The Changes of Serum Lipoprotein(a) and Plasminogen Activator Inhibitor-1 in Primary Nephrotic Syndrome

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Abstract: Objective To investigate the changes of serum lipoprotein(a) [Lp(a)] and plasminogen activator inhibitor-1 (PAI-1) in primary nephrotic syndrome (PNS). Methods 60 PNS cases whose renal pathological types were identified and 57 healthy physical examination people as normal controls (NC) were selected. Automatic biochemical analyzer detected the levels of serum Lp(a) and blood lipids. Enzyme-linked immunosorbent (Eliisa) method tested the concentration of serum PAI-1. The nonparametric Spearman rank coefficient test was used to analyze correlations between variables. Results Compared with the NC group, Lp(a) and PAI-1 were significantly higher in PNS group (P<0.05). Lp(a), PAI-1 and lipids were found negatively related with albumin in PNS group. Lp(a) was positively correlated with PAI-1 in PNS patients, and the Spearman rank coefficient was 0.381 (P=0.003). Binary logistic regression analysis results showed that Lp(a) and PAI-1 were two risk factors in group PNS. Conclusions Lp(a) and PAI-1 concentrations are increased obviously in PNS patients, and Lp(a) was positively correlated with PAI-1 in PNS patients. They could be two risk factors of PNS patients.

Keywords: Lipoprotein(a); Plasminogen Activator Inhibitor-1; Primary Nephrotic Syndrome

Introduction
Clinical diagnosis and treatments found that the damages of kidney endothelial cells, micro vascular thrombosis and blood coagulation fibrinolytic system changes are closely associated with the occurrence and development of kidney disease [1, 2]. Serum lipoprotein (a) [Lp(a)] has the homologous sequence with plasminogen. It can inhibit the fibrinolytic function on the surface of fibrin [3]. Plasminogen activator inhibitor-1 (PAI-1) is the primary inhibitor of plasminogen activators in plasma, and its high expression will also inhibit the dissolution of fibrin [4]. At present, LP (a) and PAI-1 are considered to be associated with a variety of clinical diseases closely which are based on the injury of vascular endothelial cells [5, 6]. However, the correlation of serum LP (a) and PAI-1 levels and clinical values of primary nephrotic syndrome (PNS) patients is not clearly. Therefore, this study for the first time analyzed not only the changes of Lp(a) and PAI-1 levels in PNS patients but also the correlation among plasma Lp(a), PAI-1 and albumin levels to further elucidate the possible mechanism of serum Lp(a) and PAI-1 in PNS patients.

1 Materials and methods
1.1 Study subjects
The total of 60 patients with primary nephrotic syndrome for the study was recruited from the affiliated hospital of Qingdao university chose between April 2014 and July 2014. Their renal pathological types were identified according to the renal biopsy pathology diagnosis standard. Patients with diabetes mellitus, coronary heart disease, chronic hepatic disease, cancer, recent infection and so on which could affect lipid metabolism and fibrinolytic disorders were excluded. Secondary and hereditary nephrotic syndrome was also excluded. Fifty seven healthy people who had visited the affiliated hospital of Qingdao university to a routine check-up were selected randomly at the same period constituted the control group. The study protocol had been approved by the ethical committees of the hospital, and informed consent was obtained from all the recruiters. All laboratory measurements were performed after participants completed the study protocol.
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1.2 Methods
1.2.1 Collection of specimens
Specimens of all the research objects were collected after an overnight fast for at 6 o’clock in the morning. First of all, the basic information of the research objects such as height, weight and so on were measured and recorded. Second, elbow venous blood about 3 ml was extracted into the biochemical tubes, 37°C water bath for 15 minutes, and serum was promptly separated by a 20 min centrifugation at 2500 rpm and maintained at 4°C before lipid, kidney parameters, and lipoproteins analysis. At the same time, 300μl upper serum of each specimen for PAI-1 measurement was kept at ~80 °C until analysis.

1.2.2 Detection methods
Serum levels of total protein, albumin (ALB), triglycerides (TG), total cholesterol (TC), low density lipoprotein cholesterol (LDL-c), urea nitrogen (BUN), creatinine (CREA) and Lp(a) were measured using commercial reagents by Hitachi 7600 automatic biochemical analyzer. Serum PAI-1 level was detected by a‘sandwich”Enzyme-linked Immunosorbent Assay (ELISA). The human PAI-1 Elisa kits were bought from eBioscience. Each index of the detection had a strict standard of quality control, and tested by professional individuals who was trained and operated strictly according to the operation manuals.

1.2.3 Statistical analysis
Spss version 19.0 software was used to analyze the experimental data. One Sample Kolmogorov-Smirnov Test was used to evaluate the normality of distribution of the variables. The normal distribution data were expressed as mean±standard deviation, and skewed variables were expressed as median (interquartile range). The differences of variants of the two groups were analyzed by t-test when the data were consistent with a normal distribution. The differences of variants were analyzed by Wilcoxon rank sum test when the data were skewed distribution. The nonparametric Spearman rank coefficient test was used to analyze correlations between variables. Binary linear regression analysis was used to estimate risk factors of PNS patients. A two-tailed P value less than 0.05 was considered statistically significant.

2 Results
2.1 Comparisons of general clinical information of group PNS
The general clinical information in PNS and control group were shown in Table 1. There was no significant difference of gender, age, body mass index (BMI) of the two groups. Albumin levels were significantly lower in PNS than in the controls. Urea nitrogen level was significantly higher in PNS than NC, but no significant change of creatinine level was noted between PNS and NC.

Table 1 Comparisons of general clinical information of group PNS

| Group | n | Male/Female | Age (years) | BMI (kgm⁻²) | ALB (gL⁻¹) | BUN (mmolL⁻¹) | CREA (μmolL⁻¹) |
|-------|---|-------------|-------------|-------------|------------|---------------|---------------|
| PNS   | 60 (30/30) | 46.72±13.85 | 25.21±3.36 | 27.55±8.18 | 5.72(5.05,7.57) | 88.00(75.25,106.00) |
| NC    | 57 (28/29)  | 46.93±10.64 | 24.36±3.85 | 44.86±3.13 | 4.71(4.30,5.27) | 85.00(75.00,92.00) |
| P     | 0.924 | 0.926 | 0.217 | 0.00 | 0.00 | 0.15 |

Compared with NC. *P<0.05

2.2 Lp (a), PAI-1, lipids levels in the two groups
Lp(a), PAI-1 and lipids levels including triglycerides, total cholesterol and low density lipoprotein cholesterol were significantly higher in group PNS than in NC (Table 2).

Table 2 Lp (a), PAI-1, lipids levels in group PNS and NC

| Group | TG(mmolL⁻¹) | TC(mmolL⁻¹) | LDL-c(mmolL⁻¹) | LP(a)(mgL⁻¹) | PAI-1 (ngmL⁻¹) |
|-------|-------------|-------------|----------------|--------------|----------------|
| PNS   | 1.88±0.88*  | 7.54±2.46*  | 4.28±1.90**   | 454.50(202.00,822.00) | 63.25(48.43,78.69)* |
| NC    | 0.91±0.38   | 4.87±0.57   | 2.72±0.34     | 69.00(50.50,188.50) | 40.46(32.98,48.14) |
| P     | 0.00        | 0.00        | 0.00           | 0.00         | 0.00           |

Compared with NC. *P<0.05

2.3 Lp (a), PAI-1 and lipids related with albumin in PNS patients
Albumin was found negatively correlated with Lp(a), PAI-1, total cholesterol, triglyceride and LDL cholesterol levels in PNS patients (Table 3).

Table 3 Correlation between plasma albumin and Lp(a), PAI-1, lipids in group PNS

| Lp(a) | PAI-1 | TG | TC | LDL-c |
|-------|-------|----|----|-------|
| -0.165 | -0.550 | -0.379 | -0.211 |
| 0.208 | 0.000 | 0.003 | 0.045 |

Correlated with PAI-1 in PNS patients, and the Spearman rank coefficient was 0.381 (P<0.003). No correlation was noted between Lp (a) and PAI-1 in normal control group.

2.5 The analysis of Lp(a) and PAI-1 with PNS patients

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Binary logistic regression analysis was performed to analyze whether Lp(a) and PAI-1 were risk factors in PNS patients. Consequently, we found that both Lp(a) and PAI-1 were two risk factors in PNS patients (Table 4).

Table 4  The analysis of Lp(a) and PAI-1 with PNS patients

| Variable | B    | SE   | Wald | P     | OR  | 95% CI          |
|----------|------|------|------|-------|-----|-----------------|
| PAI-1    | 0.109| 0.029| 13.735| 0.000| 1.115| 1.053 - 1.181   |
| Lp(a)    | 0.010| 0.002| 16.823| 0.000| 1.010| 1.005 - 1.015   |

3 Discussions

Due to insidious onset of kidney disease when patients went to hospital the kidney disease has become seriously. Nephrotic syndrome as severe kidney disease affects the quality and safety of patients’ life. The significant abnormalities of lipid metabolism and fibrinolytic system in patients with nephrotic syndrome are closely related to presence and severity of kidney disease.[7-9]

Lp(a) was discovered a half-century ago by Kare Berg[10]. However, it still great challenges both basic researchers and clinicians. Lp(a) resembles a LDL-like particle that is linked to apolipoprotein(a) attached to apoB-100 by a disulfide bond[11, 12]. Elevated plasma levels of Lp(a) have been shown to be a independent risk factor for atherosclerosis[13, 14]. Lp(a) is difficult to grasp because of its unique structure, its considerable heterogeneity in both gene and isoform size, its wide range of plasma concentrations and its implication in ever more numerous cellular and biochemical mechanistic pathways[11]. In our study, we found that Lp(a) and lipids levels including triglycerides, total cholesterol and low density lipoprotein cholesterol were significantly higher in PNS than in the controls. Numerous studies have reported that the concentrations of total cholesterol, LDL cholesterol and triglyceride were elevated in NS patients [9, 15]. Hyperlipidemia and higher Lp(a) level may be caused by the increased hepatic protein synthesis driven by hypoalbuminemia and reduced clearance of Lp(a) in PNS patients[12]. Because albumin was found negatively correlated with Lp(a). Kidney disease causes lipid metabolic disorder and abnormal lipid metabolism speeds up the original kidney dysfunction in PNS patients. Lp(a) concentration remain extremely stable within an individual over their lifespan, and variation within the LPA gene on chromosome 6q23 contribute independently to the interindividual variability in plasma Lp(a) concentration[11, 16]. Therefore, further researches within the LPA gene of PNS patients are needed in the future.

The fibrinolytic system constitutes a critical response mechanism to vascular injury [17]. PAI-1 is an important inhibitor of the fibrinolytic system, so elevated levels could suppress fibrinolysis and result in an increased risk of thrombosis. PAI-1 is the primary inhibitor of plasminogen activators in plasma, rapidly inactivating both tissue plasminogen activator (t-PA) and urokinase-type plasminogen activator (u-PA). So, PAI-1 inhibit t-PA converts the plasminogen into plasmin which plays a major role in fibrinolysis[4, 18]. We found that PAI-1 levels are much more higer in PNS than NC group, which showed that patients with primary nephrotic syndrome has serious fibrinolytic system disorder. In our study, PAI-1 was found negatively correlated with albumin in PNS group. PAI-1 levels also may be caused by the increased hepatic protein synthesis driven by hypoalbuminemia. PAI-1 is synthesized in the liver and by endothelial cells, and its synthesis is regulated by several physiologic mediators, including endotoxin, interleukin-1 and lipids [19], and PAI-1 gene also plays an important part in PAI-1 concentration [20]. These factors within PNS patients need to be further studies.

In conclusion, in our study the levels of Lp(a) and PAI-1 were increased in PNS group. Lp(a) and PAI-1 had already significantly changed before creatinine changes which response for renal function. They may be closely related to the presence and severity of kidney disease. Lp(a) was positively correlated with PAI-1 in PNS patients, we found that Lp(a) and PAI-1 were two risk factors in PNS patients. These findings provide a theoretical basis for the diagnosis and treatment of PNS.

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