

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 [1]. A study by Black et al. [2] reported that globally,
31 diarrhoeal diseases account for almost a fifth of all deaths of children below five years of age,
32 with an estimated 2.2 million deaths annually. Epidemiological studies of diarrhoea have been
33 reported from several African countries including Kenya [3]. In sub-Sahara Africa, an estimated
34 16% of deaths in children below 5 years of age are diarrhoea related [4]. Human
35 Immunodeficiency Virus (HIV) is also prevalent in Sub-Sahara Africa and diarrhoea can
36 exacerbate HIV related symptoms [5].

37 Studies have shown that prolonged episodes of diarrhoea in early childhood leads to stunting [6].
38 Poverty, poor sanitation and lack of balanced diet are also risk factors in diarrhoeal diseases [7].
39 In Kenya, under five year's mortality rate is seventyfour (74) deaths per 1000[8]. Sixteen per
40 cent (16%) of children under five are underweight using weight for age index [8]. In Nairobi
41 county, stunting in children increased by 4% in 2010 from an earlier survey done in 2003 [8].
42 Diarrhoea episodes increase with age peaking at six to eleven months at 30% experiencing
43 diarrhoea because during this age bracket most of the children will have started crawling while

44 others are already walking [8]. The causes of diarrhoea include a wide array of viruses, parasites
45 and bacteria. However, most of the diarrhoeal diseases are caused by the members of the family
46 **Enterobacteriaceae** [9]. Farmer [10] reported that these pathogens are named as enteric
47 pathogens which belong to the genera that initiate infection by invading the intestinal
48 epithelium. The researcher furthermore explained that the enteric pathogens belonging to
49 the family **Enterobacteriaceae** are predominantly facultative anaerobic bacterial flora of large
50 intestine of human beings. These are generally non-spore forming, non acid fast and gram
51 negative straight or curved rod.

52 The enteric disease causing members of family *Enterobacteriaceae* are *E.coli*, *Shigella*,
53 *Salmonella*, *Proteus*, *Klebsiella pneumonia*, *Citrobacter freundii*, *Enterobacter aerogenes*.
54 Some enteric organisms, for example, *Escherichia coli* are part of the normal flora and
55 incidentally cause disease while others such as salmonellae and shigellae, are regularly
56 pathogenic to humans [11;12]. The **Enterobacteraceae** are facultative anaerobes or aerobes,
57 ferment a wide range of carbohydrates, possess a complete antigenic structure, and produce a
58 variety of toxins and other virulence factors [13]. Enterobacteraceae, enteric gram-negative rods
59 and enteric bacteria may also be called **coliforms** [10]. Children living in the slums are
60 vulnerable to diarrhoeal diseases mainly due to poor sanitation. Therefore, the present study
61 seeks to study sought to determine the prevalence of **Enterobacteriaceae** isolated from childhood
62 diarrhoea in Mukuru Slums, Nairobi.

63

64 **Materials and methods**

65 **Study site**

66 The study site was the government hospital located at Mukuru Kwa Njenga slum in Nairobi
67 County. The Hospital serves the residents of Kwa Reuben, Kwa Njenga, Kayaba and Sinai slums
68 along Nairobi River. It is situated within the Industrial area of Nairobi city lying at co-ordinates
69 1°18'33"S 36°48'12"E. Mukuru Kwa Njenga is a slum in the East of Nairobi, the capital of
70 Kenya. It belongs to Embakasi Constituency. It is one of the largest slums in Nairobi. Among
71 other major slums in Nairobi are Korogocho, Kibera and Mathare. The population of the slum
72 exceeds 100,000. There have been cholera deaths in 2009 [6].

73 **Study design and population**

74 The study employed a cross-sectional laboratory based design [14]. The study population
75 comprised of children who were five years and below, attended to at the government health
76 facility in Mukuru Kwa Njenga **with signs and symptoms of diarrheal diseases.**

77 **Inclusion criteria**

- 78 • Children under five years verified by child welfare clinic records.
- 79 • Children who had diarrhoea or history of diarrhoea i.e. passage of loose or watery
80 stool more than three times a day (WHO, 1988).
- 81 • HIV negative.
- 82 • **Children** whose parents/guardians accepted to sign informed consent form

83 **Sample size determination**

84 The sample size was determined using the formula below according to [14]

$$85 \quad n = \frac{Z^2 \times P(1-p)}{d^2} \dots\dots\dots [1]$$

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

$$88 \quad n = \frac{1.96^2 \times 0.17 (0.83)}{0.05^2} \dots\dots\dots [2]$$

89 Final sample size was 178 stool samples/ anal swabs

90 N = 178; 190 participants were included in this study.

91 **Sample collection**

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

97 **Specimen processing**

98 **Culturing**

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

104 **Biochemical tests**

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

106 Colonies were selected on plate using a sterile straight wire loop. The centre of the colony was
107 lightly touched and prepared TSI medium were inoculated by stabbing the butt and streaking the
108 slants. These were then incubated at 37°C for 24 hours [15].

109 **Indole test**

110 The bacteria isolated were sub-cultured in nutrient broth and incubated for 24 hours. About 3
111 drops of Kovac's indole reagent was added and mixed gently [16].

112 **Urease test**

113 Urea agar was inoculated heavily over the entire surface of the slants in bijou bottles, incubated
114 at 37°C for 24 hours.

115 **Citrate utilization test**

116 Simmons citrate slopes were prepared in bijou bottles. The slopes were then stabbed and
117 incubated at 37°C for 48 hours.

118 **Motility test**

119 A sterile straight wire loop was used to inoculate motility indole urea media with bacterial isolate
120 and incubated overnight at 37°C. Motility was shown by diffused turbidity in the medium [16].

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

123 **Ethical Consideration**

124 The study was nested within a bigger study which was funded by The Centre for Disease Control
125 and Prevention in collaboration with the Kenya Medical Research Institute, Opportunistic

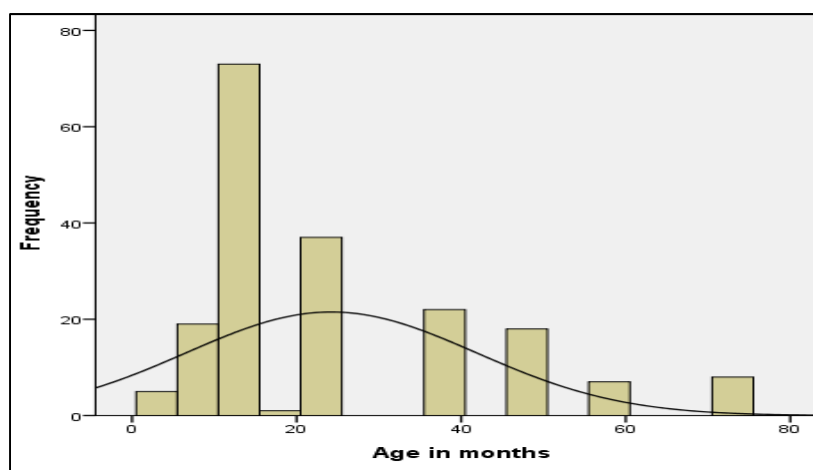
126 infection laboratories and the ministry of health central microbiology laboratories. Permission to
 127 carry out the study was granted by the investigators of the main study.

128 **Results and Discussion**

129 **Participants' characteristics**

130 A total number of 190 children below the age of five years presenting with diarrhoea in the
 131 Government health facility in Mukuru kwa Njenga slum participated in this study. The mean age
 132 of the participants was 24.21 months with the youngest child being 3 months and the oldest child
 133 being 72 months. More children who participated in the study were less than 40 months in age.
 134 The children's ages were skewed to the right of the normal curve (Figure 1). The mean age of the
 135 children was twice more than the median age with a standard deviation of 17.62. The study
 136 recorded a significant association ($p < 0.05$) between the age groups and diarrhoea among the
 137 participants.

138



139

140 **Figure 1: Distribution curve of participants ages**

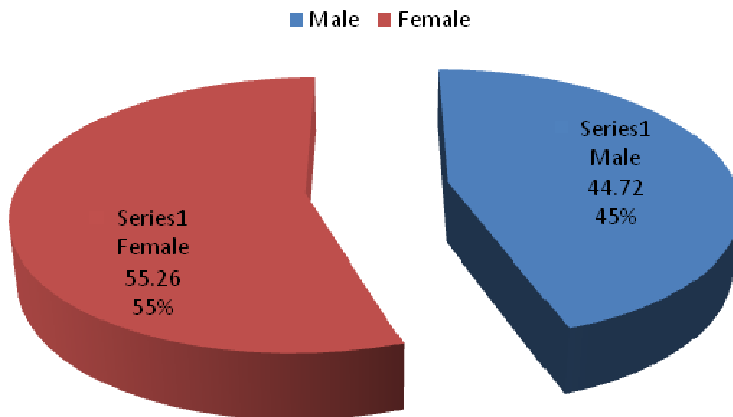
141 In this study the female children were 105(55.26%) and the males were 85(44.74%). Female
 142 children were 3.7 months older than the male children with a standard difference error mean of
 143 0.02 months as shown in Table 1 below. These results could be due to the fact that children
 144 within this age group are most often than not unaccompanied and cannot differentiate between
 145 what to eat and what not to eat; they have not learnt the rules of adherence to aseptic or hygienic
 146 practice and they can barely express themselves [13]. Those below the age of twelve months are
 147 essentially under their mothers' care, feeding mainly on breast milk thereby reducing their
 148 susceptibility to these pathogens.

149

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

Gender	N	Percentage	Mean age	Age Stdev.	Age SE. Mean	SD	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		

151 Age can be a predisposing factor to diarrhoea in children below the age of five years (WHO,
 152 2007). Living in the slums is also a predisposing factor to diarrhoeal infections because of the
 153 poor hygienic conditions coupled with poor sanitation [6]
 154 Most enteric pathogens stimulate at least partial immunity against repeated infections or illness,
 155 which helps to explain the declining incidence of diseases in older children [17].
 156 The analysis of the participants' ages versus gender revealed that there was no significant
 157 difference. The $t_{(186)}$ value was 1.458 with probability, $p = 0.146 > 0.05$, the p-value was more
 158 than 0.05 therefore there was no association between the gender in relation to diarrhoea in this
 159 study. The male participants were 85(45%) while the female were 105(55%) as shown in Figure
 160 2. There was significant association between age and diarrhoea in this study ($p=0.01$).



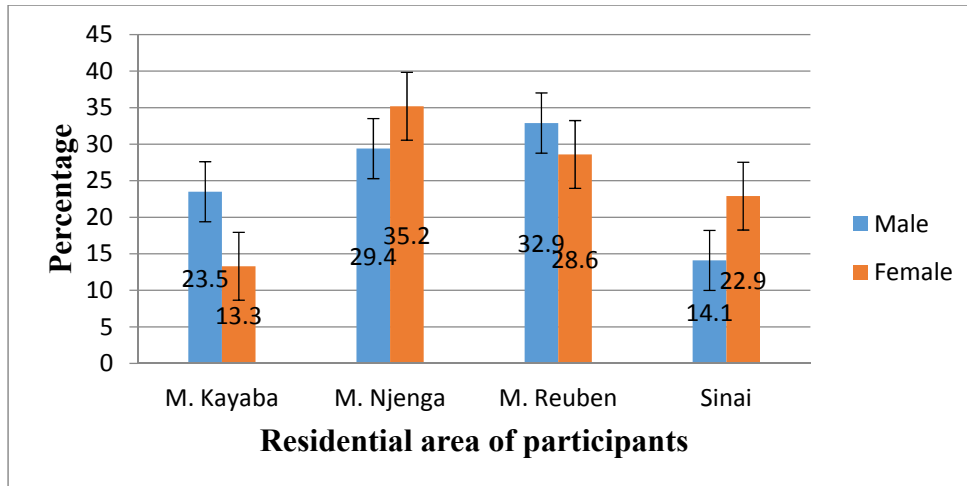
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162 **Figure 2: Gender of the participants**

163 The participants attended to at Mukuru Kwa Njenga government health facility were noted to be
 164 residents of four neighbouring slums namely; Mukuru Kwa Njenga, Mukuru Kwa Reuben,
 165 Mukuru Kayaba and Sinai. The majority of the participants were from Mukuru Kwa Njenga
 166 61(32.6%) followed by Mukuru Kwa Reuben 57(30.5%) then Sinai 35(18.9%) and the least were
 167 from Mukuru Kayaba 33(17.9%). Mukuru Kwa Njenga had the highest number of female
 168 children (35.2%) while Mukuru Kwa Reuben had the highest number of male children (32.9%).
 169 The p-values were greater than 0.05 hence there was no significance difference between the
 170 participants from different areas of residence ($\chi^2 = 5.41$, $p = 0.144$) as shown in Figure 3.

171 The results of other studies concur with the current study. Chitnis et al. [18] in their study
 172 observed that patients susceptible to Carbapenem-resistant enterobacteriaceae (CRE) were more
 173 likely to be female. The results of the current study concurs with a study done by Sule et al. [19]
 174 in Kaduna Nigeria where they found the incidence between both sexes showing female children
 175 having the highest percentage (26%) compared to males (18%). Abdullahi et al. (2010) reported
 176 that male children were more infected (22.33%) than female children (18.33%), although the
 177 difference was not statistically significant ($\chi^2 = 0.531$, $p > 0.05$) hence contradicting the finding of
 178 the current study. Most diarrhoeal episodes occur during the first two years of life due to a
 179 combination of factors; declining levels of maternal acquired antibodies, lack of active immunity
 180 in the infant, the introduction of food that may be contaminated with enteric bacteria or direct
 181 contact with human or animal faeces carrying enteric bacteria when the infant starts to crawl [3].

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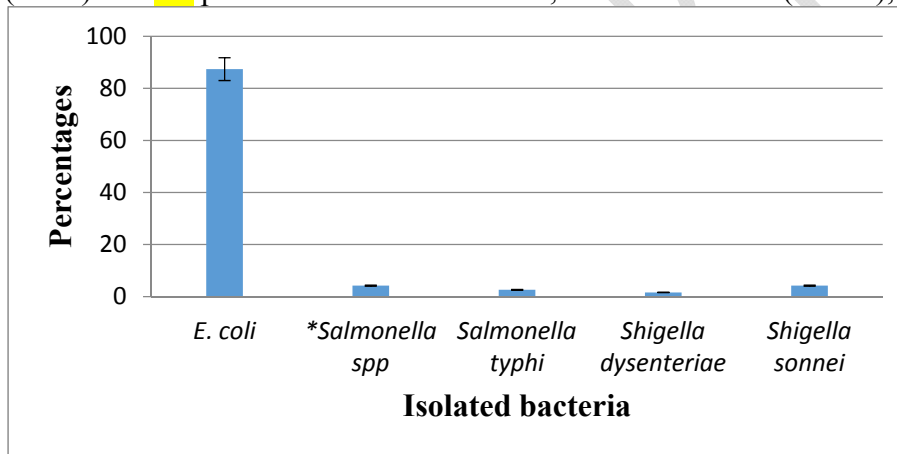


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184 **Figure 3: Residence of study participants**

185 **Isolation and identification of the bacteria**

186 The prevalence of bacteria isolated from the study were as follows: *Escherichia coli*(87.4%),
 187 *Salmonella spp*(4.2%), *Shigella sonnei* (4.2%), *Salmonella typhi* (2.6%), *Shigella dysenteriae*
 188 (1.6%) and the prevalence were as follows; *Escherichia coli* (87.4%), as shown in Figure 4.



189

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

191

192 The total percentage prevalence of bacteria species isolated among the participants by age,gender
 193 and residence in the study area were 78% *E. coli*, 4.2% *Salmonella spp*(were not identified to
 194 species level), 4.2% *Shigella sonnei*, 2.6% *Salmonella typhi* and 1.6% *Shigella dysenteriae* from
 195 all the specimens collected. At Mukuru Kwa Njenga more *E. coli*were isolated from female
 196 children (35.2%) than male children (29.4%) the rest of the isolates were uniform in both
 197 genders. At Mukuru Kwa Reuben the trend is the same in that more *E. coli*were also isolated
 198 from female children (17.5%) than from male children (13.0%). *Salmonella spp* were 4.9% from
 199 female children and 0.0% from male children while *S. typhi* were more from male children
 200 (3.0%) compared to female children (0.5%). At Sinai the percentage isolates from both male and
 201 female children were almost equal (9.0% and 9.9%, respectively).*Shigella sonnei* were more

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

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

Residence	Isolated spp	Male (% isolates)	Female (% isolates)	χ^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	

209 **Salmonella spp*- other *Salmonella* isolates which were not identified to species level, *Spp* -
 210 species, χ^2 - Chi square test, p-value- level of significance (0.05)

211
 212 Acute diarrhoea due to bacterial infections is an important cause of morbidity and mortality in
 213 infants and young children in most developing countries including Kenya especially in the slums

214 [20]. Identification of the **Enteropathogens** causing diarrhoeal diseases in the country is an
215 essential step towards the implementation of effective primary health care activities against the
216 disease [21]. Poor sanitation in the study area could have also contributed to the high prevalence
217 of bacteria isolated. The residents live in congested environments with their domesticated
218 animals which could have contributed to the high prevalence of isolated enteric bacteria.
219 According to a study done by Kariuki et al. [21], a significantly higher proportion of younger
220 children (< 3 years of age) and those from the slums presented with invasive non- typhoidal
221 *Salmonella spp* compared to older children and those from upper socio-economic groups ($p <$
222 0.001).

223 In terms of gender and area of residence, Mukuru kwa Njenga, had more *E. coli* isolated from
224 female children (35.2%) compared to male children (29.4%) the rest of the isolates were
225 uniform in both genders. In Mukuru kwa Reuben the trend was the same in that more *E. coli*
226 were also isolated from female children (17.5%) than from male children (13.0%). *Salmonella*
227 *spp* were 4.9% from female children and 0.0% from male children while *S. typhi* were more from
228 male children (3.0%) compared to female children (0.5%). At Sinai the percentages of the
229 isolates from both male and female children were almost equal (9.0% and 9.9%, respectively).
230 *Shigella sonnei* were more from (3.2) female than male children (1.0%). The rest were almost the
231 same in both male and female children. At Mukuru Kayaba *E. coli* isolates were more from
232 female (10.7%) than male children (7.2%). *S typhi* were 1.6% in females and 0.0% in males
233 while the rest were 0.0%. There was no significant association between the gender and
234 percentage isolates ($p > 0.05$). There was also no significant association between the prevalence of
235 the isolates and the area of residence of the children ($\chi^2 = 2.23$, $p = 0.693$). The results of this study
236 do not concur with what Sang et al. [6] found in their studies on the prevalence of bacteria in
237 four provinces in Kenya where they had recruited 651 participants and isolated pathogenic
238 bacteria in (17.7%) of the participants. Among the isolated bacteria were; pathogenic *E. coli*
239 (11.2%), *Salmonella* (3.5%), *Shigella* (2.3%) and *Vibrio cholera* (0.6%) [3]. The reason for the
240 different results could be because the study area was basically a slum hence the high prevalence
241 of bacteria isolated especially the *E. coli*.

242 A similar study was done by Ifeanyi et al. [22] in Abuja Nigeria among cases of diarrhoea with
243 potential bacterial pathogens detected being 65.8% of all patients screened. This was in contrast
244 to a report of the prevalence of 83.1% from similar study in Abakaliki, south –eastern Nigeria
245 [23]. Another study reported a prevalence of 63.3%-71.83% isolation of enteric bacteria in
246 ifakara Tanzania. The variation in prevalence between the two Nigerian cities might be attributed
247 to differences in infrastructural and socioeconomic [23]. In a different study, the prevalence of
248 bacterial aetiology of diarrhoea was 44% which follows the same trend with the research
249 conducted in Kano State which was found to be 40.67%. In Gabon prevalence of diarrhoea with
250 bacterial aetiology was 38% [17]. In Tanzania it was 36%. The study showed that *Shigella spp*
251 appears to be the predominant bacteria causing diarrhoea followed by *E. coli*, and *Salmonella* in
252 that order. A total of 56% of the hundred diarrhoea cases investigated had no bacterial pathogen
253 suggesting viral, protozoan or nonpathogenic factors [24].

254 *Salmonella spp* isolated in Mukuru slums could be non- typhoidal salmonella which is a zoonotic
255 strain. The children could have been contaminated with faecal matter of the domesticated
256 animals hence the acquisition of the bacteria. Occurrence of diarrhoeagenic bacteria in the
257 current study showed that gram negative bacteria (*Shigella spp*, *Salmonella spp*, *Escherichiacoli*)
258 are the main cause of bacterial diarrhoea. Sule et al. [19] in Kaduna Nigeria conducted a similar
259 study and found similar results. Generally, the aetiology of diarrhoea in young children could be

260 attributed to a wide range of factors, but one of the main causes of diarrhoea is related to
261 bacteria (such as *Salmonella spp*, *Shigella spp*, *Vibrio*, *Escherichia coli*, *Aeromonas* and
262 *Pseudomonas* [24]. Results from the current study shows that, though there are a number of
263 causative agents of diarrhoeal diseases, bacteria still remain one of the major causes with
264 *Shigella*, *Salmonella* and *Escherichia coli* being the most important pathogens among paediatric
265 patients presenting with diarrhoea in Mukuru kwa Njenga Government health facility. Judicious
266 use of antibiotic therapy requires education of health workers and patients, adequate laboratory
267 diagnostic capabilities and government regulations.

268 **Conclusion**

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

278 **Recommendation**

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

282 **Conflict of Interest**

283 The authors declare no conflict of interest

284 **References**

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