

1 **Prevalence of pulmonary tuberculosis among presumptive cases in Rivers State,**  
2 **Nigeria.**

3

4 **Abstract:**

5 **Background:** Tuberculosis (TB) is a leading cause of death in children but it is  
6 underdiagnosed and underreported in children.

7 **Objective:** To determine the prevalence of pulmonary TB in children among presumptive  
8 cases of TB and to compare the diagnostic efficacy of different screening tool for TB in  
9 children.

10 **Materials and Methods**

11 This study was a descriptive prospective study carried out over one year in nine health  
12 facilities that provide diagnosis and treatment for tuberculosis in Rivers State, Nigeria.  
13 Children aged 0 – 18 years with presumptive TB were explored. They were explored by  
14 carrying out a clinical assessment with chest radiograph, sputum or gastric aspirate for  
15 microscopy and XpertMTB/RIF screening. Sociodemographic data and results of the  
16 screening tests was retrieved from their case records as well as the National Tuberculosis  
17 registers. Ethical approval for the study was obtained from the Rivers state Ministry of  
18 Health. Those with confirmed pulmonary TB were commenced on anti TB medications and  
19 followed up for at least 6months. Obtained data was analysed by SPSS version 20 and  
20 expressed as percentages, proportions and frequencies. A test of significance (chi square and  
21 t-test) was conducted between proportions and means as appropriate. In all a p value of less  
22 than 0.05 was considered significant.

23 **Results**

24 Nine hundred and sixty three patients aged 0-18years had presumptive diagnosis of TB,  
25 394(40.9%) were males while 569(59.1%) were females. The commonest presenting  
26 symptom was chronic cough which occurred in 735 (76.5%) of the patients. The prevalence  
27 of pulmonary TB was 19.1%. Significantly more males (60.9%) than female (39.1%) had  
28 confirmed tuberculosis ( $X^2= 37.431$ , p-value <0.001). Significantly more children (54.3%)  
29 from the low socioeconomic class had confirmed pulmonary. Seventy two (39.1%) and 29  
30 (15.8%) of the patients were AFB and XpertMTB/RIF positive respectively. All (100%) the  
31 children aged 0-5years were neither AFB nor XpertMTB/RIF positive. Out of the children  
32 with confirmed TB, 170 (92.4%) had suggestive clinical features while 14(7.6%) had  
33 suggestive X-ray features. About a third of the children aged 0-5years had their TB  
34 confirmed by suggestive X-rays and Clinical features. All the patients with TB were  
35 commenced on anti TB medications, 40 (21.7%) were lost to follow up (LTFU), 21(11.4%)  
36 were transferred to other centres while 123 (66.8%) completed the treatment.

37 **Conclusion**

38 The prevalence of pulmonary TB among presumptive TB cases in this study was comparable  
39 to findings from other studies and clinical diagnosis of Pulmonary TB remains very relevant  
40 in its management. Improving the clinical skills of physicians involved in TB care and  
41 treatment and the need for community/ facility collaboration to reduce cases of LTFU is  
42 advocated.

43

44 **Introduction:**

45 Tuberculosis (TB) is a common human disease. Worldwide, it is the second leading cause of  
46 death from a single infectious disease. <sup>1</sup> In 2017, TB caused an estimated 1.3 million deaths  
47 among HIV-negative people with an additional 300 000 deaths among HIV-positive people.<sup>2</sup>  
48 Children (aged <15 years) accounted for 15% of total deaths among HIV negative people and  
49 10% of total deaths in HIV positive people.<sup>2</sup>

50 Tuberculosis affects all countries and all age groups and the World Health Organisation  
51 (WHO) has listed Nigeria as one of the countries with very high TB burden and it is hoped  
52 that this will stimulate targeted interventions and advocacy for funding to improve TB  
53 control. <sup>4</sup>

54 The actual burden of Tuberculosis in children in Nigeria is not known, however, it is  
55 estimated by the WHO that 30,000 children get TB in Nigeria each year. Study show that  
56 sputum microscopy smear-positive TB in children (<14 years old) is low.<sup>3</sup> and because the  
57 majority of children are sputum microscopy smear negative, these data underestimate the true  
58 burden of childhood TB.

59 Tuberculosis in children is underdiagnosed and underreported as microscopical examination  
60 of sputum smears is the cornerstone of diagnosis in most resource limited countries, but its  
61 usefulness is limited in young children who have paucibacillary disease and are unable to  
62 expectorate sputum. These children with undetected TB form a large pool that eventually  
63 becomes adult cases of TB. To reduce cases of missed diagnosis, children with presumptive  
64 diagnosis of TB are pooled and screened for TB by both bacteriological method and strong  
65 clinical assessment combined with chest radiograph.

66 A recent and a more reliable screening tool and the first line WHO recommended screening  
67 tool for TB the XpertMTB/RIF screening, is an important advance in rapid detection of TB

68 disease and drug resistance TB. Studies have shown that it is much more sensitive than  
69 microscopy, with sensitivity being reported from 75 to 90% on sputum samples and nearly  
70 70% on gastric aspirates, with comparable performance in HIV positive and HIV negative  
71 children.<sup>5,6</sup>

72 The study therefore was carried out to determine the prevalence of pulmonary TB disease  
73 among presumptive cases of TB and to compare the diagnostic efficacy of different screening  
74 tool for TB in children.

#### 75 **Methodology:**

76 This study was a descriptive prospective study. It was carried out in Rivers state in Southern  
77 Nigeria among nine health facilities that provide diagnosis and treatment for tuberculosis in  
78 Rivers State, Nigeria. The diagnosis and treatment of TB in these centres follow the National  
79 guidelines for Tuberculosis and Leprosy control programme and the WHO directly observed  
80 treatment short course (DOTs) strategy. Ethical approval for the study was obtained from the  
81 Rivers state Ministry of Health. The study population consisted of children aged 0 – 18 years  
82 who presented in these health facilities from 1<sup>st</sup> of April 2016 to 31<sup>st</sup> of March 2017. All  
83 children with presumptive TB were explored. Presumptive TB refers to a patient who  
84 presented with one or more of the following symptoms: cough for >2 weeks; cough not  
85 responding to adequate dose of first line antibiotics after 7days; unexplained fever for >2  
86 weeks; failure to thrive (FTT) or weight loss; a contact with a suspected or confirmed case of  
87 pulmonary TB. FTT was defined as a weight or height measurement that fall below the 3<sup>rd</sup> or  
88 5<sup>th</sup> percentile for age or sex or a downward change in growth that crosses two major growth  
89 percentile on a growth chart. They were explored by carrying out a chest x-ray, sputum or  
90 gastric aspirate for microscopy and XpertMTB/RIF screening and by noting the number of  
91 symptoms and signs that they presented with. Sociodemographic data and results of the

92 screening tests was retrieved from their hospital record files as well as the various National  
93 Tuberculosis registers kept in the DOTS clinic. All patients with extra pulmonary TB and  
94 those with Rifampicin resistance following XpertMTB/RIF test were excluded from the study  
95 while those with rifampicin resistance were referred to Drug resistant TB centres. Individual  
96 with confirmed pulmonary TB (clinically and bacteriologically) were commenced on  
97 standard first line anti TB medications and followed up for at least 6months. Follow up was  
98 done to ensure compliance to treatment and to prevent lost to follow up (LTFU) through  
99 phone calls to patients or their caregivers at least once a week (a day before their medication  
100 pick up). Obtained data was entered into an excel sheet and analysed by SPSS statistical  
101 software version 20. Data was expressed as percentages, proportions and frequencies. A test  
102 of significance (chi square and t-test) was conducted between proportions and means as  
103 appropriate. In all a p value of less than 0.05 was considered significant.

#### 104 **Result**

105 Nine hundred and sixty three patients aged 0-18years had presumptive diagnosis of TB,  
106 394(40.9%) were males while 569(59.1%) were females. The mean age was  $7.05 \pm 4.53$   
107 years, the mean age for males was  $7.3 \pm 4.6$  while the mean age for females was  $6.9 \pm 4.9$ , the  
108 age difference was not statistically significant (  $t = 0.793$  ,  $p \text{ value} = 0.120$  ). The commonest  
109 presenting symptom was chronic cough which occurred in 735 (76.5%) of the patients. Table  
110 1

111 One hundred and eighty four had confirmed TB giving a prevalence of 19.1%. More males  
112 (60.9%) than female (39.1%) had confirmed tuberculosis and the difference was statistically  
113 significant ( $X^2 = 37.431$ ,  $p\text{-value} < 0.001$ ). More children (54.3%) from the low  
114 socioeconomic class had confirmed TB and there was a statistically significant different

115 between TB and social class. There was a near equal representation of children with  
116 confirmed TB among the different age groups after one year of age. Table 2

117 Seventy two (39.1%) of the patients were AFB positive. More males (53.7%) than females  
118 (40.3%) were smear positive for AFB, however this difference was not statistically  
119 significant. Table 3. Of the patients with confirmed TB, XpertMTB/RIF was positive in only  
120 29(15.8%) of them. Seventeen (58.6%) of them were males while 12 (41.4%) were females.  
121 There was no gender difference in proportion in children with positive XpertMTB/RIF test,  
122 (p value = 0.787). Table 4. All (100%) the children aged 0-5years had undetectable TB  
123 antigen by both AFB and XpertMTB/RIF. More than half and nearly half of the children with  
124 bacteriologically detectable TB by XpertMTB/RIF and AFB respectively were between 11-  
125 15years. Table 5

126 Out of the children with confirmed TB, 98 (53.3%) had suggestive clinical features while  
127 86(46.7%) had suggestive X-ray features. More than a third (33.2%) of the children aged 0-  
128 5years had their TB confirmed by suggestive X-rays and Clinical features. Table 6

129 All the one hundred and eighty four identified TB cases were commenced on appropriate  
130 doses of standard six months first line anti TB medications (Rifampicin, Isoniazide,  
131 Pyrazinamide and Ethambutol) under DOTs strategy. One month into treatment, 17 (9.2%)  
132 were lost to follow up and by the second month, a total of 40 (21.7%) were lost to follow up.  
133 Twenty one (11.4%) relocated out of Rivers state and were transferred to centres closest to  
134 them while 123 (66.8%) completed the treatment.

### 135 **Discussion:**

136 The diagnosis of TB is protean and requires a high index of suspicion. This is because there is  
137 a wide variation in presentation and severity of disease. A presumptive TB case is any  
138 person whether adult or child, with signs and/or symptoms suggestive of TB (pulmonary or

139 extra-pulmonary, or those with chest x-ray findings suggestive of active TB disease.  
140 Presumptive diagnosis of TB increases the pool of patients with clinical risk factors for TB  
141 who are subsequently screened to confirm the presence of disease. Nine hundred and sixty  
142 three patients had a presumptive diagnosis of TB in this study with more females (59.1%)  
143 than males (40.9%) being affected. Presumptive diagnosis over diagnoses TB but however  
144 ensures that any case of TB that presents to health facilities are not left undiagnosed. Study  
145 show varying proportion of confirmed TB disease from this pool.<sup>7, 8</sup>

146 The presentation of pulmonary TB disease in children depends on many factors that include  
147 age, immunization status to the bacillus Calmette-Guerin (BCG), immune status, co-existing  
148 disease, virulence of the infecting organism and host-microbe interaction.<sup>9</sup> Chronic cough as  
149 was found in 76.3% of this study was the commonest presenting symptom in these children  
150 and this is similar to findings in other studies in children with pulmonary TB.<sup>10, 11</sup> This cough  
151 is initially unproductive but with progressive inflammation and tissue necrosis, sputum is  
152 produced. None of our patient had hemoptysis but this is occasionally a presenting symptom  
153 especially in older children but usually results from previous disease and may not indicate  
154 active tuberculosis. It may arise from rupture of a dilated vessel in the wall of a cavity  
155 (Rasmussen's aneurysm), tuberculous bronchiectasis, bacterial or fungal infection (especially  
156 *Aspergillus* mycetoma) in a cavity or erosion into an airway (broncholithiasis).<sup>10, 11</sup> The  
157 duration of cough in older children and adults has been shown to affect transmission, with  
158 other children more likely to be infected if the source case has been coughing for a longer  
159 period of time.<sup>12, 13</sup> Children who present with chronic cough especially when not responding  
160 to simple antibiotics should raise the suspicion for Pulmonary TB, this is more so when there  
161 are other suggestive symptoms like weight loss/failure to thrive (FTT) that 20.6% of our  
162 patients presented with. It is quite common not to get the history of contact with an adult with  
163 TB or chronic cough as was found in this study. This may probably be due to the fact that

164 many people still feel stigmatised by TB and so relations refuse to volunteer the history or  
165 hide it even when asked. Poor knowledge and cultural beliefs about TB may also contribute  
166 to this and these all lead to poor health seeking behaviour, delay or failure to access the health  
167 services.<sup>14, 15</sup> The fact that TB is a curable disease especially with early presentation calls for  
168 increased public health enlightenment so that any associated stigma and misconceptions  
169 could be addressed.

170 This study found a pulmonary TB disease prevalence of 19.1% among children with  
171 presumptive diagnosis of TB. This compares with 20.7% prevalence reported by Kolapo in  
172 Lagos state<sup>7</sup> but contrasts with findings from Nwachukwu et al who found a prevalence of  
173 12.3% among presumptive cases of TB in Anambra state.<sup>8</sup> These studies were among adults  
174 as there are limited studies in children. The methodology used by Nwachukwu et al may  
175 have contributed to the lower prevalence as cases of confirmed TB was by sputum  
176 microscopy only while a combination of microscopy, XpertMTB/RIF and radiological and  
177 clinical findings were used in our study.

178 Statistically significant more males than females were diagnosed with TB in this study as is  
179 reported in other studies.<sup>16, 17</sup> Males may be more predisposed due to a yet unidentified  
180 genetic risk or the fact that parents have a better health seeking behaviour for their male  
181 children especially in our part of the world. Other studies have also reported a female  
182 predominance of TB among older children aged 11 years and above.<sup>18-21</sup> This age group has  
183 been found to have a higher prevalence of TB due to hormonal and reproductive changes in  
184 this age group. Among adults however, two thirds of TB cases occur in men and globally,  
185 men are significantly more at risk of contracting and dying from TB than women.<sup>22, 23</sup>

186 It is not surprising that more children from the lower social class were diagnosed with TB as  
187 TB has been described as a disease of poverty. These individuals are more likely to live in

188 areas with high population and overcrowded environments. High population density will  
189 increase the risk of a child coming into contact with an infectious TB case, as more dense  
190 populations lead to more human encounters and encounters of more intense and prolonged  
191 physical proximity.<sup>24</sup>

192 The age of a child influences the risk of TB exposure and eventual disease. There was a near  
193 equal representation of the TB cases among the different age groups in this study, however,  
194 children aged 0-5years constituted over a third of the cases of TB. The immunity of children  
195 in this age group is poor and so progression from latent TB to TB disease is higher among  
196 them once exposed to an infectious source. However, because pre-school children interact  
197 with fewer adults, and generally come into contact with adults only in their family units, their  
198 chances of exposed to an adult with infectious TB is reduced. Older children interact with  
199 more adults in their day-to-day life, they can therefore be exposed to infectious cases of TB at  
200 home or in the community.

201 Bacteriological confirmation of TB disease in children is difficult and this is more so in  
202 younger children. This is because children have paucy bacillary disease and find it difficult to  
203 produce sputum. The skill of acquiring sputum by gastric aspirate for AFB and or  
204 XpertMTB/RIF is quite invasive and poorly done by many health workers on outpatient  
205 basis. Many do a gastric lavage instead and the required minimum 4hours fast before the  
206 procedure is not usually practiced. Sputum yield becomes poor and so the XpertMTB/RIF  
207 and AFB yield. All these contribute significantly to misdiagnosis, underdiagnoses and  
208 underreporting in children. This study found a positive yield of 39.1% and 15.8% for AFB  
209 and XpertMTB/RIF respectively. Studies have shown that XpertMTB/RIF is rapid and more  
210 sensitive to microscopy with sensitivity as high as 75-90% on sputum samples and as high as  
211 70% on gastric aspirates.<sup>25, 26</sup> Despite the low yield to both microscopy and XpertMTB/RIF  
212 in this study there was a higher sensitivity to microscopy in contrast to reported findings. This



213 may be due to poor specimen collection in younger children or poor expertise on the use of  
214 the recently introduced XpertMTB/RIF test by its users or both. The low bacteriological yield  
215 that was found in this study has been reported in other studies.<sup>27, 28</sup> Improvement on the  
216 quality of this Mycobacterium TB gene detection by XpertMTB/RIF test, improvement on  
217 the expertise of its use or use of other specimen like stool and urine specimen for detection of  
218 TB in children is advocated.

219 All children aged 0-5years had undetectable bacteriological test. This clearly suggest that  
220 either the sputum yield was poor, that the bacteriological screening methods lacked  
221 sensitivity or both. It may also depict the paucibacillary nature of TB in children and the fact  
222 that most children in this age group have primary TB which is not commonly associated with  
223 cavitatory lesions. However many children in this age group had clinical and radiological  
224 evidence of TB and responded to anti TB medications. It then follows that many children  
225 who are managed by clinicians without expertise on clinical diagnosis of TB may go  
226 undetected and progress to severe disease with poorer outcome. Over 80% of children with  
227 detectable bacteriological tests were aged 11years and above confirming the appropriateness  
228 and usefulness of these diagnostic methods in older children.

229 Nearly 50% of children with TB were diagnosed by clinical evaluation and use of chest  
230 radiograph. These children had two or more suggestive symptoms and sign and augmented  
231 with chest radiographs findings. The low diagnostic yield of sputum microscopy and  
232 XpertMTB/RIF in children especially the very young makes them an unreliable method of  
233 diagnosis in children when used solely. A high index of suspicion and clinical expertise  
234 becomes vital in making a diagnosis of pulmonary TB in children especially in children 0-  
235 5years. These very young children with low immunity are more likely to develop severe  
236 disease and to succumb to it yet diagnosis and treatment is delayed in many due to lack of  
237 experienced physicians. TB diagnosis therefore requires clinical expertise and early screening

238 of children who presents with worsening and unrelenting cough lasting for 2 weeks or more  
239 with associated fever, weight loss or poor weight gain and a positive history of contact with  
240 an adult case of TB.

241 The DOTs strategy in TB treatment requires that patients visit TB treatment centres (closest  
242 to them) on daily basis and are observed to take their medications. This strategy guarantees  
243 compliance and adherence knowing the importance of completing TB medications or  
244 achieving cure. However, this is practically difficult in the many areas as each DOTs centres  
245 covers a wide expanse of area. Therefore, Patients are given drugs on weekly basis and  
246 followed up with regular phone calls. Diagnosed patients with TB and or their caregivers are  
247 however properly counselled about the disease and drug adherence before commencement of  
248 medications and this is reinforced at every visit. Despite these measures, about a quarter of  
249 them were lost to follow up (LTFU)s. In a Kenyan study, Kibango et al reported a lower  
250 LTFU rate of 13%.<sup>29</sup> Reasons proffered for LTFU included no salaried employment, lack of  
251 family support and the male gender. Other studies proffered reasons such as financial  
252 limitation, transportation costs, lack of job, feeling better after few weeks of treatments,  
253 knowledge deficit about duration of treatment, poor patient-health worker relationship.<sup>30-35</sup>  
254 Home visits by health workers and collaboration with community health workers may  
255 overcome some of these reasons but is quite cost intensive. However, this should be explored  
256 and properly integrated into the end TB strategy milestone for 2025 that calls for a reduction  
257 in number of TB deaths by 75% and TB incidence by 50%.

## 258 **Conclusion:**

259 The prevalence of pulmonary TB among presumptive TB cases in this study was comparable  
260 to findings from other studies and clinical diagnosis of Pulmonary TB remains very relevant  
261 in its management. Improving the clinical skills of physicians involved in TB care and  
262 treatment and the need for community/ facility collaboration cannot be overemphasized.

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387  
388

389 **Tables:**

390

391

392 **Table 1: Socio demographic characteristics and presenting symptom of presumptive TB**

<b>Socio-demographic of presumptive TB cases</b>	<b>Frequency (n =963)</b>	<b>percentage ( % )</b>
<b>Gender</b>		
Males	394	40.9
Females	569	59.1
<b>Social class</b>		
Upper	210	21.8
middle	339	35.2
Lower	414	43.0
<b>Symptoms</b>		
Chronic cough> 2weeks	735	76.3
Weight loss/FTT	198	20.6
Contact with a TB case	30	3.1
<b>Age years</b>		
< 1	20	2.1
1-5	477	46.8
6- 10	201	20.8
11-15	214	22.3
16-19	51	5.3
<b>Mean age (years)</b>	<b>7.01 ± 4.53</b>	

393

394 **Prevalence of Tuberculosis:**

395 One hundred and eighty four had confirmed TB giving a prevalence of 19.1%. More males  
396 (60.9%) than female (39.1%) had confirmed tuberculosis. This difference was statistically  
397 significant ( $X^2 = 37.431$ , p-value <0.001). More children (54.3%) from the low  
398 socioeconomic class had confirmed TB and there was a statistically significant difference  
399 between TB and social class. There was a near equal representation of children with  
400 confirmed TB in the different age groups after one year of age.

401 **Table 2. Prevalence of Tuberculosis**

Variables	Unconfirmed cases	Confirmed cases	Total	$\chi^2$	df	p value
<b>Gender</b>						
Males	282 (36.2)	112(60.8)	394(40.9)	<b>37.431</b>	<b>1</b>	<b>&lt; 0.001</b>
Females	497 (63.8)	72 (39.1)	569(59.1)			
<b>Total</b>	<b>779</b>	<b>184</b>	<b>963</b>			
<b>Social class</b>						
Upper	171(22.0)	39 (21.2)	210(21.8)	<b>12.952</b>	<b>2</b>	<b>0.002</b>
Middle	293 (37.6)	46 (25.0)	339 (35.2)			
Lower	315(40.4)	99 (53.8)	414(43.0)			
<b>Total</b>	<b>779</b>	<b>184</b>	<b>963</b>			
<b>Age years</b>						
< I	11 (1.4)	9(4.9)	20(2.1)	<b>164.752</b>	<b>4</b>	<b>&lt;0.001</b>
1-5	425 (54.5)	52(28.3)	477(49.5)			
6- 10	169 (21.7)	32(17.4)	201 (20.9)			
11-15	165(21.2)	49(26.6)	214(22.2)			
16-18	9 (1.2)	42(22.8)	51(5.3)			
<b>Total</b>	<b>779 (100.0 %)</b>	<b>184(100.0%)</b>	<b>963 (100.0)</b>			

402

403 **Proportion of subjects with positive AFB test by gender**

404 Seventy two (39.1%) of the patients were AFB positive. More males (53.7%) than females  
 405 (40.3%) were smear positive for AFB, however this difference was not statistically  
 406 significant. Table 3

407

408 **Table 3: Proportion of subjects with positive AFB test by gender**

Gender	AFB		Total
	Detected	not detected	
Males	43(59.7)	69(61.6)	112(60.9)

Females	29 (40.3)	43(38.4)	72(39.1)
	72(100.0)	112(100.0)	184 (100.0)

409  $\chi^2$  0.065 , df = 1 p value 0.799

410 **Proportion of subjects with positive XpertMTB /RIF by gender**

411 Of the patients with confirmed TB, XpertMTB/RIF was positive in only 29(15.8%) of them.  
 412 Seventeen (58.6%) were males while 12 (41.4%) were females. There was no gender  
 413 difference in proportion of positive XpertMTB/RIF test, (p value = 0.787). Table 4

414

415

416 **Table 4: Proportion of subjects with positive XpertMTB/RIF test by gender**

Gender	XpertMTB/RIF		Total
	Detected	not detected	
Males	17 (58.6)	95 (61.3)	112(60.9)
Females	12 (41.4)	60(38.7)	72(39.1)
Total	29(100.0)	155(100.0)	184 (100.0)

$\chi^2$  0.073 , df = 1 p value 0.787

417

418 **Comparism between age and Microbiological results**

419 All (100%) the children aged 0-5years had undetectable TB antigen by both AFB and Gene  
 420 XpertMTB/RIF. More than half and nearly half of the children with detectable TB antigen by  
 421 XpertMTB/RIF and AFB respectively were between 11-15years.

422

423



424 **Table 5: Comparism between age and Microbiological results**

Microbiological results	XpertMTB/RIF		AFB	
	Detected( N %)	Not detected( N %)	Detected ( N %)	Not detected ( N %)
<b>Age years</b>				
< I	0(0.0)	9 (5.8)	0(0.0)	9(8.0)
1-5	0(0.0)	52 (33.5)	0 (0.0)	52(46.4)
6- 10	4(13.8)	28 (18.1)	9 (12.5)	23(20.6)
11-15	18(62.1)	31(20.0)	32(44.4)	17(15.2)
16-18	7(24.1)	35(22.6)	31(43.1)	11(9.8)
	<b>29(15.8)</b>	<b>155(84.2)</b>	<b>72(39.1)</b>	<b>112 (60.9)</b>

425

426 **Age and Clinical diagnosis of TB:**

427 Out of the children with confirmed TB, 98 (53.3%) had suggestive clinical features while  
 428 86(46.7%) had suggestive X-ray features. More than a third (33.2%) of the children aged 0-  
 429 5years had their TB confirmed by suggestive X-rays and Clinical features. Table 6

430 **Table 6: Age and Clinical diagnosis of TB:**

<b>Clinical features (CF) / x ray</b>			
<b>Age years</b>	<b>Suggestive CF</b>	<b>Suggestive Xray</b>	<b>Total</b>
< I	5(5.1)	4(4.6)	9(4.9)
1-5	30(30.6)	22(25.6)	52(28.3)
6- 10	16(16.3)	16(18.6)	32(17.4)
11-15	27(27.6)	22(25.6)	49(26.6)
16-18	20(20.4)	22(25.6)	42(22.8)
<b>Total</b>	<b>98(53.3)</b>	<b>86 (46.7)</b>	<b>184 (100.0)</b>

431  $\chi^2$  1.17 , df = 4 p value 0.883