Association between intestinal microflora and renal function in patients with chronic renal failure: A case-control analysis

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ABSTRACT

Objective: To identify the association between the changes in intestinal microflora and renal function in patients with chronic renal failure (CRF).

Methods: This retrospective case-control study included 50 patients with CRF (study group), admitted to the Clinical Laboratory Department of Shenzhen People’s Hospital from March 2021 to May 2022, and 50 healthy individuals (control group). The association between the distribution of intestinal microflora and the glomerular filtration rate (GFR), levels of serum creatinine (SCr), blood urea nitrogen (BUN), and serum cystatin C (CysC) were analyzed.

Results: Intestinal microflora of CRF patients had significantly higher levels of Enterococci compared to the control group (p-Value <0.05), while the levels of Bifidobacterium spp. and Escherichia coli were lower in the study group (p-Value <0.05). GFR was lower, and the levels of BUN, SCr, and CysC were higher in the study group compared to the control group (all p-Value <0.05). GFR, BUN, SCr and CysC levels in the study group negatively correlated with the levels of Bifidobacterium spp. and Lactobacillus spp. (r<0, P<0.05), and positively correlated with the abundance of Enterococcus spp. and Escherichia coli (r>0, P<0.05) in the intestinal microflora.

Conclusions: Changes in intestinal microbiota are associated with a significant decrease in GFR and a marked increase in serum levels of renal function indicators, and alterations in the balance of intestinal microbiota may lead to further aggravation of the renal function damage in patients with CRF.

KEYWORDS: Chronic renal failure, Intestinal microflora, Structural change, Renal function, GFR, BUN, SCr, CysC.

INTRODUCTION

Chronic renal failure (CRF) occurs with a high incidence, affecting more than 10% of the world’s population, and is caused by a variety of conditions, such as nephritis, polycystic kidney disease, and nephrotic syndrome.¹² The disease is characterized by the severe destruction of the renal parenchyma and continuous deterioration of renal function.³ Numerous studies show that the intestinal microbiota directly participates in various physiological processes including metabolism, vitamin synthesis, regulation of intestinal mucosal immunity, and nutrient absorption.⁴ The interaction between the immune system and the normal intestinal microflora can effectively prevent invasion by pathogenic bacteria,⁵ but effects on specific cardiovascular outcomes are uncertain, as are effects in people without previous cardiovascular disease (primary prevention).⁶

Furthermore, dominant populations of symbiotic bacteria in the intestinal microflora can effectively inhibit colonization with pathogenic bacteria.⁷ Changes
in the intestinal microflora may lead to immune system abnormalities that can result in disease. Intestinal microflora imbalance, or dysbiosis, may be caused by a number of factors, such as the use of antibiotics, stress, age, dietary habits and lifestyle. Dysbiosis in patients with CRF is associated with dietary restrictions, the use of specific medications, and infiltration of urinary toxins into the intestinal cavity. Moreover, studies show that the occurrence and progression of CRF directly correlate with the microecological changes in the intestinal microflora.

The results of animal studies conducted by Bakris GL et al. showed that the intestinal permeability in CRF rats significantly increased with the increase in the extent of the intestinal injury. The impact of the changes in the composition of the intestinal microflora on the metabolism of nutrients may result in further worsening of metabolic disorders that can ultimately lead to CRF aggravation. There have been many studies on gastrointestinal microbiota in patients with chronic kidney disease, but few studies explore the association between gastrointestinal microbiota changes and renal function in patients with CRF. Therefore, the aim of this study was to evaluate the association between the changes in intestinal microflora and renal function indicators, such as glomerular filtration rate (GFR), levels of serum creatinine (SCr), blood urea nitrogen (BUN), and serum cystatin C (CysC), in patients with CRF. The results of our study may further help to gain insights into the pathogenesis of CRF.

METHODS

This was a retrospective case-control study. The clinical data of the patients with CRF diagnosed at the Clinical Laboratory Department of Shenzhen People's Hospital from March 2021 to May 2022 were reviewed and finally 50 patients were screened as the study group, while the control group included 50 healthy individuals observed during the same period. The groups were matched for age and gender.

Inclusion criteria:
- The diagnoses of individuals in the study group met the clinical diagnostic criteria for CRF.
- Individuals in the control group were healthy according to the results of physical examination.
- All participants had normal cognitive and mental functions.
- All participants were ≥ 18 years.
- CCR < 80 mL/min.
- GFR < 60 mL/min/1.73m², and last for >3 months.
- Complete relevant clinical data.

Exclusion criteria:
- Patients with infectious diseases, such as viral hepatitis, AIDS, pneumonia, or the new coronavirus.
- Patients with malignancies.
- Immune system diseases, including autoimmune diseases, immunodeficiency diseases, and allergic diseases.
- Individuals with serious conditions affecting heart, lung, or stomach function, and patients with hepatobiliary disease.
- Individuals with severe coagulation dysfunction.
- Patients with severe hydronephrosis.
- Patients with intestinal infections during that occurred within three months before the study.
- Individuals using prebiotics, probiotics, or antibiotics.

Ethical Approval: The Medical Ethics Committee of our hospital approved this study (No. LL-KY-2019424, Date: 2019-07-25).

All participants provided fecal samples to assess the composition of their intestinal microflora. Specimens were collected by sterile collectors and sterile toothpicks were used to take the mid-section of the samples to prevent cross-contamination. Two weighed fecal samples were dissolved in 4mL of normal saline each, and then diluted to 10⁶ concentrations before being inoculated using a drip method. The following growth media were used for culturing: Bifidobacterium-BS, Lactobacillus-LBS, Escherichia coli-EMB, and Enterobacterium-TTC agar plates (Guangdong Huankai Microbiology Technology, China). Fecal samples were cultured in triplicates for each dilution. BS and LBS agar plates were placed in an anaerobic incubator with a temperature of 37°C for 48 hours, and EMB and TTC agar plates were placed in an aerobic biochemical incubator at 37°C for 24 hours. After the incubation period, shape, color, size, edge, and density of the colonies on the plates was observed a representative colony was selected for Gram staining, and the microorganisms were visualized using a light microscope (Nikon ECLIPSE Ni-U, Japan).

On the aerobic EMB agar plates, Escherichia coli appeared as round colonies with neat edges and smooth or slightly convex surfaces. Under the light microscope, the bacteria appeared as straight rods in pairs. Lactobacillus spp. grew on anaerobic agar LBS plates as round gray or white colonies, 0.5-2.0-mm in diameter, with regular edges, and a smooth surface. The bacteria appeared as elongated bacilli arranged in single short chains and grids. The Enterococcus spp. on the aerobic agar TTC plate presented as round, rough, flat, dark red colonies with a diameter of 0.5-1.5 mm and a lack of neat edges. Under the light microscope, the Gram-positive bacilli appeared oval or spherical (a few were globular) arranged in pairs, long or short chains, or bundles. Bifidobacterium spp. grew on the anaerobic agar BS plates and presented as round, gray and white translucent colonies of neat edges, smooth surface, and 0.7-2.4 mm in diameter. Under the light microscope, these Gram-positive bacteria had uniform cell color cells of diverse shapes (mainly curved, straight, bifurcated, or rod-shaped, without flagella). Final number of bacteria were calculated as the number of colonies per unit weight.

We collected 5 mL of fasting venous blood from each participant to measure levels of renal function indicators. Briefly, samples were centrifuged at 5000 r/min for eight minutes and the supernatants were used to measure SCr, BUN, and CysC levels using a
fully automatic biochemical analyzer (model AU5800; Beckman Kurt, USA). BUN was detected via the urease-glutamate dehydrogenase method, with a reference range of 2.8 to 7.6 mmol/L. Reference ranges for SCr levels were 40 to 90 μmol/L for women and 64 to 104 μmol/L for men. Immunoturbidimetry was used for CysC detection, with a reference value range of 0.5 to 1.07 mg/L. We calculated the GFR using the CKD-EPI formula: 1.86 × SCr -1.164 × age -0.203.

**Statistical Analysis:** Data was processed and analyzed using SPSS 25.0 software. Counting data were expressed as means and standard deviations (X±s), and inter-group comparisons were done using one-way ANOVAs. All count data were expressed as percentages (n [%]) and inter-group comparisons were performed via χ² tests. Pearson correlation analysis was used for analyzing the correlation between intestinal microflora and renal function indicators. p-Value < 0.05 was considered as statistically significant.

**RESULTS**

There was no difference in the mean basic characteristics between the two groups (p-Value >0.05; Table-I). The abundance of intestinal Lactobacillus spp. was comparable between the two groups (P>0.05). However, the abundance of Escherichia coli and intestinal Bifidobacterium spp. in the study group were significantly lower than in the control group (p-Value <0.05), while the level of Enterococcus spp. was significantly higher (p-Value <0.05; Table-II).

Glomerular filtration rate of patients in the study group was significantly lower, and the levels of SCr, BUN and CysC were significantly higher than those in the control group (P<0.05; Table-III). The levels of GFR, SCr, BUN and CysC in patients with CRF negatively correlated with the intestinal levels of Bifidobacterium spp. and Lactobacillus spp. (r<0, P<0.05), and positively correlated with the abundance of Escherichia coli and Enterococcus spp. (r>0, P<0.05; Table-IV).

**DISCUSSION**

This study showed that CRF was associated with markedly lower levels of intestinal Escherichia coli and Bifidobacterium spp., whereas the abundance of Enterococci was significantly higher, compared to healthy individuals. Serum levels of SCr, BUN, and CysC in patients with CRF were significantly higher and the GFR was significantly lower than in healthy individuals. Moreover, the levels of these renal function indicators in patients with CRF negatively correlated with the abundance of intestinal microflora.

### Table-I: Comparison of general data between the two study groups.

<table>
<thead>
<tr>
<th>Group</th>
<th>n</th>
<th>Gender (male/female)</th>
<th>Age (years)</th>
<th>BMI (kg/m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study-group</td>
<td>50</td>
<td>29/21</td>
<td>49.14±10.88</td>
<td>23.92±2.78</td>
</tr>
<tr>
<td>Control-group</td>
<td>50</td>
<td>32/18</td>
<td>50.96±10.36</td>
<td>24.61±2.40</td>
</tr>
</tbody>
</table>

χ²/t P    0.378 -0.856 -1.338 0.539 0.394 0.184

### Table-II: Comparison of intestinal flora between the two study groups (lg CFU/g, X±s).

<table>
<thead>
<tr>
<th></th>
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<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Study-group</td>
<td>50</td>
<td>8.66±2.03</td>
<td>4.88±1.01</td>
<td>3.95±1.10</td>
<td>6.58±1.49</td>
</tr>
<tr>
<td>Control-group</td>
<td>50</td>
<td>5.74±1.49</td>
<td>7.62±2.01</td>
<td>6.28±1.62</td>
<td>6.81±1.54</td>
</tr>
</tbody>
</table>

t    8.180 -8.595 -8.392 -0.758

P    <0.001 <0.001 <0.001 0.450

### Table-III: Comparison of renal function indexes between the two groups (X±s).

<table>
<thead>
<tr>
<th>Group</th>
<th>n</th>
<th>GFR (ml/min)</th>
<th>SCr (μmol/L)</th>
<th>BUN (mmol/L)</th>
<th>CCysC (mg/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study-group</td>
<td>50</td>
<td>16.02±3.03</td>
<td>385.32±28.05</td>
<td>23.42±3.29</td>
<td>4.95±1.44</td>
</tr>
<tr>
<td>Control-group</td>
<td>50</td>
<td>93.00±11.70</td>
<td>74.96±10.59</td>
<td>4.52±1.18</td>
<td>1.90±0.36</td>
</tr>
</tbody>
</table>

t    -45.040 73.192 38.234 14.477

P    <0.001 <0.001 <0.001 <0.001
individuals. Therefore, it is reasonable to hypothesize that the intestinal microflora of patients with kidney transplant or chronic kidney disease differs significantly from that of healthy individuals. Korpela K et al showed that the intestinal microflora composition can oftentimes be altered in patients with chronic kidney disease due to a variety of factors, including changes in intestinal barrier function, the presence of pathogenic bacteria, and the accumulation of toxins. These changes can ultimately lead to the worsening of symptoms of renal micro- and systemic inflammatory reactions.

Abnormal intestinal microflora in patients with CRF have been associated with the presence of intestinal endogenous toxins, such as p-cresol sulfate, indole sulfate, trimethylamine oxide, and lipopolysaccharide. Accumulation of these toxins was linked to the destruction of the intestinal barrier function, toxin invasion into the bloodstream, and the presence of symptoms of renal micro- and systemic inflammatory reactions that ultimately lead to the worsening of the disease. Moreover, intestinal microflora may be potential targets in addressing CRF-related comorbidities, which may be a significant research direction in the future.

Studies have also shown that the levels of BUN, CysC and SCr are higher in patients with severe CRF. Changes in the balance between the symbiotic and pathogenic bacteria can affect the intestinal barrier function, microecosystem stability, and ultimately damage renal function. BUN levels, which are significantly increased in all kinds of acute and chronic renal dysfunction, reflect the levels of urea nitrogen in the blood. SCr can be used to measure the glomerular filtration capacity, and is elevated in cases of renal damage.

In addition, the correlation analysis in this study showed that the levels of BUN, CysC and SCr are negatively correlated with the intestinal abundance of Lactobacillus spp. and Enterococcus spp. in patients with CRF. The correlation coefficients for these indicators are statistically significant at the 0.001 level. Therefore, it is reasonable to hypothesize that patients with CRF may have abnormal intestinal microflora composition, and this may be an important factor in disease prevention and treatment. Patients with CRF may have abnormal intestinal microflora composition, and this may be an important factor in disease prevention and treatment. Patients with CRF may have abnormal intestinal microflora composition, and this may be an important factor in disease prevention and treatment.

Table IV: Correlation between intestinal flora distribution and renal function in patients with CRF.

<table>
<thead>
<tr>
<th>Intestinal flora</th>
<th>GFR</th>
<th>SCr</th>
<th>BUN</th>
<th>CysC</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>r</td>
<td>P</td>
<td>r</td>
<td>P</td>
</tr>
<tr>
<td>Enterococcus spp.</td>
<td>0.848</td>
<td>&lt;0.001</td>
<td>0.843</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Bifidobacterium spp.</td>
<td>-0.905</td>
<td>&lt;0.001</td>
<td>-0.837</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>0.808</td>
<td>&lt;0.001</td>
<td>0.831</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Lactobacillaceae spp.</td>
<td>-0.825</td>
<td>&lt;0.001</td>
<td>-0.784</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

r: Pearson correlation coefficient.
Limitations of the study: It is a single-center retrospective study that relies on the accurate, detailed, and available diagnosis and treatment data of patients with CRF. We collected data from patients diagnosed and treated during 2022, and our sample size was relatively small. Further large-scale and multi-center studies are needed. Furthermore, intestinal microflora is a complex ecosystem containing hundreds of bacterial species, of which we only studied a few in this study.

CONCLUSION

Changes in intestinal microbiota are associated with a significant decrease in GFR and a marked increase in serum levels of renal function indicators, and alterations in the balance of intestinal microbiota may lead to further aggravation of the renal function damage in patients with CRF.

REFERENCES