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Acute Kidney Injury Severity and Long-term Readmission and Mortality Following Cardiac Surgery

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Abstract

Background—Acute kidney injury (AKI) is a common complication following cardiac surgery. While AKI severity is associated with increased risk of short-term outcomes, its long-term impact is less well understood.

Methods—Adult patients undergoing isolated coronary artery bypass graft surgery at eight centers were enrolled into the Northern New England biomarker registry (n=1,610). Patients were excluded if they had renal failure (n=15) or died during index admission (n=38). AKI severity was

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defined using the AKI Network (AKIN). We linked our cohort to national Medicare and state allpayer claims to ascertain readmissions and to the National Death Index to ascertain survival. Kaplan-Meier and multivariate Cox's proportional hazard modeling were conducted for time to readmission and death over 5 years.

Results—Within 5 years, 513 patients (33.8%) developed AKI with AKIN stage 1 (29.9%) and stage 2–3 (3.9%). There were 620 (39.9%) readmissions and 370 (23.8%) deaths. After adjustment, Stage 1 AKI patients had a 31% increased risk of readmission (95%CI: 1.10–1.57), while Stage 2–3 patients had a 98% increased risk (1.41–2.78) compared to those with no AKI. Relative to those without AKI, Stage 1 patients had a 56% increased risk of mortality (1.14–2.13) while those Stage 2–3 patients had a 3.5-times higher risk (2.16–5.60).

Conclusions—AKI severity using the AKIN stage criteria is associated with a significantly increased risk of 5-year readmission and mortality. Our findings suggest efforts to reduce AKI in the perioperative period may have significant long-term impact for patients and payers in reducing mortality and healthcare utilization.

Keywords

Coronary artery bypass graft; kidney; renal function; surgery; readmission; mortality

Readmissions account for significant increase in morbidity, mortality, costs and are potentially preventable.(1–3) Since the Hospital Readmission Reduction Program (HRRP), hospitals are facing the rapid onset of the readmission- and value-based purchasing penalties, resulting in reductions in reimbursement from the Center for Medicaid and Medicare Services (CMS, Section 3025 PPACT).(4, 5) Approximately two-thirds of the United States (US) hospitals have higher than expected readmission rates for patient populations of interest, including cardiac surgery.

Acute kidney injury (AKI) is a common complication occurring after cardiac surgery.(6) AKI has been shown to be associated with short- and long-term mortality after cardiac surgery.(7, 8) More recently, AKI has also been shown to be associated with hospital readmissions following cardiac surgery, hospitalization for heart failure, or acute myocardial infarction.(9–12) Moreover, recent evidence suggests that the severity of AKI might be an easily identifiable marker of the risk of 30-day readmission.(10) However, the long-term impacts of severity of AKI on readmission or mortality are less well understood.

Some studies have reported post-procedure AKI or renal failure as a significant risk factor for 30-day readmissions.(12–14) One study identified AKI severity as an ordinal risk factor for 30-day readmission after cardiac surgery.(10) While 30-day endpoints only provide a small window in evaluating morbidity and mortality, no report has evaluated a severity index for AKI, using the Acute Kidney Injury Network (AKIN)(15) staging criteria as a risk factor, for determining long-term risk of readmission or mortality after cardiac surgery.

Based on the evidence, we hypothesized that the severity of AKI, using the AKIN staging criteria, could serve as a risk factor for long-term readmission and mortality. Therefore, we examined whether post-cardiac surgery AKI stage (no AKI, AKI stage 1, stage 2, and stage 3) is associated with 5-year readmission and/or 5-year mortality.

Patients and Methods

Cohort: NNE Biomarker Study

Patients undergoing coronary artery bypass graft (CABG) surgery and/or valve surgery at any of 8 hospitals participating in the Northern New England Cardiovascular Disease Study Group (NNE) were prospectively enrolled into the NNE Biomarker Study from 2004 to 2007 (n = 1,690). For the present study, the sample was restricted to only those patients for whom CABG was the primary procedure (n=1610). Patients were excluded from the analysis if they had a history of renal failure (n = 15) or died during the index admission (n=38). Patients were also dropped if their recorded date of discharge occurred after their recorded date last known alive (n=1). The Committee for the Protection of Human Subjects at Dartmouth College (IRB) approved this study for both the prospective cohort with patient consent and the linkage of readmission and mortality events.

Primary Outcomes: 5-Year Readmission and Mortality

The cohort was linked to Medicare in-patient claims (completed by ResDAC) and state allpayer in-patient claims using name, gender, social security number, date of birth, and zipcode of residence at the time of surgery. Maine and Vermont completed links internally. We used probabilistic linking for New Hampshire all-payer in-patient claims. We achieved complete ascertainment for Medicare, VT, and ME. Five percent of NH patients were not matched in the NH in-patient claims. Reasons for readmission were determined using the principal diagnosis upon first readmission. Five-year mortality was ascertained by linking to the National Death Index.

Primary Exposure: AKIN Staging Criteria

The last serum creatinine prior to cardiac surgery and highest post-operative serum creatinine prior to discharge were used to classify the stage of AKI. AKI stages were defined by the AKIN definitions as follows: AKIN Stage 1: 0.3 or 50% increase in serum creatinine from baseline; AKIN Stage 2: a 2-fold increase in serum creatinine; and AKIN Stage 3: a 3-fold or 0.5 increase if the baseline serum creatinine was at least 4.0 (mg/dL), or new dialysis-dependent renal failure.(15) Due to the small proportion of patients in AKIN stage 2 and 3, we bundled stage 2 or 3 patients' outcomes in this report.

Statistical Methods

Baseline, operative, and postoperative outcomes were compared using chi-square tests, and continuous data were compared between groups using the student's t-test or Wilcoxon Rank-Sum tests, where appropriate. Kaplan-Meier time-to-event methods and log-rank tests were conducted to compare 5-year readmission (first readmission) and 5-year mortality across AKIN stages. Univariate and multivariate Cox's proportional hazard models were constructed to compare hazard ratios across AKIN stages, adjusting for age, sex, body surface area, and presence of any vascular disease for readmission and the combined readmission and mortality endpoint. Mortality models were adjusted for age and sex. Multivariate cox's proportional hazard models for 5-year readmission and mortality were repeated using multiple imputation methods to account for variables with missing values.

We then conducted time-competing event analyses. All analyses were performed using Stata 13.1 (College Station, TX).

Results

Overall, 513 patients (33.8%) developed AKI with AKIN stage 1 (29.9%) and stage 2–3 (3.9%). There were 620 (39.9%) readmissions and 370 (23.8%) deaths overall, and 620 (39.9%) readmissions and 192 (12.3%) deaths within 5 years. Patient characteristics and comorbidities are summarized in Table 1.

We stratified intra- and post-operative variables by AKIN stage in Table 2. Intra-operative factors differed by AKIN stage for intra-aortic balloon pump (IABP), number of packed red blood cells (pRBC), and lowest hematocrit on bypass. Post-operative factors differed by AKIN stage for all outcomes, except for returning to bypass, returning to the operating room for bleeding, and a new Q-wave myocardial infarction (Table 2).

5-Year Readmission

The overall incidence of readmission was 12.1 per 100 person years, while highest among those with Stage 2–3 AKI (no AKI: 9.8, Stage 1: 16.2, Stages 2–3: 32.4 per 100 person years, p<0.001, Figure 1). After adjustment, patients in AKIN Stage 1 had a 31% increased risk of 5-year readmission (95%CI: 1.10, 1.57), while those in AKIN Stage 2–3 had a 98% increased risk (95%CI: 1.41, 2.78, Table 3). Multiple imputation methods confirmed our results with AKIN stage 1 (HR: 1.32; 95%CI: 1.10, 1.57) and AKIN stage 2–3 (HR: 1.94; 95%CI: 1.39, 2.73). Accounting for death as a time competing event, results were the same for AKIN stage 1 (HR:1.29; 95%CI: 1.08, 1.55) and AKIN stage 2–3 (HR: 1.96; 95%CI: 1.42, 2.71).

The TRIBE consortium, which represents cardiac surgery data from across the United States and Canada, provided unpublished external validation of our findings, where patients with stage 1 or 2 AKI are significantly more likely to be readmitted within one year (Stage 1 adjusted OR=1.55, 95% CI: 1.13, 2.13; Stage 2 adjusted OR=3.02, 95% CI: 1.29, 7.07) compared to patients with no AKI.

5-Year Mortality

The incidence of 5-year mortality was 2.6 per 100 person-years. The rate increased across AKI severity (no AKI: 1.9, AKIN Stage 1: 3.5, AKIN Stages 2–3: 9.4 per 100 person years, p<0.001, Figure 2). After adjustment, relative to those without AKI, patients in AKIN Stage 1 had a 56% increased risk of mortality (95%CI: 1.14, 2.13) while those in Stage 2 or 3 had a 348% higher risk (95%CI: 2.16, 5.60). Multiple imputation confirmed our results with AKIN stage 1 (HR: 1.56; 95%CI: 1.14, 2.12) and AKIN stage 2–3 (HR: 3.48; 95%CI: 2.16, 5.61).

5-Year Readmission or Mortality

The overall incidence of 5-year combined readmission or mortality was 13.1 per 100 personyears. Incidence rates increased across AKI severity (no AKI: 10.7, AKIN Stage 1: 17.6,

AKIN Stages 2–3: 34.0 per 100 person years, p<0.001, Figure 3). After adjustment, patients in AKIN Stage 1 had a 32% increased risk of mortality (95%CI: 1.11, 1.56) relative to those without AKI, while those in Stage 2 or 3 had a 93% higher risk (95%CI: 1.39, 2.69). Multiple imputation methods confirmed our results for AKIN stage 1 (HR: 1.32; 95%CI: 1.12, 1.56) and AKIN stage 2–3 (HR: 1.90; 95%CI: 1.37, 2.64).

Reasons for readmission and death

The reasons for hospital readmission are shown in Figure 4. The primary causes of death are illustrated in Figure 5. The most common causes of readmission were infection and angina or coronary artery disease. The most common cause of death was low output heart failure.

Comment

Using AKIN stage criteria, our study confirms similar findings from past literature that AKI severity is associated with increased hospital readmission and mortality.(16, 17) We discovered AKIN stage 1 and stages 2–3 were associated with a significant 31% to 98% increased hazard of 5-year readmission. In addition, AKIN stage 1 and stages 2–3 were associated with a significant 56% to 3.5-fold increased hazard of 5-year mortality. Our findings suggest that current and novel therapeutic interventions to reduce AKI may have a significant long-term impact for patients and payers in reducing readmission and mortality up to five years.

The effects of creatinine elevation on long-term patient readmission and mortality are significant, even when the creatinine returns to baseline before the patient is discharged from the hospital.(7) We evaluated the association of an AKI severity index (AKIN staging criteria) and risk of patients dying or being readmitted up to five years after cardiac surgery. We found AKIN stage to be a simple risk factor for risk of long-term readmission or mortality. Although we are the first to report on the predictive ability of AKIN stage as a severity index for mortality and readmission out to five years among patients alive at discharge, recent studies have supported the association between AKI and readmission, predominately in patients hospitalized for heart failure. These reports have identified that worsening renal function, among patients hospitalized for heart failure, has been predictive of mortality and 30-day heart failure readmissions.(18, 19) Others have investigated AKI or the RIFLE (Risk, Injury, Failure, Loss of function, End-stage renal disease) criteria and their association with mortality and readmission. Among patients hospitalized for heart failure, those that developed AKI were at a significant increased risk of readmission within 30 days (9.2% versus 6.0%, p=0.0003).(12) In another investigation, the RIFLE criteria was used as a significant independent risk factor for mortality or heart failure readmissions up to one year.(20) However, when heart failure readmissions were abstracted from the one-year events, the RIFLE criteria did not demonstrate a trend in rates of one-year heart failure readmissions. Contrary to this report, our research supports the use of the AKIN staging criteria for prediction of 5-year readmissions or five-year mortality after live discharge from cardiac surgery in a multi-center regional collaborative.

Strengths and Limitations

There are limitations to consider for our investigation of AKIN staging criteria and the association with 5-year readmission and mortality. First, our prospective cohort was developed to enroll patients into a biomarker cohort. While enrollment was consistent at all medical centers, this investigation is limited to patients consented into the NNE biomarker cohort and does not reflect consecutive patients. However, Table 4 demonstrates patients enrolled were comparable to non-enrolled patients in the NNE. External validation from the TRIBE consortium supports generalizability of our findings to other centers.

Second, serum creatinine used to define AKI and AKIN staging criteria were measured daily postoperatively at each medical center, rather than at a single laboratory. Therefore, there may be variability in the measurement of AKI. However, we have no reason to believe there was differential ascertainment of creatinine data between patients with AKI versus those without AKI.

Third, as with any observational trial, the issue of unmeasured confounding is a concern. We did not capture post-hospitalization risk factors for readmission or mortality. However, the NNE registry has some of the most extensive patient and procedural data for cardiac surgery, and we are confident we have thoroughly evaluated patient, procedural, and perfusion risk factors for readmission. In addition, we adjusted for variables used in the age, sex, body surface area, and vascular disease in our models.(21)

Currently, we have a limited understanding on the causal relationship between AKI and long-term readmission or mortality. It is likely that AKIN stage is acting as a severity index for risk of near-term adverse events, such as readmissions and mortality, and acting as a surrogate for the physiologic vulnerability of patients. In our previous research, we identified that AKI duration, after cardiac surgery, impacts 5-year survival.(16) This effect on survival may not be directly related to the kidney injury. AKI may be a surrogate for other undetected organ injuries, an epiphenomenon, thereby being useful as a signal for increased likelihood of readmission and long-term survival. While the renal injury may not be substantial enough to significantly impair toxin excretion and electrolyte balance, AKI, as an epiphenomenon, may be contributing to adverse outcomes by virtue of other affected organ systems.(10, 16) It is unknown whether this notion is valid. However, it is also likely that injuring the kidney during or after cardiac surgery may cause more rapid progression of kidney disease(22) and result in injury to other organs, therefore supporting the need of biomarker evaluation in readmission.(23, 24)

Our study also has notable strengths. Ascertainment of readmission is commonly a problem in epidemiological investigations, often limited to readmission occurring to the same site. Readmission to other facilities is often missing in most analyses.(3) We, however, linked our cohort with both Medicare in-patient claims and state all-payer in-patient claims to provide the most complete ascertainment of readmission possible. We also report on the association of AKI severity and long-term outcomes using a multi-centered regional collaborative. We identified and evaluated the association of AKIN stage as a severity index for 5-year readmission and 5-year mortality following adult cardiac surgery. These efforts provide the cardiac surgical community with a simple tool for pre-discharge screening of long-term risk

of readmission and mortality. Further research is needed in this area to understand the pathophysiology of decompensated renal function and long-term events, such as readmission and mortality occurring well after the index cardiac surgery. In addition, novel approaches to prevent renal injury in the peri- and post-operative setting must continue to be developed and evaluated in cardiac surgery.

Conclusions

We discovered that AKI severity, using the AKIN stage criteria, is associated with a significantly increased risk of 5-year readmission and 5-year mortality following cardiac surgery. Our findings suggest that efforts to reduce AKI may have significant long-term impact for patients and payers on survival and health care utilization.

Acknowledgments

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List of Abbreviations

AKI	Acute Kidney Injury
AKIN	Acute Kidney Injury Network
CABG	Coronary Artery Bypass Grafting
CI	Confidence Interval
CMS	Centers for Medicare & Medicaid Services
HR	Hazard Ratio
HRRP	Hospital Readmission Reduction Program
IABP	Intra-Aortic Balloon Pump
NNE	Northern New England Cardiovascular Disease Study Group
OR	Odds Ratio
pRBC	packed Red Blood Cells
ResDAC	Research Data Assistance Center (at the Centers for Medicare & Medicaid Services)
RIFLE	Risk, Injury, Failure, Loss of function, End-stage renal disease (criteria for classifying AKI)
STS	Society of Thoracic Surgeons

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Figure 1. 5-Year Readmission

We plot time from alive cardiac surgery discharge to the first readmission to any hospital stratified by AKIN stage: No AKI (blue line), AKIN Stage 1 (red line), and AKIN Stages 2–3 (green line). Log-rank test chi2 46.43, p-value <0.001. AKI= acute kidney injury. AKIN= Acute Kidney Injury Network.

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Figure 2. 5-Year Mortality

We plot time from alive cardiac surgery discharge to all-cause death stratified by AKIN stage: No AKI (blue line), AKIN Stage 1 (red line), and AKIN Stages 2–3 (green line). Log-rank test chi2 56.23, p-value <0.001. AKI= acute kidney injury. AKIN= Acute Kidney Injury Network.

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Figure 3. 5-Year Readmission or Mortality

We plot time from alive cardiac surgery discharge to readmission or all-cause death stratified by AKIN stage: No AKI (blue line), AKIN Stage 1 (red line), and AKIN Stages 2–3 (green line). Log-rank test chi2 48.49, pvalue <0.001. AKI= acute kidney injury. AKIN= Acute Kidney Injury Network.

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Figure 4. Reasons for Hospital Readmission

This Pareto chart depicts the reasons for readmission in our cohort. The category "Other" is not included in this chart accounting for 2.5% of readmissions. Blue line shows the cumulative percentage of readmissions, and the black bars indicate the number of readmissions in this cohort.

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Figure 5. Primary Causes of Death

Pie chart illustrates the most frequent causes of death and their percentages of occurrence in the study cohort.

Table 1

Patient Characteristics

Total N = 1,690	Percent/Mean ± SD
AKIN	
No AKIN	66.25%
Stage 1	29.87%
Stage 2	2.30%
Stage 3	1.58%
Age (mean, SD)	65.18 ± 10.06
Female	22.37%
BMI (mean, SD)	29.72 ± 5.50
BSA (mean, SD)	2.04 ± 0.24
Smoker	23.23%
Atrial fibrillation	6.75%
CHF	9.46%
Last pre-op serum creatinine (mean, SD)	1.11 ± 0.90
Diabetes	37.72%
Ejection fraction <40%	10.85%
Hypertension	80.89%
IABP pre-op	4.63%
Prior MI	
No	55.85%
< 24 hours pre-op	1.74%
> 24 hours & < 7 days pre-op	18.57%
>7 days & <365 days pre-op	9.96%
> 365 days pre-op	13.88%
Vascular disease	27.12%
Unstable angina	54.34%
COPD	13.11%
Left main, 50% stenosis	34.19%
Prior CABG	2.08%
Prior PCI	20.50%

Total N = 1,690	Percent/Mean ± SD
Priority	
Emerge	nt 1.93%
Urge	nt 67.42%
Non-urge	nt 30.66%
Received pRBC uni	ts 36.21%
Number of pRBC units given pre-op	
	0 97.81%
	1 0.64%
	2 0.97%
3 or mo	re 0.58%

AKI=acute kidney injury, AKIN=Acute Kidney Injury Network, BMI=body mass index (kg/m²), BSA=body surface area (m²), CABG=coronary artery bypass graft, CHF=congestive heart failure, COPD=chronic obstructive pulmonary disease, IABP=intra-operative balloon pump, MI=myocardial infarction, PCI=percutaneous coronary intervention, pRBC=packed red blood cell

Table 2

Procedural Characteristics by AKI Status

	No AKI	AKIN Stage 1	AKIN Stage 2 or 3	p-value
	Percent/Mean ± SD	Percent/Mean ± SD	Percent/Mean ± SD	
Pump time (minutes; mean, SD)	98.86±28.71	105.60±30.54	115.46±37.16	< 0.001
Intra- or post-operative IABP	1.39%	1.76%	8.47%	< 0.001
Total Number of pRBC units given during admission				
0	71.04%	50.11%	30.51%	< 0.001
1	6.17%	10.15%	13.56%	
2	12.14%	17.66%	15.25%	
3 or more	10.65%	22.08%	40.68%	
Number of pRBC units given intra-op				
0	87.48%	77.53%	62.71%	< 0.001
1	3.98%	7.05%	11.86%	
2	6.56%	10.13%	5.08%	
3 or more	1.99%	5.29%	20.34%	
Low hematocrit <20%	13.28%	21.45%	25.93%	< 0.001
Number of pRBC units given post-op				
0	77.73%	62.69%	44.07%	< 0.001
1	6.06%	9.71%	16.95%	
2	10.83%	16.11%	15.25%	
3 or more	5.37%	11.48%	23.73%	
Renal failure outcome	0.00%	0.00%	13.56%	< 0.001
Atrial fibrillation, new post-op	16.29%	27.53%	52.54%	< 0.001
Low output	3.54%	5.15%	15.79%	< 0.001
Any Mediastinitis/sternal discharge req. re-op	0.30%	1.10%	6.78%	< 0.001
Return to OR for bleeding	1.09%	1.76%	1.69%	0.563
Q wave post-op	0.44%	1.23%	1.85%	0.180
Pneumonia post-op	0.70%	2.20%	13.56%	< 0.001
Return to bypass	2.32%	2.89%	1.69%	0.751
Outcomes				
Dead at 30 days	0.30%	1.54%	3.39%	0.003

	No AKI	AKIN Stage 1	AKIN Stage 2 or 3	p-value
	Percent/Mean ± SD	Percent/Mean ± SD	Percent/Mean ± SD	
Dead at 1 year	1.19%	3.96%	10.17%	<0.001
Dead at 5 years	9.14%	16.08%	37.29%	<0.001
Readmitted at 30 days	9.54%	13.59%	15.52%	0.039
Readmitted at 1 year	24.05%	36.10%	50.00%	<0.001
Readmitted at 5 years	35.79%	49.20%	68.42%	<0.001
Readmitted or dead at 30 days	9.63%	14.54%	16.95%	0.009
Readmitted or dead at 1 year	24.43%	37.22%	50.85%	<0.001
Readmitted or dead at 5 years	37.84%	51.32%	69.49%	<0.001

AKI=acute kidney injury, AKIN=Acute Kidney Injury Network, IABP=intra-operative balloon pump, OR=operating room, pRBC=packed red blood cell

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Table 3

Unadjusted and adjusted hazard ratios for 5-year readmission, 5-year mortality, and 5-year readmission or mortality.

		Unadjusted			Adjusted	
	Hazard Ratio	Confidence Interval	p-value	Hazard Ratio	Confidence Interval	p-value
5-year Readmission ^d						
No AKI	1.00	(1.00-1.00)	1	1.00	(1.00-1.00)	ł
AKIN Stage 1	1.53	(1.29–1.81)	<0.001	1.31	(1.10–1.57)	0.002
AKIN Stage 2 or 3	2.51	(1.80–3.50)	<0.001	1.98	(1.41–2.78)	<0.001
5-year Readmission ^d (with death as a competing risk)						
No AKI	1.00	(1.00-1.00)	ł	1.00	(1.00-1.00)	I
AKIN Stage 1	1.51	(1.27–1.79)	<0.001	1.29	(1.08–1.55)	0.004
AKIN Stage 2 or 3	2.46	(1.79–3.38)	<0.001	1.96	(1.42–2.71)	<0.001
5-year Readmission ^d (with missing observations imputed)						
No AKI	1.00	(1.00-1.00)	:	1.00	(1.00-1.00)	1
AKIN Stage 1	1.53	(1.29–1.82)	<0.001	1.32	(1.10–1.57)	0.002
AKIN Stage 2 or 3	2.49	(1.79–3.47)	<0.001	1.94	(1.39–2.73)	<0.001
5-year Mortality b						
No AKI	1.00	(1.00-1.00)	1	1.00	(1.00-1.00)	I
AKIN Stage 1	1.85	(1.36–2.51)	<0.001	1.56	(1.14–2.13)	0.005
AKIN Stage 2 or 3	4.89	(3.07–7.79)	<0.001	3.48	(2.16–5.60)	<0.001
5-year Mortality ^b (with missing observations imputed) No AKI	1.00	(1.00-1.00)	:	1.00	(1.00-1.00)	

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	Hazard Ratio	Confidence Interval	p-value	Hazard Ratio	Confidence Interval	p-value
AKIN Stage 1	1.82	(1.34–2.48)	<0.001	1.56	(1.14–2.12)	0.005
AKIN Stage 2 or 3	4.79	(3.01–7.62)	<0.001	3.48	(2.16–5.61)	<0.001
5-year Readmission or Mortality ^d No AKI	1.00	(1.00–1.00)	1	1.00	(1.00–1.00)	
AKIN Stage 1	1.53	(1.30–1.80)	<0.001	1.32	(1.11–1.56)	0.001
AKIN Stage 2 or 3	2.46	(1.78–3.39)	<0.001	1.93	(1.39–2.69)	<0.001
5-year Readmission or Mortality ^d (with missing observations imputed)						
No AKI	1.00	(1.00-1.00)	I	1.00	(1.00-1.00)	1
AKIN Stage 1	1.53	(1.30–1.81)	<0.001	1.32	(1.12–1.56)	0.001
AKIN Stage 2 or 3	2.44	(1.77–3.37)	<0.001	1.90	(1.37–2.64)	<0.001
^a Adjusted for age, sex, body surface area,	, and vascul	ar disease diagr	losis.			
b Adjusted for age and sex.						

Table 4

Patient Characteristics by NNE Biomarker Cohort status

	Not in biomarker cohort	In biomarker cohort	
	Percent/Mean ± SD	Percent/Mean ± SD	p-value
Age (mean, SD)	65.37 ± 10.56	65.18 ± 10.06	0.528
Female	24.66%	22.37%	0.053
BMI (mean, SD)	29.68 ± 5.90	29.72 ± 5.50	0.789
BSA (mean, SD)	2.03 ± 0.25	2.04 ± 0.24	0.206
Smoker	24.71%	23.23%	0.214
CHF	11.93%	9.46%	0005
Last pre-op serum creatinine (mean, SD)	1.10 ± 0.88	1.11 ± 0.90	0.678
Diabetes	35.54%	37.72%	0.107
Vascular disease	25.26%	27.12%	0.131
Unstable angina	55.45%	54.34%	0.290
COPD	12.28%	13.11%	0.374
Left main, 50% stenosis	34.94%	34.19%	0.593
Prior CABG	3.56%	2.08%	0.003
Priority			
Emergent	8.70%	1.93%	< 0.001
Urgent	63.96%	67.42%	
Non-urgent	27.34%	30.66%	

BMI=body mass index (kg/m²), BSA=body surface area (m²), CABG=coronary artery bypass graft, CHF=congestive heart failure, COPD=chronic obstructive pulmonary disease, MI=myocardial infarction, pRBC=packed red blood cell