Research Article

Environmental Lead Exposure Accelerates Progressive Diabetic Nephropathy in Type II Diabetic Patients

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Whether environmental lead exposure has a long-term effect on progressive diabetic nephropathy in type II diabetic patients remains unclear. A total of 107 type II diabetic patients with stage 3 diabetic nephropathy (estimated glomerular filtration rate (eGFR) range, 30–60 mL/min/1.73 m²) with normal body lead burden (BLB) (<600 μg/72 hr in EDTA mobilization tests) and no history of exposure to lead were prospectively followed for 2 years. Patients were divided into high-normal BLB (>80 μg) and low-normal BLB (<80 μg) groups. The primary outcome was a 2-fold increase in the initial creatinine levels, long-term dialysis, or death. The secondary outcome was a change in eGFR over time. Forty-five patients reached the primary outcome within 2 years. Although there were no differences in baseline data and renal function, progressive nephropathy was slower in the low-normal BLB group than that in the high-normal BLB group. During the study period, we demonstrated that each 100 μg increment in BLB and each 10 μg increment in blood lead levels could decrease GFR by 2.2 mL/min/1.72 m² and 3.0 mL/min/1.72 m² (P = 0.005), respectively, as estimated by generalized equations. Moreover, BLB was associated with increased risk of achieving primary outcome. Environmental exposure to lead may have a long-term effect on progressive diabetic nephropathy in type II diabetic patients.

1. Introduction

Over the past 25 years, the prevalence of type II diabetes in the USA has almost doubled, with 3- to 5-fold increases in developing countries [1]. Diabetes is now the major cause of end-stage renal disease and the primary diagnosis causing kidney disease in 20–40% of patients starting treatment for end-stage renal disease worldwide [2, 3]. However, few studies have investigated the relationship between environmental exposure to lead and diabetic nephropathy.

Previous epidemiological studies [4–6] showed that blood lead levels (BLL) are related to renal function [4, 5] and exacerbated age-related decreases in renal function [6] in the general population, suggesting that environmental exposure to lead influences renal function in healthy individuals. Because BLL only indicates recent lead exposure [4, 7], body lead burden (BLB) is usually assessed by X-ray fluorescence to determine bone lead content and calcium disodium ethylenediaminetetraacetic acid (EDTA) mobilization tests [7]. A BLB greater than 600 μg, as determined by EDTA mobilization tests, is considered lead poisoning. Previous investigations that used EDTA mobilization tests to assess BLB in nondiabetic chronic kidney disease (CKD) patients with normal BLB [8–12] suggested that environmental lead exposure is associated with progressive CKD.

A 6-year study [13] indicated that bone lead content is related to progressive elevation of serum creatinine in persons with diabetes. However, these values were not adjusted for daily urinary protein excretion or daily protein intake. A short-term 1-year observational study [14] of type II diabetic patients with diabetic nephropathy suggested that environmental lead exposure might influence progressive diabetic nephropathy. However, the observation period was too short to demonstrate the long-term toxic effect of environmental
lead exposure; therefore, the estimated glomerular filtration rate (eGFR) was calculated from the American Modification of Diet in Renal Disease (MDRD) formula for CKD patients [15] rather than the Chinese formula for type II diabetic patients [16]. Hence, the relationship between low-level environmental exposure to lead and progressive diabetic nephropathy remains unclear. This 2-year prospective study was performed to clarify the relationship in type II diabetic patients.

2. Study Population and Methods

2.1. Subjects. The Institutional Review Board Committee of Chang Gung Memorial Hospital approved the study protocol. Each patient provided written informed consent.

Patients aged from 30 to 83 years who had type II diabetes mellitus with nephropathy and who received followup care at Chang Gung Memorial for more than 1 year were eligible for inclusion in this study if they met all the following criteria [14]: abnormal serum creatinine (>1.4 mg/dL); stage 3 CKD (eGFR between 30 mL/min/1.73 m² and 60 mL/min/1.73 m²); diabetic retinopathy treated with or without laser therapy; daily urinary protein excretion of more than 0.5 g/day; no microhematuria in urine tests; normal-sized kidneys as determined by echograms; history of diabetes for more than 5 years; no known history of exposure to lead or other heavy metals; and a BLB of less than 600 µg as measured by EDTA mobilization testing and 72-hour urine collection. Diabetic nephropathy diagnoses were also based on renal histological examination findings in cases where renal biopsies were performed.

The exclusion criteria were as follows: type I diabetes; renal insufficiency with a potentially reversible cause such as malignant hypertension, urinary tract infection, hypercalcemia, or drug-induced nephrotoxic effects; presence of other systemic diseases such as connective tissue diseases; use of drugs that might alter the course of renal disease such as nonsteroidal anti-inflammatory agents, steroids, immunosuppressive drugs, or Chinese herbal drugs; having joined a previous study [14]; drug allergies; and the absence of informed consent. The blood pressure of each patient was maintained at less than 140/90 mm Hg with diuretics and angiotensin-converting-enzyme inhibitors (ACEI) or angiotensin II receptor antagonists (ARA), with or without calcium-blocking agents and/or vasodilators [17]. Calcium carbonate was employed to maintain patients' phosphate levels. No patients received vitamin D3 supplements because their parathyroid hormone was below 200 pg/mL. Each patient received dietary consultation. A diabetic diet (35 Kcal/kg of body weight per day) with normal-protein intake (0.8–1.0 g of high biological value protein per kilogram of body weight per day) was recommended to each patient. A nutritionist reviewed the dietary intake of each patient every 3 to 6 months. A 24-hour urea excretion analysis was performed every 3 months to determine nitrogen balance and dietary compliance [18].

2.2. Measurements of Blood Lead Levels and Body Lead Burdens. BLL and BLB were measured as described previously [7–12]. BLB was measured using EDTA mobilization tests as modified by Behringer et al. [19]. Urinary excretion measured 72 hours after the intravenous infusion of 1 g of calcium disodium EDTA (Abbott Laboratories, North Chicago, IL, USA) was used to measure BLB. Blood and urine lead levels were determined by electrothermal atomic-absorption spectrometry (SpectrAA-200Z; Varian, CA, USA) with Zeeman background correction and a L'ovot platform. Both internal and external quality-control procedures were applied throughout this study and achieved consistently satisfactory results. A certified commercially prepared product (Seronorm Trace Elements, Sero AS, Billingstad, Norway) was utilized to monitor intrabatch accuracy and ensure interbatch standardization. The coefficient of variation for lead measurement was <5.3%. The detection limit was 0.01 µg/dL. External quality control was maintained via participation in the governmental National Quality-Control Program. Low-normal BLB was defined as <80 µg and high-normal BLB was defined as >80 µg and <600 µg [9–12, 14].

2.3. Study Protocol. Serum creatinine, glycosylated hemoglobin (HbA1c), daily urine protein excretion, daily protein intake, mean arterial pressure, cholesterol, and triglyceride levels were measured with an autoanalyzer system (model 736; Hitachi, Tokyo, Japan) at the beginning and end of the study and every 3 months during the 24-month clinical observation period. Blood pressure and body mass index were also measured at 3-month intervals. At the end of this period, we compared the changes in renal function between the 2 groups and assessed the relationship between BLB and the progressive decline of diabetic nephropathy. Renal function was assessed by creatinine clearance and eGFR (both in mL/min/1.73 m² of body surface area). A modified eGFR equation for Chinese patients with type II diabetes was employed [16] (D-GFR) (mL/min/1.73 m²) \( R^2 = 0.95 \): 313 \( \times \) (age) \( ^{-0.094} \) × (years) \( \times \) [SCr] \( ^{-1.059} \) (mg/dL) \( \times \) [Alb] \( ^{0.485} \) (g/dL) for men, and 783 \( \times \) (age) \( ^{-0.480} \) (years) \( \times \) [SCr] \( ^{-0.077} \) (mg/dL) \( \times \) [SUN] \( ^{-0.130} \) (mg/dL) for women. A total of 85 patients completed the initial study period (Figure 1).

2.4. Outcome Measures. The primary endpoint was a 2-fold elevation in serum creatinine (measured twice, 1 month apart) from baseline values, need for long-term dialysis, or death during the 24-month observation period. The secondary endpoint was temporal changes in renal function during the study period.

2.5. Statistical Analysis. The differences in variables and renal function between the 2 groups were analyzed by the Chi-square test and Student's t-test. All P values were two-tailed, and all results are presented as means ± SD. The Mann-Whitney U test was employed for data not normally distributed. We performed a sensitivity analysis that assigned the mean eGFR value of the treatment group to controls lost to followup and assigned the mean eGFR value of the control group to treated patients lost to followup. Generalized estimating equations (GEE) with linear analysis were employed in longitudinal multivariate analyses using SAS statistical
Type II diabetic patients \((n=198)\) with abnormal serum creatinine levels, eGFR between 30 and 60 mL/min/1.73 m\(^2\), and without previous lead exposure were enrolled.

The following patients were excluded: 21 who participated in a previous study, 47 who did not receive laser therapy for diabetic neuropathy, 12 with microhematuria, 10 with urine protein excretion <0.5 g/day, 10 with small-sized kidneys, 2 with BLB >600 \(\mu\)g, and 7 without informed consents.

Inclusion of 89 patients with diabetic neuropathy and normal BLB into the 24-month observation period.

Eventually, 85 patients completed the 24-month observation study.

Two patients were lost to follow-up, and 2 died of acute myocardial infarction.

**Figure 1:** Flow chart showing the enrollment and status of patients.

software (version 6.12) to further assess the temporal changes in variables and associations with progressive renal function (eGFR) during the observation period. Moreover, multivariate Cox analyses were used to determine the significance of the baseline variables for predicting the primary endpoint during the study period. These models included all variables identified in the literature as related to the progression of diabetic nephropathy [12–16]. A value of \(P < 0.05\) was considered statistically significant. Data were analyzed using SPSS, version 18.0 for Windows 95 (SPSS Inc., Chicago, IL).

3. Results

3.1. Study Subjects. A total of 89 patients participated in the study and 85 completed the 24-month observation period (58 men and 31 women) (Figure 1). The following baseline data were obtained: patient mean age, 60.1 ± 9.5 years (range, 33–83); body-mass index (weight in kilograms divided by the square of height in meters), 24.9 ± 3.3 (range, 14.9–33.4); serum creatinine level, 1.9 ± 0.3 mg/dL (range, 1.5–2.8 mg/dL); eGFR, 41.3 ± 6.9 mL/min/1.73 m\(^2\) of body surface area (range, 30.3–59.9 mL/min/1.73 m\(^2\) of body surface area); daily protein excretion, 3.0 ± 2.5 g (range, 0.5–12.2 g); daily protein intake, 0.97 ± 0.18 g/kg (range, 0.58–1.63 g/kg); HbA1c, 8.3 ± 1.9% (range, 5.7–14.7%); BLL, 4.3 ± 1.1 \(\mu\)g/dL (range, 0.8–10.4 \(\mu\)g/dL); and BLB, 109.9 ± 52.3 \(\mu\)g (range, 14.4–316.8 \(\mu\)g). Sixty-two patients (70.0%) had hyperlipidemia. Eighty-four patients (95.5%) had hypertension, and they were treated with ACEI or ARA. Fourteen patients (15.7%) smoked. Seventy-six patients (85.4%) had retinopathy, which was treated with laser therapy. Among all the study patients, 29 (32.6%) had a history of cardiovascular diseases, including myocardial infarction, congestive heart failure, stroke, and diabetic foot. BLL was associated with BLB in all study patients \((r = 0.274, P = 0.009)\).

3.2. Longitudinal Followup for a 24-Month Period. Table 1 summarizes demographic data, baseline chronic disease condition, use of ACEI or ARA, daily urinary urea and protein levels, and body lead burden for participants in each group. No significant differences in these baseline values were noted between the 2 groups on initial assessment or during the observation period. Table 2 compares the progression of diabetic nephropathy between the high-normal BLB and low-normal BLB groups during the observation period. Creatinine clearance and eGFR were higher in the low-normal BLB group than in the high-normal BLB group during months 18 to 24 of the observation period. Similar results were obtained in the sensitivity test (Table 3).

3.3. Outcome Measures. Thirty-nine patients had a 2-fold elevation in serum creatinine from the baseline values during the 24-month observation period; 5 patients in the high-normal BLB group required hemodialysis; 1 patient with high-normal and 1 with low BLB died of acute myocardial
| Variable                                      | Low-normal BLB group | High-normal BLB group | P value |
|-----------------------------------------------|----------------------|-----------------------|---------|
| Age (yr)                                       |                      |                       |         |
| Mean ± SD                                     | 60.7 ± 8.3           | 59.8 ± 10.1           | 0.698   |
| Range                                         | 46–79                | 33–83                 |         |
| Sex (no. of patients)                         |                      |                       |         |
| Men/women                                     | 17/10                | 41/21                 | 0.812   |
| Education yrs > 9 yrs (no. of patients)       |                      |                       |         |
| Mean ± SD                                     | 24.5 ± 3.0           | 25.1 ± 3.7            | 0.459   |
| Range                                         | 17.6–32.3            | 14.9–33.4             |         |
| Hyperlipidemia (no. of patients) (%)          |                      |                       |         |
| Mean ± SD                                     | 16 (60.6)            | 46 (68.9)             | 0.210†  |
| Range                                         | 15 (57.6)            | 41 (60.8)             | 0.352†  |
| Hypertension (no. of patients) (%)            |                      |                       |         |
| Mean ± SD                                     | 26 (97.0)            | 58 (94.6)             | 0.999†  |
| Use of angiotensin-converting-enzyme inhibitors or angiotensin-receptor antagonists (no. of patients) (%) | | |         |
| Mean ± SD                                     | 27 (100.0)           | 60 (98.6)             | 0.999†  |
| Use of nondihydropyridine calcium channel blockers (no. of patients) (%) | | |         |
| Mean ± SD                                     | 10 (48.5)            | 22 (33.8)             | 0.999†  |
| Use of dihydropyridine calcium channel blockers (no. of patients) (%) | | |         |
| Mean ± SD                                     | 9 (27.3)             | 24 (40.5)             | 0.812†  |
| Smoking (no. of patients) (%)                 |                      |                       |         |
| Mean ± SD                                     | 5 (15.2)             | 9 (24.3)              | 0.753†  |
| History of cardiovascular disease (no. of patients) (%) | | |         |
| Mean ± SD                                     | 8 (36.4)             | 21 (29.7)             | 0.808†  |
| Use of insulin at entry (no. of patients) (%) |                      |                       |         |
| Mean ± SD                                     | 6 (18.2)             | 16 (24.3)             | 0.795†  |
| HbA1c (%)                                     |                      |                       |         |
| Mean ± SD                                     | 8.2 ± 2.1            | 8.6 ± 1.8             | 0.367   |
| Range                                         | 5.8–14.0             | 5.7–14.7              |         |
| Mean arterial pressure (mmHg)                 |                      |                       |         |
| Mean ± SD                                     | 96.6 ± 12.0          | 98.6 ± 11.9           | 0.477   |
| Range                                         | 68.7–120             | 75–126                |         |
| Cholesterol (mg/dL)                           |                      |                       |         |
| Mean ± SD                                     | 211.0 ± 38.2         | 217.2 ± 54.5          | 0.596   |
| Range                                         | 124–278              | 105–414               |         |
| Triglycerides (mg/dL)                         |                      |                       |         |
| Mean ± SD                                     | 211.3 ± 107.2        | 206.2 ± 187.6         | 0.697   |
| Range                                         | 91–635               | 56–1185               |         |
| Serum creatinine (mg/dL)§                     |                      |                       |         |
| Mean ± SD                                     | 1.79 ± 0.21          | 1.88 ± 0.30           | 0.170   |
| Range                                         | 1.5–3.7              | 1.5–3.8               |         |
| Creatinine clearance rate (mL/min/1.73 m²)    |                      |                       |         |
| Mean ± SD                                     | 44.9 ± 10.8          | 41.1 ± 12.0           | 0.123   |
| Range                                         | 29.2–61.4            | 24.6–70.9             |         |
| Glomerular filtration rate§ (mL/min/1.73 m²)  |                      |                       |         |
| Mean ± SD                                     | 42.4 ± 6.2           | 40.8 ± 7.1            | 0.380   |
| Range                                         | 31.9–58.3            | 30.3–59.9             |         |
| Blood lead (µg/dL)                            |                      |                       |         |
| Mean ± SD                                     | 3.8 ± 3.0            | 4.6 ± 3.1             | 0.278   |
| Range                                         | 1.6–10.4             | 0.8–10.3              |         |
variable
Low-normal BLB group High-normal BLB group

| Variable                          | Low-normal BLB group | High-normal BLB group | P value |
|-----------------------------------|----------------------|-----------------------|---------|
| Body lead burden (µg)†            |                      |                       |         |
| Mean ± SD                         | 58.1 ± 16.7          | 132.4 ± 46.1          | 0.001   |
| Range                             | 14.4–79.8            | 82.8–316.8            |         |
| Daily protein excretion (g)       |                      |                       |         |
| Mean ± SD                         | 2.8 ± 2.5            | 3.2 ± 2.4             | 0.364   |
| Range                             | 0.5–10.5             | 0.5–12.2              |         |
| Daily protein intake (g/kg)       |                      |                       |         |
| Mean ± SD                         | 0.99 ± 0.16          | 0.96 ± 0.18           | 0.569   |
| Range                             | 0.60–1.55            | 0.53–1.68             |         |

* A high-normal body lead burden was defined as a lead value of at least 80 µg (0.39 µmol) but less than 600 µg (2.9 µmol) and a low-normal body lead burden as a lead value less than 80 µg (0.39 µmol).
† P values were calculated by Fisher’s Chi-square test, except in the comparisons of age, body-mass index, serum creatinine, creatinine clearance, glomerular filtration rate, blood lead level, and body lead burden, which were calculated by Student’s t-test.
‡ To convert values for serum creatinine to micromoles per liter, multiply by 88.4.
§ Modified equation for glomerular filtration rate of Chinese diabetic patients.
∧ To convert values for lead to micromoles per liter, multiply by 0.04286.
* Hyperlipidemia was defined as a serum cholesterol level above 240 mg per deciliter (6.2 mmol per liter) after diet control.
|| Hypertension was defined by the presence of at least two blood-pressure measurements above 140/90 mmHg in patients.
Cardiovascular diseases included ischemic heart disease, congestive heart failure, stroke, and diabetic foot.

infarction; and 2 patients with high BLB were lost to followup. A total of 45 (50.6%) patients reached the primary endpoint. Only 9 (9/27, 33.3%) patients had a body lead burden <80 µg, and 36 (36/62; 58.1%) of these subjects had body lead burdens >80 µg (Logrank tests, P = 0.023) (Figure 2). In addition, GEE with linear analysis showed that BLB or BLL were significant variables for predicting the progression of eGFR, after adjusting for other variables (Tables 4 and 5). Each 1 µg increase in BLB led to a decrease of 0.022 mL/min/1.73 m² in eGFR (P = 0.009) and each 1 µg/dL increase in BLL led to a 0.298 mL/min/1.73 m² decrease in eGFR (P = 0.010) during the 2-year study period. Moreover, multivariate Cox regression analysis demonstrated that BLB was a significant risk factor (hazard ratio [HR] = 1.01, 95% confidence interval [CI]: 1.01–1.02; P < 0.001) for achieving primary outcome in type II diabetic patients, even after adjustment for other factors (Table 6). Similarly, multivariate Cox regression analysis demonstrated that BLL >80 µg was a significant risk factor (HR = 2.79, 95% CI: 1.25–6.25; P = 0.012) for achieving primary outcome in these patients.

4. Discussion

The results of the present study indicate that BLB and BLL, even at low levels, are important risk factors for progressive diabetic nephropathy. These associations were strong, dose dependent, and consistent, even after comprehensive adjustments for other covariates. Our result is similar to those of previous reports showing that increased BLL is associated with a progressive decline in renal function in the general population [4, 5].

In comparison with our previous work [14], this study enrolled a different study cohort and showed several novel findings. First, patients with a high-normal BLB showed a higher incidence of progressive diabetic nephropathy than those with low-normal BLB, although the corresponding variables were not different between the 2 groups during the 2-year followup period. Moreover, similar results were
Table 2: Means of renal function during the 24-month observation period (n = 89).

| Renal function (mL/min/1.73 m²) | Low-normal BLB group (n = 27) | High-normal BLB group (n = 62) | P (95% CI) |
|---------------------------------|-----------------------------|-------------------------------|-----------|
| **Month 0**                     |                             |                               |           |
| Ccr                             | 44.9 ± 10.8                 | 41.1 ± 12.0                   | 0.123 (−9.1–1.0) |
| D-GFR                           | 42.4 ± 6.2                  | 40.8 ± 7.1                    | 0.300 (−1.5–4.8) |
| **Month 6**                     | (n = 27)                    | (n = 62)                      |           |
| Ccr                             | 41.5 ± 15.2                 | 33.1 ± 13.0                   | 0.010 (−14.6–2.1) |
| D-GFR                           | 38.1 ± 9.2                  | 32.9 ± 8.3                    | 0.010 (−9.1–1.3) |
| **Month 12**                    | (n = 27)                    | (n = 62)                      |           |
| Ccr                             | 41.0 ± 19.2                 | 26.2 ± 12.2                   | <0.001 (−21.6–8.1) |
| D-GFR                           | 38.1 ± 9.5                  | 27.2 ± 8.9                    | <0.001 (−13.6–5.2) |
| **Month 18**                    | (n = 26)                    | (n = 59)                      |           |
| Ccr                             | 33.1 ± 13.7                 | 24.5 ± 10.8                   | 0.003 (−14.1–3.0) |
| D-GFR                           | 32.7 ± 11.2                 | 23.7 ± 8.0                    | <0.001 (−13.3–4.7) |
| **Month 24**                    | (n = 24)                    | (n = 55)                      |           |
| Ccr                             | 34.2 ± 17.3                 | 19.8 ± 9.8                    | <0.001 (−20.5–8.2) |
| D-GFR                           | 31.4 ± 11.2                 | 20.2 ± 7.0                    | <0.001 (−15.3–7.1) |
| **Total decrease of renal function (mL/min/1.73 m²) during the 2-year observation period** | | | |
| Ccr                             | 12.1 ± 15.5                 | 21.2 ± 8.7                    | 0.002 (3.7–14.5) |
| D-GFR                           | 11.3 ± 11.7                 | 20.5 ± 7.7                    | 0.001 (4.8–13.7) |

Data were measured by the Student’s t-test except * data by Mann-Whitney method. P < 0.05 means significant differences. Ccr: creatinine clearance; D-GFR: estimated GFR for Chinese patients with type II diabetes.

Table 3: Sensitivity analysis of renal function from month 18 to month 24 of the observation period (n = 89).

| Renal function (mL/min/1.73 m²) | Low BLB group (n = 27) | High BLB group (n = 62) | P (95% CI) |
|---------------------------------|------------------------|-------------------------|-----------|
| **Month 18**                    |                        |                         |           |
| Ccr                             | 32.8 ± 13.6             | 24.9 ± 10.7             | 0.005 (−13.2–2.5) |
| D-GFR                           | 32.4 ± 11.1             | 24.2 ± 8.1              | <0.001 (−12.4–4.1) |
| **Month 24**                    | (n = 24)                | (n = 55)                |           |
| Ccr                             | 32.6 ± 16.1             | 21.5 ± 9.8              | <0.001 (−16.9–5.4) |
| D-GFR                           | 30.2 ± 11.1             | 21.5 ± 7.5              | <0.001 (−12.7–4.7) |
| **Total decrease of renal function (mL/min/1.73 m²) during the 2-year observation period** | | | |
| Ccr                             | 12.2 ± 14.6             | 18.6 ± 11.2             | 0.010 (0.7–12.0) |
| D-GFR                           | 12.3 ± 11.5             | 19.3 ± 8.4              | 0.006 (2.7–11.3) |

Data were measured by the Student’s t-test except * data by Mann-Whitney method. P < 0.05 means significant differences. Ccr: creatinine clearance; D-GFR: estimated GFR for Chinese patients with type II diabetes.

obtained in the sensitivity test. Second, each increment of 10 μg/dL of BLL was determined to potentially decrease GFR by 3.0 mL/min/1.73 m² after adjustment for covariates. In addition to BLL, BLB is a strong predictor of progressive diabetic nephropathy and can be easily monitored in clinical practice. Importantly, there were no safe limits of lead indices in our study. Consistent with our results, previous studies of healthy populations indicated a high correlation between measured BLL and BLB [20, 21]. Therefore, one can assume that under conditions of constant environmental lead exposure, a higher BLL should correspond to a higher BLB. Third, the present study included a more homogenous population than our previous study [14]. Only patients with stage 3 CKD were included in the present study, whereas patients with stages 2, 3, and 4 CKD were included in our previous work [14]. Achieving primary outcome in patients with different stages of CKD is associated with confounding effects. Moreover, patients with stage 4 CKD may have hyperparathyroidism, which can cause osteopathy; increase BLB, as measured by EDTA tests [22]; and result in selection bias in the classification of high-normal or low-normal BLB groups. Fourth, the eGFR was calculated from the Chinese-modified MDRD formula for CKD patients with type II diabetes rather than the formula used for American CKD patients. Lastly, because the present study used stricter definitions of the primary outcome (a 2-fold versus a 1.5-fold
increase in serum creatinine level from that of the baseline) and a longer followup period (24 months versus 12 months) than previous studies [13, 14], a more definitive conclusion regarding the long-term effect of environmental exposure to lead on progressive diabetic nephropathy can be drawn.

The mean BLL of our patients was only 4.3 μg/dL, which is lower than observed in our previous study [14] and slightly higher than that reported by nationwide surveys in Taiwan (3.0 μg/dL) [23], Europe (2.57 μg/dL) [24], and the USA (3.5 μg/dL) [25]. This difference could be the result of the older age (mean, 60.1 years old) of our study patients. The mean BLL of our patients was only 109.9 μg, which is much lower than subtle lead poisoning (>600 μg) levels [7]. Although there were no differences in baseline data, Kaplan-Meier analysis showed that patients with high-normal BLB were more likely (58.1%) to achieve the primary outcome than those (33.3%) with low-normal BLB during the 24-month followup. Multivariate Cox analysis indicated each 100-μg increase of BLB could lead to a 100% increase in the risk of achieving primary outcome. Consistent with this result, EDTA chelation therapy has shown benefits in retarding progressive diabetic nephropathy in type II diabetic patients with high-normal BLB [14, 26]. Hence, environmental exposure to lead may accelerate progressive diabetic nephropathy in these patients, and it is reasonable to suggest chelation therapy for patients with high-normal BLB, who accounted for 70% (62/89) of the current study patients.

The mechanism underlying the effect of environmental exposure to low-levels of lead on accelerating the development of progressive diabetic nephropathy remains unclear. Low-level lead exposure in a rat CKD model was found to hasten progressive CKD by accelerating microvascular and tubulointerstitial injury [27]. Studies performed on animals [28, 29] have shown that chronic exposure to low-dose lead results in the generation of reactive oxygen species, reduces nitric oxide availability and the expression of angiotensin II, and increases blood pressure [29]. It also promotes hydroxyl radical generation and lipid peroxidation [30], enhances vascular reactivity to sympathetic stimulation, and decreases DNA repair capacity, which might be relevant for rapidly dividing cells in the inflamed arterial wall [31]. Moreover, chronic exposure to low-level lead-induced oxidative stress and reduced nitric oxide availability were successfully treated with a lead chelating agent or antioxidants in rats [29, 32].

### Table 4: Longitudinal multivariate analysis of body lead burden and other predictors of progressive change in the estimated glomerular filtration rate (D-GFR), using generalized estimating equations, during the 24-month longitudinal study period (n = 89).

| Variable                                      | Estimate (interactive effect)* | P value |
|-----------------------------------------------|--------------------------------|---------|
| Age (each increment of 1 yr)                  | −0.271                         | <0.001  |
| Gender (female versus male)                   | −3.575                         | <0.001  |
| Smoking (no versus yes)                       | −0.259                         | 0.813   |
| Body-mass index (each increment of 1 kg/m²)   | 0.020                          | 0.852   |
| History of cardiovascular diseases (no versus yes) | −0.375                        | 0.686   |
| MAP (mmHg) (each increment of 1 mmHg)         | −0.054                         | 0.033   |
| Cholesterol (mg/dL) (each increment of 1 mg/dL) | −0.002                        | 0.811   |
| Triglycerides (mg/dL) (each increment of 1 mg/dL) | −0.004                        | 0.075   |
| HbA1c (%) (each increment of 1%)              | 0.013                          | 0.793   |
| Serum creatinine (mg/dL) (each increment of 1 mg/dL) | −6.997                        | <0.001  |
| Body lead burden (μg) (each increment of 1 μg) | −0.022                         | 0.009   |
| Daily protein intake (g/kg) (each increment of 1 g/kg) | 1.287                         | 0.460   |
| Daily protein excretion (g) (each increment of 1 g) | −0.417                         | 0.035   |

The interactive effect of variables was calculated by a generalized estimating equation. Negative values for the interactive effect indicate a decline in the glomerular filtration rate, and positive values indicate an increase. Cardiovascular diseases included ischemic heart disease, congestive heart failure, stroke, and diabetic foot. MAP: mean arterial pressure.

### Table 5: Longitudinal multivariate analysis of blood lead level and other predictors of progressive change in the estimated glomerular filtration rate (D-GFR), using generalized estimating equations, during the 24-month longitudinal study period (n = 89).

| Variable                                      | Estimate (interactive effect)* | P value |
|-----------------------------------------------|--------------------------------|---------|
| Age (each increment of 1 yr)                  | −0.268                         | <0.001  |
| Gender (female versus male)                   | −3.261                         | <0.001  |
| Smoking (no versus yes)                       | −0.631                         | 0.604   |
| Body-mass index (each increment of 1 kg/m²)   | −0.018                         | 0.861   |
| Previous cardiovascular diseases (no versus yes) | −0.220                        | 0.818   |
| MAP (mmHg) (each increment of 1 mmHg)         | −0.057                         | 0.025   |
| Cholesterol (mg/dL) (each increment of 1 mg/dL) | −0.006                        | 0.406   |
| Triglycerides (mg/dL) (each increment of 1 mg/dL) | −0.005                        | 0.024   |
| HbA1c (%) (each increment of 1%)              | −0.002                         | 0.976   |
| Serum creatinine (mg/dL) (each increment of 1 mg/dL) | −7.550                        | <0.001  |
| Blood lead level (μg/dL) (each increment of 1 μg/dL) | −0.298                        | 0.010   |
| Daily protein intake (g/kg) (each increment of 1 g/kg) | 1.143                         | 0.539   |
| Daily protein excretion (g) (each increment of 1 g) | −0.400                         | 0.045   |

The interactive effect of variables was calculated by a generalized estimating equation. Negative values for the interactive effect indicate a decline in the glomerular filtration rate, and positive values indicate an increase. Cardiovascular diseases included ischemic heart disease, congestive heart failure, stroke, and diabetic foot. MAP: mean arterial pressure.
show not be related to low eGFR in levels, is associated with CKD in the general population [33, 34]. Moreover, higher BLL in the range below 10 μg/dL was shown to be related to lower cystatin-estimated GFR [35] in adolescents. These previous studies support the current study results. However, much remains to be explored regarding the mechanisms of lead-induced progressive diabetic nephropathy.

The use of eGFR to assess altered renal function is one of the limitations of the present study. However, a study on eGFR in Chinese patients with type II diabetes conducted by Barbosa et al. [36] demonstrated a strong correlation between eGFR and isotopic GFR ($r^2 = 0.95$) in the Chinese population. Another limitation of this study was that BLB was not assessed using X-ray fluorescence methods. However, there are several important limitations associated with X-ray fluorescence-based methods [36], such as lack of precision, nonhomogenous lead distribution in cortical bone, and a low turnover rate with low biological activity of lead in cortical bone. However, lead that can be chelated by EDTA predominantly reflects lead concentrations in the blood and soft tissues. Because the kidneys are included among soft tissues, EDTA mobilization may reflect the lead content of the kidney [37], which may influence progressive CKD.

**Table 6:** Cox regression analysis of the overall risk of the primary outcome of progressive renal insufficiency, according to baseline prognostic factors ($N = 89$).

| Variable                                      | Hazard ratio (95% CI) | $P$ value |
|-----------------------------------------------|-----------------------|-----------|
| Age (each increment of 1 yr)                  | 0.98 (0.94–1.02)      | 0.337     |
| Female sex                                    | 1.83 (0.87–3.83)      | 0.111     |
| Smoking (no versus yes)                       | 0.75 (0.26–2.12)      | 0.582     |
| Baseline body-mass index (each increment of 1 kg/m$^2$) | 0.90 (0.81–0.99)      | 0.023     |
| Previous cardiovascular diseases (no versus yes) | 0.55 (0.24–1.29)      | 0.170     |
| MAP (mmHg) (each increment of 1 mmHg)         | 1.03 (1.00–1.06)      | 0.088     |
| Cholesterol (mg/dL) (each increment of 1 mg/dL) | 1.00 (0.99–1.01)      | 0.822     |
| Triglycerides (mg/dL) (each increment of 1 mg/dL) | 1.00 (1.00–1.00)      | 0.937     |
| HbA1c (%) (each increment of 1%)              | 0.98 (0.83–1.16)      | 0.806     |
| Baseline serum creatinine (each increment of 1 mg/dL) | 0.29 (0.06–1.29)      | 0.104     |
| Body lead burden (each increment of 1 μg)     | 1.01 (1.01–1.02)      | $<0.001$  |
| Baseline daily protein intake (each increment of 1 g/kg) | 0.41 (1.06–1.44)      | 0.462     |
| Baseline daily protein excretion (each increment of 1 g) | 1.24 (1.12–1.42)      | 0.008     |

Cardiovascular diseases included ischemic heart disease, congestive heart failure, stroke, and diabetic foot. MAP: mean arterial pressure.

5. Conclusion

The results of this prospective study indicate that environmental exposure to lead may accelerate progressive diabetic nephropathy in type II diabetic patients despite the control of treatable factors during long-term followup. These results suggest that avoiding exposure to any sources of lead in the environment and chelation therapy are important in patients with BLB >80 μg. The findings of the current study are important because diabetic nephropathy is the major cause of end-stage renal disease in the world.

**Conflict of Interests**

All authors report that there is no conflict of interests.

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