HUMAN CONTAMINATION AND SCHISTOSOME INFECTION INTENSITY IN BULINID AND PLANORBID SNAIL VECTORS IN KADAWA IRRIGATION AREA, KANO STATE, NIGERIA

4

5 ABSTRACT

An Epidemiological Research was conducted to determine the magnitude of human 6 contamination of irrigation canal perimeter as it relates to the prevalence and intensity of 7 schistosome cercarial infection in snail vectors. The study area was categorized into Zone of 8 Heavy Contamination (ZHC), Zone of Light Contamination (ZLC) and Zone of Free 9 Contamination (ZFC) based on the density of faecal lumps observed along the canal perimeter 10 using $1m^2$ quadrat sampling technique. Snail vectors of schistosomiasis were collected from 11 these zones, identified and subjected to cercarial shedding between January and June, 2012. Of 12 the 827 snails collected 28.54% shed schistosome cercariae. The breakdown of infection 13 prevalence was 31.37%, 27.69% and 26.26% for ZHC, ZLC and ZFC respectively. Three snail 14 species recovered in the study area, Bulinus globosus, B. rohlfsi and Biomphalaria pfeifferi had 15 infection intensity of 8.6, 5.67 and 3.94 respectively, with total mean intensity of 4.67. A Chi-16 squared analysis did not show any significant difference in infection prevalence in the three 17 zones ($\chi^2_{cal.}$ 0.025, $\chi^2_{2, 0.05} = 5.99$). However, infection intensity was significantly different in the 18 three zones and among the three snail species using analysis of variance (P < 0.05). Thus, human 19 environmental contamination with faeces and urine around irrigation canals remains the source 20 21 of infection to snail vectors and then to humans. It is therefore presumed that contact control through avoidance of defaecation in the open and building of pit latrines near water contact 22 points along irrigation canals will be an effective means of drawing a barrier to infection with 23 schistosomes in epidemiological sense. 24

Key words: Human Contamination, Schistosome Cercaria, Infection Intensity, Snail Vectors,
 Irrigation Canal

27 INTRODUCTION

Human schistosomiasis is a water-based disease and one of the neglected tropical diseases that is 28 more prevalent where there is high frequency of human contact with infested water. Water 29 resource schemes for power generation and irrigation have resulted in the increase in the 30 transmission and outbreaks of schistosomiasis in several African countries [1]. In sub-Saharan 31 Africa schistosomiasis is widespread with foci of high prevalence and high morbidity found 32 adjacent to rivers, lakes and irrigation schemes [2]. The disease epidemiology is attributable to 33 water contact pattern, biology and distribution of the potential snail vectors and the local 34 geographical, geological and climatic conditions [3, 4]. Contamination of surface waters or their 35

36 surrounding with faeces and urine containing schistosome eggs is essential for transmission of the parasite [4]. Humans become infected with schistosome following contact with contaminated 37 water through various water contact activities [5]. A combination of environmental and 38 anthropogenic parameters controls the distribution of schistosomes within a surface water 39 network [6]. Heavy rains aid contamination by carrying the schistosome eggs to water bodies 40 where they can successfully hatch into viable miracidia [7]. A wet climate is an important 41 contributor to water contamination as seen in the decreased viability of S. mansoni eggs exposed 42 to the sun within a few days after fecal deposition. The level of contamination is thus dependent 43 44 on both direct factors such as defaecation patterns and indirect factors such as rain events, overflowing latrines, and level of community sanitation [7]. Faecal contamination of surface 45 water with schistosome eggs occurs in rural endemic regions with low sanitation infrastructure 46 [7,8]. The continuum of infection is linked to the continuous water contact activities and 47 anthropogenic faecal and urine contamination, coupled with prevailing snail vector population 48 [9]. The bulinid and planorbid snail vectors, implicated in the transmission of human 49 schistosomiasis, have been reported in Kano State and many parts of Nigeria [10,11,12,13,14]. 50 Human activity of gross contamination of the water body perimeter is a major factor for infection 51 of the snails with human schistosome species. Even though snail infection rates may be low, the 52 presence of infected snails portends potential transmission of schistosomiasis. However, marked 53 seasonal fluctuation in snail infections may occur [14]. In the North-western parts of Nigeria, 54 55 comprising Sokoto, Katsina and Kebbi States, there are about 16 large and many small-scale formal irrigation and many private ones. Here, the general prevalence of urinary schistosomiasis 56 was shown to be 22.3% [15]. In Kano State, Nigeria, a considerable amount of water 57 development projects has been carried out and more are being proposed which will enhance 58 transmission [13]. In a prevalence study in Katsina State, Idris et al. [16] reported infection rates 59

60 of 12% and 3.3% for S. haematobium and S. mansoni among primary school pupils. Tukur and Galadima [17] reported a prevalence and intensity of S. haematobium infection of 50.9% and 61 151.0 eggs/10ml urine respectively, in Bakolori irrigation project area of Zamfara State. They 62 also found that persons aged 10-19 years had the highest prevalence rate of 70.3% and mean 63 intensity of 324.33 eggs/10ml urine, while those aged 40 years and above had the least 64 prevalence of 20.8%. Adamu et al. [18] reported 41% prevalence for urinary schistosomiasis in 65 Wurno district of Sokoto State, with intensity of 310 eggs/10ml urine. However, low prevalence 66 and intensity of 5% and 10 eggs/gm stools were recorded for intestinal schistosomiasis. In Kano 67 and Bauchi States, where a number of irrigation schemes and other water projects have been 68 executed and still more are expected, there was high rate of schistosomiasis recorded from these 69 water projects [19]. Schistosoma haematobium infection prevalence rates in some parts of Kano 70 State and its neighbours have been monitored. Umar [20] recorded as high as 28.4% prevalence 71 rate for S. haematobium among pupils of 8-10 years in Kura Local Government Area of Kano 72 State which is an area that is extensively irrigated as well as being very rich in ponds and rivers. 73 Betterton et al. [10] showed the presence of S. haematobium among the 813 school children and 74 adults from Tomas and Rimin Gado dam areas of Kano State, with prevalence of 26.6% and 75 36.8% respectively. They observed that the prevalence and intensity of S. haematobium were low 76 and similar in both study areas and no cases of S. mansoni infection were found. The study area is 77 sandwiched between tow village communities, Dakasoye and Dorawar Sallau, with a reported 78 overall prevalence of 32.8% and 16.8% for S. haematobium and S. mansoni infections 79 respectively [14]. Ali and Ndams [14] further reported an association between infection 80 prevalence and water contact activities in both communities. This research work reports an 81 82 investigation on the magnitude of human environmental contamination and its epidemiological implication in relation to the prevalence and intensity of schistosome infections in snail vector 83

population in the study area, with a view to highlighting the role of defaecation in the open in
maintaining schistosome infection in susceptible snail vectors.

86 MATERIALS AND METHODS

87 Study Area

The study area is an irrigated area lying about 35km southwest of Kano City (Lat. 11°59'N, Long. 88 8°30'E) on both sides of Kano-Zaria trunk road. The irrigation water is conveyed from Tiga Dam 89 to the project site through an 18km-long main canal, which splits into East and West branches 90 of canals and earthen field channels from where water is finally abstracted for crops 91 irrigation using plastic siphon tubes. Canals are designed such that the west branch canal and 92 the lateral canal are lined with the side slopes kept at $1:1\frac{1}{2}$ and maximum velocity of 1.8 m/s 93 while the earthen distributary canals have side slopes kept at 1:2, with a velocity below 0.3m/s to 94 prevent erosion [21]. The study area is bordered by two villages, Dakasoye (Lat. 11°44'N, Long. 95 8°25'E) and Dorawar Sallau (Lat. 11°39'N, Long. 8°23'E) within the Kano River Project Phase I 96 (KRP I), which is one of the largest and successful irrigation projects in Nigeria. The study area 97 98 comprised established communities with the irrigation agriculture-based economy and whose lives are directly or indirectly linked to the water that is constantly present in the irrigation canals. 99

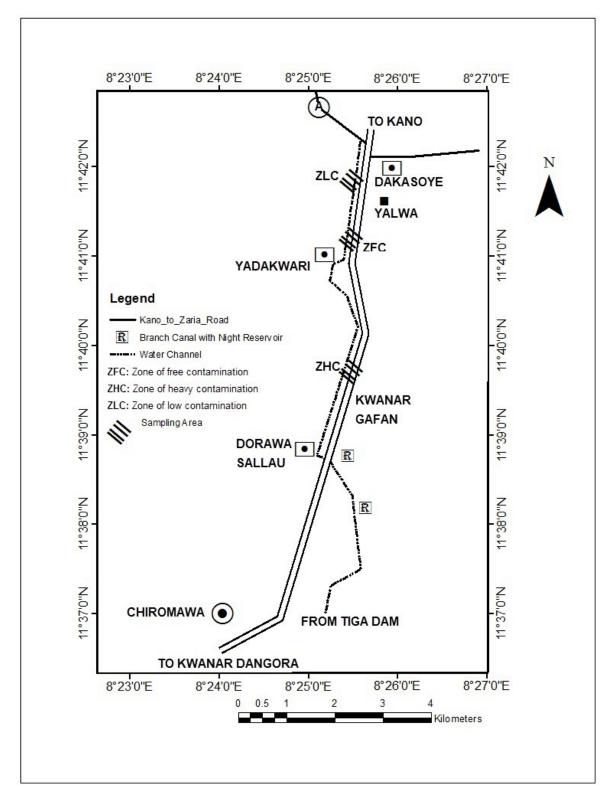




Figure 1: Map of the study area

103 Study Design

The study area was categorized into three (3) zones: Zone of Heavy Contamination (ZHC), Zone 104 of Light Contamination (ZLC) and Zone of Free Contamination (ZFC). The categorization was 105 on the basis of the observed level of human faecal and urine contamination during pre-sampling 106 107 visits to the study area, and the established presence of snail intermediate hosts in the water canals reported in previous studies [9,14]. The degree of contamination was determined by the 108 density of faecal lumps in each zone, using 1m² quadrat sampling technique. The quadrat was 109 thrown three times at random, on each landing the area covered by it was observed. The number 110 of visible faecal lumps within the quadrat was recorded and the average number of faecal lumps 111 calculated as lumps/m². Selection of contamination zones in the study area was made on the 112 basis of faecal density as: ZHC (> 3 lumps/m²), ZLC (1-3 lumps/m²) and ZFC (<1 lump/m²). 113 The distance between ZHC and ZFC was about 2.6km and that between ZFC and ZLC was 114 roughly 1.2km. The distance between points of faecal contamination and the edge of the water 115 canal was also determined using meter rule. The study area covers a distance of about 4km along 116 the water canal and Kano-Zaria Trunk Road, with the direction of the water course from ZHC to 117 ZFC to ZLC. The source of water in the irrigation canal was Tiga Dam. The topography of the 118 three zones, in particular, the vegetation covers and the nature of gradient around the perimeter 119 of the water canal, were also noted. All the three zones were measured approximately 8m by 120 150m to obtain an approximate canal perimeter area of 1200m² along the water canal. ZHC was 121 located proximal to Kwanar Gafan seasonal Vegetable Market. The people attending the market 122 come from various parts of Nigeria transacting in green vegetables which were harvested from 123 124 the surrounding irrigation area; although majority were from the neighboring communities. ZLC is located near the town of Dakasoye, where it forms a partial open latrine to some members of 125 126 the village community and visiting irrigation farmers, who do not have access to standard latrines during water exposure for occupational or recreational purposes. ZFC interspersed ZHC 127

and ZLC. Throughout the research period, rain boots, protective and disposable hand gloves, andnose cover, were worn during each sampling.

130 131

Snail Collection and Identification

Collection of snails was done between January and June, 2012 directly from 3 or 4 points 132 adjacent to the respective zones of contamination. The peak period of snail abundance and 133 anthropogenic environmental contamination in the hot dry seasons as well as water contact 134 activities reported earlier by [12] and [14], informed the decision of confining the study to six 135 months. The snails were searched from aquatic substrata such as macrophytes, plant twigs, rock 136 surfaces and floating objects and collected by hand picking with the aid of tea strainer from the 137 three zones (ZHC, ZLC and ZFC), during the study period, taking into cognizance of the 138 substrates to which the snails were attached. Protective hand gloves were worn during each 139 sampling. The snails were then transferred to labeled plastic beakers containing the canal water 140 141 and transported to the laboratory for identification and cercarial shedding. Identification of the snail was based on gross morphology of snail shells as in Brown [22]. 142

143 Snail Cercarial Shedding and Counting

Snails were examined for schistosome infection by immersing each snail in 5ml of dechlorinated 144 water in a Petri dish after exposure to light from a lamp-bulb for about 2-3 hours according to 145 [11]. The cercariae observed in the water contained in each Petri dish were counted by adopting 146 the method of [23] as follows. Water sample in each Petri dish was passed through 7cm Whatman 147 No.1 filter paper in a Buchner funnel under partial vacuum. Dechlorinated water was used to 148 149 rinse the Petri dishes to ensure washing out of all shed cercariae. Cercariae trapped on the filter paper were stained and immobilized with Lugol's Iodine and counted systematically under low 150 power (×10 objective) of dissecting microscope. Only the heads of cercariae were counted since 151

tails may become detached during sample preparation [23].

153 **RESULTS**

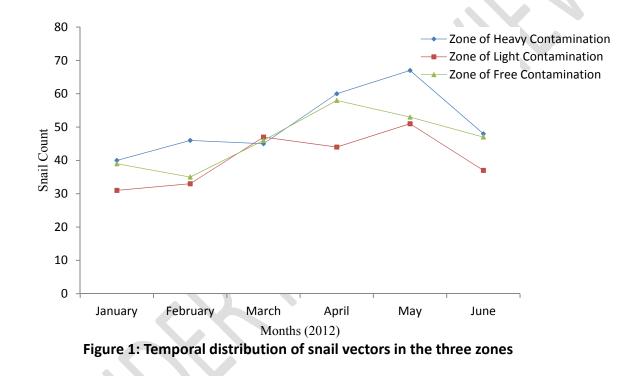
154 Snail Vector Abundance and Temporal Distribution

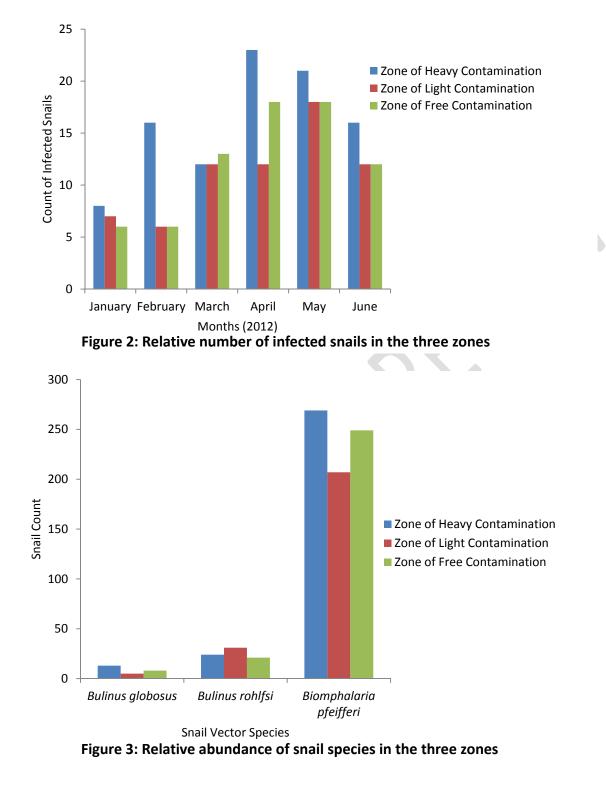
The results for the abundance and temporal distribution of snail vector species in the three zones 155 of contamination have been presented in Figures 1 and 3. There was a monthly variation in snail 156 abundance in the three zones of contamination. Snail count was generally low in the months of 157 January and February, high between the months of March and May and highest in May in the 158 zones of heavy and light contamination. However, snail count dropped in all the three zones in 159 160 June, although the highest snail count was recorded in April in ZHC, during the research period. Only three species of snail intermediate hosts of human schistosomiasis were recovered in the 161 study area; viz.: Biomphalaria pfeifferi, Bulinus globosus and B. rohlfsi, the former species being 162 predominant in all the three zones. 163

164 Snail Infection Prevalence and Intensity

The prevalence of infection in the snail intermediate hosts was presented in Table 1 and Figures 2 165 and 4. The prevalence of schistosome cercarial infection in the snail vectors in the three zones 166 was in the following order: ZHC, 31.37%; ZLC, 27.69% and ZFC, 26.26%, with overall infection 167 prevalence of 28.54% in the study area (Table 1). Figure 2 showed that the rate of infection with 168 schistosome cercariae followed a spatio-temporal pattern. Infection was highest in the month of 169 May, followed by April and January. The ZHC has the highest infection prevalence in 5 out of 6 170 171 months of the study. This is followed by ZFC and ZLC. All the three snail species were infected with schistosome cercariae (Figure 4). Infection prevalence was highest in *Biomphalaria pfeifferi* 172 and lowest in Bulinus globosus. Infection in B. pfeifferi was highest in ZHC followed by ZFC. 173 Conversely, in B. globosus, infection was highest in ZFC, followed by ZHC. The order of 174 increasing infection prevalence in B. rohlfsi was: ZC, ZLC and ZHC. However, there was no 175

statistically significant difference in infection prevalence in the three zones ($\chi^2 = 0.025$). Tables 2and 3 revealed the results of the mean intensity of schistosome cercarial infection in the snail species. The mean infection intensities for *Bulinus globosus*, *B. rohlfsi* and *Biomphalaria pfeifferi* were 8.6, 5.67 and 3.94, respectively; with total mean intensity of 4.67. Moreover, infection intensity was significantly different in the three zones and among the three snail species using analysis of variance at P<0.05 (Table 3).







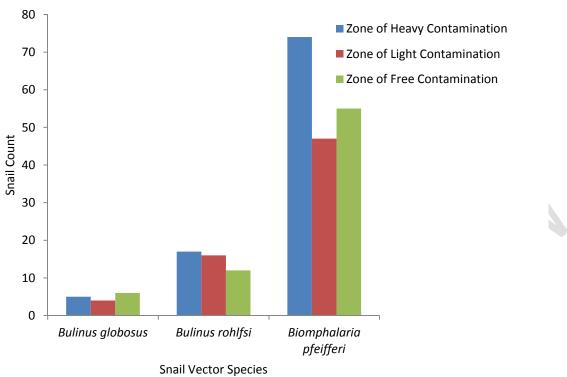


Figure 4: Relative abundance of snail vector species shedding schistosome cercariae.

186

187

188 Table 1: Schistosome Cercarial Infection Prevalence in Snail Vectors

	Zone of	No. of Snail Vectors	No. of Infected Snails	% Infection Prevalence					
	Contamination								
	ZHC	306	96	31.37					
	ZLC	242	67	27.69					
	ZFC	278	73	26.26					
	Total	827	236	28.54					
189	$\chi^2 = 0.025$								
190									
191									
192									
193									
194									
195									

	Snail Vector Species			
				199
Zone of Contamination	Bulinus globosus	Bulinus rohlfsi	Biomphalaria pfeifferi	Mean 200
				201
				202
ZHC	12.2	8.24	5.19	6.203
ZLC	6.75	3.94	3.96	4.2024
ZFC	6.83	4.33	2.67	3.205
				206
Mean	8.6	5.67	3.94	4.2077
				208

197 Table 2: Mean Intensity of Schistosome Cercariae Infection

209

Table 3: Analysis of Variance for Snail Cercarial Infection Intensity in the Three
 Contamination Zones

212

Source of Variation	Degree of	Sum of	Mean	F _{calculated}
	freedom	Squares	Squares	
Snail Species	2	33.65	16.83	13.304 ^s
Zones of Contamination	2	28.94	14.47	11.439 ^s
Error	4	5.06	1.265	
Total	8	67.65		

F_{0.05} (2, 4) = 6.94; **S** = Significant at 5% level of Statistical Significance

214 DISCUSSION

The temporal distribution and abundance of snail vectors of schistosomiasis show variation in 215 the three contamination zones. This was similarly observed by [14] who inferred that variation in 216 snail vector population in time and space was due to fluctuating seasonal temperature as water 217 218 constancy in the canal has neutral influence on snail abundance throughout the study period. Moreover, the snail vectors were observed to be mainly spatially distributed in the littoral zone 219 of the water canal where water flow velocity was lowest, usually attached to submerged and 220 221 floating objects as is typical of periphytonic communities. There was no marked variation in species distribution in the three zones of contamination. However, Biomphalaria pfeifferi has far 222

outnumbered the other two species; *Bulinus globosus* and *B. rohlfsi*, recovered in the study area.
The predominance of *Biomphalaria pfeifferi* in the study area was earlier reported by [14].
Although, the overall snail vector population was highest in ZHC and lowest in ZLC during the research survey.

The observed anthropogenic faecal and urine contamination of the irrigation water canal 227 perimeter has been documented by several researchers [13,14,25]. The lack of standard pit 228 latrines in the study area was the major cause of human contamination activities around the water 229 canal. The contribution of environmental contamination to the spread of schistosomiasis is 230 immense epidemiologically since human urine and faecal matter are the sources of infection to 231 snail vectors in which the juvenile stages of schistosomes perpetuate to release the human 232 infective cercariae. This observation was in agreement with that of Akullian [7] who reported 233 that the perpetual contamination of waterways with human waste, and subsequent exposure to 234 contaminated water is essential for the parasite's continued asexual reproduction in the snail host 235 and sexual reproduction within the mammalian host. Amadou et al. [24] and WHO [1] have 236 linked schistosomiasis to very low standard of hygiene coupled with inadequate potable water 237 supply that may lead to unprotected water contact activities. The prevalence of schistosome 238 infection in the snail vectors is indicated by the mature patent infection by cercarial shedding 239 which was 28.54% altogether. This finding was slightly higher than that of [14] who recorded an 240 overall infection prevalence of 20.9% in the snail vectors. This may be attributed to small sample 241 size in this research, and the varying environmental conditions which are never static. In 242 addition, Li et al. [8] attributed schistosomiasis prevalence to levels of local surface water 243 contamination contributed by sanitation levels and faecal contamination patterns in humans and 244 domestic animals. They further observed that faecal contamination of surface water with 245

schistosome eggs occurs in rural endemic regions with low sanitation infrastructure. These findings were further corroborated by [7] who indicated that geographic distribution of schistosomes along waterways might have more to do with human behavior and the geographic extent of human travel than environmental factors alone.

The significant difference in infection prevalence in the three zones observed in this study 250 indicates the pivotal role of anthropogenic activity of faecal and urine contamination of the edges 251 along water canals in the epidemiology of schistosomiasis; maintenance of infection in snail host 252 reliant upon infected urine and faecal matter that slip into the canal. Similar findings have been 253 reported by [14] who observed that human activity of grossly contaminating the canal periphery 254 contributed to increased infection of snail vectors with human schistosome species. This 255 promiscuous contamination is the sole source of human-to-snail transmission which has an 256 attendant effect of maintaining schistosome infection in humans in the study area as a result of 257 water contact with cercariae-infested water, hence, the endemicity of schistosomiasis in the study 258 area, as reported by several researchers [9,13,14,20,25]. Moreover, Akullian [7] further observed 259 that in many endemic areas humans contribute heavily to both the parasite's survival and the 260 resulting burden of disease within the human population through continued faecal and urinary 261 contamination of heavily used waterways. This study revealed a significant difference in 262 infection intensity in the three zones and among the three snail species, namely Biomphalaria 263 pfeifferi, Bulinus globosus and B. rohlfsi. The presence of infected snail vectors in the ZFC might 264 be attributed to the influence of water currents in horizontal transportation of snail infective 265 larval forms, miracidia, thereby seeding the near and distant snail colonies along the water 266 course, as well as the greater chance of the surrounding contaminated soil to be blown into the 267 water canal especially by the whirling wind during the hot dry season, precisely the months of 268

269 April and May, when water contact and contamination activities of the surrounding communities were highest, and when the water canal accommodates a higher population of the thriving 270 susceptible snail vectors and at the advent of the rains due to a slight slanted topography of the 271 water canal perimeter; a triple tragedy in epidemiological point of view. This finding thus, 272 strengthens the epidemiologic importance of contamination activity in schistosomiasis 273 transmission, in particular human-to-snail transmission. Moreover, the infectivity of the three 274 snail vector species indicates their competence in hosting and nurturing the developing juveniles 275 of schistosomes, with bulinid species surpassing in vectorial competence, therefore connoting a 276 higher prevalence of urinary schistosomiasis in the study area, as reported by [9,13,20]. 277

278 **Recommendations**

For effective and lasting control of schistosomiasis, contact control strategies should be employed as a preventative tool drawing a barrier between human definitive host and schistosomiasis. Moreover, it is recommended based on the findings in this research that:

i. Mass drug administration (MDA) of anti-schistosomal regimen, Praziquantel, 282 following mass screening should be implemented once a year, targeting children of 283 school age in all schistosomiasis-endemic areas with the intention of providing mass 284 prevention. This exercise should be a sole responsibility of health department under 285 state and local government authorities. However, the WHO [1] criterion for MDA is a 286 primary school prevalence of \geq 50% of infection. Moreover, the WHO Control 287 Strategy for urinary schistosomiasis states that the major control plans of urinary 288 schistosomiasis are provision of Praziguantel to primary school children, provision of 289 safe tap water to the whole community and health education. 290

291 ii. Government should enact sanitation laws targeting schistosomiasis-endemic communities to include components as follows: banning any form of anthropogenic 292 contamination of the environment around canal perimeters; building a reasonable 293 number of public convenience in the irrigation area near water contact points along 294 the canals by the local authorities; establishing community sanitation clubs (CSC) to 295 curb any form of faecal and urine environmental contamination through vigilance and 296 awareness campaign; inclusion of public health education in the curricula of primary 297 and secondary schools which will lay emphasis on the health-risk associated with 298 299 unprotected exposure to water that is laden with susceptible snail vectors.

300 iii. Periodic community awareness campaign on the health-risk of unprotected water301 contact activities through community health and agricultural extension workers.

The aforesaid recommendations, though not exhaustive, would proffer a tremendous impact in our dream of eradicating schistosomiasis, or at least halting the progression of its transmission in endemic areas.

305 Conclusion

The human 'contaminatory' behavior of the endemic communities around the study area and the lack of measures to improve sanitary conditions will continue to predispose the inhabitants to the risk of infection, and re-infection with schistosome parasites so long that the wet climate remains, as the irrigation scheme provides for subsistence agriculture and water contact for domestic purposes to the majority of the local populace, and so long that faecal and urine contamination of the canal perimeter continues, thereby seeding the surface water that harbors susceptible snail population. 314 Disclaimer: - This manuscript was presented in a Conference.

315 Conference name: 37th PPSN Annual Conference

316 Available link: -

<u>a</u>

- 317 <u>https://www.researchgate.net/publication/329813851_Human_Contamination_and_Schistosome_Infec</u>
- 318 tion_Intensity_in_Bulinid_and_Planorbid_Snail_Vectors_in_Kadawa_Irrigation_Area_Kano_State_Nigeri
- 319
- 320

321

322 **REFERENCES**

- 323 1. WHO (2006). Neglected Tropical Diseases. WHO Annual report
 324 2006.WHO/CDS/NTD/2007.4
- Alan, F., Lorenzo, S., Dirk, E., Robert, N.B. and Matthew, H.T. (2003). Drugs for the control of parasitic diseases: Current status and development in schistosomiasis. *Trends in Parasitology*. 19(11):509-515.
- 328 3. Ugbomoiko, U.S. (2000). The prevalence, incidence and distribution of human urinary
 329 schistosomiasis in Edo State, Nigeria. *The Nigerian Journal of Parasitology*. 21:3-14.
- 4. Jozef, V., Darren, J.S. and Jan, D.B. (2001). Index of potential contamination for schistosomiasis. *Trends in Parasitology*. **17**(6):256-261.
- 5. Cheesbrough, M. (1998). *District laboratory practice in Tropical countries part 1*. Cambridge
 University Press. pp.236-239.
- 334 6. Jordan, P. and Webbe, G. (1993). "Epidemiology" from Human Schistosomiasis Edited by
 335 Jordan, Webbe and Sturrock. CAB International. Wallingford, UK.
- 7. Akullian, A.N. (2007). Spatial Patterns of Schistosomiasis: A Watershed approach to
 measuring *S. japonicum* environmental risk and human and animal disease outcomes. B.Sc.
 Thesis, Brown University, Rhode Island, USA. Pp.5.
- 8. Li, B.G. (1998). Relationship between number and quality of private lavatory and social
 economic situation in schistosomiasis endemic areas. *J. Pract. Parasit. Dis.* 6:185-186.
- 9. Ali, M.U. and Ndams, I.S. (2013). Human *Schistosoma* Infections: Intensity versus Water
 Contact Patterns among Irrigation Communities in Kano State, Nigeria. *Biological and Environmental Sciences Journal for the Tropics*. 10(3):120-124.
- Betterton, C., Ndifon, G.T., Bassey, S.E., Tan, R.M. and Oyeyi, T. (1988). Schistosomiasis in
 Kano State, Nigeria. I. Human infections near dam sites and the distribution and habitat
 preferences of potential snail intermediate hosts. *Annals of Tropical Medicine and Parasitology*. 82(6):561-570.

- 348 11. Opaluwa, S.A., Galadima, M., Odama, L.E. and Nock, I.H. (1999). Prevalence of
 349 schistosomiasis in some rice-farming communities of Niger State, Nigeria. *The Nigerian* 350 *Journal of Surgical Research*. 1(2):56-61.
- Idris, H.S. and Ajanusi, O.J. (2002). Snail intermediate hosts and the etiology of human
 schistosomiasis in Katsina State, Nigeria. *The Nigerian Journal of Parasitology*. 23:145-152.
- 13. Bassey, S.E. and Umar, Z. (2004). A re-assessment of schistosomiasis infection in GarunBabba, Kadawa and Kura in Kano State, Nigeria. *The Nigerian Journal of Parasitology*.
 25:107-109.
- 14. Ali, M.U. and Ndams, I.S. (2012). Distribution of Snail Intermediate Hosts of
 Schistosomiasis in Human Water Contact Points along Kadawa Irrigation Canals, Kano State,
 Nigeria. *Biological and Environmental Sciences Journal for the Tropics*. 9(2):169-175.
- 15. Ofoezie, I.E.; Bolton, P.; Imevbore, A.M.A. and Christensen, N.O. (1996). Schistosomiasis
 and other helminth infections in irrigation schemes in Sokoto, Katsina and Kebbi States of
 Nigeria. *The Nigerian Journal of Parasitology*. 1(17):31-38
- Idris, H.S., Ajanusi, J.O., Umoh, J.U., Galadima, M. and Ogbugu, V.C. (2001). Prevalence of
 schistosomiasis among pupils in some Local Government Areas of Katsina State, Nigeria. The
 Nigerian Journal of Parasitology. 22:75-80
- Tukur, A. and Galadima, M. (1998). Epidemiological study of schistosomiasis in the Bakalori
 Irrigation Project Area of Zamfara State, Nigeria: Prevalence and intensity of *Schistosoma haematobium* infection. The Nigerian Journal of Parasitology. 19:73-75
- 18. Adamu, T., Abubakar, U. and Danga, C. (2001). Schistosomiasis in Wurno District of Sokoto
 State, Nigeria. *The Nigerian Journal of Parasitology*. 22(1&2):81-84
- 19. Oyeyi, T.I.; Ndifon, G.T.; Betterton, C. and Bassey, S.E. (1988). Field and Laboratory observations on the aestivation pattern of *Bulinus senegalensis* (Muller) snail intermediate hosts of schistosomiasis in Kano State, Nigeria. In: 13th Annual conference, *Nigerian Soc. Parasitol*. September, 24-27. Pp.23
- 20. Umar, Z.O. (2001). A re-assessment of the Epidemiology of schistosomiasis at Kura Local
 Government Area, Kano State, Nigeria. An M.Sc thesis, Department of Biological Sciences,
 Bayero University, Kano. Pp. 5-21
- HJRBDA. (2004). Brief on Achievements of Hadejia-Jama'are River Basin Development
 Authority, Kano, presented to the distinguished senators of the Senate Committee on Water
 Resources, Federal Ministry of Water Resources, September, 2004. Pp.1-2.
- Brown D.S. (1980). Freshwater snails of Africa and their medical importance. London:
 Taylor and Francis.
- Oyeyi, T.I. (2000). Studies on the Aestivation, Growth, Susceptibility and Cercarial
 production in some snail hosts of Schistosomiasis in Kano. A PhD Thesis, Department of
 Biological Sciences, Bayero University, Kano, Nigeria. Pp.46.

- 24. Ali, M.U. (2011). Vector Abundance and Prevalence of *Schistosoma* Infections in Two
 Irrigation Communities in Kura Local Government Area, Kano State. An M.Sc Thesis,
 Department of Biological Sciences, Ahmadu Bello University, Zaria, Nigeria. Pp. 91.
- 388 25. Amadou, G., Seydou, T., Robert, D., Elisa, B. and Alan, F. (2006). Implementation of
 389 National schistosomiasis control programmes in West Africa. *Trends in Parasitology*.
 390 22(7):322-326
- 391