Thyroid hormone ameliorates diabetic nephropathy in a mouse model of type II diabetes

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Abstract

Conventional therapies for diabetic patients, such as strict glycemic control, do not completely stop the progression of diabetic nephropathy. Serum-free tri-iodothyronine ($T_3$) levels were lower in patients with type II diabetes. The purpose of this study was to test a hypothesis that treatment with $T_3$ would improve diabetic nephropathy in db/db mice, a model of type II diabetes. Male db/db mice (16 weeks) were treated with $T_3$ for 4 weeks. Urinary excretions of albumin and blood glucose levels were measured. Kidneys were collected for histological examination and molecular assays of transforming growth factor-$\beta_1$ (TGF-$\beta_1$) expression and phosphatidylinositol 3-kinase (PI3K). $T_3$ attenuated albuminuria in db/db mice, suggesting an improved kidney function. $T_3$ significantly decreased accumulation of collagenous components in cortical interstitium (interstitial fibrosis) and expansion of mesangial matrix in glomeruli (glomerulosclerosis) and prevented the loss of glomeruli in db/db mice. Therefore, $T_3$ improved the renal structural damage seen in diabetic mice. Notably, diabetic nephropathy was accompanied by a significant decrease in PI3K activity and an increase in TGF-$\beta_1$ expression in kidneys. $T_3$ restored renal PI3K activity, attenuated hyperglycemia, and decreased renal TGF-$\beta_1$ expression in db/db mice. These effects of $T_3$ were abolished by simultaneous treatment with PI3K inhibitor (LY294002). These data suggest that $T_3$ prevented progressive kidney damage and remodeling in db/db mice by improving insulin signaling (e.g. PI3K activity).

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Introduction

Diabetic nephropathy is now the most common cause of end-stage renal disease (ESRD), which significantly contributes to the high mortality in type II diabetic patients (Ruggenenti & Remuzzi 1998). It is projected that 30–40% of patients with type I diabetes and 5–10% of patients with type II diabetes eventually develop ESRD (Ruggenenti & Remuzzi 1998, Lin & Sun 2010a). Conventional therapies such as strict glycemic control and antihypertensive treatment do not completely stop the progression of diabetic nephropathy in diabetic patients (Ruggenenti & Remuzzi 1998).

The db/db mouse is a useful animal model of type II diabetes with a point mutation in leptin receptor gene, which develops hyperglycemia, insulin resistance in peripheral tissues, and obesity by 8 weeks of age (Breyer et al. 2005). The db/db mice exhibit renal glomerular mesangial matrix expansion, increases in renal collagen and serum creatinine, albuminuria, and a decrease in glomerular filtration rate (GFR), which are comparable to advanced human diabetic nephropathy with renal insufficiency (Sharma et al. 2003).

Glomerular mesangial matrix expansion has been considered the essential characteristic of diabetic nephropathy in human (Sharma et al. 2003).

Hypothyroidism is accompanied by a decrease in GFR and renal blood flow (Iglesias & Diez 2009). The low level of tri-iodothyronine ($T_3$) is associated with a survival disadvantage of chronic kidney disease (Iglesias & Diez 2009). Type II diabetic patients with subclinical hypothyroidism are associated with an increased risk of diabetic nephropathy (Chen et al. 2007). In contrast, hyperthyroidism increases GFR and renal blood flow (Iglesias & Diez 2009).

Clinical studies indicated that serum-free $T_3$ level was about 47% lower in patients with type II diabetes compared with non-diabetic patients (Islam et al. 2008). There were no significant differences in serum levels of free thyroxine ($T_4$) and TSH between the control and the study subjects (Islam et al. 2008). The serum level of $T_4$ was lower in db/db mice and the tissue level of $T_3$ was lower in liver and brain due to impaired deiodination in these tissues of db/db mice (Kaplan & Young 1987). The purpose of this study was to test our hypothesis that chronic administration of $T_3$ would attenuate diabetic nephropathy in db/db mice.
Materials and Methods

Animals

BKS.Cg-+Lepr<sup>db</sup>/+Lepr<sup>db</sup>/OlaHsd mice (db/db mice) and BKS.Cg-m+ /+Lepr<sup>db</sup>/OlaHsd (lean mice) (all males, 12 weeks) were purchased from Harlan (Indianapolis, IN, USA). All mice were housed at room temperature (25 ± 1°C) and were provided with Purina laboratory chow (No. 5001) and tap water which was made available ad libitum. This study was approved by the Institutional Animal Care and Use Committee at the University of Oklahoma Health Sciences Center.

Experimental protocol

The kidneys used for this study were from the same groups of animals used in our previous publication on the effects of T<sub>3</sub> on diabetes in db/db mice (Lin & Sun 2010b). Briefly, three groups of db/db and three groups of lean mice were used (all 12 weeks, five mice/group). Following a 4-week control period, three groups of each strain (16 weeks) received i.p. injections of vehicles (35% DMSO and PBS), T<sub>3</sub> (7 ng/g b.w. in PBS, Sigma), and LY294002 (3 mg/g dissolved in 35% DMSO, Sigma) followed by T<sub>3</sub> respectively. The same doses of T<sub>3</sub> and LY294002 were given twice daily (0900 and 1700 h) for 18 days (14 ng/g per day for T<sub>3</sub> and 6 mg/g per day for LY294002). LY294002 was given 20 min before injection of T<sub>3</sub>. Animals were further treated with the increased dose of T<sub>3</sub> (28 ng/g per day) and LY294002 (9 mg/g per day) for another 10 days. A group treated with LY294002 alone was not included because the db/db mice died within 2–3 days after treatments, probably due to severe hyperglycemia. At 4 weeks prior to the treatment and during week 4 of the treatment, 24-h urine samples were collected for measurements of urinary albumin.

At the end of week 4 of the treatments with T<sub>3</sub>, animals were killed and blood was collected in EDTA for measuring blood glucose and plasma T<sub>3</sub>. Blood glucose was measured using a Reli On Ultima glucose reader (Solartek Products, Inc., Alameda, CA, USA). Plasma T<sub>3</sub> was measured using a T<sub>3</sub> ELISA kit (Alpha Diagnostic International, San Antonio, TX, USA). Animals were then perfused transcardially using heparinized saline. Following perfusion, kidneys were isolated and weighed. The left kidney from each mouse was fixed with 4% paraformaldehyde in PBS for 24 h. The kidneys were set in a plane perpendicular to the long axis and were embedded in paraffin for histological and immunohistochemical analysis.

Measurements of albumin

Urinary albumin excretion was measured with a mouse-specific microalbuminuria ELISA kit (Albuwell M; Exocell, Philadelphia, PA, USA) according to the manufacturer’s instruction.

Morphological investigations

A series of cross sections of the left kidney (3–5 μm) were cut at the vascular poles. Tissue sections (four to five specimens per group) were stained with periodic acid Schiff (PAS), Masson’s trichrome, or H&E. For sections stained with Masson’s trichrome (5μm thickness), images of cortex from three consecutive sections for each animal were collected at equal exposure conditions under Nikon Eclipse Ti microscopy (magnification ×100). The fraction area for collagenous components in cortex was obtained with NIS-Elements BR 3.0 (Nikon, Melville, NY, USA). Images of sections for each animal were also collected at equal exposure conditions under the microscopy at the magnification of 400.

For sections stained with PAS (3μm), images of 20 glomeruli for each animal were collected at equal exposure conditions and at the magnification of 400 under a microscope (Nikon Eclipse Ti). Mesangial matrix area was defined by

![Figure 1](https://example.com/figure1.png)

**Figure 1** Treatments with T<sub>3</sub> decreased albuminuria in db/db mice. T<sub>3</sub> decreased urinary albumin excretion in db/db mice (A). T<sub>3</sub> increased kidney mass in lean and db/db mice (B). Data = mean ± S.E.M. n=5. *P<0.05, ***P<0.001 versus lean-DMSO-PBS group; ++*P<0.001 versus db/db-DMSO-PBS; ***P<0.001 versus db/db-DMSO-T<sub>3</sub>.
PAS-positive and nuclei-free area in the mesangium. The glomerular area was defined by tracing along the borders of the capillary loop. Relative mesangial area (defined as fraction area of mesangial matrix area over glomerular area) was obtained using Image J (NIH freeware, Bethesda, MD, USA).

For sections stained with H&E (5 μm), the number of glomeruli was quantified using Nikon Eclipse Ti microscopy at a magnification of 100. Briefly, the kidneys were sectioned in a plane perpendicular to the long axis, beginning at the vascular poles. The kidneys were cut at 250 μm intervals. The number of glomeruli was quantified in three randomly chosen fields in each section and averaged based on five consecutive sections for each animal. The area (mm²) of each field was measured. Images of cortex for each animal were also collected at equal exposure conditions at a magnification of 400.

Phosphatidylinositol 3-kinase activity

Lysates of mouse kidney were immunoprecipitated with antibody against phosphatidylinositol 3-kinase (PI3K; Millipore, Temecula, CA, USA). PI3K activity in the immunocomplexes was determined using a PI3K ELISA kit (Echelon Biosciences, Salt Lake City, UT, USA).

Western blotting

Lysates of mouse kidney under reduced conditions were directly subjected to SDS-PAGE followed by western blotting with an antibody against transforming growth factor-β1 (TGF-β1; Santa Cruz Biotechnology, Santa Cruz, CA, USA, TGF-β1 and its precursor) and then with an antibody against β-actin (Santa Cruz) after stripping the blot.

Statistical analysis

Data were analyzed using a two-way ANOVA (strain and treatments). The Newman–Keuls procedure was used to assess differences between means. Data were expressed as mean ± S.E.M. A value with \( P<0.05 \) was considered significant.

Results

\( T_3 \) decreased urinary albumin excretion in diabetic mice

At the age of 12 weeks, the db/db mice had a significantly higher level of urinary albumin than that of the lean mice (Fig. 1A). No significant difference of urinary albumin excretion was found between any two groups of db/db mice or lean mice at the age of 12 weeks (Fig. 1A). Albuminuria of the db/db mice at the age of 20 weeks was significantly increased compared with that at the age of 12 weeks (Fig. 1A). Treatment with \( T_3 \) significantly decreased albuminuria in db/db mice (Fig. 1A). Treatment with PI3K inhibitor (LY294002) abolished the beneficial effect of \( T_3 \) on albuminuria in db/db mice (Fig. 1A), suggesting that this effect may be mediated by PI3K.

There was a significant decrease in kidney mass in the db/db-DMSO-PBS group (control) versus the lean-DMSO-PBS group at the age of 20 weeks (Fig. 1B). Treatment with \( T_3 \) significantly increased the kidney mass in lean mice (Fig. 1B). Body weight was significantly greater in the db/db-DMSO-PBS group than in the lean groups (Lin & Sun 2010b). Treatment with \( T_3 \) did not affect body weights in either lean or db/db mice (Lin & Sun 2010b).

Figure 2  Treatment with \( T_3 \) attenuated collagen deposition in renal cortex of db/db mice. Representative photomicrographs of Masson’s trichrome-stained kidney sections with original magnification ×400 (A). Arrows indicate trichrome-positive collagenous components in cortical interstitium (blue). Kidney sections were cut at their vascular poles. Quantitative analysis of collagenous components in renal cortex (B). Images of cortex from three consecutive sections for each animal, collected at equal exposure conditions under Nikon Eclipse Ti microscopy (×100), were used in the analysis. Data = mean ± S.E.M. \( n=4–5 \). ** \( P<0.01 \) versus the lean-DMSO-PBS; + \( P<0.05 \) versus the db/db-DMSO-PBS group; ++ \( P<0.01 \) versus the db/db-DMSO-T3 group.
**T3 attenuated collagen accumulation in kidney cortex in diabetic mice**

The Masson trichrome staining showed that the 20-week-old db/db mice had significantly higher levels of collagenous components in cortical interstitium compared with those of the lean mice (Fig. 2A and B). Treatments with T3 significantly decreased collagenous components in db/db mice (Fig. 2A and B). PI3K inhibitor abolished the beneficial effect of T3 (Fig. 2A and B). These data indicated that T3 prevented interstitial fibrosis in db/db mice and that PI3K was important in mediating this beneficial effect in db/db mice.

**T3 decreased glomerular mesangial matrix expansion in diabetic mice**

The db/db mice showed significant glomerular mesangial matrix expansion compared with the lean mice (Fig. 3A and B), indicating glomerular remodeling. Treatments with T3 abolished glomerulosclerosis in db/db mice (Fig. 3A and B). PI3K inhibitor partially blocked the effect of T3 on the mesangial matrix expansion.

**T3 attenuated kidney damage in diabetic mice**

The db/db mice showed renal tubule atrophy, some lost typical structures of cortical tubules (Fig. 4A), suggesting renal remodeling in diabetic mice. T3 restored tubular structures (Fig. 4A). The number of glomeruli was significantly decreased in the db/db-DMSO-PBS group compared with the lean-DMSO-PBS groups (Fig. 4B), suggesting a loss of glomeruli at the age of 20 weeks. The glomeruli were replaced by massive cellular infiltrates in the diabetic mice. T3 prevented the loss of glomeruli in the db/db mice (Fig. 4B). These beneficial effects of T3 may be mediated by PI3K because they could be abolished by the PI3K inhibitor (Fig. 4B).

**T3 increased PI3K activities in kidneys and attenuated hyperglycemia in diabetic mice**

The PI3K activity (PIP3 products) was significantly lower in kidneys of the db/db-DMSO-PBS mice than that in the lean mice (Fig. 5). T3 significantly increased the PI3K activity in kidneys of both db/db and lean mice (Fig. 5). The PI3K inhibitor abolished the stimulating effects of T3 on PI3K activities.

The blood glucose level was significantly higher in the db/db-DMSO-PBS mice than in the lean mice (Lin & Sun 2010b). T3 (14 ng/g per day) significantly attenuated the blood glucose level in the db/db mice but not in the lean mice. An increase in the dose of T3 (28 ng/g per day) further attenuated hyperglycemia (Lin & Sun 2010b). The PI3K inhibitor abolished the anti-hyperglycemic effects of T3 (Lin & Sun 2010b).

The basal level of plasma T3 in db/db mice (61.7 ± 5.2 ng/dl) was significantly lower than that of the lean mice (86.5 ± 7.2 ng/dl). Daily administration of T3 (28 ng/g per day) for 7 days increased the plasma levels of T3 by 2.7-folds in db/db mice (Lin & Sun 2010b). As we reported recently (Lin & Sun 2010b), the basal level of plasma insulin was not significantly different between the lean and the db/db mice, although the latter had severe hyperglycemia. Treatment with T3 significantly increased the plasma level of insulin (Lin & Sun 2010b).

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**Figure 3** Treatment with T3 prevented mesangial matrix expansion in db/db mice. Representative photomicrographs of PAS-stained kidney sections with original magnification of ×400 (A). Arrows indicate PAS-positive mesangial material (pink). Quantitative measurements of mesangial matrix expansion (B). Relative mesangial matrix area is expressed as PAS-positive mesangial matrix per total glomerular tuft cross-sectional area. Kidney sections were cut at their vascular poles. An average value was obtained from analyses of 20 glomeruli per mouse.

Data = mean ± s.e.m. n = 4–5. ***P < 0.001 versus the lean-DMSO-PBS group; ++P < 0.01 versus the db/db-DMSO-PBS group; ++P < 0.001 versus the db/db-DMSO-T3 group.
Renal fibrosis, including tubulointerstitial and glomerular fibrosis, is an important feature of diabetic nephropathy (Brosius 2008). Interstitial fibrosis seems to contribute to the deterioration of renal function, especially during the late stage of diabetic nephropathy (Brosius 2008). Diabetic glomerular fibrosis is mainly due to accumulation of extracellular matrix proteins such as collagen and fibronectin in the mesangial interstitial space, which results in progressive diabetic nephropathy by reducing the surface area of glomerular capillaries for filtration (Steffes et al. 1989, Brosius 2008). This study showed that there was renal structural remodeling as evidenced by a loss of typical structures of renal tubules and glomeruli in db/db mice (Figs 2 and 4). The loss of glomeruli in db/db mice is surprising as it has not been observed in human and other experimental models of type II diabetes. The loss of glomeruli and cellular infiltration in the late stage of diabetes may be due to hyperglycemia-related vascular damages (Lehmann & Schleicher 2000, Koya et al. 2003). It is known that long-term hyperglycemia causes vascular oxidative stress resulting in vascular damage and cellular infiltration (Suzuki & Miyata 1999, Lehmann & Schleicher 2000, Koya et al. 2003), which may be eventually replaced by fibrosis (Lehmann & Schleicher 2000). Interestingly, treatments with T3 prevented accumulation of collagenous components in cortical interstitium and glomerular mesangial matrix expansion in diabetic mice (Figs 2 and 3).

TGF-β has been shown to be linked to renal fibrosis in diabetic nephropathy in animals and humans (Lehmann & Schleicher 2000, Zhu et al. 2007, Chiarelli et al. 2009). Suppression of TGF-β1 inhibited hyperglycemia-induced collagen synthesis and prevented glomerular fibrosis and renal insufficiency in db/db mice (Lehmann & Schleicher 2000, Ziyadeh et al. 2000). This study revealed that T3 significantly attenuated the increase in TGF-β1 expression in kidneys of db/db mice (Fig. 6). This effect may be mediated by an increase in PI3K activity because inhibition of PI3K abolished
the attenuating effect of T3 on the up-regulation of TGF-β1 in diabetic kidneys (Fig. 6). This finding further suggested that PI3K may be involved in the up-regulation of TGF-β1 in diabetic nephropathy. Further studies are required to determine whether PI3K regulates TGF-β1 via its direct action on TGF-β1 or through its effect on glycemia in diabetic kidneys. Therefore, the beneficial effects of T3 on diabetic nephropathy may be mediated, at least in part, by its attenuating effect on TGF-β1 expression. These findings also support a notion that thyroid hormone is an important regulator of renal structure in diabetic mice.

Although renal hypertrophy is found in the early stage of diabetes (Sharma et al. 2003), this study showed that kidney mass did not remain increased in diabetic mice at the age of 20 weeks (Fig. 1B). This finding is consistent with a report that renal hypertrophy disappears in db/db mice at the age of 21 and 25 weeks (Koya et al. 2000). The decrease in kidney mass is probably due to the loss of glomeruli and tubules and structural remodeling at a late stage of diabetes.

The db/db mice showed progressive renal functional damage as evidenced by a significant increase in urinary excretion of albumin (Fig. 1A), which is consistent with the published data (Ziyadeh et al. 2000, Cohen et al. 2001). Notably, treatments with T3 markedly ameliorated renal functional damage. Plasma T3 levels were significantly decreased in db/db mice (Lin & Sun 2010b). It was reported that female db/db mice aged 7 weeks have lower T4 levels in serum and lower T3 levels in the liver and brain (Kaplan & Young 1987). An epidemiological study indicated that type II diabetic patients with subclinical hypothyroidism are associated with an increased risk of diabetic nephropathy (Chen et al. 2007). The GFR in hypothyroid patients is approximately one-third lower than that of the euthyroid individuals (Singer 2001). Suher et al. (2005) showed that subjects with hypothyroidism are associated with increased urinary albumin excretion rate and decreased creatinine clearance. In corroboration with our findings, T3 was reported to have beneficial effects on renal function in a diabetic patient with end-stage renal failure and severe hypothyroidism (van Welsem & Lobatto 2007). Renal function was improved by T3 in 32 patients with hypothyroidism (den Hollander et al. 2005). Therefore, treatments with thyroid hormone may be an effective therapeutic approach for delaying progressive renal damage or even renal failure in diabetic patients with hypothyroidism or subclinical hypothyroidism.

Given that hyperglycemia may be involved in the development of diabetic nephropathy (Brosius 2008) and that T3 decreased blood glucose levels in db/db mice (Fig. 5A and B), the anti-hyperglycemic effect of T3 may contribute to its protective effects on diabetic nephropathy. On the other hand, strict glycemic control does not completely stop the progression of diabetic nephropathy in diabetic patients (Ruggenenti & Remuzzi 1998). It is noted that T3 restored the decreased activity of PI3K in diabetic kidneys (Fig. 5C). The present data suggest that T3 ameliorated diabetic nephropathy in db/db mice, probably by increasing PI3K activity because inhibition of PI3K activity abolished the beneficial effect of T3. Therefore, the beneficial effects of T3 on diabetic nephropathy may be mediated by the increased insulin synthesis and release, enhanced insulin signaling, and improved insulin resistance by T3 (Lin & Sun 2010b). The limitation of this study is that it cannot determine the relative importance of the increased renal PI3K activity and the attenuated hyperglycemia in the beneficial effect of T3 on diabetic nephropathy. A further study is needed to address this important question by controlling glucose levels in all groups.

The PI3K/AKT signaling was activated in renal cortex of db/db mice in the early phase of diabetes (at ages of 6–8 weeks) likely due to an increase in insulin receptor activities (Feliers et al. 2001). Activation of the PI3K/AKT signaling has been considered an important factor for diabetic renal hypertrophy, which contributes to diabetic nephropathy (Price 2007). On the other hand, the PI3K/AKT signaling was found to be lower in glomeruli of 12-week-old db/db mice, which has been linked to the death of podocytes of db/db mice (Tejada et al. 2008). Most recent data indicated that the down-regulation of PI3K/AKT signaling contributes...
to renal tubular apoptosis in kidneys of streptozocin-induced diabetic mice and apoptosis in renal proximal tubular cells (Rane et al. 2010). Apoptosis has been considered as an initiator of diabetic nephropathy (Kumar et al. 2004). Therefore, the PI3K/AKT signaling may play different roles at different stages of diabetic nephropathy.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement

Z. S conceived and designed the experiment and wrote the paper. Y. L designed and performed the experiments, analyzed the data, and wrote the paper.

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