, a baseline creatinine on the day of contrast exposure and for 5 days following contrast administration. Data from patients on any form of renal replacement therapy were excluded from the study. Information regarding diabetes mellitus, congestive heart failure, peripheral vascular disease, prophylactic medications received, dose and type of intravenous contrast medium, acute myocardial infarction, unstable angina, hypotension (systolic blood pressure < 80 mm Hg), and body mass index was gathered. Serum creatinine values were recorded before contrast exposure and for 5 days after coronary angiography. All creatinine values were measured at one central laboratory by dry slide rate enzymatic methodology. The primary endpoint was development of CIN, which was defined by 25% increase of serum creatinine from baseline or an increase in serum creatinine by 0.5 mg/dL from baseline.

Statistical Analysis

For the calculation of the sample size, we assumed an incidence of CIN of 20% in the control group and 40% in the group on RAAS blockade. The inclusion of 39 patients in the RAAS group allowed for a two-sided significance level of 5% and 80% power, assuming a normal distribution. The odds ratio of acute kidney failure (AKF) and associated 95% confidence intervals (CIs) were calculated for the ACE inhibitor and ARB groups. Multivariable analysis was performed to determine any significant covariates that could affect the incidence of CIN in these groups. The variable included in the multivariable model consisted of age, ethnicity, comorbidities, type and volume of contrast medium, and prophylactic strategies.

RESULTS

Between January 1, 2007 and December 31, 2008, a total of 398 patients with CKD Stage 3 or 4 underwent coronary angiography. Of these, 220 were excluded due to lack of sufficient follow-up data, resulting in inclusion of 178, of whom 35% were on ACE inhibitors and 7% were on ARBs. Review of the electronic medical records confirmed
the diagnosis of CIN by the definition noted above. Characteristics of the patients at baseline are shown in Table 1. The mean age was $71.3 \pm 10.7$ years and the mean estimated GFR was $44 \pm 11.5$ mL/min. There were no significant differences in the baseline characteristics of groups on RAAS blockade agents and the control group (Table 2).

On day 2, 13% of patients had experienced AKF. The peak incidence of acute kidney failure was on day 3 at 19%. Figure 1 illustrates the percentage of patients with AKF relative to days after contrast exposure. The odds ratio of ARF on day 5 was 0.73 (95% CI, 0.31 to 1.69) for ACE inhibitors (Figure 2) and 0.46 (95% CI, 0.06 to 3.70) for ARBs (Figure 3). Multivariable analysis revealed the findings to be independent of demographic variables, comorbidities, type of contrast medium, and the prophylactic strategies utilized.

**DISCUSSION**

We performed a retrospective study of patients with CKD Stage 3 or 4 who were on or off RAAS blockade at the time of coronary angiography. A total of 178 patients were included in the study, of whom 35% were on ACE inhibitors and 7% were on ARBs. The results showed a trend towards a

![Figure 1](image1)

**Figure 1.** Percentage of patients with acute kidney failure relative to days after contrast exposure.

![Figure 2](image2)

**Figure 2.** Odds ratios of acute kidney failure in patients on angiotensin-converting enzyme inhibitors (95% confidence interval) relative to days after contrast exposure.
protective effect with RAAS blockade, however, the results did not achieve statistical significance. Multivariable analysis revealed the findings to be independent of demographic variables, comorbidities, type of contrast medium, and the prophylactic strategies utilized.

The incidence of CIN varies widely between 10% to 50% among patients, depending on the presence of risk factors such as CKD, the definition of CIN, the amount and type of the contrast medium administered, and the exact radiologic procedure. Overall, the rate of AKF requiring dialysis from CIN appears to be low. In a retrospective study of 58,000 coronary procedures, only 10 and 49 patients required dialysis within one week and one month after contrast administration, respectively.16

Several risk factors for the development of CIN have been established. Studies have shown that underlying renal insufficiency (serum creatinine greater than 1.5 mg/dL or estimated GFR less than 60 mL/min/1.73 m² by the MDRD), diabetic nephropathy with renal insufficiency, heart failure, hypovolemia or states of reduced perfusion, percutaneous coronary intervention, and high total doses of the contrast medium all predispose the patient to development of CIN.2,17,18 A few scoring systems including well-established risk factors for CIN have been developed to estimate the individual risk of CIN and dialysis, the most familiar of which has been developed by Mehran and coworkers.7

Whether certain medications, specifically RAAS blockade agents, are risk factors for development of CIN remains controversial. A randomized prospective study was published by Gupta and colleagues over a decade ago, in which patients undergoing coronary angiography were randomized to receive captopril or placebo. Renal scintigraphy scans were used to measure GFR and it was concluded that patients who had received captopril had a mean increase in the GFR of 13 mL/min compared to placebo.19 Other investigators have reported conflicting findings. In a recent study, Kiski and colleagues performed a retrospective analysis of patients who participated in a single-center prospective study to compare different prophylactic strategies for CIN prevention. The impact of RAAS blockade on the frequency of CIN was assessed. Patients treated with RAAS blockade before exposure to contrast media developed significantly more CIN episodes within 72 hours (11.9% versus 4.2%, \( P = .006 \)). Using multivariable analysis, RAAS blockade was found to be an independent predictor of CIN (odds ratio, 3.082; 95% CI, 1.23 to 7.70, \( P = .02 \)).20

The limitations of this study include the single-center nature and the retrospective design. In addition, baseline creatinine was noted on the day of contrast administration; however, not all patients had serum creatinine values documented months prior to determine true baseline. Lastly, only patients who had remained in the hospital and had creatinine values 5 days after contrast exposure were included in this analysis. Thus, the results may not be generalizable to stable outpatients undergoing coronary angiography.

CONCLUSIONS
This retrospective study demonstrates that patients on ACE inhibitor or ARB while undergoing cardiac catheterization are not at a higher risk of developing CIN. Prospective randomized trials are needed to help determine the effect of RAAS blockade on CIN.

CONFLICT OF INTEREST
Dr Nasrollah Ghahramani is supported by Award Number K23DK084300 from the National Institute of Diabetes and Digestive and Kidney Diseases. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institute of Diabetes and Digestive and Kidney Diseases or the National Institutes of Health.
REFERENCES


Correspondence to:
Nasrollah Ghahramani, MD, MS, FACP, FASN
Division of Nephrology, Mail Code H040, Pennsylvania State University College of Medicine, Hershey, PA 17033, USA
Tel: +1 717 531 8156
Fax: +1 717 531 6776
E-mail: nghahramani@yahoo.com

Received January 2012
Revised June 2012
Accepted July 2012