Spirometric Indices in Type 2 Diabetes Mellitus Patients in a Nigerian Tertiary Institution

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ABSTRACT

Background: Diabetes mellitus is a non-communicable disease of global health importance. It is a metabolic disorder caused by increased levels of blood glucose over a prolonged period of time. Type 2 diabetes mellitus (T2DM) is usually associated with obesity and insulin resistance. Several studies have also revealed that diabetes mellitus hampers pulmonary functions. This study was aimed at estimating the spirometric indices in type 2 diabetes mellitus patients.

Methods: A cross-sectional study of T2DM patients and apparently healthy control attending the medical outpatient clinic in a tertiary institution in south western Nigeria.

Results: A total of 146 participants with 73 patients with T2DM and 73 control groups. There were no significant differences in the age, body mass index, and gender distribution of the diabetics and control. However, patients with diabetes had higher SBP (133.2±20.17 mmHg vs 111.6±6.5 mmHg p<0.0001), and DBP (78.4±11.8mmHg vs 73.7±6.3 mmHg, p=0.003) when compared to the control. The mean FEV (1.98±0.5 vs 2.09±1.2, p=0.033), FVC (2.35±0.6 vs 2.53±1.3, p=0.045) and FEV/FVC ratio (83.61±7.2 vs 81.14±10.7, p=0.029) were significantly lower in diabetic patients when compared to matched controls. There was no significant difference in the PEF of both groups.

Conclusion: Type 2 diabetes mellitus patients had significant decrease in their spirometric indices, hence pulmonary function should be included in the periodic comprehensive diabetic check for holistic management.

Keywords: Pulmonary function, Spirometric indices, Diabetes Mellitus, Non-communicable diseases.

I. INTRODUCTION

Diabetes mellitus is a metabolic non-communicable disease, one of the most serious public health challenges worldwide [1], [2]. Globally, it was estimated that 451 million people developed diabetes mellitus in 2017, and this is expected to increase to 693 million by 2045 [3]. Diabetes mellitus is one of the non-communicable diseases (NCD) with global rising burden and it is associated with various microvascular and macrovascular complications which contribute to increased morbidity and mortality [4].

Type 2 Diabetes Mellitus (T2DM) comprises 90% of the global burden of diabetes mellitus, interfacing the increase burden of other NCD like obesity, hyperlipidemia, and systemic hypertension as well as communicable diseases like tuberculosis [5]. There is a global rise in the incidence of T2DM with 80% of people with DM living in low- and middle-income countries and an observed increase in the burden in the African population [2], [6]. Nigeria has the highest burden of diabetes mellitus in Africa, with every 1 in 17 adults having this disease [7], [8]. The multi-systemic effects of DM and the consequent microvascular damage affect virtually every organ by the glycation of connective tissue and collagen in most tissues [9]-[12]. The abundant microvascular circulation and connective tissue in the lungs raises the possibility of lung affection. The histopathologic evidence of lung involvement in subjects with DM has included thickened alveolar epithelial and pulmonary capillary basal laminae, the latter suggestive of existing pulmonary microangiopathy [13]. Studies have shown impaired lung function among patients with diabetes mellitus [14]-[17]. On the contrary, some investigators reported no decline in lung function among patients [18].

Despite these conflicting reports regarding diabetes mellitus and lung function, few studies have been conducted on this subject in Nigeria and none in our center. Thus, this study was conducted to assess the spirometric indices and their determinants in patients with type 2 diabetes mellitus attending the medical outpatient department of Ekiti State University Teaching Hospital (EKSUTH), Ado-Ekiti, Nigeria.

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II. METHODS

A. Study Setting

This study was conducted at the medical outpatients’ clinic of the Ekiti State University Teaching Hospital (EKSUTH), Ado-Ekiti, Nigeria. The hospital is a tertiary facility that serves both general and specialized healthcare needs of the inhabitants of Ekiti State and border communities of neighbouring states.

B. Study Design and Data Collection

A cross-sectional analytic study among 73 diabetic patients and 73 matched controls. The subjects who participated gave both verbal and written informed consent. The anthropometric measurement, spirometric indices, smoking history and blood pressures of all participants were measured. Biochemical investigations such as fasting plasma glucose and haemoglobin A1C (for patients with diabetes mellitus only) were determined.

C. Sample Size Considerations

The sample size of the study was 146 adult patients comprising 73 diabetes mellitus and 73 controlled group. This was determined using the select statistical services software formula for sample size in a comparative study and setting the study power at 80% [19]. To derive our sample size, we used prevalence data from a previously published local study on prevalence of abnormal spirometric indices in T2DM in Nigeria [20].

D. Anthropometry

Participants’ weight was measured with a digital weighing scale (OMRON 511, OMRON healthcare Co., Ltd. Japan) to the nearest 0.1 kg. Height was determined with a stadiometer to the nearest 0.1 m while standing without shoes, head gear or cap. Body mass index was taken as weight (kg)/height (m)^2. Body mass index was defined according to the WHO classification: <18.5 kg/m^2, underweight; 18.5 to 24.9 kg/m^2, normal; 25.0 to 29.9 kg/m^2, overweight; >30 kg/m^2, obese. [21].

E. Spirometric Indices

The spirometric indices (FEV1, FVC, FEV%, PEFR, and FEF25–75%) were measured using spirometer (spirolab®) in accordance with recommendations of the American Thoracic Society (ATS). The apparatus was calibrated for each procedure and three maneuvers were performed per subject. The maximum value of three trials was used for analysis. The American Thoracic Society/European Respiratory Society (ATS/ERS) guideline was used to categorize spirometry values into obstructive, restrictive, and mixed patterns. The percentage predicted value was calculated dividing observed value by predicted value and multiplying by 100 and this value was automatically inputted and calculated by the spirometer. [22]

F. Blood Pressure Measurement

The blood pressure was measured on the left upper arm with the participants in sitting position. Accussons® mercury sphygmomanometer attached to appropriate cuff sizes was used. The first and fifth Korokoff sounds were taken as the systolic and diastolic blood pressures respectively. Systemic hypertension was defined as persistent elevation of blood pressure ≥140/90 mmHg. [23]

G. Biochemical Analysis

After an overnight fast, plasma venous blood was obtained aseptically into fluoride oxalate bottle and plasma glucose was analysed by the glucose oxidase method. Glycosylated haemoglobin was analyzed on capillary blood using the Clover HbA1c analyser (Infopia Co Ltd., Korea) which utilizes a fully automated boronate affinity assay.

H. Statistical Analysis

Data were coded, entered, cleaned, and analyzed using Statistical Package for Social Science (SPSS) version 22. Descriptive statistics and independent samples t-test were conducted in the data analysis and interpretation. Categorical variables were represented as proportions and compared using chi-square while continuous variables were represented as mean± standard deviation and compared using t-test.

I. Ethical Consideration

Approval was obtained from the Ethical and Research committee of Ekiti State University Teaching Hospital, Ado-Ekiti. Verbal and written informed consent were also obtained from all participants and confidentiality was ensured.

III. RESULTS

In this study, 73 patients with DM and 73 matched controls were recruited. There is female preponderance in both the DM and control groups. The mean age of the diabetic patients and the matched controls were 69.9±9.8 and 59.2±12.8 years respectively with no significant statistical difference (p=0.14).

| TABLE I: DEMOGRAPHICS OF THE STUDY POPULATION |
|---------------------------------------------|
| Variables                | T2DM N (%) | Control N (%) |
| Gender                   |            |              |
| Male                     | 23 (31.5)  | 25 (34.2)    |
| Female                   | 50 (68.5)  | 48 (65.8)    |
| Age (years)              |            |              |
| 69.9±0.8                 | 1 (1.4)    | 5 (6.8)      |
| 59.2±12.8 (p=0.14)      | 40-60      | 32 (43.8)    | 39 (53.4) |
| Above 60                 | 40 (54.8)  | 29 (39.7)    |
| Smoking History          |            |              |
| Ex-smokers               | 13 (17.8)  | 6 (8.2)      |
| Never                    | 60 (82.2)  | 67 (91.8)    |

As presented in Table I, there was no significant difference in the smoking history of both groups. Majority of the subjects with diabetes mellitus (52.3%) were diagnosed between the period of less than 5 years while 3.4 % were diagnosed more than 10 years ago.

Table II shows that there was no significant difference in the anthropometric profile of the subjects. However, the results also indicated that T2DM patients had a significantly higher systolic (p < 0.001) and diastolic (p = 0.003) blood pressure when compared to that of the control.

The spirometric indices as presented in Table II showed significant reduction in forced expiratory volume (p=0.033), forced vital capacity (p=0.045) and forced expiratory ratio (p=0.029) compared to the control group. However, there was no significant difference in peak expiratory flow rate and the...
forced expiratory flow rate of the diabetic patients and their matched controls. The pattern of lung function amongst diabetic and non-diabetic patients was significantly different (p=0.0238).

**TABLE II: BIOCHEMICAL, ANTHROPOMETRIC AND SPIROMETRIC INDICES**

| Variables                        | Diabetic (mean±sd) | Non-diabetics (mean±sd) | P-value |
|----------------------------------|--------------------|-------------------------|---------|
| **Biochemical and Clinical profile** |                    |                         |         |
| SBP (mmHg)                       | 133.2 ± 20.17      | 111.6 ± 6.5             | < 0.0001** |
| DBP (mmHg)                       | 78.4 ± 11.8        | 73.7 ± 6.3              | 0.0033*  |
| **Anthropometric profile**       |                    |                         |         |
| Height (m)                       | 1.74 ± 0.17        | 1.74 ± 0.23             | 0.859   |
| Weight (kg)                      | 160.1 ± 13.2       | 158.4 ± 20.1            | 0.547   |
| BMI (kg/m²)                      | < 18.51            | 1 (1.4)                 | 0.96    |
|                                  | 18.5 – 24.9        | 22 (30.1)               |         |
|                                  | 25 – 29.9          | 26 (35.6)               |         |
|                                  | > 30               | 23 (31.5)               |         |
| **Pulmonary function tests**     |                    |                         |         |
| FEV1 (L)                         | 1.98 ± 0.5         | 2.09 ± 1.2              | 0.033** |
| FVC (L)                          | 2.35 ± 0.6         | 2.53 ± 1.3              | 0.045** |
| FEV1/FVC (%)                     | 83.61 ± 7.2        | 81.14 ± 10.7            | 0.029** |
| PEF (L)                          | 4.92 ± 1.8         | 4.88 ± 2.1              | 0.65    |
| FEF25-75 (L)                     | 4.64 ± 0.5         | 4.54 ± 0.5              | 0.965   |

Keys: BMI, body mass index; DBP, diastolic blood pressure; SBP, systolic blood pressure; FPG, fasting plasma glucose; HbA1c, glycosylated haemoglobin.

Significant p-values were asterisk (*).
In contrast, Sreeja et al., and Keerthi et al., reported obstructive lung impairment characterized by reduced FEV1/FVC ratio [36], [40]. Albeit the FEV1/FVC% is a more sensitive indicator of airway obstruction than FVC or FEV1 alone. The decrease in FEV1/FVC% in diabetes mellitus subjects may be related with the poor mechanical properties of the lung, like lung compliance and elastic recoil of lungs. Loss of elastic recoil leads to dynamic collapse of small airways during expiration [40]. Consistent with these observations, some authors demonstrated reduction in alveolar gas exchange among patients with T2DM [34]. Furthermore, insulin resistance which is typical of T2DM is also associated with decline in pulmonary function [16].

V. CONCLUSION

The lungs are prime targets for diabetes mellitus as pulmonary dysfunction may be one of the earliest and early measurable derangements in T2DM patients. Type 2 Diabetes Mellitus poses an underlying reduction in pulmonary functions alongside other known complications. It is advisable therefore, that diabetic mellitus patients should have periodic spirometry tests to assess the severity of lung function impairment.

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