

1 Prevalence of Tuberculosis among Children with Severe Acute Malnutrition at Ola During
2 Children's Hospital in Freetown Sierra Leone

3

4 **Abstract**

5 **Prevalence of Tuberculosis among Children with Severe Acute Malnutrition at Ola**
6 **during Children's Hospital in Freetown Sierra Leone. Worldwide, pediatric tuberculosis**
7 **account for about 1million cases, annually, accounting for 10-15% of all tuberculosis;**
8 **with more than 100,000 estimated mortality annually, it is also one of the 10 most**
9 **common causes of childhood mortality. Aim of this study was to determine the**
10 **prevalence of tuberculosis among children with severe acute malnutrition at Ola During**
11 **Children's Hospital in Freetown Sierra Leone. It was a descriptive cross-sectional**
12 **study, carried out at the therapeutic feeding center (TFC) of Ola During Children's**
13 **Hospital in 2018. An opportunistic sampling method in which every next patient whose**
14 **mother gave consent was recruited until the number 74 was reached.** Patients who met
15 the World Health Organisation (WHO) criteria for diagnosis of severe acute malnutrition
16 **were** admitted into the TFC **and consecutively** selected and interviewed using a
17 structured questionnaire after obtaining written informed consent, from their mothers
18 or caregivers. **All the mothers approached during the study period consented for the**
19 **study.** Diagnosis of tuberculosis was both clinically and by laboratory investigations, 74
20 **children whose mothers/caregivers** consented for the study were recruited. Data was
21 entered into an excel spread sheet and analyzed using Epi info version 7. There were 74
22 children with a median age of 11months \pm 9.9SD. Forty (54.1%) Males and 34(45.9%)
23 Females, with a M:F ratio of 1.18:1. Prevalence of tuberculosis was 20%. Diagnosis of
24 Tuberculosis was based on clinical findings of extreme weight loss or failure to gain

25 weight, Chest x-ray findings of perihilar infiltrates. Gene Xpert MTB RIF results were all
26 negative 0(0%). Most of the mothers 59 (79.7%) were aged between 20-29years,
27 45(60.9%) of them were petty traders, while 15(20.3%) had no formal education. **The chi**
28 **square was used to determine the statistical difference,** there was no statistically
29 significant difference between gender and TB, $P= 0.3415$, there is a statistically
30 significant difference between no formal education and occurrence of tuberculosis in
31 their children $P= 0.0467$.

32 **CONCLUSIONS/RECOMMENDATIONS:** Prevalence of Tuberculosis is still high among
33 children with severe acute malnutrition. Gene Xpert MTB RIF was unable to make a
34 bacteriological confirmation. **There are difficulties with making bacteriological**
35 **confirmation of tuberculosis in resource poor settings. Guidelines requiring mainly**
36 **clinical parameters need to be developed for use in resource limited countries.**

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39 **KEY WORDS:** Prevalence, Tuberculosis, Severe acute malnutrition. Paediatrics.

40 1. INTRODUCTION

41 Ten to twenty percent of deaths in children under the age of 15years in tuberculosis
42 (TB) endemic countries are alleged to be associated with tuberculosis. [1,2] The World
43 health Organisation reported a total of 140,000 mortalities in Paediatric age in their
44 2015 global TB report from vital registration data. [3] In 2012, TB accounted for 2% of
45 total deaths in children. [4] In Southeast Asia and Sub-Saharan Africa tuberculosis in
46 children accounted for less than 4% among the notified new tuberculosis cases[4]
47 Tuberculosis continues to be a major cause of morbidity and mortality in children

48 globally especially in those from resource limited settings.[5] Globally there are about
49 9million new TB cases each year and 11% of these occur in Paediatric patients.[5]
50 Children living in areas where TB is endemic are also plagued with malnutrition and it
51 accounts for 2.2million deaths in children less than 5years all over the world.[6]
52 Malnutrition and poor infection control have blossomed in an environment of poverty,
53 overcrowding, food insecurity, human immunodeficiency Virus [7] Malnutrition is deadly
54 when coexisting with tuberculosis, social and economic factors that cause malnutrition
55 to thrive such as poverty, illiteracy, ignorance, overcrowding and poor sanitation also
56 contribute to the prevalence of tuberculosis.[8] Hence we tried to look at the
57 prevalence of tuberculosis among children with severe acute malnutrition at Ola During
58 Children's hospital in Freetown and some of its socioeconomic factors, since there has
59 been no known study in this subject matter in Freetown.

60 2. Materials and Methods

61 a. Study Area

62 Therapeutic feeding center (TFC) of the Ola During children's hospital in
63 Freetown, Sierra Leone. A place where children with severe acute
64 malnutrition are admitted and managed. Ola During children's hospital is the
65 only Paediatric tertiary hospital in Freetown Sierra Leone and as such
66 receives referrals from all across the country.

67 b. Study Population

68 Under-five children admitted in to TFC during the period of the study whose
69 parents or care giver consented for the study were recruited. The children
70 were admitted in to TFC, if they met the WHO criteria for severe acute
71 malnutrition. Severe acute malnutrition criteria was met as defined by WHO, if
72 there was very low weight for height (Below -3zscores of the median

73 NCHS/WHO growth standards), or visible severe wasting, or presence of
74 nutritional oedema. [9]

75 **c. Selection and Inclusion criteria**

76 **i. Inclusion criteria**

77 All children on admission at TFC ward during the period of the study
78 whose parent or caregiver consented for the study.

79 **2.3.2 Exclusion criteria**

80 All patients on admission in TFC during the period of the study whose
81 parents or caregivers refused to consent for the study.

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84 **d. Sampling Method**

85 This was a descriptive cross-sectional study. A non-probability
86 sampling method (opportunistic sampling) was used, in which every
87 next child admitted into TFC, whose parents and caregivers consented
88 for the study was recruited into the study until we got 74 subjects. All
89 parents or caregivers whose children were on admission at TFC during
90 the period of the study, who were approached by the researcher
91 consented for the study. The study was collected over a six months
92 period in 2018. A structured questionnaire was used to collect data on
93 parents or caregiver's biodata, child's biodata, clinical and laboratory
94 results. All children were to have a chest X-ray and a gene X-pert MTB
95 -RIF test done for the diagnosis of tuberculosis. Laboratory results
96 were obtained from patient's case note by the researcher. But often

97 times the diagnosis of TB was made clinically as most times the chest
98 X-ray machine was not working or the gene X-pert machine was not
99 functioning. The diagnosis was most times based on failure of the child
100 to gain weight despite appropriate feeding. On one occasion we had
101 history of contact with mother who was on treatment for tuberculosis,
102 as often times in paediatric, patients it is difficult to find history of
103 contact. Rarely diagnosis was based on therapeutic trial on those with
104 chronic cough and failure to gain weight despite appropriate feeding.

105 Data was entered into Microsoft excel spread sheet and analyzed
106 using Epi-info version 7

107 3. Results and Discussions

108 There were 74 subjects, 40(54.05%) Males and 34(45.95%) females giving a M:F
109 ratio of 1.17:1. Their median age was 11 months SD \pm 9.9. 48(64.9%) were aged
110 between 12-59 months. The prevalence of tuberculosis was high 20(27%).
111 Tuberculosis was highest in the age group 12-59 months 14(70.0%) Diagnosis
112 was mostly clinical and with chest X-ray 15(20.27%) as all the gene X-pert test
113 done 13(17.57%) came out negative. The chest X-rays showed pulmonary
114 infiltrates and perihilar opacities All 20(27%) of the children with TB had received
115 BCG at birth. 59(79.7%) of the parents/caregivers were aged between 20-
116 29 years., they were mostly traders 45(60.9%) while 8(10.45) were unemployed,
117 15(20.3%) had no formal education. All the patients (100%) got well and were
118 discharged home to the nearest moderate acute malnutrition (MAM) clinic to
119 their home. The chi square was used to test for statistical significance. There was
120 a statistically significant difference between no formal education in

121 parents/caregivers and occurrence of TB in their children P-value 0.046. There
122 was no significant difference in sex P-value 0.341

123 **Discussions**

124 This study found a 20% prevalence of TB among patients with SAM. This in contrast to
125 the finding of Munthali et al [10] working in Lusaka, Zambia who had a prevalence of
126 1.58% in Zambia among malnourished children. The number of patients (74) in this
127 study was much smaller than the Zambian study that was reported among 9540, this
128 may have accounted for the difference in the prevalence. [11] **Christi et al in**
129 **Bangladesh who reported a prevalence of 7% among children with SAM and signs of**
130 **Pneumonia, and the work of Bhat et al in Karnataka, also in India who found a 4%**
131 **prevalence among children with SAM following the diagnostic algorithm and a 0.3%**
132 **prevalence among children with SAM who did not follow the diagnostic alogorithm.**
133 These three studies had similar values which are lower than what we found in this
134 study. However, ours is comparable to the work of Veeraraja et al [8] in India who found
135 a prevalence of tuberculosis of 22% among children with severe acute malnutrition.
136 There was a 0 % bacteriological confirmation in this study which is also in contrast to
137 the Zambian study that had a 25% bacteriological confirmation among the 151patients
138 with tuberculosis in their study. Although the method of bacteriological confirmation
139 employed in their study was a smear microscopy performed on gastric aspirates. This
140 study used a more sensitive Xpert MTB/ RIF which gave a 0% yield, however, **Christi et**
141 **al [11] in Bangladesh reported that tuberculosis was microbiologically confirmed in 7%**
142 **(27/396) of the children who provided sputum. Twenty-one was by Xpert MTB/RIF while**
143 **10 was by culture and 4 was by both methods.** [11] Using more sensitive X pert MTB RIF
144 was also of no additional value among severely malnourished children in Malawi [13]
145 This however shows that there is a low yield of Mycobacterium tuberculosis. The

146 bacteriological isolation of mycobacterium tuberculosis in children is said to be very
147 difficult due to the pauci bacillary nature of childhood tuberculosis [8] This study found
148 a median of age of 11 ± 9.9 months this is in keeping with the finding of other workings
149 with a peak incidence of pulmonary tuberculosis among malnourished children of 1-
150 3 years [8,11,14,15] However Veeraraja et al [8] found incidence of pulmonary
151 tuberculosis at a younger age of 6-12 months in severely malnourished children. While
152 Munthali et al had a higher median age of 16 months. However, they are all among
153 preschool children. Just like other workers [8,14,15] this study did not find any sex
154 predilection. No child in this study had the severe forms of tuberculosis such as
155 disseminated TB and neuro tuberculosis. as they were all vaccinated with BCG. This is
156 not surprising as BCG is known to protect from the very severe forms of tuberculosis
157 such as disseminated TB and neuro tuberculosis [15] 20.27% of the parents in this study
158 had no formal education, 10% were unemployed and 79.7% of them were young adults.
159 These features conform to the description of poverty, illiteracy, ignorance that
160 constitute risk factors for the formation of tuberculosis [6] There was an epileptic
161 functioning of the chest x-ray machine and the Gene Xpert machine this made making a
162 diagnosis of tuberculosis very difficult and making a diagnosis had to be done
163 clinically, following failure of the patients to gain weight despite adequate therapeutic
164 feeding and therapeutic trials. For some other patients a history of contact with a
165 confirmed case was used among other criteria. This is no different from what is
166 obtained in other resource limited countries like India. Bhat et al [12] reported that full
167 current electricity required for the x-ray machine was only available for 3 hours during
168 working hours resulting in long waiting hours for patients and as such not all patients
169 could have the X-ray done for their diagnosis. The diagnostic algorithm in their protocol
170 places importance in detecting AFB in the sputum, broncho-alveolar specimen or

171 gastric lavage but it was difficult to carry out on all patients in their setting. This also
172 made bacteriological confirmation difficult in their setting like we had in this study.
173 All the children in this study recovered and were discharged home to be followed up at
174 the moderate acute malnutrition centers closest to them. similar to the work of Christi
175 et al [11]in Bangladesh where the patients were discharged and followed up for six
176 months in case of tuberculosis. While Bhat et al [12] had patients, who may have died
177 who they were unable to account for in Karma kata.

178

179 **Conclusion**

180 The prevalence of tuberculosis is high among children with severe acute
181 malnutrition. One interesting thing that was found in this study is the fact that
182 although Xpert MTB/ RIF is said to be highly sensitive in diagnosis of
183 mycobacterium tuberculosis there was a zero percent yield with it in this study. T
184 he high technology machines used in more advanced countries were not very
185 helpful in diagnosis of TB in this setting because of its repeated break down and
186 lack of uninterrupted power supply. Clinical guidelines for diagnosis of
187 tuberculosis, which do not require such high technology machines should be
188 developed for resource poor countries. They should be used to compliment this
189 high technology machines.

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191 **Limitations**

192 This was a hospital-based study and so needs to be replicated in a rural
193 community. Also, the Xpert MTB /RIF test should be done in a community-based
194 study as it was epileptic in its function during the period of this study and the

195 results obtained in this study need to be validated in a larger sample in the
196 community.

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199 4. Tables and Figures

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Table. 1 Socio-demographic information of children

Age-Groups (months)	Frequency (n = 74)	Percent (%)
1 – 11	26	35.1
12 – 59	48	64.9
Median \pmSD	11.0 \pm9.9	
Gender		
Male	40	54.1
Female	34	45.9

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Table 2. Cross-Tabulation of TB with age groups

Age-Groups (months)	TB positive	TB Negative	Chi-square (p-value)
1 – 11	6 (30.0)	20 (37.04)	0.31
12 – 59	14 (70.0)	34 (62.96)	(0.5733)**
Total	20 (100.0)	54 (100.0)	

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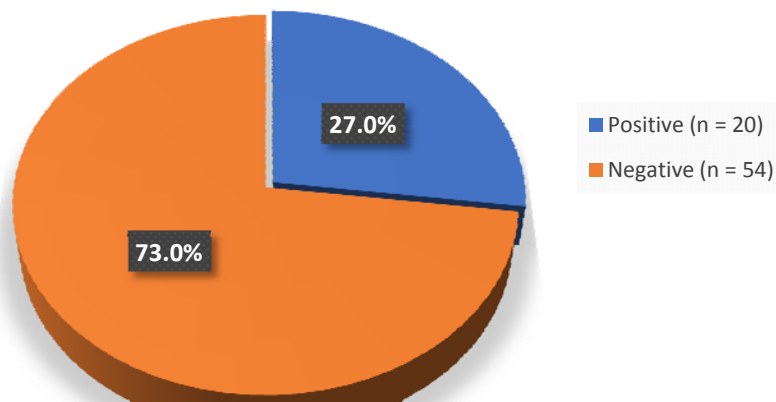
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Figure 1.

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Table 3. Distribution of Nutritional Status of Patients

Nutritional Status Of Patients	Frequency (%)
Marasmus	65 (87.80)
Kwashiokor	9(12.20)
Total	74(100)

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Table 4. Chest X-ray and GeneXpert MTB RIF

	CXR (%)	GeneXpert MTB RIF (%)
Test Done	15 (20.27)	13 (17.57)
Not Done	59(79.73)	61 (82.43)
Total	74 (100.0)	74 (100.0)

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Table 5. Socio-demographic information of parents

Variables	Frequency (n = 74)	Percent (%)
Age Groups (years)		
20 - 29	59	79.7
30 - 39	11	14.9
≥ 40	4	5.4
Mother's Occupation		
Unemployed	8	10.80
Trader	45	60.80
Student	13	17.60
Teacher	1	1.35
Tailor	1	1.35
Electrician	1	1.35
Hair dressing	3	4.05
Driver	1	1.35
Caterer	1	1.35
Education		
No Formal Education	15	20.27
Primary	5	6.76
Secondary	50	67.57
Tertiary	4	5.4

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Table 6. Cross tabulation of Mother's education and TB Prevalence

Education	TB Positive	TB Negative	Chi-square (p-value)
No Formal Education	1 (5.0)	14 (25.9)	3.95 (0.0467)*
Primary	2 (10.0)	3 (5.6)	0.45 (0.4987)**
Secondary	15 (75.0)	35 (64.8)	0.85 (0.3559)**

Tertiary	2 (10.0)	2 (3.7)	1.13 (0.2874)**
Total	20 (100.0)	54 (100.0)	

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Table 7. Cross tabulation of Gender and TB among children

Gender	TB Positive	TB Negative	Chi-square (p-value)
Male	9 (45.0)	31 (57.4)	0.90 (0.3415)**
Female	11 (55.0)	23 (42.6)	
Total	20 (100.0)	54 (100.0)	

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**Difference between both groups is not statistically significant (p > 0.05)

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Table 8. BCG Vaccination Status of Patients

Vaccine	TB POSITIVE (%)	TB NEGATIVE (%)	TOTAL (%)
BCG	20(27.0)	54(73.0)	74(100)

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245 **Competing Interest** There was no competing interest in this study

246 **Consent.** Written informed consent was obtained from participants

247 **Ethical Approval.** There no ethical Issues in this study, given that it was a descriptive
248 cohort study, WHO exempts it from ethical approval.

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