



PREVALANCE OF CHRONIC KIDNEY DISEASE STAGES 3-5 AMONG PATIENTS WITH TYPE 2 DIABETES MELLITUS: EXPERIENCE FROM A TERTIARY CARE HOSPITAL OF BANGLADESH

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Introduction and aims

Prevalence of diabetes mellitus (DM) is increasing and mostly contributed by type 2 diabetes mellitus (T2DM). DM is the most common cause of chronic kidney disease (CKD) and end-stage renal disease in developed as well as in developing countries [1-6]. This study was designed to evaluate the frequency of CKD stages 3-5 among selected group of T2DM subjects and to evaluate its relation with selected socio-demographic, clinical and laboratory parameters.

Methods

This cross-sectional study was done in Department of Nephrology, Bangladesh Institute of Research and Rehabilitation in Diabetes, Endocrine and Metabolic Disorders (BIRDEM) General Hospital, Dhaka, Bangladesh from July to December 2015. Adult male and non-pregnant female T2DM subjects were consecutively and purposively included in this study. Patients with acute kidney injury (AKI), AKI on CKD, non-diabetic kidney disease and on renal replacement therapy were excluded. Enrolled patients were evaluated clinically and by laboratory tests. Diagnosis of CKD and CKD stages were determined according to Kidney Disease: Improving Global Outcomes (KDIGO) Clinical Practice Guidelines (2012), using estimated glomerular filtration rate (eGFR) [7] which were calculated by Modification of Diet in Renal Disease (MDRD), Cockcroft-Gault (C-G) and Chronic Kidney Disease Epidemiology (CKD-EPI) creatinine based formula.

Results

Total patients were 400 including 231 females. Base-line characteristics of the study participants are shown in Table I.

Table I. Base-line characteristics of the study subjects (N=400)

| Characteristics | Value |
|-----------------------------------|-----------|
| Mean age (years) | 55.2±11.2 |
| Mean duration of diabetes (years) | 11.6±7.6 |
| Male: Female | 1:1.4 |
| Rural: Urban | 1:3.1 |
| Hypertension (%) | 82 |
| Dyslipidaemia (%) | 37.5 |
| Mean BMI (Kg/m ²) | 24.6±4.5 |
| Mean HbA1c (%) | 9.1±2.0 |

[BMI= body mass index; HbA1c=glycated haemoglobin]

Out of 400 cases, 254 (63.5%), 259 (64.75%) and 218 (54.5%) cases had CKD stages 3-5 according to MDRD, C-G and CKD-EPI GFR estimating equations respectively (Table II). There was high burden of other classic complications of diabetes among the study subjects (Table III).

Table II. Frequency of different stages of CKD according to different equations among the study subjects (N=400)

| CKD stages | MDRD n (%) | C-G n (%) | CKD-EPI n (%) |
|----------------------|-------------|-------------|---------------|
| Stage 3a (GFR 45-59) | 111 (27.75) | 87 (21.75) | 101 (25.25) |
| Stage 3b (GFR 30-44) | 77 (19.25) | 106 (24.50) | 54 (13.5) |
| Stage 4 (GFR 15-29) | 45 (11.25) | 51 (12.75) | 42 (10.5) |
| Stage 5 (GFR <15) | 21 (5.25) | 15 (3.75) | 21 (5.25) |

GFR was measured in ml/min/1.73 m²

[GFR=glomerular filtration rate; MDRD= Modification of Diet in Renal Disease; C-G=Cockcroft-Gault; CKD-EPI=Chronic Kidney Disease Epidemiology]

Table III. Microvascular and macrovascular complications among the study subjects (N=400)

| Complications | Frequency (%) |
|-----------------------------------|---------------|
| Microvascular | |
| Nephropathy/CKD* stages 3-5 | 218 (54.5) |
| Neuropathy | 119 (29.8) |
| Retinopathy | 151 (37.8) |
| Macrovascular | |
| Coronary artery disease | 103 (25.8) |
| Peripheral vascular disease | 56 (14) |
| Stroke/Transient ischaemic attack | 44 (11) |

[*CKD stages 3-5 as per CKD-EPI creatinine based formula]

CKD was more among female patients, with long duration of diabetes and hypertension (Table IV).

Table IV. Presence of CKD stages 3-5 in relation to different risk factors (N=400)

| Risk factors | Total number | CKD stages 3-5 n (%) | p value |
|--------------------------|--------------|----------------------|---------|
| Sex | Male | 71 (42.0) | 0.001 |
| | Female | 147 (63.6) | |
| Family history of DM | Present | 110 (53.9) | 0.810 |
| | Absent | 108 (51.1) | |
| Duration of DM | ≥ 5 years | 196 (59.6) | 0.001 |
| | < 5 years | 22 (30.0) | |
| Hypertension | Present | 186 (57.2) | 0.022 |
| | Absent | 32 (42.7) | |
| Dyslipidaemia | Present | 84 (55.6) | 0.726 |
| | Absent | 134 (53.8) | |
| BMI (Kg/m ²) | ≥ 25 | 94 (58.4) | 0.200 |
| | <25 | 124 (51.9) | |
| HbA1c (%) | ≥ 7 | 185 (52.1) | 0.007 |
| | < 7 | 33 (73.3) | |

Conclusions

More than half of the T2DM subjects had CKD stages 3-5 in this study. CKD was more in female T2DM subjects with long duration of diabetes and hypertension. Issues like glycaemic control and control of hypertension and weight management should be emphasized for a better renal and patient outcome.

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References

- Bailey RA, Wang Y, Zhu V, Rupnow MFT. Chronic kidney disease in US adults with type 2 diabetes: an updated national estimate of prevalence based on Kidney Disease: Improving Global Outcomes (KDIGO) staging. BMC Res Notes. 2014; 7: 415.
- The UK Renal Registry. The Sixth Annual Report 2003.
- ANZ Data Registry. The twenty-sixth report. Adelaide: Australia and New Zealand Dialysis and Transplant registry. 2003.
- Singh AK, Farag YMK, Mittal BV, Subramanian KK, Reddy SRK, Acharya VN et al. Epidemiology and risk factors of chronic kidney disease in India – results from the SEEK (Screening and Early Evaluation of Kidney Disease) study. BMC Nephrology. 2013; 14: 114.
- Ahmed ST, Rahim MA, Ali MZ, Iqbal MM. Prevalence of primary renal diseases among patients on maintenance haemodialysis: a hospital based study. KYAMC Journal. 2012; 2(2): 182-86.
- Fiseha T, Kassim M, Yemane T. Prevalence of chronic kidney disease and associated risk factors among diabetic patients in southern Ethiopia. Am J Health Res. 2014; 2(4): 216-21.
- Kidney Disease: Improving Global Outcomes (KDIGO) CKD Work Group. KDIGO 2012 Clinical Practice Guideline for the Evaluation and Management of Chronic Kidney Disease. Kidney Int. (Suppl.) 2013; 3: 1–150.

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