Hypogonadism and Erectile Dysfunction in Male Patients of Type 2 DM without CAD and With CAD in Indian Scenario

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ABSTRACT
AIM:- The present study is carried out to investigate hypogonadism using serum testosterone levels in male Type 2 diabetes mellitus (T2DM) subjects with and without coronary artery disease (CAD) and prevalence of erectile dysfunction in type 2 DM with and without CAD

MATERIAL AND METHODS: This was a cross-sectional study, conducted among 80 men (aged 30-70 years) with type 2 diabetes and 40 nondiabetic (aged 30-70 years) who visited medicine OPD of mahatma Gandhi hospital, Jaipur between Feb 2015 to March 2016. The patients’ demographic characteristics were collected using a pre structured questionnaire. Duration of diabetes, smoking habits, family history, drug history, trauma history, any tumour or radiation history or history of chronic liver and kidney disease were collected. Venous blood sample was collected to test for total testosterone (TT), free testosterone (FT), serum lipids, and glycosylated haemoglobin (HbA1c). In all men, the morning (08.00–10.00 hours) TT and FT levels were measured after an overnight fast. Normal levels of TT were taken as 300-1000 ng/dl and normal levels of FT as 9-40 pg/ml. Presence and degree of ED was assessed by the validated international index of erectile Function-5 (IIEF-5) questionnaire. Erectile dysfunction was considered present when the IIEF–5 score was ≤ 21.

RESULTS: 36.11 % of type 2 DM with CAD subjects had low total testosterone as compared to type 2 DM without CAD (22.72%) subjects. 61.11% type 2 DM with CAD subjects had low free testosterone than 54.54% type 2 DM without CAD subjects. Total testosterone (298.63±24.75) in patients with type 2 DM were significantly lower as compared to control (383.81±58.36) p<0.001. Free testosterone (7.61±2.12) in patients with type 2 DM were significantly lower as compared to control (11.17±2.34) p<0.001.

CONCLUSION: We observed hypogonadism as indicated by low testosterone levels in a significant proportion of male T2DM subjects with CAD. Erectile dysfunction was found to be more severe in DM2 with CAD. Thus erectile dysfunction can be a indicator of CAD in type 2 DM and correcting them early may prevent coronary artery disease.

KEYWORDS: Cardiovascular risk, coronary artery disease, diabetes, hypogonadism, testosterone.

INTRODUCTION
In recent years, low testosterone levels have been recognized to be closely associated with increased cardiovascular (CV) risk.¹,² Studies in the past have reported that testosterone deficiency is inversely associated with coronary intima-media
thickness (CIMT), a surrogate marker for atherosclerosis which leads to the development of coronary artery disease (CAD). Inverse relationships between the serum testosterone level and cardiovascular risk factors, such as obesity, hypertension, dyslipidemia and insulin resistance, have been observed. Cardiovascular disease is an important cause of morbidity and mortality in men with type 2 diabetes. In the population-based Osteoporotic Fractures in Men Study cohort from Sweden, men in the highest quartile of serum testosterone level had the lowest risk of cardiovascular events compared with men in the other three quartiles. When men with known CVD at baseline were excluded, the hazard remained. Similar observations had been reported in Asians. In a prospective cohort of 171 middle-aged Japanese men with coronary risk factors without a previous history of CVD, low serum total testosterone was associated with a significant fourfold higher risk of cardiovascular events when comparing men from the lowest testosterone tertile with those in the highest tertile (P < 0.01), independent of coronary risk factors and endothelial dysfunction. Decrease in testosterone levels is also influenced by age, obesity, and insulin resistance (IR). Subjects with diabetes have been reported to have low serum testosterone levels as compared with the healthy individuals. Subjects with diabetes also possess a greater CV risk. This study is done to determine the relationship of low testosterone and erectile dysfunction in DM2 with CAD and DM2 without CAD.

MATERIALS AND METHODS
The study was conducted in 80 male subjects with diabetes mellitus type 2 and 40 non diabetic subjects age and sex matched, attending Diabetic OPD and admitted in medicine ward Mahatma Gandhi Hospital between March 2015 and February 2016 with official permission and ethical clearance given by ethical committee of Mahatma Gandhi Medical College and Hospital, Jaipur, Rajasthan.

Patients with a known history of hypogonadism, panhypopituitarism, patients not willing for study, patients taking exogenous testosterone, patients with hyperthyroidism, or chronic debilitating disease, such as renal failure, cirrhosis, or HIV, were excluded from the study. The mean level of serum testosterone was calculated in various age and BMI groups and compared with controls. TT was measured by fully automated bi-directionally interfaced chemiluminescent immunoassay (CLIA) and FT was measured by radio immune assay (RIA), with an intra-assay coefficient of variation of 2.6% and an inter assay coefficient of variation of 4.3%. Normal levels of TT were taken as 300-1000 ng/dl and normal levels of FT as 9-40 pg/ml. Presence of diabetes mellitus was defined as per American Diabetes Association Criteria.

Presence and degree of ED was assessed by the validated international index of erectile Function-5 (IIEF-5) questionnaire. Erectile dysfunction was considered present when the IIEF–5 score was ≤ 21.

Statistical Method
For different qualitative parameters mean and standard deviation calculated. To compare the means between two groups, student unpaired ‘t’ test is used. Level of significance is taken as p<0.05. Chi square test is used to find the association between two qualitative variables. Statistical analysis
The data was coded and entered into Microsoft Excel spreadsheet. Analysis was done using SPSS version 20 (IBM SPSS Statistics Inc., Chicago, Illinois, USA) Windows software program. The variables were assessed for normality using the Kolmogorov Smirnov test. Descriptive statistics included computation of means and standard deviations. Level of significance was set at P≤0.05.
RESULTS

Table 1: Distribution According to Age

| Age in Years | Case N (%) | Control N (%) |
|--------------|------------|---------------|
| 30-40        | 15 (18.75%) | 10 (25%)      |
| 41-50        | 32 (40%)    | 15 (37.5%)    |
| 51-60        | 28 (35%)    | 12 (30%)      |
| >60          | 5 (6.25%)   | 3 (7.25%)     |
| Total        | 80          | 40            |

Table shows age distribution among case and control groups. Most of subjects were from 41-60 year age groups among both case (75%) and control (67.5%) groups.

Table 2 Distribution of case subject

| No. of Cases | Diabetes Mellitus Type2 With Coronary Artery Disease | Diabetes Mellitus Type2 Without Coronary Artery Disease |
|--------------|------------------------------------------------------|-------------------------------------------------------|
|              | 36                                                   | 44                                                    |

Table revealed only Diabetes Mellitus Type 2 cases in which 36 subjects were Diabetes Mellitus Type 2 with coronary artery disease and 44 were from Diabetes Mellitus Type 2 without coronary artery disease.

Table 3 General Characteristics of the Study Variables

| Variables                                      | Case          | Control        | P value |
|------------------------------------------------|---------------|----------------|---------|
| AGE (mean in years)                           | 48.32±8.25    | 47.02±9.54     | 0.44    |
| Body mass index (mean in kg/m^2)              | 27.11±4.603   | 27.07±3.608    | 0.96    |
| Systolic Blood Pressure (mean in mm Hg)       | 127.75±11.202 | 120.4±7.24     | <0.001 (S) |
| Diastolic Blood Pressure (mean in mm Hg)      | 80.25±5.63    | 78.6±3.62      | 0.09    |
| Dyslipidemia (HDL<40 mg/dl, TG>150mg/dl)      | n=72          | n=14           | 0.01 (S) |
| Smoking                                        | n=24          | n=8            | 0.01 (S) |
| Alcohol                                        | n=9           | n=4            | 0.22    |
| Family history                                 | n=40          | n=6            | 0.01 (S) |

Table provided general characteristics of the study variables. Systolic blood pressure (127.75, 120.4), dyslipidemia (72, 14), smoking (24, 8) and family history (40, 6) showed statistically significant results among subjects with type 2 DM and control groups respectively.

Table 4 Risk factor for CAD

| Risk factor                          | Case n (%) | Control n (%) | P value |
|--------------------------------------|------------|---------------|---------|
| Age (>45 years)                      | 46 (67.6)  | 22 (32.4)     | 0.79    |
| BODY MASS INDEX (>30kg/m^2)          | 26 (72.2)  | 10 (27.8)     | 0.39    |
| HYPERTENSION                         | 13 (16.2)  | 1 (2.5)       | 0.002 (S) |
| Smoking                              | 1 (100)    | 8 (88.9)      | 0.01 (S) |
| Alcohol                              | 9 (69.2)   | 4 (30.8)      | 0.01 (S) |
| Family history                       | 40 (87)    | 6 (13)        | <0.001 (S) |
| DYSLIPIDEMIA (HDL<40mg/dl, TG>150mg/dl) | 72(90)    | 14(35)        | 0.001 (S) |
Table reveals the risk factors for CAD among the study groups. Cases showed more risk with hypertension (16.2), alcohol (69.2), dyslipidemia (90) family history (87) as compared to control which is statistically significant. Age (67.6) and BMI (72.2) didn’t have effect on risk factors for CAD which provide statistically non-significant results.

Table 5 Relationship of DM Patients with and without CAD with BMI (kg/m²)

| BMI No. of participants (%) | P value |
|-----------------------------|---------|
| <18.5 | 18.5-24.9 | 25-29.9 | >30 |
| DM2 without CAD | 3 (3.75%) | 15 (18.75%) | 14 (17.5%) | 12 (15%) | 0.348 |
| DM2 with CAD | 1 (1.25%) | 7 (8.75%) | 14 (17.5%) | 14 (17.5%) |
| Total | 4 (5%) | 22 (27.5%) | 28 (35%) | 26 (32.5%) |

Table showed that percentage of obese (15%, 17.5%), overweight (17.5%, 17.5%) in DM2 with CAD and DM2 without CAD were almost similar.

Table 6 Comparison of Testosterone Profile

| Case | Control | P value |
|------|---------|---------|
| Mean± std Deviation | Mean± std Deviation | |
| Total Testosterone (ng/dl) | 298.63±24.75 | 383.81±58.36 | <0.001 (S) |
| Free Testosterone (pg/dl) | 7.61±2.12 | 11.17±2.34 | <0.001 (S) |
| International Index Erectile Function -5 Score | 14.62±5.39 | 23.15±3.70 | <0.001 (S) |

Table showed comparison of testosterone profile among case and control group. Total testosterone, free testosterone and International Index Erectile Function Score (IIEF-5) in case (298.63, 7.61, 14.62) and control (383.81, 11.17, 23.15) group respectively showed statistically significant results.

Table 7 Comparison of testosterone profile in DM2 with CAD, DM2 without CAD and nondiabetic controls

| | DM2 WITH CAD | DM2 WITHOUT CAD | NONDIABETIC (CONTROL) |
|-----------------------------|---------------|-----------------|----------------------|
| TOTAL TESTOSTERONE | 263 | 302 | 383.81 |
| FREE TESTOSTERONE | 7.07 | 7.65 | 11.17 |

Table shows that total and free testosterone were significantly lower in DM2 with CAD as compared to non diabetic and DM2 without CAD.

Table 7 Comparison of Study of Erectile Dysfunction in DM2 without CAD and DM2 with CAD among Cases

| | IIEF-5 Score |
|-----------------------------|-------------|-------------|-------------|-------------|-------------|
| | 22-25 | 17-21 | 12-16 | 8-11 | 5-7 |
| DM2 without CAD | 15 | 12 | 12 | 3 | 2 |
| DM2 with CAD | 0 | 1 | 14 | 14 | 7 |
| Total | 15 | 13 | 26 | 17 | 9 |

Table showed Comparison of study of erectile dysfunction in DM2 and DM2 with CAD among cases. Case of DM2 with CAD had more severe erectile dysfunction as compared to DM2 without CAD which had statistically significant results.
DISCUSSION

Distribution According to Age
It has been documented in the literature that serum testosterone levels decreased with age. After age of 30 years 1-2% of serum testosterone levels decrease with every year as a part of normal aging process. But few recent studies have shown that decrease in serum testosterone level with age is not normal (Guay et al, 2003). The mean age in the present study was 48.32±8.20 in case group and 47.02±9.42 in control group. Majority of the cases (40.1%) and controls (37.5%) were in age groups of 41-50 yrs. The mean age in the study by Koopman et al (2005) was 46.01±1.27yrs. According to Gale et al (2010), the average age of onset of diabetes mellitus type 2 in Indian population is 42.5 yrs. The present study was comparable to that seen in Koopman et al 2005 study (46.01 years).

Risk Factor of CAD and their Relationship with Diabetes Mellitus Type 2
In our study significant correlation with HTN (16.2%), alcohol (68.2%), dyslipidemia (90%), and family history of CAD and DM2 (87%) was seen with case as compared to control groups. Age and BMI shown no effect as risk factor for CAD which provide statistically non significant results. In our study, it was observed that total cardiovascular risk was less in patients without ED. Moreover, as severity of ED increased, total cardiovascular risk also increased. Various other workers have reported the correlation between ED, cardiovascular risk and other risk factors such as HT, dyslipidemias and obesity.

In 2002, Jackson et al, concluded that erectile dysfunction and cardiovascular disease share several risk factors that are similar and commonly coexist. ED in asymptomatic man may be a marker for underlying coronary artery disease.

In 2003, Roth et al, studied 1412 Israeli men and found that ED and cardiovascular disease share common risk factors and may be aggravated by medical treatment for reducing them. They concluded that ED is common among patients who are at high risk for cardiovascular disease because of diabetes and/or hypertension.

In 2003, Mota et al, studied 310 male diabetic patients and observed that ED showed a positive correlation with obesity, stroke, arteriopathy, retinopathy and smoking, but it was not correlated to type of diabetes mellitus, duration of diabetes <10 years, hypertension, IHD, nephropathy, dyslipidemia and alcohol consumption.

Correlation of Dyslipidemia and CAD and Erectile Dysfunction
Epidemiologic data also suggest that hypercholesterolemia and perhaps coronary atherosclerosis itself are risk factors for ischemic stroke. Specific indices of dyslipidemia are elevated levels of plasma triglyceride (>150 mg/dl), low levels of HDL-C (<40 mg/dl) and normal LDL-C levels but smaller and denser particles which increase their atherogenic potential. The present finding found that cholesterol and triglycerides were elevated in DM2 patients. Also there were a statistically significant differences in triglycerides and HDL-C among DM2 patients and controls (p<0.05). The observed dyslipidemia among DM2 patients may be one of the risk factors that increases incidence of ED among those patients.

Hyperlipidemia may impair erectile function by affecting endothelial and smooth muscle cells of the penis. It was clear from the study that Dyslipidemia is linked to ED. Epidemiological studies have shown that elevated serum cholesterol and reduced HDL-C levels are associated with an increased risk of ED. According to present findings, dyslipidemia was more prevalent among DM2 patients than controls. These results coincide with previous studies in that dyslipidemia is a risk factor of ED and is linked to it. In addition, ED was correlated with duration of DM2 and its complications (p<0.05), this may be attributed to the fact that endothelial dysfunction which characterizes diabetic patients is leading cause of ED among DM2 patients.
Distribution According to Serum Testosterone
The Mean serum total testosterone in DM2 with CAD and without CAD was (263,302) respectively and in control group was 383.81. The difference was statistically significant. The Mean serum free testosterone in DM2 with and without CAD was 7.07,7.65 and in control group was 11.17. In present study, 23 out of 80 type 2 diabetes mellitus patients (28.75%) and 8 out of 40 controls (25%) had low level of total testosterone in serum, whereas 46 out of 80 diabetes mellitus patients (57.5%) and 8 out of 40 controls (25%) had low levels of free testosterone. The difference was statistically significant.

In the study by Ernani et al28 (2005), low levels of serum total testosterone was found in patients with diabetes mellitus (34%) when compared to healthy control subjects (23%). In the study by Kapoor et al29 (2007), low levels of total testosterone was found in (25%) of patients with diabetes mellitus. In the study by Ernani et al28(2005) level of free testosterone was decreased in 46% of diabetics as compared to 24% in non-diabetics. In the study by Koopar et al29 (2007), 42% of patients with DM type 2 had low levels of serum free testosterone.

In the present study the low levels of serum total and free testosterone in DM2 patients and control group were comparable to other previous studies. A meta-analysis study showed that low testosterone levels are associated with increased CV risk and mortality.30 This analysis also showed that testosterone replacement therapy in subjects with hypogonadism moderates metabolic components associated with CV risk.30 Significant testosterone deficiency observed in diabetes subjects with CAD raises the important issue whether these subjects should be given testosterone replacement therapy in an attempt to reduce CV risk. This notion is also based on the findings of a recent retrospective study involving over 83,000 veterans.31 Results of this study showed that normalization of TT levels with testosterone replacement therapy leads to a significant lowering of all causes of mortality, MI risk, and stroke.31

Study of Erectile Dysfunction in Patient of DM2
In the present study, out of 80 diabetic male, the erectile dysfunction was present in 65 (81.25%) patients. Out of which, 14 (17.5%) had mild ED and 30 (37.5%) patients had moderate and (26.25%) severe ED. Erectile dysfunction was found to be more severe in DM2 with CAD than without CAD.

Schiavi et al32, studied 40 diabetic men, free from other illness or drugs that could affect sexual capacity and 40 age-matched healthy control subjects. ED was present in 77% of patients. Sundaram et al33, reported that in diabetic patients, the prevalence of ED was 66%. Ledda et al34, reported that ED was very common among diabetic patients.

CONCLUSION
The present study finds testosterone deficiency in a significant proportion of male T2DM subjects particularly those with evidence of CV disease. Low testosterone levels could contribute to a significantly higher cardiovascular risk in subjects with T2DM.

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