

1 **Prevalence of Enterobacteriaceae Isolated from Childhood Diarrhoea in Mukuru Slums,**  
2 **Nairobi- Kenya**

3  
4 **Abstract**

5 Diarrhoea in young children continues to be a major public health concern in developing  
6 countries, including Kenya. Poor sanitation among other factors can predispose a child to  
7 diarrhoea. Therefore, the present study sought to determine the prevalence of enterobacteriaceae  
8 isolated from childhood diarrhoea in Mukuru Slums, Nairobi. . It employed a cross-sectional  
9 design targeting children below 5 years of age. Stool specimens were obtained aseptically and  
10 cultured on MacConkey agar and Salmonella Shigella agar. Biochemical tests were used to  
11 identify the isolated bacteria to genus and species using biochemical characterization scheme and  
12 the Analytic Profile Index 20E. Drugs sensitivity tests were done using standard techniques.  
13 *Escherichia coli* ATCC 25922 was included as a control strain. Analysis of gender verses  
14 diarrhoea revealed that ( $p = 0.146 > 0.05$ ) there was no statistical significant association between  
15 the gender (male and female) and area of residence in relation to diarrhoea in this study. There  
16 was no statistical significant difference between the participants characteristics and their area of  
17 residence ( $p= 0.144$ ). Age of the participants had significant association with the prevalence of  
18 diarrhoea ( $p=0.00$ ). The *E. coli* bacteria showed the highest percentage of enteric pathogens  
19 isolated (35.2%) from female children at Mukuru kwa Njenga and 29.4% from male children,  
20 *Salmonella spp* being second (4.9%) from female at Mukuru kwa Reuben and the least was  
21 *Shigellasonnei* (3.2%) from female children at Sinai. Emphasis should therefore be placed on  
22 primary preventive measures such as ensuring good sewerage management and safe supply of  
23 drinking water in the study area and Kenya at large especially in the slums.

24  
25 **Keywords: Enterobacteriaceae, Diarrhoea, prevalence, Mukuru slums**

26  
27 **Introduction**

28  
29 Diarrhoea is a significant health problem globally, particularly in the developing world where  
30 adequate sanitation facilities are lacking (Okeke et al., 2000). A study by Black et al. (2003)  
31 reported that globally, diarrhoeal diseases account for almost a fifth of all deaths of children  
32 below five years of age, with an estimated 2.2 million deaths annually. Epidemiological studies  
33 of diarrhoea have been reported from several African countries including Kenya (Sang et al.,  
34 2012). In sub-Sahara Africa, an estimated 16% of deaths in children below 5 years of age are  
35 diarrhoea related (Bryce et al., 2005). Human Immunodeficiency Virus (HIV) is also prevalent in  
36 Sub-Sahara Africa and diarrhoea can exacerbate HIV related symptoms (Obimbo et al., 2004).  
37 Studies have shown that prolonged episodes of diarrhoea in early childhood leads to stunting  
38 (FAO, 2008; WHO, 2009). Poverty, poor sanitation and lack of balanced diet are also risk factors  
39 in diarrhoeal diseases (MOH, 2010). In Kenya, under five year's mortality rate is seventy four  
40 (74) deaths per 1000(KDHS, 2010). Sixteen per cent (16%) of children under five are  
41 underweight using weight for age index (KDHS, 2010). In Nairobi county, stunting in children  
42 increased by 4% in 2010 from an earlier survey done in 2003 (KDHS, 2010). Diarrhoea episodes  
43 increase with age peaking at six to eleven months at 30% experiencing diarrhoea because during

44 this age bracket most of the children will have started crawling while others are already walking  
45 (KDHS, 2010). The causes of diarrhoea include a wide array of viruses, parasites and bacteria.  
46 However, most of the diarrhoeal diseases are caused by the members of the family  
47 enterobacteriaceae (Lakshmi *et al.*, 2014). Farmer (2003) reported that these pathogens are  
48 named as enteric pathogens which belong to the genera that initiate infection by invading the  
49 intestinal epithelium. The researcher furthermore explained that the enteric pathogens  
50 belonging to the family enterobacteriaceae are predominantly facultative anaerobic  
51 bacterial flora of large intestine of human beings. These are generally non-spore forming, non  
52 acid fast and gram negative straight or curved rod.

53 The enteric disease causing members of family *Enterobacteriaceae* are *E.coli*, *Shigella*,  
54 *Salmonella*, *Proteus*, *Klebsiella pneumonia*, *Citrobacter freundii*, *Enterobacter aerogenes*.  
55 Some enteric organisms, for example, *Escherichia coli* are part of the normal flora and  
56 incidentally cause disease while others such as salmonellae and shigellae, are regularly  
57 pathogenic to humans (Abbott, 2003; Kariuki *et al.*, 2013). The enterobacteraceae are facultative  
58 anaerobes or aerobes, ferment a wide range of carbohydrates, possess a complete antigenic  
59 structure, and produce a variety of toxins and other virulence factors (Sang *et al.*, 2012).  
60 Enterobacteraceae, enteric gram-negative rods and enteric bacteria may also be called coli forms  
61 (Farmer, 2003). Children living in the slums are vulnerable to diarrhoeal diseases mainly due to  
62 poor sanitation. Therefore, the present study seeks to study sought to determine the prevalence of  
63 enterobacteriaceae isolated from childhood diarrhoea in Mukuru Slums, Nairobi.

64

## 65 **Materials and methods**

### 66 **Study site**

67 The study site was the government hospital located at Mukuru Kwa Njenga slum in Nairobi  
68 County. The Hospital serves the residents of Kwa Reuben, Kwa Njenga, Kayaba and Sinai slums  
69 along Nairobi River. It is situated within the Industrial area of Nairobi city lying at co-ordinates  
70 1°18'33"S 36°48'12"E (KBS, 2009). Mukuru Kwa Njenga is a slum in the East of Nairobi, the  
71 capital of Kenya. It belongs to Embakasi Constituency. It is one of the largest slums in Nairobi.  
72 Among other major slums in Nairobi are Korogocho, Kibera and Mathare. The population of the  
73 slum exceeds 100,000. There have been cholera deaths in 2009 (WHO, 2010).

### 74 **Study design and population**

75 The study employed a cross-sectional laboratory based design (Fischer *et al.*, 1986). The study  
76 population comprised of children who were five years and below, attended to at the government  
77 health facility in Mukuru Kwa Njenga.

### 78 **Sample size determination**

79 The sample size was determined using the formula below according to Fischer *et al.* (1986)

80 
$$n = \frac{Z^2 \times P(1-p)}{d^2} \dots\dots\dots [1]$$

81 Where n is the sample size, z is the confidence interval at 95% and p is the prevalence got from  
82 Kenya Demographic health survey (KDHS), 2010, d is the margin of error at 5%

83 
$$n = \frac{1.96^2 \times 0.17 (0.83)}{0.05^2} \dots\dots\dots [2]$$

84 Final sample size was 178 stool samples/ anal swabs

85 N = 178; 190 participants were included in this study. The study sampled 190 children to take  
86 care of specimen that might get spoiled on the way before reaching the laboratory for processing.

87 **Sample collection**

88 Stool samples were collected into sterile, wide-mouthed, screw cap containers and preserved in  
89 cool boxes. Anal swabs were collected from participants who were unable to produce stool  
90 samples and the specimens were labelled and assigned unique code numbers during the time of  
91 sample collection. Specimens once collected were taken to the centre for microbiology research  
92 laboratory (CMR)-KEMRI within the shortest time possible for processing.

93 **Specimen processing**

94 **Culturing**

95 The specimens were enriched in selenite F media overnight at 37°C. After enrichment,  
96 inoculations were done both on MacConkey Agar and Shigella Salmonella Agar (Oxoid,  
97 Basingstoke, United Kingdom). Lactose fermenters and non-lactose fermenters that had grown  
98 colonies were inoculated onto biochemically impregnated API 20E strips (BioMerieux,  
99 Basingstoke, United Kingdom) for identification.

100 **Biochemical tests**

101 **Triple sugar iron agar (TSI)**

102 Colonies were selected on plate using a sterile straight wire loop. The centre of the colony was  
103 lightly touched and prepared TSI medium were inoculated by stabbing the butt and streaking the  
104 slants. These were then incubated at 37°C for 24 hours (Cowan and Steel, 2002).

105 **Indole test**

106 The bacteria isolated were sub-cultured in nutrient broth and incubated for 24 hours. About 3  
107 drops of Kovac’s indole reagent was added and mixed gently (Cheesbrough, 2005).

108 **Urease test**

109 Urea agar was inoculated heavily over the entire surface of the slants in bijoux bottles, incubated  
110 at 37°C for 24 hours.

111 **Citrate utilization test**

112 Simmons citrate slopes were prepared in bijoux bottles. The slopes were then stabbed and  
113 incubated at 37°C for 48 hours.

114 **Motility test**

115 A sterile straight wire loop was used to inoculate motility indole urea media with bacterial isolate  
116 and incubated overnight at 37°C. Motility was shown by diffused turbidity in the medium  
117 (Cheesbrough,2005).

118 N/B: All these tests mentioned above were used for the purpose of identification of  
119 Enterobacteriaceae. The results were either positive or negative for a particular entero pathogen.

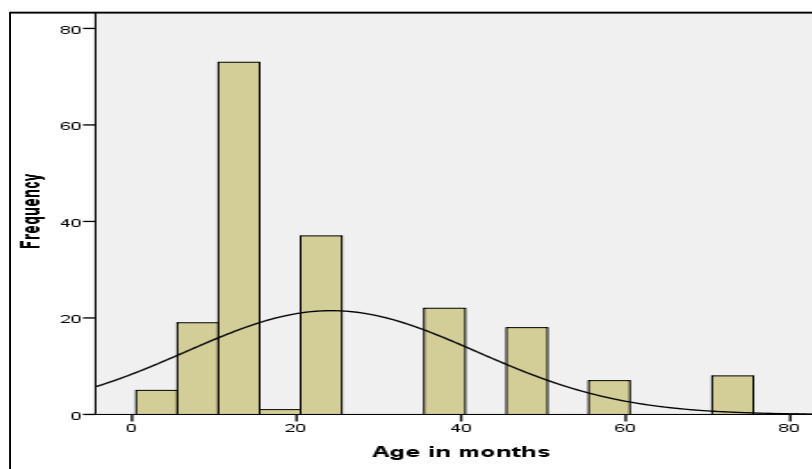
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121 **Results and Discussion**

122 **Participants’ characteristics**

123 A total number of 190 children below the age of five years presenting with diarrhoea in the  
124 Government health facility in Mukuru kwa Njenga slum participated in this study. The mean age

125 of the participants was 24.21 months with the youngest child being 3 months and the oldest child  
 126 being 72 months. More children who participated in the study were less than 40 months in age.  
 127 The children's ages were skewed to the right of the normal curve (Figure 1). The mean age of the  
 128 children was twice more than the median age with a standard deviation of 17.62. The study  
 129 recorded a significant association ( $p < 0.05$ ) between the age groups and diarrhoea among the  
 130 participants.  
 131



132  
 133 **Figure 1: Distribution curve of participants ages**

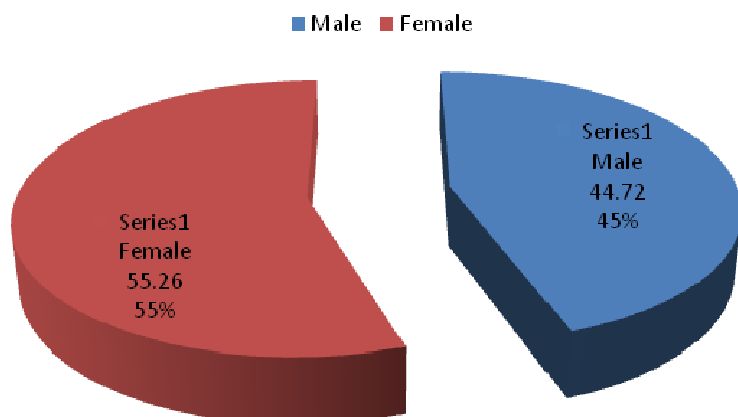
134 In this study the female children were 105(55.26%) and the males were 85(44.74%). Female  
 135 children were 3.7 months older than the male children with a standard difference error mean of  
 136 0.02 months as shown in Table1 below. These results could be due to the fact that children  
 137 within this age group are most often than not unaccompanied and cannot differentiate between  
 138 what to eat and what not to eat; they have not learnt the rules of adherence to aseptic or hygienic  
 139 practice and they can barely express themselves (Sang *et al.*, 2012). Those below the age of  
 140 twelve months are essentially under their mothers' care, feeding mainly on breast milk thereby  
 141 reducing their susceptibility to these pathogens.  
 142

143 **Table 1: Analysis of age of the participants**

Gender	N	Percentage	Mean age	Age Stdev.	Age SE. Mean	Std. Diff	P-value
Male	85	44.74%	16.414	1.780	16.414	0.02	0.00
Female	105	55.26%	18.449	1.800	18.449		

144 Age can be a predisposing factor to diarrhoea in children below the age of five years (WHO,  
 145 2007). Living in the slums is also a predisposing factor to diarrhoeal infections because of the  
 146 poor hygienic conditions coupled with poor sanitation (WHO, 2010).  
 147 Most enteric pathogens stimulate at least partial immunity against repeated infections or illness,  
 148 which helps to explain the declining incidence of diseases in older children (Patwari *et al.*, 1993).  
 149 The analysis of the participants' ages verses gender revealed that there was no significant  
 150 difference. The  $t_{(186)}$  value was 1.458 with probability,  $p = 0.146 > 0.05$ , the p-value was more  
 151 than 0.05 therefore there was no association between the gender in relation to diarrhoea in this

152 study. The male participants were 85(45%) while the female were 105(55%) as shown in Figure  
153 2. There was significant association between age and diarrhoea in this study (p=0.01).



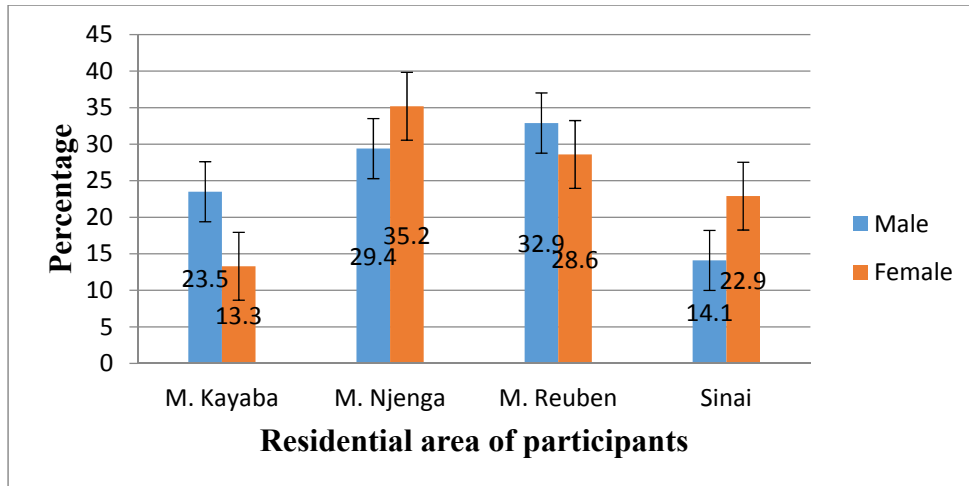
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155 **Figure 2: Gender of the participants**

156 The participants attended to at Mukuru Kwa Njenga government health facility were noted to be  
157 residents of four neighbouring slums namely; Mukuru Kwa Njenga, Mukuru Kwa Reuben,  
158 Mukuru Kayaba and Sinai. The majority of the participants were from Mukuru Kwa Njenga  
159 61(32.6%) followed by Mukuru Kwa Reuben 57(30.5%) then Sinai 35(18.9%) and the least were  
160 from Mukuru Kayaba 33(17.9%). Mukuru Kwa Njenga had the highest number of female  
161 children (35.2%) while Mukuru Kwa Reuben had the highest number of male children (32.9%).  
162 The p-values were greater than 0.05 hence there was no significance difference between the  
163 participants from different areas of residence ( $\chi^2=5.41$ ,  $p=0.144$ ) as shown in Figure 3.

164 The results of other studies concur with the current study. Chitnis et al. (2012) in their study  
165 observed that patients susceptible to Carbapenem-resistant enterobacteriaceae (CRE) were more  
166 likely to be female. The results of the current study concurs with a study done by Sule et  
167 al.(2011) in Kaduna Nigeria where they found the incidence between both sexes showing female  
168 children having the highest percentage (26%) compared to males (18%). Abdullahi et al. (2010)  
169 reported that male children were more infected (22.33%) than female children (18.33%),  
170 although the difference was not statistically significant ( $\chi^2 = 0.531$ ,  $p>0.05$ ) hence contradicting  
171 the finding of the current study. Most diarrhoeal episodes occur during the first two years of life  
172 due to a combination of factors; declining levels of maternal acquired antibodies, lack of active  
173 immunity in the infant, the introduction of food that may be contaminated with enteric bacteria  
174 or direct contact with human or animal faeces carrying enteric bacteria when the infant starts to  
175 crawl (Sang, 2007).

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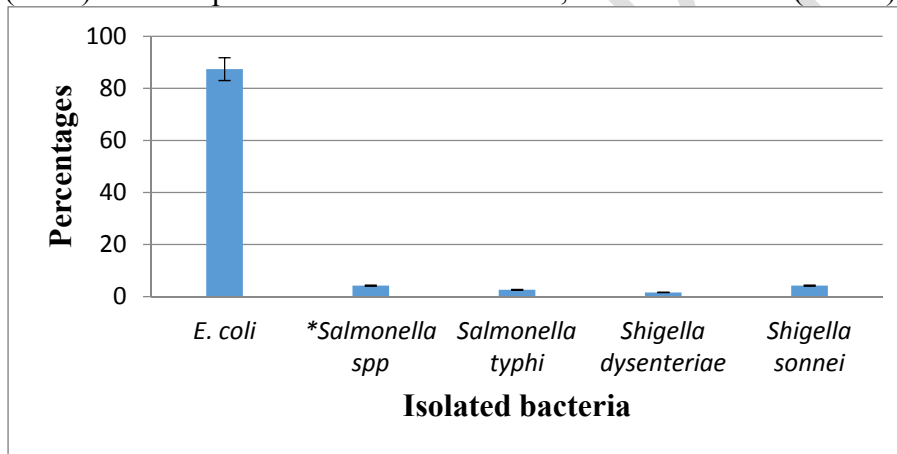


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178 **Figure 3: Residence of study participants**

179 **Isolation and identification of the bacteria**

180 The prevalence of bacteria isolated from the study were as follows: *Escherichia coli*(87.4%),  
 181 *Salmonella spp*(4.2%), *Shigella sonnei* (4.2%), *Salmonella typhi* (2.6%), *Shigella dysenteriae*  
 182 (1.6%) and. The prevalence were as follows; *Escherichia coli* (87.4%), as shown in Figure 4.



183

184 **Figure 4: Bacteria species isolated from the stool samples**

185

186 The total percentage prevalence of bacteria species isolated among the participants by age,gender  
 187 and residence in the study area were 78% *E. coli*, 4.2% *Salmonella spp*(were not identified to  
 188 species level), 4.2% *Shigella sonnei*, 2.6% *Salmonella typhi* and 1.6% *Shigella dysenteriae* from  
 189 all the specimens collected. At Mukuru Kwa Njenga more *E. coli*were isolated from female  
 190 children (35.2%) than male children (29.4%) the rest of the isolates were uniform in both  
 191 genders. At Mukuru Kwa Reuben the trend is the same in that more *E. coli*were also isolated  
 192 from female children (17.5%) than from male children (13.0%). *Salmonella spp* were 4.9% from  
 193 female children and 0.0% from male children while *S. typhi* were more from male children  
 194 (3.0%) compared to female children (0.5%). At Sinai the percentage isolates from both male and  
 195 female children were almost equal (9.0% and 9.9%, respectively).*Shigella sonnei* were more

196 from (3.2) female than male children (1.0%). The rest were almost the same in both male and  
 197 female children. At Kayaba *E. coli* isolates were more from female (10.7%) than from male  
 198 children (7.2%). *S. typhi* were 1.6% in females and 0.0% in males while the rest were 0.0%.  
 199 There was no significant association between the gender and percentage isolates ( $p > 0.05$ ). There  
 200 was also no significant association between the prevalence of the isolates and the area of  
 201 residence of the children ( $\chi^2 = 2.23$ ,  $p = 0.693$ ). The results are as shown in Table 2.

202 **Table 2: Prevalence of bacteria isolated by gender and residence of participants**

Residence	Isolated spp	Male (% isolates)	Female (% isolates)	$\chi^2$ (p-value)
M. Njenga	<i>E. coli</i>	29.4	35.2	2.23 (0.693)
	* <i>Salmonella spp</i>	1.0	1.1	
	<i>S. typhi</i>	0.0	0.0	
	<i>S. dysenteriae</i>	1.5	1.5	
	<i>Shigella sonnei</i>	1.1	1.0	
M. Reuben	<i>E. coli</i>	13.0	17.5	
	* <i>Salmonella spp</i>	0.0	4.9	
	<i>S. typhi</i>	3.0	0.5	
	<i>S. dysenteriae</i>	1.0	1.1	
	<i>Shigella sonnei</i>	2.9	2.0	
Sinai	<i>E. coli</i>	9.0	9.9	
	* <i>Salmonella spp</i>	2.0	2.2	
	<i>S. typhi</i>	1.6	1.0	
	<i>S. dysenteriae</i>	0.0	1.6	
	<i>Shigella sonnei</i>	1	3.2	
M. Kayaba	<i>E. coli</i>	7.2	10.7	
	* <i>Salmonella spp</i>	0.0	0.0	
	<i>S. typhi</i>	0.0	1.6	
	<i>S. dysenteriae</i>	0.0	0.0	
	<i>Shigella sonnei</i>	0.0	0.0	

203 \**Salmonella spp*- other *Salmonella* isolates which were not identified to species level, *Spp* -  
 204 species,  $\chi^2$  – Chi square test, p-value- level of significance (0.05)

205  
 206 Acute diarrhoea due to bacterial infections is an important cause of morbidity and mortality in  
 207 infants and young children in most developing countries including Kenya especially in the slums

208 (Adegunloye, 2005). Identification of the enteropathogens causing diarrhoeal diseases in the  
209 country is an essential step towards the implementation of effective primary health care activities  
210 against the disease (Olowe et al., 2003). Poor sanitation in the study area could have also  
211 contributed to the high prevalence of bacteria isolated. The residents live in congested  
212 environments with their domesticated animals which could have contributed to the high  
213 prevalence of isolated enteric bacteria. According to a study done by Kariuki et al. (2006), a  
214 significantly higher proportion of younger children (< 3 years of age) and those from the slums  
215 presented with invasive non- typhoidal *Salmonella spp* compared to older children and those  
216 from upper socio-economic groups ( $p < 0.001$ ).

217 In terms of gender and area of residence, Mukuru kwa Njenga, had more *E. coli* isolated from  
218 female children (35.2%) compared to male children (29.4%) the rest of the isolates were  
219 uniform in both genders. In Mukuru kwa Reuben the trend was the same in that more *E. coli*  
220 were also isolated from female children (17.5%) than from male children (13.0%). *Salmonella*  
221 *spp* were 4.9% from female children and 0.0% from male children while *S. typhi* were more from  
222 male children (3.0%) compared to female children (0.5%). At Sinai the percentages of the  
223 isolates from both male and female children were almost equal (9.0% and 9.9%, respectively).  
224 *Shigella sonnei* were more from (3.2) female than male children (1.0%). The rest were almost the  
225 same in both male and female children. At Mukuru Kayaba *E. coli* isolates were more from  
226 female (10.7%) than male children (7.2%). *S typhi* were 1.6% in females and 0.0% in males  
227 while the rest were 0.0%. There was no significant association between the gender and  
228 percentage isolates ( $p > 0.05$ ). There was also no significant association between the prevalence of  
229 the isolates and the area of residence of the children ( $\chi^2 = 2.23$ ,  $p = 0.693$ ). The results of this study  
230 do not concur with what Sang et al. (2013) found in their studies on the prevalence of bacteria in  
231 four provinces in Kenya where they had recruited 651 participants and isolated pathogenic  
232 bacteria in (17.7%) of the participants. Among the isolated bacteria were; pathogenic *E. coli*  
233 (11.2%), *Salmonella* (3.5%), *Shigella* (2.3%) and *Vibrio cholera* (0.6%) (Sang, 2007). The reason  
234 for the different results could be because the study area was basically a slum hence the high  
235 prevalence of bacteria isolated especially the *E. coli*.

236 A similar study was done by Ifeanyi et al. (2010) in Abuja Nigeria among cases of diarrhoea  
237 with potential bacterial pathogens detected being 65.8% of all patients screened. This was in  
238 contrast to a report of the prevalence of 83.1% from similar study in Abakaliki, south –eastern  
239 Nigeria (Ogbu et al., 2008). Another study reported a prevalence of 63.3%-71.83% isolation of  
240 enteric bacteria in ifakara Tanzania (Vargas et al., 2004). The variation in prevalence between  
241 the two Nigerian cities might be attributed to differences in infrastructural and socioeconomic  
242 indices (Ogbu et al., 2012). In a different study, the prevalence of bacterial aetiology of  
243 diarrhoea was 44% which follows the same trend with the research conducted in Kano State  
244 which was found to be 40.67% (Tsang et al., 2009; Abdullahi et al., 2010). In Gabon prevalence  
245 of diarrhoea with bacterial aetiology was 38% (Patwari et al., 1993). In Tanzania it was 36%  
246 (Molbak et al., 1997). The study showed that *Shigella spp* appears to be the predominant  
247 bacteria causing diarrhoea followed by *E. coli*, and *Salmonella* in that order. A total of 56% of  
248 the hundred diarrhoea cases investigated had no bacterial pathogen suggesting viral, protozoan or  
249 nonpathogenic factors (Abdullahi et al., 2010).

250 *Salmonella spp* isolated in Mukuru slums could be non- typhoidal salmonella which is a zoonotic  
251 strain. The children could have been contaminated with faecal matter of the domesticated  
252 animals hence the acquisition of the bacteria. Occurrence of diarrhoeagenic bacteria in the  
253 current study showed that gram negative bacteria (*Shigella spp*, *Salmonella spp*, *Escherichiacoli*)



254 are the main cause of bacterial diarrhoea. Sule et al. (2011) in Kaduna Nigeria conducted a  
255 similar study and found similar results. Generally, the aetiology of diarrhoea in young children  
256 could be attributed to a wide range of factors, but one of the main causes of diarrhoea is related  
257 to bacteria (such as *Salmonella spp*, *Shigella spp*, *Vvibrio*, *Escherichia coli*, *Aeromonas* and  
258 *Pseudomonas* (Abdullahi et al., 2010). Results from the current study shows that, though there  
259 are a number of causative agents of diarrhoeal diseases, bacteria still remain one of the major  
260 causes with *Shigella*, *Salmonella* and *Escherichia coli* being the most important pathogens  
261 among paediatric patients presenting with diarrhoea in Mukuru kwa Njenga Government health  
262 facility. Judicious use of antibiotic therapy requires education of health workers and patients,  
263 adequate laboratory diagnostic capabilities and government regulations.

#### 264 **Conclusion**

265 In this study the female participants were more than the males. Mukuru Kwa Njenga had the  
266 highest (35.2%) number of female children while Mukuru Kwa Ruben had the highest (32.9)  
267 number of male children. There was no statistical significant difference between the participants  
268 characteristics and their area of residence ( $p=0.144$ ). Age of the participants had significant  
269 association with the prevalence of diarrhoea ( $p=0.00$ ). The total prevalence of isolated bacteria  
270 among the participants was very high (90.6%). The *E. coli* bacteria showed the highest  
271 percentage of enteric pathogens isolated (35.2%) from female children at Mukuru Kwa Njenga  
272 and 29.4% from male children, *Salmonella spp* being second (4.9%) from female at Mukuru  
273 Kwa Reuben and the least was *Shigellasonnei* (3.2%) from female children at Sinai.

#### 274 **Recommendation**

275 Further studies should investigate social demographic characteristics of children, parents and  
276 their households in order to understand more the causes and predisposing factors of diarrhoea in  
277 the slums.

#### 278 **Conflict of Interest**

279 The authors declare no conflict of interest

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