Serum neutrophil gelatinase-associated lipocalin as an early biomarker of acute kidney injury in subjects undergoing percutaneous transluminal coronary angiogram with normal EGFR

Shylaja.T.V1,*, Amit Sonagra2, Asmabi Makandar3, Patnaik4, Shanthinaidu5

1,3Assistant Professor, Dept. of Biochemistry, East Point College of Medical Sciences, Bengaluru, Karnataka, 2Assistant Professor, Dept. of Biochemistry, G.M.E.R.S. Medical College, Patan, Gujarat, 4Professor, Dept. of Cardiology, Nizam’s Institute Of Medical Sciences, Hyderabad, Andhra Pradesh, 5Consultant Biochemist and Head of Laboratory Medicine, Care Hospitals, Hyderabad, Andhra Pradesh, India

*Corresponding Author:
Email: shylajavishal@gmail.com

Abstract
Introduction: Detecting AKI in a timely fashion with the current AKIN staging criteria is a challenge because the diagnosis of AKI is usually based on changes in serum creatinine (SCr) which is a poor marker of early renal dysfunction.

Aim: To study the role of Serum Neutrophil Gelatinase-Associated Lipocalin (NGAL) as early biomarker for the detection of Contrast induced acute kidney injury (CIAKI) in subjects undergoing Percutaneous transluminal coronary angiogram (PTCA) with normal eGFR.

Materials and Methods: Prospective cohort study was conducted, where SCr and serum NGAL were serially measured in a heterogeneous group of subjects (n=60) presenting to cardiology department.

Results: The study population consisted of 60 subjects. All subjects were divided into 2 groups “CIAKI group” and “no-CIAKI group” according to predefined definition. The serum NGAL increased and reached its peak at 4 hours after contrast media (CM) administration and did not returned to baseline by 24 hours while the SCr increased at 24 hours and reached peak at 48 hours respectively (P<0.001). Thus, 4 hours after CM administration were considered to be appropriate time point for NGAL measurement and there was no significant correlation between serum NGAL with SCr at 0 hours, 4 hours, 24 hours and 48 hours.

Conclusion: In our study, we found that serum NGAL promises to be a simple, safe, non-invasive and reliable early biomarker for predicting possible onset of CIAKI following contrast administration.

Keywords: Contrast induced acute kidney injury, Percutaneous Trans luminal Coronary angiogram, Serum Creatinine, Estimated glomerular filtration rate.

Introduction

Acute kidney injury (AKI) Previously referred to as Acute Renal Failure (ARF) is a heterogeneous entity associated with various clinical presentations, treatments, and procedures. The incidence of hospital-acquired AKI varies from 5% in patients with normal preoperative renal function to 25% in intensive care unit (ICU) patients.1,2 In developed countries, AKI is common in the elderly and hospital-acquired causes dominate, where as in developing countries, AKI is the disease of younger subjects and community-acquired cases are common. Acute diarrheal diseases, acute glomerulonephritis, tropical infections (mainly malaria and leptospirosis), environmental agents and snakebite are the common causes of AKI.3 Acute radio contrast nephropathy (ACN) is an important cause of AKI in hospitalized patients undergoing contrast-based procedure.4 SCr has been the predominant marker of renal function in clinical practice for more than half a century and its limitations are well documented. As a marker of renal function rather than injury, the nonlinear relationship between Glomerular Filtration Rate (GFR) and SCr, means GFR may decrease by more than 50% from normal before a significant rise in SCr occurs, making SCr insensitive to small but significant reductions in GFR.5,6 A particularly acute diagnostic problem is the lack of biochemical markers for detecting early kidney injury that are sensitive and easily applicable in clinical practice and may help to predict the development of AKI. NGAL, as a member of the lipocalin super family, is produced from the nephron in response to tubular epithelial damage.7 It has been identified as one of the earliest and potentially one of the most indicative biomarkers of AKI from a diverse array of conditions and can differentiate between prerenal and intrinsic causes.8,9 Downstream proteomic analysis also revealed NGAL to be one of the most highly induced protein in the
kidney after ischemic or nephrotoxic AKI in animal models and studies implicated NGAL as an early diagnostic biomarker for AKI in common clinical situations. Because of the limitations regarding use of SCr for the early detection of AKI, and the importance of early detection, this study is being taken up to know the role of serum NGAL that would allow earlier detection of AKI. And it has been conducted in subjects undergoing contrast related procedure as a cause of AKI, since the baseline values were also available in all these subjects.

Materials and Methods

Study design and patient population: Prospective observational cohort study of 30 cases and 30 controls conducted in the department of Biochemistry, Nizam’s Institute of Medical Sciences, Hyderabad, India.

Study material: Subjects admitted in the department of Cardiology to undergo emergency or elective PTCA.

Inclusion criteria: Subjects aged between 18-70 years of either gender who were undergoing elective and emergency PTCA by using Iohexol contrast, with normal SCr levels and normal eGFR $\geq$60 ml/min/1.73 m$^2$ (MDRD formula) admitted to hospital during the study period and willing to participate in this study after obtaining informed consent and ethical committee approval has been taken.

Exclusion criteria:
1. Subjects with non availability of normal SCr levels
2. Subjects with non availability of normal eGFR $\geq$60 ml/min/ 1.73 m$^2$ (MDRD) at admission
3. Subjects underwent contrast related procedure within 1 week or less from the index procedure

Sample Size calculated: Based on baseline data available on serum NGAL the anticipated difference in mean between two groups is 147 and anticipated standard deviation (SD) is 100, to obtain a power of 99% with type 1 error of 0.001, the required sample size is 21. In this study a total of 30 cases and 30 controls are enrolled.

Sample collection: Venous blood samples were drawn from all participants after 12-hour overnight fast before PTCA, and other samples were taken at 4, 24 and 48 hours after the procedure. Blood samples were centrifuged at 2000 $\times$g for 10 minutes, serum separated SCr, and fasting plasma glucose were estimated in overnight fasting sample. And in after procedure samples taken at 4 hours, 24 hours and 48 hours same centrifugation procedure repeated, only SCr concentrations estimated in all these samples. Remaining serum was stored at $-20^\circ$C.Subjects were observed for primary outcome variable which was the development of CI AKI, defined as an absolute increase in SCr of $>25\%$ or 0.5mg/dl from baseline values occurring within 24 to 48 hours after the coronary procedure. Finally by looking at the primary outcome.

Cases: 30 Subjects (an increase in SCr by $>0.5$ mg/dl or $>25\%$ within 3 days after intravascular administration of CM without an alternative aetiology). Controls: 30 Subjects (who did not showed $>25\%$ increase in the SCr at end of 48 hours).

All the data were analysed after completion of study period.

Methods
1. eGFR is Calculated by applying the MDRD formula,$^{11}$
2. SCr estimated by Jaffe method(IDMS-traceable version)$^{12}$
3. Glucose estimated by HK (hexokinase)method$^{13}$
4. All the parameters were analysed by using fully automated analyzer(Hitachi-912)
5. Serum NGAL levels were measured using a commercially available ELISA kits (BIOPORTO Diagnostics [Gentofte, Denmark]) where serum samples were diluted to 1in 100 dilution before performing an ELISA assay to fit the concentration of respective NGAL protein in the linear range of the standard curve ,the inter and intra assay CV% $<10\%$, the measurement were made in duplicate and in blinded fashion.

Statistical analysis
Continuous variables were expressed as mean $\pm$ SD and tested with Student’s t test and categorical variables were expressed as frequencies and compared with the use of the Pearson chi-square test (Fisher’s exact test was applied if the number of observations per cell was fewer than five). The changing trends of serum NGAL was analysed by taking repeated measures at 0 hours, 4 hours, 24 hours and 48 hours before and after procedure. All the analysis was performed using the SPSS 12.0 for Windows.

Results
Total 60 subjects were included in the present study. All subjects were divided into 2
groups: “CIAKI group” and “no-CIAKI group” according to predefined definition.
The “no-CIAKI group” included a total of 30 controls, 27 males (90%) and 3 females (10%) with average age 60.20 ± 05.73 years.

Table 1: The clinical characteristics of the patient population

<table>
<thead>
<tr>
<th>Variables</th>
<th>No CIAKI (n=30)</th>
<th>CIAKI (n=30)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Demographic Characteristics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean age</td>
<td>57.60±05.4</td>
<td>60.20±05.73</td>
<td>&lt;0.081</td>
</tr>
<tr>
<td>Male n (%)</td>
<td>27(90%)</td>
<td>26(86.7%)</td>
<td>1.0000</td>
</tr>
<tr>
<td>Clinical Characteristics n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diabetes Mellitus (DM)</td>
<td>06(20.0%)</td>
<td>28(93.3%)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>eGFR ml/min/1.73 m2</td>
<td>64.90±03.07</td>
<td>62.80±02.68</td>
<td>0.0049</td>
</tr>
<tr>
<td>Type of contrast media (CM)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n (%)</td>
<td>30(100%)</td>
<td>30(100%)</td>
<td>-</td>
</tr>
<tr>
<td>Dose of contrast (ml)</td>
<td>117.70±33.70</td>
<td>280.00±64.17</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

Continuous values are expressed as mean ±SD; categorical values are expressed as total number and percentage of the global population (in parentheses).

As shown in Table 1, that patients with CIAKI group were found no difference in age than controls 60.20 ± 5.73 vs 57.60 ± 05.4 yrs respectively (P<0.081), DM (P<0.0001), eGFR (P=0.004) compared with patients of “no-CIAKI group” and in subjects who developed CIAKI, a higher dose of CM was administered (P<0.0001), while the compared analysis of the patient groups in relation to the molecule type chosen to make coronaries opaque (Iohexol) did not show any significant statistical difference.

Table 2: Changing trends of novel biomarker NGAL after CM exposure in both the subjects (0, 4, 24 and 48 hours)

<table>
<thead>
<tr>
<th>Time (Hours)</th>
<th>NGAL(ng/ml) no CIAKI</th>
<th>NGAL(ng/ml) CIAKI</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>71.2</td>
<td>71.2</td>
</tr>
<tr>
<td>4</td>
<td>71.3</td>
<td>156.1</td>
</tr>
<tr>
<td>24</td>
<td>69.9</td>
<td>99.4</td>
</tr>
<tr>
<td>48</td>
<td>69.6</td>
<td>71.2</td>
</tr>
</tbody>
</table>

Table 3: Changing trends of Conventional marker (SCr) after CM exposure in both the subjects (0, 4, 24 and 48 hours)

<table>
<thead>
<tr>
<th>Time(Hours)</th>
<th>S Cr (mmol/L) No CIAKI</th>
<th>S Cr (mmol/L) CIAKI</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>94.7</td>
<td>95.3</td>
</tr>
<tr>
<td>4</td>
<td>94.0</td>
<td>92.7</td>
</tr>
<tr>
<td>24</td>
<td>92.7</td>
<td>93.6</td>
</tr>
<tr>
<td>48</td>
<td>93.0</td>
<td>177.0</td>
</tr>
</tbody>
</table>

The changing trends of Serum NGAL and SCr were showed in Table 2, 3 and Fig. 1 & 2 respectively. The serum NGAL increased and reached its peak at 4 hours after CM administration and did not returned to baseline by 24 hours while the SCr increased at 24 hours and reached peak at 48 hours respectively. Thus, 4 hours after CM administration were considered to be appropriate time point for NGAL measurement.
Fig. 1: Shows changing trends of Serum NGAL after CM exposure

Fig. 2: Shows changing trends of SCr after CM exposure

Table 4: Changes in Serum NGAL and SCr before and after PTCA in CIAKI group

<table>
<thead>
<tr>
<th>Variables</th>
<th>Before PTCA</th>
<th>4 hrs after PTCA</th>
<th>24 hrs after PTCA</th>
<th>48 hrs after PTCA</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>NGAL(ng/ml)</td>
<td>71.2±3.14</td>
<td>156.0±24.9 a</td>
<td>99.4±9.7 b</td>
<td>71.1±4.0</td>
<td>&lt;0.0001a,b</td>
</tr>
<tr>
<td>SCr(mmol/l)</td>
<td>94.6±5.7</td>
<td>92.6±4.4</td>
<td>96.6±4.3</td>
<td>177±16.8 c</td>
<td>&lt;0.0001 c</td>
</tr>
</tbody>
</table>

Results expressed as Mean ± SD

a,b=P<0.0001 - NGAL Baseline vs 4 hrs and 24 hrs

c=P<0.0001 - SCr- Baseline vs 48 hrs

Discussion

It is important to detect AKI in an initial stage to start early interventions. DM is one of the strongest predictors of AKI after coronary intervention. In our study, we found that occurrence of CIAKI is significantly higher among subjects with DM compared to other subjects. This is in accordance with other study ToprakOet-al. eGFR found to have significant difference (P<0.004) in CIAKI subjects compared to no CIAKI subjects. Renal function deterioration after exposure to radiographic contrast agents is common in patients with impaired renal function. Volume of contrast showed that the CIAKI risk increases proportionally to the dose of CM like in CIAKI than no CIAKI this is in accordance with other study Kane GC et al.15

The discrepancies in the above findings may be due to contributions from several systemic factors like prolonged vasoconstriction, alterations in nitric oxide metabolism that lead to renal vasoconstriction, and impaired auto
regulation induced by CM predisposing to medullary hypoxia, in combination with direct cytotoxicity to the renal tubular epithelium.\textsuperscript{16}

Numerous studies have demonstrated that the treatment of AKI should be started well before the rise of SCr and immediately after the injury.\textsuperscript{17,18} Sensitive biologic markers of renal tubular injury are needed to detect early AKI because currently AKI diagnosing and staging criteria are entirely based on an increase in SCr or decrease in urine output. SCr is insensitive and increases too slow, and urine output is affected by the use of diuretic or prerenal azotemia. In this study, we demonstrated that, the serum NGAL increased and reached its peak at 4 hours after CM administration and did not returned to baseline by 24 hours, while the SCr increased at 24 hours and reached peak at 48 hours respectively in subjects undergoing PTCA. Bachorzewska-Gajewska H \textit{et al} in their cohort study found a significant rise in serum NGAL after 2 and 4 hrs and a significant rise in urinary NGAL after 4 and 8 hours after coronary angiography, while SCr and creatinine clearance remained unchanged after procedure.\textsuperscript{19} Similar, but not identical, results were observed by Mishra J \textit{et al} who reported a significant rise in serum and urinary NGAL in samples taken after 2 hrs or at the first available sample after cardiopulmonary bypass in children who developed ARF.\textsuperscript{20} Another study by Wagener G \textit{et al} in patients after cardiac surgery demonstrated higher urinary NGAL in patients developing ARF compared to patients without ARF.\textsuperscript{21} Recent studies have provided clues toward understanding the role of NGAL in the kidney. In the post ischemic kidney, NGAL is markedly up regulated in proximal tubules and distal nephron segments. The similar pattern of proximal tubule NGAL expression was observed following nephrotoxic injury after cisplatin administration.\textsuperscript{22} The reason for NGAL up regulation is not clear, but it may represent a defence mechanism to protect against tubular cell death and preserve renal function, because exogenous NGAL protects against murine ischemia reperfusion injury.\textsuperscript{23} This may be due to marked up regulation of NGAL mRNA and protein levels in the early post-ischemic kidney.

Limitation of our study, since we estimated serum NGAL at 4 hours, so the role of early detection of CIAKI (within 2 hours) in preventing and improving clinical outcomes of high risk patients was not discussed in the current study due to the original study design and limited sample size. Further study with large sample size and intervention may be needed to identify the role of early detection of CIAKI in clinical care of patients with high risk of CIAKI.

Conclusion

We showed in our study that serum NGAL promises to be a simple, safe, non-invasive and reliable early biomarker for predicting possible onset of CIAKI following contrast administration overriding the conventional renal failure markers.

References


