Diagnostic accuracy of plasma neutrophil gelatinase-associated lipocalin (NGAL) as an inflammatory biomarker for low-grade inflammation.

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Abstract

Neutrophil gelatinase-associated lipocalin (NGAL) is an acute phase reactant in certain inflammatory conditions. The aim of this study was to investigate whether plasma NGAL is a sensitive biomarker of low-grade inflammation. The severity of inflammation was determined by a scoring system using high-sensitivity C-reactive protein (hsCRP) and corrected erythrocyte sedimentation rate. The sensitivity of NGAL in low-grade inflammation was 27.3%, lower than hsCRP (73.8%, P<0.001). Plasma NGAL concentration was significantly associated with hsCRP, tumor necrosis factor-alpha (TNF-α), and interleukin-6 (IL-6) in high-grade inflammation, but not in low-grade inflammation. In a receiver operating characteristic curve analysis, the diagnostic value of NGAL to identify an increase in IL-6 and TNF-α was similar to that of hsCRP in high-grade inflammation. However, the area under the curve of NGAL was significantly lower than that of hsCRP in low-grade inflammation (0.579 (95% CI, 0.449-0.708) versus 0.691 (95% CI, 0.557-0.826), P<0.001). Plasma NGAL may provide helpful information when monitoring patients with high-grade inflammation but does not seem to accurately reflect the severity of inflammation in low-grade inflammation.

Keywords: Neutrophil gelatinase-associated lipocalin, Tumor necrosis factor-alpha, Interleukin-6, Sensitivity, Inflammatory biomarker, Low-grade inflammation.

Introduction

Inflammation is an adaptive response to infection, noxious stimuli, and cellular injury that initiates elimination of toxic and foreign agents and repair of damaged tissue. Inflammation is an essential component of host defense and immune response. However, unresolved low-grade chronic inflammation is generally viewed as a core perturbation in a range of chronic diseases [1]. Low-grade inflammation is defined as a two- to three-fold increase in circulating levels of acute phase proteins and inflammatory cytokines and other markers of immune system activity, including C-reactive protein (CRP), tumor necrosis factor-alpha (TNF-α), and interleukin-6 (IL-6). Low-grade systemic inflammation is associated with a number of chronic conditions, e.g. metabolic syndrome, atherosclerosis, type 2 diabetes mellitus, and non-alcoholic fatty liver disease [2,3].

Neutrophil gelatinase associated lipocalin (NGAL) is a 25 kDa glycoprotein of the lipocalin family, which was originally identified as a novel protein isolated from secondary granules of activated neutrophils. In contrast to serum creatinine, NGAL is specifically induced in the damaged nephron. Because NGAL is up-regulated shortly after damage in renal tubular cells, it is used for early detection of acute kidney injury (AKI). NGAL is a promising marker of renal epithelial injury; however, it is highly induced in a number of inflammatory diseases [4]. Recently, some investigators reported that plasma NGAL is a novel inflammatory marker for low-grade inflammatory conditions, such as Alzheimer’s disease and late-life depression [5].

Previous studies of NGAL have generally focused on its ability to predict worsening kidney function. There have been few studies that closely examined the role of NGAL as an inflammatory marker in low-grade inflammation. In the present study, we investigated whether plasma NGAL accurately reflects the severity of inflammation in low-grade inflammation. In the present study, we investigated whether plasma NGAL accurately reflects the severity of inflammation in low-grade inflammation, particularly based on the inflammatory markers of high-sensitivity CRP (hsCRP), TNF-α, IL-6, and corrected erythrocyte sedimentation rate (cESR).

Materials and Methods

Study population

The study included 184 patients under clinical investigation of systemic inflammation who had no renal dysfunction. Their
Sample collection and laboratory investigation

Blood samples were collected from patients at admission, immediately centrifuged, and stored in aliquots at -80°C until assay. All specimens were obtained before treatment. Plasma NGAL concentrations were measured by a fluorescence immunoassay using the Triage NGAL assay (Alere, Inc., San Diego, CA, USA), which analyzes plasma NGAL with a measurable range from 15 ng/mL to 1300 ng/mL. The intra-assay CVs (n=20) for three samples (mean NGAL, 68-547 ng/mL) were 4.2-6.5%; the inter-assay CVs calculated from duplicate results in 10 subsequent assays were 4.6-7.1%. A medical decision point was regarded as 150 ng/mL [6].

Serum hsCRP level was determined by the particle-enhanced immunonephelometry assay (Dade Behring, Inc, Deerfield, IL, USA). A value of P<0.05 was considered statistically significant.

Results

NGAL, TNF-α, and IL-6 in study population

The baseline characteristics of the study population are summarized in Table 1. Of the 184 patients, 114 (61.9%) had an elevated NGAL level, and 159 (86.4%) had an increase in the levels of IL-6 and TNF-α in patients with inflammation. Data were analyzed using SPSS software (IBM SPSS Statistics for Windows, Version 19.0, Armonk, NY, USA). A value of P<0.05 was considered statistically significant.
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The median levels of NGAL, TNF-α, IL-6, and hsCRP in high-grade inflammation were significantly elevated compared to those in low-grade inflammation. The positive rate of NGAL in low-grade inflammation was 27.3%, prominently lower than that of hsCRP (73.8%, P<0.001). However, no significant difference was observed between the positive rate of NGAL and hsCRP in high-grade inflammation (93.8% versus 97.9%, P=0.154). Median hsCRP level in high-grade inflammation was 8.8-times higher than that in low-grade inflammation, in contrast to plasma NGAL levels (3.4-times) between the corresponding groups (Table 2).

Table 2. NGAL, cytokines, and inflammatory markers by grades of inflammation.

<table>
<thead>
<tr>
<th>Grades of inflammation</th>
<th>Low grade (inflammation index; &gt;1.0 to &lt;2.5; n=88)</th>
<th>High grade (inflammation P value index; ≥ 2.5 to 4.0; n=96)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NGAL (ng/mL)</td>
<td>103.5 (69.5-159.0)</td>
<td>351.9 (198.5-852.4)</td>
</tr>
<tr>
<td>Prevalence (n, %)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NGAL&gt;150 ng/mL</td>
<td>24 (27.3)</td>
<td>90 (93.8)</td>
</tr>
<tr>
<td>hsCRP&gt;0.3 mg/dL</td>
<td>65 (73.8)</td>
<td>94 (97.9)</td>
</tr>
<tr>
<td>cESR&gt;15 mm/h</td>
<td>41 (46.6)</td>
<td>92 (95.8)</td>
</tr>
</tbody>
</table>

Data are expressed by mean ± SD, median (interquartile range), or frequency (%).

Regression analysis

In high-grade inflammation, plasma NGAL levels were significantly correlated with IL-6 (r=0.413, P<0.001), TNF-α (r=0.325, P<0.001), and hsCRP (r=0.432, P<0.001) after adjusting for potential confounders, such as age, sex, BMI, systolic blood pressure, and eGFR. However, in low-grade inflammation, no significant associations were noted between plasma NGAL concentrations and the levels of cytokines and inflammatory markers (Table 3).

Table 3. Multivariate regression analysis of plasma NGAL levels and physiologic parameters by severity of inflammation.

<table>
<thead>
<tr>
<th></th>
<th>Univariate</th>
<th>Multivariate*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Low grade</td>
<td>High grade</td>
</tr>
<tr>
<td>hsCRP (mg/dL)</td>
<td>0.152</td>
<td>0.495</td>
</tr>
<tr>
<td>(0.103)</td>
<td>(&lt;0.001)</td>
<td>(0.520)</td>
</tr>
<tr>
<td>cESR (mm/h)</td>
<td>0.126</td>
<td>0.467</td>
</tr>
<tr>
<td>(0.195)</td>
<td>(&lt;0.001)</td>
<td>(0.613)</td>
</tr>
<tr>
<td>TNF-α (pg/mL)</td>
<td>0.124</td>
<td>0.392</td>
</tr>
<tr>
<td>(0.208)</td>
<td>(&lt;0.001)</td>
<td>(0.264)</td>
</tr>
<tr>
<td>IL-6 (pg/mL)</td>
<td>0.138</td>
<td>0.481</td>
</tr>
<tr>
<td>(0.141)</td>
<td>(&lt;0.001)</td>
<td>(0.245)</td>
</tr>
</tbody>
</table>

Data are expressed as standard β (P value).

*Adjusted for age, sex, BMI, systolic BP, and eGFR.

ROC curve analysis

The diagnostic values of NGAL and hsCRP to identify an increase in IL-6 (>4.8 pg/mL) and TNF-α (>10.5 pg/mL) in patients with inflammation were investigated. The area under
the curve (AUC) of NGAL was significantly lower than that of hsCRP in low-grade inflammation (0.579 (95% CI, 0.449-0.708) versus 0.691 (95% CI, 0.557-0.826), P<0.001). However, there were no significant differences between the AUCs of NGAL and hsCRP in high-grade inflammation (0.703 (95% CI, 0.608-0.799) versus 0.726 (95% CI, 0.635-0.816), P=0.184) (Figure 1).

**Discussion**

In this study, the relationship between plasma NGAL levels and inflammatory parameters was investigated in patients with different grade of inflammation. The grade of inflammation was determined by a new scoring system of an inflammation index using hsCRP and cESR. A significant association was observed between NGAL and the levels of TNF-α, IL-6, and hsCRP in high-grade inflammation, but not in low-grade inflammation. The diagnostic ability of NGAL to identify an increase in IL-6 and TNF-α did not outperform that of hsCRP in low-grade inflammation. The results suggest that plasma NGAL does not accurately reflect the severity of inflammation in low-grade inflammation. NGAL is an early predictor for acute kidney damage; however, a wide heterogeneity in its predictive value is reported [6,9]. Several researchers observed that plasma NGAL can increase in the absence of tubular damage [10]. Moreover, it is unclear whether NGAL is a more crucial as an indicator of AKI than as a marker for inflammation. A group of investigators recently reported that measurement of plasma NGAL is helpful for assessing patients with low-grade inflammatory diseases [11].

Low-grade inflammation is a term for conditions in which cytokine-induced acute-phase response is ongoing. However, the scope of the condition is extremely broad and ambiguous. Clinically, low-grade systemic inflammation, mainly characterized by an elevated CRP level, is commonly associated with developing cardiovascular disease [12,13]. Duncan et al. [14] designed a classification method to indicate low-grade inflammation using several parameters including leukocytes, CRP, fibrinogen, and IL-6. In our study, we defined low-grade inflammation as the score (>1.0 to <2.5) of an inflammation index, based on the levels of hsCRP and cESR. In the current study, 47.8% of patient populations had low-grade inflammation, 2.4- to 3.2-times higher levels in TNF-α and IL-6 compared to healthy individuals.

Catalán et al. [15] reported that low-grade inflammation leads to an increased gene expression of NGAL in patients with obesity. In our study, there were no significant associations between plasma NGAL concentrations and the levels of IL-6, TNF-α, hsCRP, and cESR in patient with low-grade inflammation. These findings may reflect the differences in study populations, severity of disease, and detection methods of NGAL, particularly between gene expression and circulating plasma level of NGAL. Additionally, in the present study, the positive rate of NGAL in patients with low-grade inflammation was significantly lower than that of hsCRP. These results imply that plasma NGAL does not correctly represent the extent of inflammation in patients with low-grade inflammation.

Proinflammatory cytokines, such as TNF-α and IL-6, are crucial in inflammatory reaction. TNF-α stimulates the secretion of IL-6, which contributes to ischemia-reperfusion injury as a multifunctional cytokine [16]. In the present study, the ability of NGAL to detect an increase in IL-6 and TNF-α was investigated using an ROC curve analysis. The hsCRP exhibited a better diagnostic performance than NGAL in
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patients with low-grade inflammation, but the AUC of hsCRP was similar to that of NGAL in high-grade inflammation. These observations indicate that plasma NGAL is not superior to hsCRP for assessing low-grade inflammation, although the diagnostic efficacy of NGAL is comparable to that of hsCRP in high-grade inflammation. A measure of plasma NGAL does not seem to offer any clinical advantage over hsCRP, at least for screening low-grade inflammation. Inflammation is a risk factor for decreased renal function [17]. Systemic inflammation is frequently accompanied by kidney injury owing to its pathogenesis of hypoperfusion, apoptosis, microvascular thrombosis, and infiltration of immune cells [18]. In our study, to minimize the influence of kidney function on plasma NGAL levels, only subjects without renal impairment were enrolled. Our data obtained in patients with high-grade inflammation, who had well-preserved renal function, suggest that NGAL is a potential indicator for inflammation, but NGAL may be confined to the advanced degree of inflammation.

There are several limitations in this study. We did not measure plasma NGAL level in serial samples to assess the changes in NGAL in relation to the progression of disease. There may be unmeasured confounders for which we could not adjust during statistical analysis. Additionally, we emphasize that plasma NGAL level might be influenced by possibly missing information on renal or other organ injuries. Despite these limitations, our data may provide additional information for the diagnostic competence of NGAL in different grades of inflammation. In conclusions, this study demonstrates that plasma NGAL has sensitivity comparable to hsCRP in high-grade inflammation, but does not surpass the diagnostic accuracy of hsCRP in low-grade inflammation. The clinical relevance of NGAL as an inflammatory marker appears to be limited according to the intensity of inflammation. Thus plasma NGAL should be used with care as a biomarker to screen for patients with low-grade inflammation.

Acknowledgment

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