

## SHORT REPORT

# CHRONIC KIDNEY DISEASE SCREENING AND RENOPROTECTION IN TYPE 2 DIABETES

<sup>1</sup>E. I. Agaba, <sup>2</sup>F. H. Puepet, <sup>2</sup>S. O. Ugoya, <sup>3</sup>P. A. Agaba, <sup>1</sup>R. Adabe,  
<sup>1</sup>M. Duguru and <sup>2</sup>A. I. Rowland

<sup>1</sup>Renal Unit, Department of Medicine, Jos University Teaching Hospital, Jos, Nigeria

<sup>2</sup>Endocrinology Unit, Department of Medicine, Jos University Teaching Hospital, Jos, Nigeria

<sup>3</sup>AIDS Preventive Initiative, Nigeria Center, Jos University Teaching Hospital, Jos, Nigeria

Reprint requests to: Dr. Emmanuel I. Agaba, Department of Medicine, Jos University Teaching Hospital,  
P.M.B. 2076, Jos, Nigeria E-mail: [eiagaba@unijos.edu](mailto:eiagaba@unijos.edu); [eiagaba@yahoo.com](mailto:eiagaba@yahoo.com)

Accepted: 29<sup>th</sup> August 2008

### Abstract

**Background:** Type 2 diabetes (T2D) is a major cause of chronic kidney disease. Control of hypertension and the use of angiotensin converting enzyme inhibitors (ACEI) and/or angiotensin II receptor blockers especially in those with proteinuria have been shown to protect against chronic kidney disease and delay its progression to kidney failure.

**Methods:** We reviewed the medical records of 169 patients at 12 months with a view of auditing the screening for chronic kidney disease and the use of renoprotective measures.

**Results:** Urinalysis was done in 49.1% and serum creatinine in 50.3%. No patient had glomerular filtration rate estimated. Seventy nine (67.6%) of the hypertensive patients were on anti-hypertensives. ACEI was used in 49 (45.8%) of these patients BP control was optimal in 29.1%.

**Conclusion:** There is poor adherence to guidelines on chronic kidney disease screening and renoprotection in T2D.

**Keywords:** Chronic kidney disease, diabetes, renoprotection

### Résumé

**Fond:** Le type 2 le diabète (T2D) est une cause importante de la maladie chronique de rein. La commande de l'hypertension et l'utilisation des inhibiteurs d'enzyme convertissants d'angiotensine (ACEI) et/ou des dresseurs de récepteur de l'angiotensine II particulièrement dans ceux avec le proteinuria ont été montrées pour se protéger contre la maladie chronique de rein et pour retarder sa progression à l'échec de rein.

**Méthodes:** Nous avons passé en revue les disques médicaux de 169 patients à 12 mois avec une vue d'auditer le criblage pour la maladie chronique de rein et l'utilisation des mesures renoprotective.

**Résultats:** L'analyse d'urine a été faite en 49.1% et créatinine de sérum dans 50.3%. Aucun patient n'a fait estimer le taux de filtrage glomérulaire. Soixante-dix-neuf (67.6%) des patients hypertendus étaient sur des antihypertensifs. ACEI a été employé dans 49 (45.8%) de ces patients que la commande de BP était optimale dans 29.1%.

**Conclusion:** Il y a d'adhérence pauvre aux directives sur le criblage et le renoprotection chroniques de la maladie de rein dans T2D.

**Mots-clés:** La maladie chronique de rein, diabète, renoprotection

Chronic kidney disease (CKD) is of pandemic proportion and a major cause of morbidity and mortality worldwide. Current clinical practice guidelines emphasize the need for prevention of end

stage renal disease (ESRD) largely by the screening of persons at increased risk of and early detection and treatment of CKD.<sup>1</sup> Urine examination for markers of kidney damage (like proteinuria and haematuria) and an estimation of glomerular filtration rate (GFR) by the Cock-Croft Gault (CG) formula<sup>2</sup> or the Modification of Diet in Renal Disease (MDRD) study derived equation<sup>3</sup> using serum creatinine form the mainstay of this screening.

Type 2 Diabetes (T2D) is a major cause of CKD.<sup>4, 5</sup> Once overt nephropathy occurs there is a relentless progression to ESRD. Control of hypertension and the use of Angiotensin converting enzyme inhibitors (ACEI) and/or Angiotensin II receptor blockers (ARB) especially in those with proteinuria have been shown to delay this progression.<sup>6-8</sup> This report describes screening for CKD and renoprotection in T2D in a teaching hospital in Nigeria. This audit would contribute immensely to fighting the global scourge of CKD.

## Materials and Methods

We analyzed records of T2D patients being followed up at the Diabetes Clinic of the Jos University Teaching Hospital (JUTH) in a cross-sectional study between June and September 2004. Assessment of records was done at 12 months of continued care at the clinic.

Clinical data analyzed included age, gender, duration of diabetes, documentation of urinalysis and estimation of GFR (eGFR) from serum creatinine measurement. Other variables of interest were treatment of hypertension and the use of ACEI/ARB.

### Statistical analysis

Continuous variables were reported as mean±standard deviation and categorical variables as proportions. The Chi-Squared statistic was used to compare proportions. A p value of less than 0.05 was considered significant.

## Results

### Study characteristics

Records of 169 consecutive type 2 diabetics (68 males and 101 females) were reviewed. One hundred and seven (63.3%) patients were hypertensive and 62 (36.7%) non-hypertensive. The mean age of the patients was 51±12 years. The duration of T2D ranged from one (1) to 27 years with a median of four (4) years.

### Screening for CKD

Eighty three patients (49.1%) had urinalysis done. Proteinuria was present 19.7% of those who had urinalysis carried out. Serum creatinine was assayed in 85 (50.3%) patients. None had eGFR documented.

Analysis of eGFR in those who had serum creatinine showed that eight percent had GFR less than 60ml/mi/1.73m<sup>2</sup>.

Urinalysis was done in 47.7% of those with hypertension and 52.3% of those without (OR 1.77, 95% CI 0.92- 3.40; p = .11). Similarly, serum creatinine was done in 51.4% of those with hypertension and 48.6% of those without (OR 1.67, 95% CI 0.88- 3.16; p = .11).

### Renoprotection

Seventy nine (67.6%) of the hypertensive patients were on treatment for hypertension and 28 (26.2%) were not. ACEI was used in 49 (45.8%) of these patients. BP control was satisfactory (<130/80 mm Hg) only in 23 (29.1%) of the patients on treatment for hypertension. Only two non-hypertensives were on ACEI therapy. No patient (hypertensive or not) was on ARB. The treatment of hypertension was ACEI based in four (19.0%) of the patients with proteinuria and four (16.7%) without (OR 1.17, 95% CI 0.25- 5.42).

## Discussion

### Main finding of the study

The main findings of our study were; 1) less than half of the patients with T2D are screened for CKD; 2) only two-thirds of T2D patients with hypertension are on hypertensive medication. In addition, only a third of those with hypertension had optimal BP control.

### Screening for CKD

An earlier study in Nigeria showed that only 16.9% of diabetic patients are screened for CKD.<sup>9</sup> Harzallah and colleagues<sup>10</sup> in a similar study of Tunisian diabetics demonstrated that only 19.8% had urinalysis done. The low screening rate for CKD is not only limited to the "diabetic" population as a recent survey revealed that only 26% of patients in primary care had their GFR estimated.<sup>11</sup> The National Kidney Foundation Kidney Disease Outcomes Quality Initiative (KDOQI) clinical practice guidelines recommend screening for CKD in at risk individuals using proteinuria and the eGFR.<sup>1</sup>

### Renoprotection

As cardinal as BP control is to the prevention of and slowing the progression of CKD, only a third of our patients had optimal BP control. A similar figure has been reported from Australia.<sup>12</sup> This corroborates previous reports. The National Health and Nutrition Examination Survey revealed that only 40% of US hypertensive diabetics achieved blood pressure of <130/80 mm Hg.<sup>13</sup> Lower figures (between 13% and 23%) have been reported in Europe.<sup>14-16</sup>

Though ACEIs was used in nearly half of the diabetics with hypertension in this study, the use of ACEIs was scanty in those with proteinuria (19%).

There was no patient on an ARB. Blockade of the Renin-Angiotensin-Aldosterone system has been shown to reduce proteinuria and retard the decline of GFR in CKD. A recent survey in the UK showed that 32% of diabetics with proteinuria at CKD stage 3 were on an ACEI and as much as 26% of diabetics with hypertension were not on any BP lowering medication regardless of the CKD stage.<sup>17</sup> In clinical practice renoprotection is sub-optimal in T2D.

Adherence to international guidelines is no doubt generally poor. The gap between guidelines and clinical practice needs to be bridged. Regular audit, continuing medical education, local adaptation of international guidelines and action checklists have been shown to improve the care of diabetics.<sup>15, 18, 19</sup>

In conclusion, this study demonstrates a low screening rate for CKD and an inadequate utilization of renoprotection in T2D. The burden of CKD, especially that due to T2D will remain unchecked unless measures are instituted to improve adherence to existing guidelines.

#### Acknowledgment

This data was presented at the 4th World Congress of Nephrology organized by the International Society of Nephrology, 21-25 April, 2007, Rio de Janeiro, Brazil.

#### References

1. National Kidney Foundation. K/DOQI Clinical Practice Guidelines for Chronic Kidney Disease: Evaluation, classification, and stratification. *Am J Kidney Dis.* 2002; 39:S1-S246.
2. Cockcroft DW, Gault MH. Prediction of creatinine clearance from serum creatinine. *Nephron.* 1976; 16:31-41.
3. Levey AS, Bosch JP, Lewis JB, et al. A more accurate method to estimate glomerular filtration rate from serum creatinine: A new prediction equation. Modification of Diet in Renal Disease Study Group. *Ann Intern Med.* 1999; 130:461-470.
4. United States Renal Data System. USRDS. The United States Renal Data System. *Am J Kidney Dis.* 2003; 42:1-23.
5. Ritz E, Rychlik I, Locatelli F, et al. End-stage renal failure in type 2 diabetes: a medical catastrophe of worldwide dimension. *Am J Kidney Dis.* 1999; 34:795-808.
6. Kasiske BL, Kalil RS, Ma JZ, et al. Effect of antihypertensive therapy on the kidney in patients with diabetes: A meta-regression analysis. *Ann Intern Med.* 1993; 118:129-138.
7. Marre M, Chatellier G, Leblanc H, et al. Prevention of diabetic nephropathy with enalapril in normotensive diabetics with microalbuminuria. *BMJ.* 1988; 297:1092-1095.
8. Remuzzi G, Perico N, Macia M, et al. The role of renin-angiotensin-aldosterone system in the progression of chronic kidney disease. *Kidney Int.* 2005 ;( Suppl 99):S57-S65.
9. Okoro EO, Adejumo AO, Oyejola BA. Diabetic care in Nigeria: report of a self-audit. *J Diabetes Complications.* 2002; 16:159-164.
10. Harzallah F, Kanoun F, Elhouch F, et al. Quality of ambulatory care of non-insulin dependent diabetic patients. *East Mediterr Health J.* 2006; 12:98-104.
11. de Lusignan S. Chronic kidney disease. *Br J Gen Pract.* 2006; 56:885.
12. Bryant W, Greenfield JR, Chisholm DJ, et al. Diabetes guidelines: easier to preach than to practice. *Med J Aust.* 2006; 185:305-309.
13. Resnick HE, Foster GL, Bardsley J, et al. Achievement of American Diabetes Association clinical practice recommendations among U.S. adults with diabetes, 1999-2002: the National Health and Nutrition Examination Survey. *Diabetes Care.* 2006; 29:531-537.
14. Eliasson B, Cederholm J, Nilsson P, et al. The gap between guidelines and reality: Type diabetes in a National Diabetes Register 1996-2003. *Diabet Med.* 2005; 22:1420-1426.
15. Madsbad S, Larsen ML, Adeler HF, et al. Implementation of clinical guidelines in general practice. The effect of journal audit and continuing education for the treatment of cardiovascular risk factors in patients with and without type 2 diabetes. *Ugeskr Laeger.* 2006; 168:1640-1645.
16. Mallion JM, Kahn JC, Poncelet P, et al. Differences between management guidelines and global health strategies for arterial hypertension with metabolic disorders in France in 1999. Ohara study. *Arch Mal Coeur Vaiss.* 2001; 94:834-838.
17. New JP, Middleton RJ, Klebe B, et al. Assessing the prevalence, monitoring and management of chronic kidney disease in patients with diabetes compared with those without diabetes in general practice. *Diabet Med.* 2007; 24:364-369.
18. Kirkman MS, Williams SR, Caffrey HH, et al. Impact of a program to improve adherence to diabetes guidelines by primary care physicians. *Diabetes Care.* 2002; 25:1946-1951.
19. Schectman JM, Schorling JB, Nadkarni MM, et al. The effect of physician feedback and an action checklist on diabetes care measures. *Am J Med Qual.* 2004; 19:207-213.