

Effect of Anti-tubercular Drugs and Impact of Glycemic Status in Patients with Tuberculosis and Concomitant Diabetic- Tuberculosis Patients

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ABSTRACT

India is a key nation which is endemic and epidemic to tuberculosis and diabetes. An association between diabetes and TB is biological plausible, where diabetes impairs the immune system and making it harder for the body to fight against infection. **Methodology:** Patients on DOTS therapy in continuation phase of anti-TB therapy, Patients with TB and Diabetes and Patients with TB alone were included in the study. Blood samples were collected and Fasting Blood Glucose (FBG), Post Prandial Glucose (PPG) and Glycated Hemoglobin (HbA1c) were measured to understand the glycemic status and its impact on TB treatment outcomes. **Results:** Our study confirms the fact that type II diabetes is a strong risk factor for tuberculosis and is associated with a slower response to TB treatment and a higher mortality rate. Incidence of TB is greatest among those with conditions impairing immunity such as DM. Diabetes impairs the immune system, making it harder for the body to fight against infection.

Keywords: Tuberculosis, Glycemic status, Lung field, Diabetes Mellitus, Glycated Hemoglobin.

INTRODUCTION

The global burden of disease from diabetes and tuberculosis is immense.¹ India is a key nation which is endemic and epidemic to tuberculosis and diabetes.^{1,2} Tuberculosis is a major public health problem where India accounts for one fifth of the global burden of TB incident cases. Each year nearly 2 million people in India develop TB, of which around 0.87 million are infectious cases. It is estimated that annually around 3,30,000 Indians die due to TB.³

An association between diabetes and TB is biological plausible, for diabetes impairs the immune system, making it harder for the body to fight against infection.⁴ Clinicians have observed an association between DM and TB, although they were often unable to determine whether DM caused by TB or whether TB led to the clinical manifestations of DM. Multiple rigorous epidemiological studies, have demonstrated that DM is indeed positively associated with TB.

A causal link between DM and TB does not bode well for the future, as the global burden of DM is expected to rise from an estimated 180 million prevalent cases currently to a predicted 366 million by 2030. Experts have raised concerns about the merging epidemics of DM and TB. Prevalence of tuberculosis among diabetic is 2.5 times higher than in the non-diabetic population, and a higher rate of lower lung field involvement seen in diabetic patients.⁵ In 85% of the patients, tuberculosis had developed after the onset of diabetes. The association is more common among those who are above 40 years of age and males appear to be at a greater risk compared with females. Diabetes may also complicate the management of tuberculosis.⁶

TB is more common in patients with diabetes, especially in those with poor glucose control. Diabetes has been associated with increased risk of TB treat-

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ment failure or relapse, and diminished 2 month and 6 month culture conversion rates.

Hence this study was aimed to find out the effect of anti-TB drugs in patients with TB alone and TB with Diabetes groups.

AIM: To find out the effect of anti-TB drugs in patients with TB alone and TB with Diabetes groups.

MATERIAL AND METHODS

The protocol was accepted by Institutional Ethics Committee. It was a Prospective case control study conducted at a 1000 bedded tertiary care hospital at Tamil Nadu. Patients on DOTS therapy in continuation phase of anti-TB therapy. Patients with TB and Diabetes and Patients with TB alone were included in the study whereas patients who are HIV positive, pregnant and lactating women, abnormal renal functions and abnormal liver functions were excluded from our study. Patients underwent tests for Blood glucose and lipid profile. Blood samples were collected and Fasting Blood Glucose (FBG), Post Prandial Glucose (PPG) and Glycated Hemoglobin (HbA1c) were measured to understand the glycemic status and its impact on TB treatment outcomes. Similarly, Total Cholesterol, Triglycerides, HDL and LDL were also measured to find whether any variation in these among the test and control groups.

RESULT

In the study, no statistically significant difference was found in exposure of age, vital studies and lipid profile, while the statistically significant increase was seen in FBG, PPG and HbA1c in diabetic patients with TB than in patients with TB and without diabetes (Table 1).

The Table 2 explains the difference in glycemic status among the test and control groups. The TBDM group had shown an extremely significant increase in FBG (P value 0.0231)

From the Table 3, it is observed a slight variation in lipid profile among the TBDM and TB groups. Total cholesterol was slightly higher in TB group where the triglycerides, HDL and LDL values were greater in TBDM group. But no significant changes were found.

Considering the category of treatment taken, the patients with diabetes along with tuberculosis were found with a higher fasting and post prandial glucose levels in Cat. II patients than in Cat. I patients, where, HbA1c (glycated hemoglobin) values were

Table 1: Effect of anti-TB drugs in patients with TB alone and TB with Diabetes group.

	TBDM group	TB group	P value
Age (28-60)	49	46	0.2742
Karnofsky Performance Status Scale (60-90)	72.5	75	0.3093
Temperature (F)	98.6	98.6	0.1426
Pulse rate (bts/min)	74	74	0.3906
Systolic BP (mmHg)	123	125	0.2696
Diastolic BP (mmHg)	84	80	0.0447
FBG (mg/dl)	179	72	0.0231
PPG (mg/dl)	318	115	0.0009
HbA1c (%)	7.7	5.5	<0.0001
Total Cholesterol (mg/dl)	176	182	0.2616
Triglycerides (mg/dl)	164	158	0.1610
HDL (mg/dl)	47	44	0.2414
LDL (mg/dl)	109	104	0.2906

Table 2: Comparison of Glycemic status among the test (TBDM) and control (TB) group

	TBDM Mean \pm SD (SEM)	TB Mean \pm SD (SEM)
FBG (mg/dl)	179.38 \pm 138.03 (48.80)	71.750 \pm 17.393 (6.149)
PPG (mg/dl)	317.75 \pm 148.25 (52.415)	115.13 \pm 21.263 (7.518)
HbA1c (%)	7.738 \pm 0.5397 (0.1908)	5.475 \pm 0.2866 (0.1013)

(P value 0.0009) and HbA1c (P value <0.0001) levels than the TB alone group

Table 3: Comparison of Lipid profile among TBDM and TB groups

	TBDM Mean \pm SD (SEM)	TB Mean \pm SD (SEM)
Total Cholesterol (mg/dl)	175.88 \pm 23.333 (8.249)	181.88 \pm 11.269 (3.984)
Triglycerides (mg/dl)	164 \pm 8.106 (2.866)	157.75 \pm 15.192 (5.371)
HDL (mg/dl)	46.750 \pm 8.812 (3.115)	43.500 \pm 9.212 (3.257)
LDL (mg/dl)	108.75 \pm 16.202 (5.728)	103.75 \pm 19.092 (6.750)

found higher in Cat. I patients than in Cat. II patients. There was not much marked inter-individual variability was observed in patients with tuberculosis and without diabetes, based on category of treatment taken.

Table 4: Comparison of Glycemic status of patients in TBDM and TB group

	TBDM		TB	
	Median(Range)		Median(Range)	
	Cat. I	Cat. II	Cat. I	Cat. II
FBG (mg/dl)	124 (57-158)	323 (107-455)	67 (57-105)	65 (57-73)
PPG (mg/dl)	280 (166-338)	448 (218-611)	121 (105-136)	101.5 (71-132)
HbA1c (%)	8.1 (6.9-8.5)	7.4 (7.1-7.8)	5.65 (5.1-5.9)	5.3 (5.3-5.3)

DISCUSSION

Our study confirms the fact that type II diabetes is a strong risk factor for tuberculosis and is associated with a slower response to TB treatment and a higher mortality rate.^{7,8,9} Incidence of TB is greatest among those with conditions impairing immunity such as DM. Diabetes impairs the immune system, making it harder for the body to fight against infection.^{1,10}

Earlier studies shows that tuberculosis and diabetes mostly affects males than females, and also the patients with DM above 40 years are vulnerable to TB.^{8,9} In our study too males were found to be affected more than females and they were of age above 30 year old with DM along with TB.

In our study it was painful to note that all the patients recruited in the study were alcoholic and smokers at the onset of TB. Since they were advised to stop alcohol and smoking in order to make treatment effective they had stopped. The tobacco smoke and alcohol abuse are major risk factors which reduces the effectiveness of TB treatment and increases prevalence of TB.^{3,7,9}

Among the patients with DM in anti-TB therapy around 90% cases were pulmonary tuberculosis with 70% sputum positive cases. The studies done earlier had shown a high percentage of sputum positivity and a higher rate of pulmonary TB than extra pulmonary tuberculosis in diabetic patients.^{1,6,9,10} It is because glucose stimulates the mycobacterial growth⁵ and uncontrolled DM is responsible for poor clinical response to anti-TB therapy⁶ which also increases the susceptibility to mycobacterial growth.^{1,8}

The patients with diabetes along with TB will be having a lower health performance as these disease affects the host defenses and immunity of body, and thereby unable to resist against infections and other comorbidities.^{1,8}

CONCLUSION

India is a developing country which is endemic and epidemic to tuberculosis and diabetes. Our study also had shown the need to improve the care of patients with concomitant DM and TB. Enhanced medical vigilance, especially in patients with diabetes is required in tuberculosis population. It seems to be rational to screen for DM in patients with TB and vice versa, as these may have implications for control and treatment of both diseases. Therapeutic monitoring of anti-TB drugs along with regular monitoring of glucose levels using insulin can be recommended in patients with diabetes and tuberculosis.

Our study also recommends TB control program should consider targeting patients with diabetes for interventions such as active case finding and treatment of latent TB and conversely that effort to diagnose detect and treat diabetes may have a beneficial impact on TB control.

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CONFLICT OF INTEREST

Nil

ABBREVIATIONS USED

TB: tuberculosis; DM: Diabetes mellitus; TBDM: tuberculosis and Diabetes mellitus; DOTS: directly observed treatment short course; HIV: human immunodeficiency virus; FBG: fasting blood glucose; PPG: post prandial glucose; HDL: High density lipoprotein; LDL: low density lipoprotein; Hb A1c: Glycated hemoglobin.

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