

Assesment of Health-Related Quality of Life in Post-Tuberculosis Lung Disease Patients: A Cross Sectional Comparative Study

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Submitted: 04 Oct 2025; Accepted: 11 Nov 2025; Published: 22 Nov 2025

Citation: Odinaka, C.V., Onyedum, C.C., Udeh, C.F., Mbata, G.C., Eke, C.O.U., et al. (2025). Assesment of Health-Related Quality of Life in Post-Tuberculosis Lung Disease Patients: A Cross Sectional Comparative Study. *Med Clin Res*, 10(11), 01-10.

Abstract

Background: Tuberculosis is a disease of public health importance because of its high morbidity and mortality and also its communicable nature. Pulmonary tuberculosis is an airborne respiratory disease caused by *Mycobacterium tuberculosis*. It is one of the oldest diseases known to affect man and is a major cause of death globally. Patients treated for pulmonary tuberculosis and cured can come down with poor health-related quality of life and impaired lung function although this has not been extensively studied. This study assessed the quality of life of post-tuberculosis lung disease patients.

Aim: The study was done to assess health-related quality of life in post-tuberculosis lung disease patients.

Methods: This was a hospital based cross sectional comparative study of post-pulmonary tuberculosis patients attending Federal Teaching Hospital, Owerri, who had completed treatment for tuberculosis at the point of treatment completion down/up to 2 years previously. Interviewer based questionnaire was administered to study participant to obtain data regarding socio-demographics. Other relevant clinical data including health-related Quality of Life of participants were collected using St George's respiratory Questionnaire.

All participants were subjected to spirometry procedure to measure the ventilatory function parameters using spirolab III. Data was analyzed using IBM SPSS statistics version 25.0. Chi-square was used to determine association between categorical variables. The independent student t-test was used to compare continuous data while one-way ANOVA test was used to compare mean lung volumes of participants across the patterns of lung function with post hoc analysis conducted as appropriate. The Holmes-Bonferonni method was used to correct the analysis for multiple testing. Spearman rank correlation analysis was conducted between the lung volumes and median values of the symptom, impact and activity domains of the St George's Respiratory Questionnaire. A two-sided level of significance of 5% was used in all the analyses.

Results: Out of the 200 participants recruited, consisting of 100 post-tuberculosis lung disease patients (18 years and above) and 100 age-, sex- and height-matched apparently healthy controls; mean age: 48.0±15.2 for cases and 46.7±13.9 for controls. The most frequent respiratory symptom was cough and sputum production which was present in 56 (56%) and 51 (51%). Respiratory lung function impairment occurred in 71% of cases and 12% of the control. The SGRQ component score for symptoms was 17.1 (IQR 16.8-40.8) followed by impact 1.05 (IQR 0-30.26) and total components 7.48 (IQR 1.20-34.51) for post-TB individuals and none for controls, Also, the current health status were significantly lower in post-TB cases ($p = <0.001$) compared to controls.

Conclusion: Overall, health-related quality of life was significantly reduced across the symptom and impact domains in post-TB individuals compared to healthy controls, highlighting the significant impact of pulmonary tuberculosis on affected individuals even after successful treatment.

Keywords: Tuberculosis, Post-tuberculosis, Quality of life, Lung function

Abbreviations: TB: Tuberculosis; PTB: Pulmonary Tuberculosis; BMI: Body Mass Index; QOL: Quality of Life; HRQOL: Health-Related Quality of Life; FEV1: Forced Expiratory Volume in one Second; FVC: Forced Vital Capacity; CI: Confidence Interval; WHO: World Health Organisation

1. Introduction

Tuberculosis disease is spread when people who are infected with tuberculosis expel bacteria into the air by coughing, sneezing, talking, and singing [1]. In the immunocompetent host, the alveolar macrophages ingest the *M. tuberculosis* organisms; whether or not the macrophages destroy the bacteria depends on the degree to which they are activated, the host genetic factors and resistance mechanisms in the bacteria [2]. When innate macrophage microbicidal activity is inadequate to destroy the initial few bacteria of the droplet nucleus, they replicate doubling every 24 hours until the macrophage bursts to release its bacterial progeny. The bacilli spread from the initial lesion via lymphatic or circulatory systems to other parts of the body [2,3,4,6]. It is during this stage of infection that seeding to lung apices occur which is critical to the later development of pulmonary tuberculosis.

Pulmonary tuberculosis is an important risk factor for chronic respiratory diseases because of lasting lung damage [4,5,7]. Post-tuberculosis sequelae add substantially to the overall disease burden caused by tuberculosis and associated adverse patient outcomes including poor health related quality of life as well as increased lung function reduction after completion of treatment [10,12,13,15].

According to WHO (1999), Health-related quality of life (HRQoL) was defined as individuals' perception of their position in life in the context of the culture and value systems in which they live and in relation to their goals, expectations, standards and concerns [19, 20].

Health-related quality of life measures the physical, social, psychosocial and environmental functional status of an individual [20-22]. It is a multi-faceted concept that reflects how a person's health impacts their physical, mental, and social well-being and their ability to function in daily life [19,20]. Chronic diseases such as pulmonary tuberculosis have a strong influence on both physical health and quality of life which collectively make up the health-related quality of life [23,25,26]. Pulmonary tuberculosis is a challenging public health problem associated with poor health related quality of life [12,19,31]. A patient's quality of life is an important measure of effective treatment particularly for PTB [28,29,32,34].

Irreparable lung damage visible as scarring, fibrosis, cavitation, or other types of damage on radiological imaging may occur following pulmonary tuberculosis which may impact on health-related quality of life [13,37,38]. Data on the quality of life of post-TB lung disease patients are limited. This study evaluated patients who have completed treatment for pulmonary tuberculosis

for assessment of health-related quality of life with a view to identifying individuals who might require further respiratory function evaluation as part of their post treatment monitoring program.

2. Methods

2.1 Study Design

This was a hospital based cross sectional comparative study that spanned over a period of 7 months that assessed the health-related quality of life of patients treated and cured for pulmonary tuberculosis compared with apparently healthy control who have not been previously treated for PTB. Socio-demographic information included age, gender, marital status, occupation, tribe, level of education and religion. Detailed history was obtained from the participants by the researcher about participants' medical history, and quality of life using St Georges' respiratory questionnaire. Physical examination was carried out by the researcher including general and systemic examinations.

2.2 Inclusion Criteria

For the post-TB patients

1. Patients who gave written informed consent
2. Patients aged 18 years and above
3. Patients who had completed TB treatment and certified cured at six months of treatment completion up to two years.

For the controls

1. Persons who gave written informed consent
2. Persons with no historical evidence of lung disease
3. Persons aged 18 years and above who had not been previously treated for TB

2.3 Exclusion Criteria (For Both Cases and Controls)

1. Persons with active pulmonary tuberculosis
2. Persons with massive haemoptysis and those whose spirometry results fail to meet acceptability and repeatability criteria
3. Individuals with recent myocardial infarction, recent thoracic, abdominal or eye surgery, spine or chest deformities eg kyphoscoliosis, pectus deformities
4. Individuals with pre-existing bronchial asthma, chronic obstructive lung disease, interstitial lung disease, congestive cardiac failure, stroke or neuromuscular disease
5. Individuals with elevated blood pressure $\geq 180/100$ mmHg and those on long term treatment with drugs eg Bleomycin, Methotrexate, Amiodarone.

2.4 Data Analysis

Data were analysed using the IBM Statistical Package for Social Science (SPSS) version 25.0. Results were expressed as mean \pm standard deviation at 95% confidence interval (CI).

Categorical variables (such as gender, symptoms, etc) were expressed as percentages. The independent Student t-test was used to compare continuous data while one-way ANOVA test was used to compare mean lung volumes of participants across the patterns of lung function with post hoc analysis conducted as

appropriate. The Holmes-Bonferonni method was used to correct the analysis for multiple testing. Spearman rank correlation analysis was conducted between the lung volumes and median values of the symptom, impact and activity domains of the St George's Respiratory Questionnaire. A P value of less than 0.05 was considered statistically significant.

3. Results

One hundred post-TB individuals (54 females and 46 males) and 100 controls (56 females and 44 males) were recruited for this

study. There was no statistically significant difference in mean age for cases (48.0±15.2 years) relative to the controls (46.7±13.9 years, $t=0.660$; $p=0.510$). Of the 100 cases, 9 (9%) had no formal education, 39 (39%) had only primary education, 36(36%) had secondary education while 16 (16%) had up to tertiary education. This was statistically significant ($p<0.001$) when compared with that of controls. The predominant comorbidity was HIV in the cases 19 (19%) compared to the control group 6 (6%) and this was statistically significant ($p=0.005$) (Table 1). Thirty-nine (39%) of the cases had been previously treated for tuberculosis.

Table 1: Distribution of sociodemographic characteristics of study participants

	Cases n = 100, (%)	Control n = 100, (%)	Chi-square (<i>p-value</i>)
Gender			
Female	54 (54.0)	56 (56.0)	0.08 (0.776)
Male	46 (46.0)	44 (44.0)	
Mean age ± SD (years)	48.0 ±15.2	46.7 ± 13.9	0.660 (0.510) [†]
Age groups (in years)			
<25	8 (8.0)	6 (6.0)	1.97 (0.894)
25 – 34	11 (11.0)	14 (14.0)	
35 – 44	18 (18.0)	24 (24.0)	
45 – 54	31 (31.0)	28 (28.0)	
55 – 64	18 (18.0)	15 (15.0)	
≥65	14 (14.0)	13 (13.0)	
Education			
No formal education	9 (9.0)	0 (0)	39.54 (<0.001)
Primary	39 (39.0)	25 (25.0)	
Secondary	36 (36.0)	19 (19.0)	
Tertiary	16 (16.0)	56 (56.0)	
Marital status			
Single	24 (24.0)	25 (25.0)	3.81 (0.432)
Married	47 (47.0)	49 (49.0)	
Divorced	5 (5.0)	1 (1.0)	
Separated	1 (1.0)	0 (0)	
Widowed	23 (23.0)	25 (25.0)	
Smoking	8 (8.0)	2 (2.0)	3.79 (0.052)
Alcohol	16 (16.0)	0 (0)	17.39 (<0.001)
Comorbidities			
Diabetes	1 (1.0)	1 (1.0)	0 (1.000)
Hypertension	5 (5.0)	1 (1.0)	2.75 (0.097)
HIV	19 (19.0)	6 (6.0)	7.73 (0.005)

[†] independent Student-t test

Those who had been previously treated for tuberculosis more than once had significant cough ($p=0.003$), sputum production ($p=0.004$), breathlessness ($p=0.013$), chest pain ($p=0.009$) and haemoptysis ($p=0.034$) as shown in table 2.

Table 2: Association of sociodemographic characteristics with respiratory symptoms in post TB study participants

Symptoms	Cough n = 56, % (p-value)	χ^2	Sputum n = 51, % (p-value)	χ^2	Breathlessness n=34, % (p-value)	χ^2	Chest pain n=31, % (p-value)	χ^2	Wheezing n=13, % (p-value)	χ^2	Haemoptysis n=10, % (p-value)	χ^2
Gender												
Male	29 (51.8)	1.72 (0.190)	25 (49.0)	0.38 (0.537)	20 (58.8)	3.41 (0.065)	14 (45.2)	0.01 (0.910)	4 (30.8)	1.40 (0.237)	7 (70.0)	2.58 (0.108)
Female	27 (48.2)		26 (51.0)		14 (41.2)		17 (54.8)		9 (69.2)		3 (30.0)	
Odds Ratio (95% CI)	1.71 (0.77 – 3.80)		1.28 (0.58 – 2.82)		2.20 (0.95 – 5.11)		0.95 (0.41 – 2.23)		0.48 (0.14 – 1.66)		3.05 (0.74 – 12.57)	
Age group												
≤48 years	26 (46.4)	1.58 (0.208)	24 (47.1)	1.02 (0.313)	16 (47.1)	0.50 (0.478)	14 (45.2)	0.84 (0.359)	7 (53.8)	0.02 (0.886)	2 (20.0)	4.56 (0.033)
>48 years	30 (53.6)		27 (52.9)		18 (52.9)		17 (54.8)		6 (46.2)		8 (80.0)	
Odds Ratio (95% CI)	0.60 (0.27 – 1.33)		0.67 (0.30 – 1.47)		0.74 (0.32 – 1.70)		0.67 (0.29 – 1.57)		1.09 (0.34 – 3.51)		0.20 (0.04 – 0.99)	
Level of education												
Up to primary	32 (57.1)	4.26 (0.039)	27 (52.9)	1.02 (0.313)	22 (64.7)	5.76 (0.016)	20 (64.5)	4.91 (0.027)	7 (53.8)	0.21 (0.651)	5 (50.0)	0.02 (0.894)
At least secondary	24 (42.9)		24 (47.1)		12 (35.3)		11 (35.5)		6 (46.2)		5 (50.0)	
Odds Ratio (95% CI)	2.33 (1.04 – 5.24)		1.50 (0.68 – 3.30)		2.82 (1.19 – 6.67)		2.66 (1.11 – 6.41)		1.31 (0.41 – 4.22)		1.09 (0.30 – 4.03)	
Marital status												
Currently married	22 (39.3)	3.04 (0.081)	18 (35.3)	5.73 (0.017)	16 (47.1)	0.00 (0.993)	14 (45.2)	0.06 (0.805)	4 (30.8)	1.58 (0.209)	3 (30.0)	1.29 (0.256)
Not currently married	34 (60.7)		33 (64.7)		18 (52.9)		17 (54.8)		9 (69.2)		7 (70.0)	
Odds Ratio (95% CI)	0.49 (0.22 – 1.10)		0.38 (0.17 – 0.84)		1.00 (0.44 – 2.30)		0.90 (0.38 – 2.11)		0.45 (0.13 – 1.59)		0.45 (0.11 – 1.84)	
Body Mass Index												
Not overweight/obese	43 (76.8)	0.22 (0.642)	41 (80.4)	1.61 (0.204)	30 (88.2)	4.81 (0.028)	24 (77.4)	0.14 (0.708)	10 (76.9)	0.03 (0.864)	6 (60.0)	1.33 (0.248)
Overweight/obese	13 (23.2)		10 (19.6)		4 (11.8)		7 (22.6)		3 (23.1)		4 (40.0)	
Odds Ratio (95% CI)	1.24 (0.50 – 3.08)		1.81 (0.72 – 4.55)		3.50 (1.09 – 11.24)		1.21 (0.45 – 3.29)		1.13 (0.28 – 4.84)		0.46 (0.12 – 1.77)	
Smoking												
Yes	6 (10.7)	1.27 (0.259)	4 (7.8)	0.00 (0.953)	2 (5.9)	0.31 (0.575)	4 (12.9)	1.47 (0.226)	1 (7.7)	0.00 (0.965)	1 (10.0)	0.06 (0.806)
No	50 (89.3)		47 (92.2)		32 (94.1)		27 (87.1)		12 (92.3)		9 (90.0)	
Odds Ratio (95% CI)	2.52 (0.48 – 13.15)		0.96 (0.23 – 4.06)		0.63 (0.12 – 3.28)		2.41 (0.56 – 10.33)		0.95 (0.11 – 8.44)		1.32 (0.15 – 11.95)	
Alcohol												
Yes	13 (23.2)	4.93 (0.026)	11 (21.6)	2.40 (0.121)	7 (20.6)	0.81 (0.369)	5 (16.1)	0.00 (0.981)	3 (23.1)	0.56 (0.456)	2 (20.0)	0.13 (0.716)
No	43 (76.8)		40 (78.4)		27 (79.4)		26 (83.9)		10 (76.9)		8 (80.0)	
Odds Ratio (95% CI)	4.13 (1.10 – 15.56)		2.42 (0.77 – 7.57)		1.64 (0.55 – 4.88)		1.01 (0.32 – 3.22)		1.71 (0.41 – 7.05)		1.36 (0.26 – 7.07)	
Comorbidities												
Present	11 (19.6)	1.33 (0.250)	10 (19.6)	1.10 (0.294)	7 (20.6)	0.33 (0.566)	9 (29.0)	0.62 (0.430)	1 (7.7)	2.18 (0.140)	2 (20.0)	0.10 (0.755)
Absent	45 (56)		41 (80.4)		27 (79.4)		22 (71.0)		12 (92.3)		8 (80.0)	
Odds Ratio (95% CI)	0.58 (0.23 – 1.47)		0.61 (0.24 – 1.54)		0.75 (0.28 – 2.03)		1.47 (0.56 – 3.86)		0.23 (0.03 – 1.88)		0.77 (0.15 – 3.91)	
Previous tuberculosis treatment												
Yes	29 (51.8)	8.75 (0.003)	27 (52.9)	8.50 (0.004)	19 (55.9)	6.17 (0.013)	18 (58.1)	6.86 (0.009)	6 (46.2)	0.32 (0.571)	7 (70.0)	4.49 (0.034)
No	27 (48.2)		24 (47.1)		15 (44.1)		13 (41.9)		7 (53.8)		3 (30.0)	
Odds Ratio (95% CI)	3.65 (1.52 – 8.79)		3.47 (1.48 – 8.13)		2.91 (1.24 – 6.86)		3.17 (1.32 – 7.62)		1.40 (0.43 – 4.53)		4.23 (1.02 – 17.49)	

χ^2 – Chi-square; CI – Confidence interval

The prevalence of lung function impairment was 71% among the post-TB cases compared to 12% in the controls; as shown in figure 1.

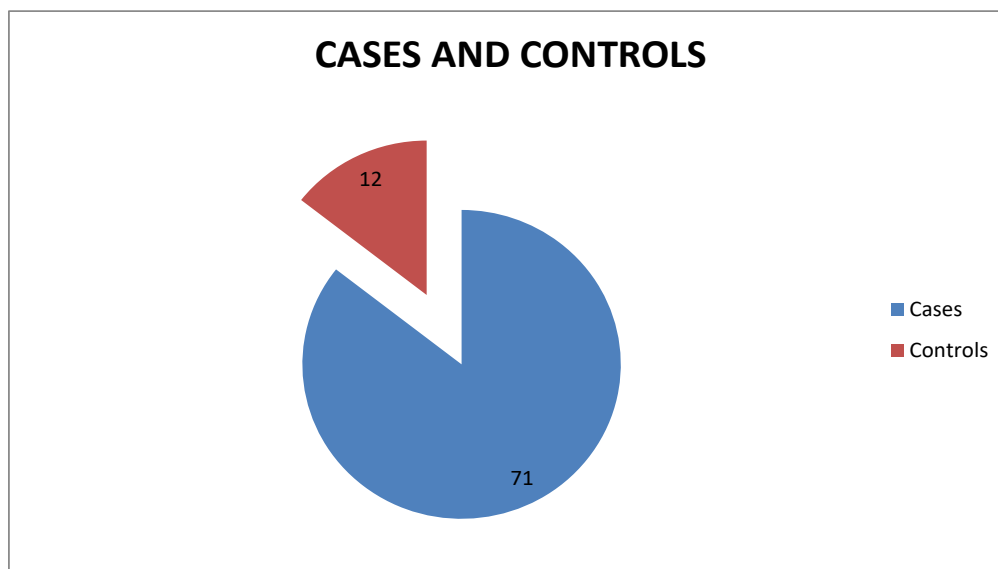


Figure 1: Prevalence of abnormal spirometry in study participants

The mean FEV₁ among the post-TB cases was 1.93 ± 0.73L (median z-score -2.06) comparatively lower to that of the control group with mean FEV₁ of 2.40 ± 0.64L (median z-score -0.7) and the difference was statistically significant (t -4.79, p<0.001). Similarly, the mean FVC among the post-TB cases was 2.72 ± 0.87L (median z-score -2.27) which was lower compared to the control group with mean FVC of 3.12 ± 0.88L (median z-score -0.31) and the difference was statistically significant (t -3.22,

p=0.001). Also the mean FEV₁/FVC of 69 ± 12.4 among the post-TB cases (t -5.62, p<0.001), mean FEF₂₅₋₇₅ of 1.57 ± 0.83 (t -4.67, p<0.001) and mean PEF_R of 4.66 ± 1.92 (t -5.34, p<0.001) were statistically significantly lower in the post-TB cases compared to the control group (Table 3). Obstructive ventilatory pattern was the predominant pattern followed by restrictive and mixed patterns (Table 4).

Table 3: Lung function indices of study participants with their z-scores

	Cases n = 100 mean ± SD	Control n = 100 mean ± SD	Student t (p-value)
FEV ₁ (litres)	1.93 ± 0.73	2.40 ± 0.64	-4.79 (<0.001)
FEV ₁ z-score	-2.06 (-2.80 – -0.55) [†]	-0.71 (-0.96 – -0.27) [†]	2769 (<0.001) [‡]
FVC (litres)	2.72 ± 0.87	3.12 ± 0.88	-3.22 (0.001)
FVC z-score	-2.27 (-2.30 – -0.24) [†]	-0.31 (-0.84 – -0.26) [†]	3417 (<0.001) [‡]
FEV ₁ /FVC (%)	69.5 ± 12.4	77.7 ± 7.9	-5.62 (<0.001)
FEV ₁ /FVC z-score	-1.38 (-2.33 – -0.94) [†]	-1.34 (-1.38 – -1.21) [†]	4137 (0.035) [‡]
FEF ₂₅₋₇₅ (l/sec)	1.57 ± 0.83	2.09 ± 0.75	-4.67 (<0.001)
PEFR (l/sec)	4.66 ± 1.92	6.16 ± 2.04	-5.34 (<0.001)

[†] median (interquartile ranges), [‡] Mann-Whitney U test

Table 4: Distribution of lung function abnormalities in study participants

Lung Function Pattern	Cases (n, %)	Control (n, %)	Chi-square (p-value)
Normal	29 (29.0)	88 (88.0)	73.10 (<0.001)
Obstructive	29 (29.0)	8 (8.0)	
Restrictive	28 (28.0)	3 (3.0)	
Mixed	14 (14.0)	1 (1.0)	
Total	100 (100.0)	100 (100.0)	

The FEV₁ (p<0.001), FVC (p=0.002), FEF₂₅₋₇₅ (p<0.001) and FEV₁/FVC (p<0.001) of the post-TB individuals were significantly reduced across the patterns of lung function impairment as shown in Table 5.

On the average, the mean FEV₁ values were about 720ml lower in those with restrictive lung pattern compared to those with a normal pattern (95% CI: 250 to 1,180ml) and 990ml (95%CI: 430 to 1,560ml) lower in those with a mixed pattern compared to the normal. The difference between the mean FEV₁ in those with a normal pattern and those with an obstructive pattern was no longer significant after adjusting for multiple testing by the Bonferroni method (p=0.112). There was no significant difference in mean FEV₁ values between those with obstructive, restrictive and mixed patterns (Table 6).

The mean FVC was 770ml higher (95%CI: 180 to 1340ml) in the normal pattern than in those with a restrictive pattern and 710ml higher (95%CI: 10 to 1,430ml) than in those with a mixed

pattern. There was no significant difference in the mean FVC values between those with a normal and an obstructive pattern, or between those with obstructive, restrictive and mixed patterns (p>0.05) (Table 6).

Similarly, the mean FEF₂₅₋₇₅ was significant only between those with a normal lung function pattern and those with obstructive, restrictive and mixed patterns; there were no statistically significant differences in the mean FEF₂₅₋₇₅ between those with obstructive, restrictive or mixed patterns. Those with a normal pattern had a higher mean FEF₂₅₋₇₅ value compared to those with an obstructive pattern (0.67l/s, 95%CI: 0.13 to 1.21l/s), those with a restrictive pattern (0.68l/s, 95%CI: 0.16 to 1.21l/s) and those with a mixed pattern (1.09l/s, 95%CI: 0.44 to 1.73l/s) (Table 6).

The mean FEV₁/FVC of the cases differed between those with normal and obstructive and mixed patterns, but not in those with normal and restrictive patterns. Also, the mean FEV₁/FVC values of the cases were different between those with obstructive and

restrictive patterns and between those with restrictive and mixed patterns. On the average, the mean FEV₁/FVC ratio was 14.8 percentage points (95%CI: 8.6 to 21.0%) higher in those with normal patterns than in those with obstructive patterns and 21.6 percentage points (95%CI: 13.8 to 29.3%) higher than in those with mixed patterns. While it was lower in those with a normal pattern compared to those with a restrictive pattern, this was not statistically significant (-0.6%, 95%CI: -6.9 to 5.6%). For those

with a restrictive pattern, the mean ratio was also higher than in those with an obstructive pattern by 15.4 percentage points (95%CI: 9.1 to 21.7%) as well as in those with a mixed pattern by 22.2 percentage points (95%CI: 14.4 to 29.9%) (Table 6). The Lung function patterns showed significant association in those with primary level of education (p=0.043) and underweight individuals (p=0.008) (table 7) while participants above 55 years of age had more of obstructive ventilatory pattern as shown in figure 3.

Table 5: Comparison of lung volumes of post-TB cases by pattern of lung function impairment

Pattern of lung function	FEV ₁ (litres) mean±SD	FVC (litres) mean±SD	FEF ₂₅₋₇₅ (l/sec) mean±SD	FEV ₁ /FVC (%) mean±SD
Normal (n = 29)	2.42 ± 0.83	3.10 ± 0.81	2.11 ± 0.92	76.3 ± 9.1
Obstructive (n = 29)	1.93 ± 0.49	2.89 ± 0.76	1.44 ± 0.78	61.8 ± 6.1
Restrictive (n = 28)	1.71 ± 0.62	2.32 ± 0.93	1.42 ± 0.54	77.3 ± 11.1
Mixed (n = 14)	1.43 ± 0.57	2.39 ± 0.74	1.02 ± 0.70	55.1 ± 7.6
F	9.462	5.356	7.965	33.54
p-value	<0.001	0.002	<0.001	<0.001

F: One-way analysis of variance (ANOVA) test

Table 6: Post-hoc analysis of lung volumes of cases across patterns of lung function impairment

	FEV ₁ (l) Mean difference (95% CI)	FVC (l) Mean difference (95% CI)	FEF ₂₅₋₇₅ (l/s) Mean difference (95% CI)	FEV ₁ /FVC (%) Mean difference (95% CI)
Normal				
Obstructive	0.49 (0.03 to 0.95)	0.20 (-0.38 to 0.79)	<i>0.67 (0.13 to 1.21)</i>	<i>14.8 (8.6 to 21.0)</i>
Restrictive	<i>0.72 (0.25 to 1.18)</i>	<i>0.77 (0.18 to 1.34)</i>	<i>0.68 (0.14 to 1.22)</i>	-0.6 (-6.9 to 5.6)
Mixed	<i>0.99 (0.43 to 1.56)</i>	0.71 (-0.01 to 1.42)	<i>1.08 (0.42 to 1.75)</i>	<i>21.6 (13.8 to 29.3)</i>
Obstructive				
Normal	-0.49 (-0.95 to -0.03)	-0.20 (-0.79 to 0.38)	<i>-0.67 (-1.21 to -0.13)</i>	<i>-14.8 (-21.0 to -8.6)</i>
Restrictive	0.22 (-0.24 to 0.69)	0.57 (-0.02 to 1.15)	0.01 (-0.53 to 0.53)	<i>-15.4 (-21.7 to -9.1)</i>
Mixed	0.50 (-0.07 to 1.07)	0.50 (-0.22 to 1.23)	0.41 (-0.25 to 1.07)	6.8 (-0.9 to 14.5)
Restrictive				
Normal	-0.72 (-1.18 to -0.25)	<i>-0.77 (-1.36 to -0.18)</i>	<i>-0.68 (-1.22 to -0.14)</i>	0.6 (-5.6 to 6.9)
Obstructive	-0.22 (-0.69 to 0.24)	-0.57 (-1.15 to 0.02)	-0.01 (-0.55 to 0.53)	15.4 (9.1 to 21.7)
Mixed	0.28 (-0.29 to 0.85)	-0.06 (-0.79 to 0.66)	0.40 (-0.25 to 1.07)	<i>22.2 (14.4 to 29.9)</i>
Mixed				
Normal	<i>-0.99 (-1.56 to -0.43)</i>	<i>-0.71 (-1.42 to -0.01)</i>	<i>-1.08 (-1.75 to -0.42)</i>	<i>-21.6 (-29.3 to -13.8)</i>
Obstructive	-0.50 (-1.07 to 0.07)	-0.50 (-1.23 to 0.22)	-0.41 (-1.08 to 0.25)	-6.8 (-14.4 to 0.9)
Restrictive	-0.28 (-0.85 to 0.29)	0.06 (-0.66 to 0.79)	-0.40 (-1.07 to 0.26)	<i>-22.2 (-29.9 to -14.4)</i>

values in italics are significant (p<0.05)

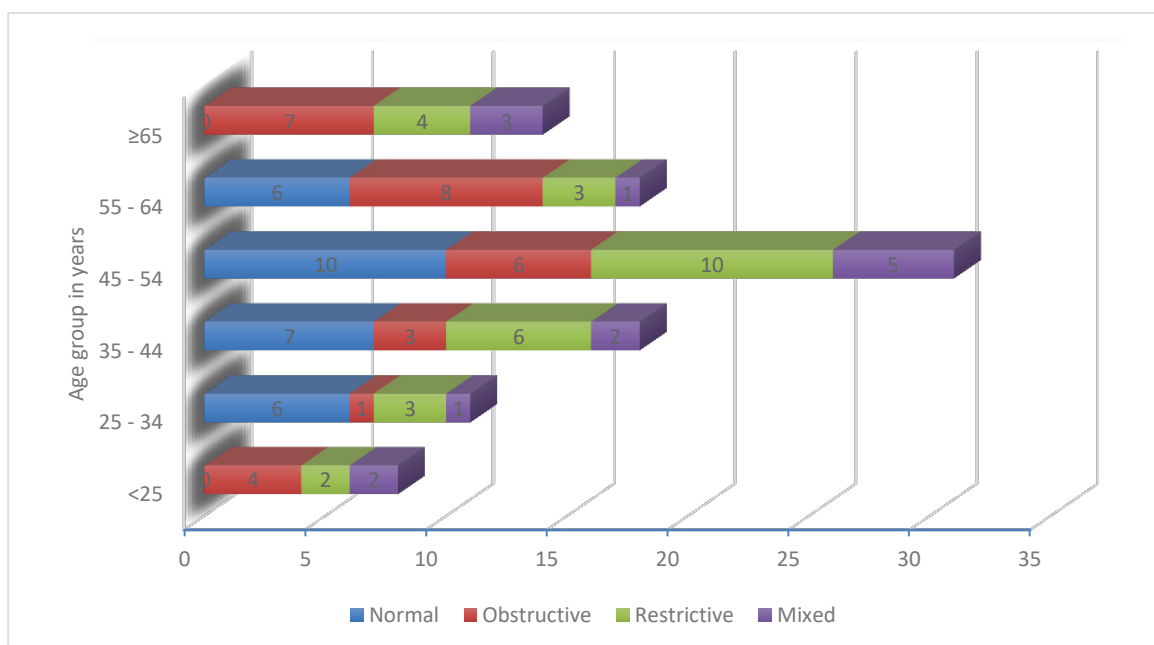


Figure 2: Distribution of lung function by age group in post-TB cases.

Table 7: Association of lung function patterns with sociodemographic factors in post TB individuals

Variable	Normal (n=29) N, %	Obstructive (n=29) N, %	Restrictive (n=28) N, %	Mixed (n=14) N, %	χ^2	<i>p-value</i>
Age						
≤48 years	18 (62.1)	11 (37.9)	16 (57.1)	7 (50.0)	3.80	0.284
>48 years	11 (37.9)	18 (62.1)	12 (42.9)	7 (50.0)		
Gender						
Male	10 (34.5)	12 (41.4)	15 (53.6)	8 (57.1)	3.47	0.325
Female	19 (65.5)	17 (58.6)	13 (46.4)	6 (42.9)		
Level of education						
≤Primary	13 (44.8)	18 (62.1)	8 (28.6)	9 (64.3)	8.14	0.043
≥Secondary	16 (55.2)	11 (37.9)	20 (71.4)	5 (35.7)		
Marital status						
Married	15 (51.7)	13 (44.8)	15 (53.6)	4 (28.6)	2.71	0.439
Not married	14 (48.3)	16 (55.2)	13 (46.4)	10 (71.4)		
Smoking						
Yes	1 (3.4)	3 (10.3)	3 (10.7)	1 (7.1)	1.33	0.723
No	28 (96.6)	26 (89.7)	25 (89.3)	13 (92.9)		
Comorbidities						
Present	9 (31.0)	5 (17.2)	8 (28.6)	2 (14.3)	2.56	0.465
Absent	20 (69.0)	24 (82.8)	20 (71.4)	12 (85.7)		
Previous TB treatment						
Yes	10 (34.5)	12 (41.4)	11 (39.3)	6 (42.9)	0.406	0.939
No	19 (65.5)	17 (58.6)	17 (60.7)	8 (57.1)		
BMI						
Normal/underweight	15 (51.7)	25 (86.2)	23 (82.1)	12 (85.7)	11.94	0.008
Overweight/obese	14 (48.3)	4 (13.8)	5 (17.9)	2 (14.3)		

χ^2 – Chi-square

The current health status of post-TB clients was statistically significantly lower than their healthy counterparts. Quality of life was lower across the symptom 17.1 (IQR 6.8-40.8) and impact

1.05 (IQR 0 – 30.26) domains in post-TB participants compared to controls who never had Pulmonary tuberculosis. This difference was statistically significant with *p* value < 0.001 as shown in table

8. Reduced lung volume indices (FEV_1 , FEF_{25-75} and FEV_1/FVC) were associated with worsened health-related quality of life (table 9)

Table 8: Health status of cases and controls

	Cases N=100, %	Controls n=100, %	Total n=200, %	χ^2 (p-value)
Current health status				
Poor	10 (10.0)	0 (0.0)	10 (5.0)	27.60 (<i><0.001</i>)
Fair	29 (29.0)	10 (10.0)	39 (19.5%)	
Good	58 (58.0)	78 (78.0)	136 (68.0)	
Very Good	3 (3.0)	12 (12.0)	15 (7.5)	

χ^2 – Chi-square

Table 9: Spearman correlation coefficient of St. Georges Respiratory Questionnaire scores with lung volumes of cases

	Total	Symptom	Activity	Impact
FEV_1	-0.25*	-0.27**	-0.26*	-0.19
FVC	-0.14	-0.17	-0.13	-0.10
FEF_{25-75}	-0.29**	-0.24**	-0.34	-0.24*
FEV_1/FVC	-0.28**	-0.28	-0.24	-0.31**

*p<0.05, ** p<0.01

4. Discussion

This study showed that Health-related quality of life was significantly lower across the symptom and impact domains in post-TB patients than in the control group. Median symptom domain score was 17.1 for cases and none for control while median impact domain score was 1.05 for cases and none for control. This is similar with findings by Ozoh et al¹² and Ojuawo et al.¹⁹ who noted significant component score for symptom followed by activity and impact domains. In India, Kim et al⁶ also noted significant decline in HRQoL among post-TB individuals with jeopardised physical and mental health. This was similar to finding in this work which noted compromised current health status. This could be because, in addition to the impact of PTB on the patient’s physical health and fitness, the infectious nature of the disease could cause the individual to be stigmatized in the society even after successful treatment causing significant psychological trauma.

The current health status of Post-tuberculosis individuals was significantly lower than the healthy controls. This can be explained by adverse health-related impact of the disease on patients as well as improved socioeconomic standards among the healthy controls. In this study, a greater percentage of the post-TB patients; 58% (58) had good overall quality of life perception compared to 10% (10) of them who had poor perception. This is similar to what was reported by Kastien-Hilka et al²⁰ who found a significant improvement during the treatment course of TB relative to the beginning of TB treatment. These findings point to the favourable impact of successful treatment for tuberculosis on the patients.

5. Conclusion

There is diminished quality of life following treatment for pulmonary tuberculosis in Owerri. This underscores the need for incorporation of health-related quality of life assessment as an essential tool in the evaluation of post-tuberculosis lung disease.

Declarations

Ethics and Consent to Participate: The research was done in compliance with the Helsinki Declaration. The study was approved by the Ethics committee of the Federal Teaching Hospital, Owerri (FMC/OWR/HREC/VOL11./78). Informed consent to participate was obtained from all of the participants.

Consent for Publication: Not applicable.

Availability of Data and Materials: Data are available from the corresponding author upon reasonable request

Competing Interests: The authors declare no conflict of interests
Funding: There was no external funding for the research Authors’ contributions

Contributions

All authors made meaningful contributions to the study
CV and CC– conceptualised the study and wrote the manuscript
NI and PA – assisted with interpretation of results, revised and supervised the writing of the manuscript
CF and AG– assisted with data collation
GC and COU- assisted with interpretation of results
JC and FC – assisted with data analysis
Acknowledgements: Nil

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References

1. Chakaya, J., Khan, M., Ntoumi, F., Aklillu, E., Fatima, R., Mwaba, P., ... & Zumla, A. (2021). Global Tuberculosis Report 2020—Reflections on the Global TB burden, treatment and prevention efforts. *International journal of infectious diseases*, 113, S7-S12.
2. Kwaghe, A. V., Umeokonkwo, C. D., & Aworh, M. K. (2020). Evaluation of the national tuberculosis surveillance and response systems, 2018 to 2019: National Tuberculosis, Leprosy Buruli Ulcer control programme, Abuja, Nigeria. *Pan African Medical Journal*, 35(1).
3. van Kampen, S. C., Wanner, A., Edwards, M., Harries, A. D., Kirenga, B. J., Chakaya, J., & Jones, R. (2018). International research and guidelines on post-tuberculosis chronic lung disorders: a systematic scoping review. *BMJ global health*, 3(4).
4. Ravimohan, S., Kornfeld, H., Weissman, D., & Bisson, G. P. (2018). Tuberculosis and lung damage: from epidemiology to pathophysiology. *European respiratory review*, 27(147).
5. Allwood, B. W., Stolbrink, M., Baines, N., Louw, E., Wademan, D. T., Lupton-Smith, A., ... & Rylance, J. (2021). Persistent chronic respiratory symptoms despite TB cure is poorly correlated with lung function. *The International Journal of Tuberculosis and Lung Disease*, 25(4), 262-270.
6. Kim, S. H., Lee, H., & Kim, Y. (2021). Health-related quality of life after pulmonary tuberculosis in South Korea: analysis from the Korea National Health and Nutrition Examination Survey between 2010 and 2018. *Health and Quality of Life Outcomes*, 19(1), 195.
7. Nightingale, R., Chinoko, B., Lesosky, M., Rylance, S. J., Mnesa, B., Banda, N. P. K., ... & Rylance, J. (2022). Respiratory symptoms and lung function in patients treated for pulmonary tuberculosis in Malawi: a prospective cohort study. *Thorax*, 77(11), 1131-1139.
8. Menzies, N. A., Quaife, M., Allwood, B. W., Byrne, A. L., Coussens, A. K., Harries, A. D., ... & Cohen, T. (2021). Lifetime burden of disease due to incident tuberculosis: a global reappraisal including post-tuberculosis sequelae. *The Lancet Global Health*, 9(12), e1679-e1687.
9. Chakaya, J., Khan, M., Ntoumi, F., Aklillu, E., Fatima, R., Mwaba, P., ... & Zumla, A. (2021). Global Tuberculosis Report 2020—Reflections on the Global TB burden, treatment and prevention efforts. *International journal of infectious diseases*, 113, S7-S12.
10. Allwood, B. W., Byrne, A., Meghji, J., Rachow, A., Van Der Zalm, M. M., & Schoch, O. D. (2021). Post-tuberculosis lung disease: clinical review of an under-recognised global challenge. *Respiration*, 100(8), 751-763.
11. Pandey, A., Agrawal, R., Agarwal, R., Kumar, A., Gupta, U., & Sharma, D. (2020). Assessment of symptomatic post tuberculosis patients by spirometry and chest X ray. *Int J Contemp Med Res*, 7(1), 2454-7379.
12. Ozoh, O. B., Ojo, O. O., Dania, M. G., Dede, S. K., Adegboyega, O. A., Irurhe, N. K., ... & Adeyeye, O. O. (2021). Impact of post-tuberculosis lung disease on health-related quality of life in patients from two tertiary hospitals in Lagos, Nigeria. *African Journal of Thoracic and Critical Care Medicine*, 27(2), 46-52.
13. Alami, B., Lamrani, Y. A., Amraoui, T., Boubbou, M., & Maaroufi, M. (2015). Pulmonary tuberculosis sequelae: spectrum of radiologic findings and review of literature. *J Med Surg Res*, 2(2), 144-153.
14. Mkocho, P., Naidoo, S., Mbanga, L. C., Nomvete, F., Muloiwa, R., & Dlamini, S. (2019). Chronic lung disease and a history of tuberculosis (post-tuberculosis lung disease): Clinical features and in-hospital outcomes in a resource-limited setting with a high HIV burden. *South African Medical Journal*, 109(3), 169-173.
15. Mpagama, S. G., Msaji, K. S., Kaswaga, O., Zurba, L. J., Mbelele, P. M., Allwood, B. W., ... & Mortimer, K. (2021). The burden and determinants of post-TB lung disease. *The International Journal of Tuberculosis and Lung Disease*, 25(10), 846-853.
16. Allwood, B. W., Stolbrink, M., Baines, N., Louw, E., Wademan, D. T., Lupton-Smith, A., ... & Rylance, J. (2021). Persistent chronic respiratory symptoms despite TB cure is poorly correlated with lung function. *The International Journal of Tuberculosis and Lung Disease*, 25(4), 262-270.
17. Mancuzo, E. V., Martins, E., Sulmonetti, N., Viana, V. D. S., Croda, J., Kritski, A. L., ... & Miranda, S. S. D. (2020). Spirometry results after treatment for pulmonary tuberculosis: comparison between patients with and without previous lung disease: a multicenter study. *Jornal Brasileiro de Pneumologia*, 46(02), e20180198.
18. Meghji, J., Lesosky, M., Joekes, E., Banda, P., Rylance, J., Gordon, S., ... & Squire, S. B. (2020). Patient outcomes associated with post-tuberculosis lung damage in Malawi: a prospective cohort study. *Thorax*, 75(3), 269-278.
19. Ojuawo, O. B., Fawibe, A. E., Desalu, O. O., Ojuawo, A. B., Aladesanmi, A. O., Opeyemi, C. M., ... & Salami, A. K. (2020). Spirometric abnormalities following treatment for pulmonary tuberculosis in Ilorin, Nigeria. *Nigerian Postgraduate Medical Journal*, 27(3), 163-170.
20. Kastien-Hilka, T., Rosenkranz, B., Sinanovic, E., Bennett, B., & Schwenkglens, M. (2017). Health-related quality of life in South African patients with pulmonary tuberculosis. *PloS one*, 12(4), e0174605.
21. Abdelaleem, N. A., Ahmed, M. K., Mohamed, M. N., & Bayoumi, H. A. (2022). Lung health after tuberculosis: clinical

- and functional assessment in post-pulmonary tuberculosis Egyptian patients. *The Egyptian Journal of Bronchology*, 16(1), 23.
22. Nidoi, J., Muttamba, W., Walusimbi, S., Imoko, J. F., Lochoro, P., Ichtho, J., ... & Kirenga, B. (2021). Impact of socio-economic factors on Tuberculosis treatment outcomes in north-eastern Uganda: a mixed methods study. *BMC Public Health*, 21(1), 2167.
 23. Patil, S., Patil, R., & Jadhav, A. (2018). Pulmonary functions' assessment in post-tuberculosis cases by spirometry: Obstructive pattern is predominant and needs cautious evaluation in all treated cases irrespective of symptoms. *The International Journal of Mycobacteriology*, 7(2), 128-133.
 24. Pandey, A., Agrawal, R., Agarwal, R., Kumar, A., Gupta, U., & Sharma, D. (2020). Assessment of symptomatic post tuberculosis patients by spirometry and chest X ray. *Int J Contemp Med Res*, 7(1), 2454-7379.
 25. Meghji, J., Lesosky, M., Joekes, E., Banda, P., Rylance, J., Gordon, S., ... & Squire, S. B. (2020). Patient outcomes associated with post-tuberculosis lung damage in Malawi: a prospective cohort study. *Thorax*, 75(3), 269-278.
 26. Ngahane, B. H. M., Nouyep, J., Motto, M. N., Njankouo, Y. M., Wandji, A., Endale, M., & Ze, E. A. (2016). Post-tuberculous lung function impairment in a tuberculosis reference clinic in Cameroon. *Respiratory medicine*, 114, 67-71.
 27. Manji, M., Shayo, G., Mamuya, S., Mpembeni, R., Jusabani, A., & Mugusi, F. (2016). Lung functions among patients with pulmonary tuberculosis in Dar es Salaam—a cross-sectional study. *BMC pulmonary medicine*, 16(1), 58.
 28. Pasipanodya, J. G., McNabb, S. J., Hilsenrath, P., Bae, S., Lykens, K., Vecino, E., ... & Weis, S. E. (2010). Pulmonary impairment after tuberculosis and its contribution to TB burden. *BMC public health*, 10(1), 259.
 29. Fiogbe, A. A., Agodokpessi, G., Tessier, J. F., Affolabi, D., Zannou, D. M., Adé, G., ... & Marcy, O. (2019). Prevalence of lung function impairment in cured pulmonary tuberculosis patients in Cotonou, Benin. *The International Journal of Tuberculosis and Lung Disease*, 23(2), 195-202.
 30. Nihues, S. D. S. E., Mancuzo, E. V., Sulmonetti, N., Sacchi, F. P. C., de Souza Viana, V., Netto, E. M., ... & Croda, J. (2015). Chronic symptoms and pulmonary dysfunction in post-tuberculosis Brazilian patients. *The Brazilian Journal of Infectious Diseases*, 19(5), 492-497.
 31. Khosa, C., Bhatt, N., Massango, I., Azam, K., Saathoff, E., Bakuli, A., ... & Rachow, A. (2020). Development of chronic lung impairment in Mozambican TB patients and associated risks. *BMC Pulmonary Medicine*, 20(1), 127.
 32. Nightingale, R., Chinoko, B., Lesosky, M., Rylance, S. J., Mnesa, B., Banda, N. P. K., ... & Rylance, J. (2022). Respiratory symptoms and lung function in patients treated for pulmonary tuberculosis in Malawi: a prospective cohort study. *Thorax*, 77(11), 1131-1139.
 33. van Kampen, S. C., Wanner, A., Edwards, M., Harries, A. D., Kirenga, B. J., Chakaya, J., & Jones, R. (2018). International research and guidelines on post-tuberculosis chronic lung disorders: a systematic scoping review. *BMJ global health*, 3(4).
 34. Mancuzo, E. V., Martins, E., Sulmonetti, N., Viana, V. D. S., Croda, J., Kritski, A. L., ... & Miranda, S. S. D. (2020). Spirometry results after treatment for pulmonary tuberculosis: comparison between patients with and without previous lung disease: a multicenter study. *Jornal Brasileiro de Pneumologia*, 46(02), e20180198.
 35. Blamed, E. (2012). *Transmission & Pathogenesis of Tuberculosis*.
 36. Smith, I. (2003). *Mycobacterium tuberculosis pathogenesis and molecular determinants of virulence*. *Clinical microbiology reviews*, 16(3), 463-496.

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