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

ABSTRACT

Aims: An Epidemiological Research was conducted to determine the magnitude of human contamination of irrigation canal perimeter as it relates to the prevalence and intensity of schistosome cercarial infection in snail vectors.

Study design: The study area was categorized into Zone of Heavy Contamination (ZHC), Zone of Light Contamination (ZLC) and Zone of Free Contamination (ZFC) based on the density of faecal lumps observed along the canal perimeter using $1m^2$ quadrat sampling technique. Snail vectors of schistosomiasis were collected from these zones, identified and subjected to cercarial shedding between January and June 2012.

Results: Of the 827 snails collected 28.54% shed schistosome cercariae. The breakdown of infection prevalence was 31.37%, 27.69% and 26.26% for ZHC, ZLC and ZFC respectively. Three snail species recovered in the study area, Bulinus globosus, B. rohlfsi and Biomphalaria pfeifferi had infection intensity of 8.6, 5.67 and 3.94 respectively, with a total mean intensity of 4.67. A Chi-squared analysis did not show any significant difference in infection prevalence in the three zones ($\chi^2_{cal.}$ 0.025, $\chi^2_{2,\ 0.05}$ = 5.99). However, infection intensity was significantly different in the three zones and among the three snail species using analysis of variance ($P<0.05$).

Conclusion: It is concluded that human environmental contamination with faeces and urine around irrigation canals remains the source 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 points along irrigation canals will be an effective means of drawing a barrier to infection with schistosomes in an epidemiological sense.

***Keywords**: Human Contamination, Schistosome Cercaria, Infection Intensity, Snail Vectors, Irrigation Canal*

INTRODUCTION

Human schistosomiasis is a water-based disease and one of the neglected tropical diseases that are more prevalent where there is a high frequency of human contact with infested water. Water resource schemes for power generation and irrigation have resulted in an increase in the transmission and outbreaks of schistosomiasis in several African countries [1]. In sub-Saharan Africa, schistosomiasis is widespread with foci of high prevalence and high morbidity found adjacent to rivers, lakes and irrigation schemes [2]. The disease epidemiology is attributable to

water contact pattern, biology and distribution of the potential snail vectors and the local geographical, geological and climatic conditions [3, 4]. Contamination of surface waters or their 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 water through various water contact activities [5]. A combination of environmental and anthropogenic parameters controls the distribution of schistosomes within a surface water network [6]. Heavy rains aid contamination by carrying the schistosome eggs to water bodies where they can successfully hatch into viable miracidia [7]. A wet climate is an important contributor to water contamination as seen in the decreased viability of *S. mansoni* eggs exposed to the sun within a few days after fecal deposition. The level of contamination is thus dependent 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 water with schistosome eggs occurs in rural endemic regions with low sanitation infrastructure [7,8]. The continuum of infection is linked to the continuous water contact activities and anthropogenic faecal and urine contamination, coupled with prevailing snail vector population [9]. The bulinid and planorbid snail vectors, implicated in the transmission of human schistosomiasis, have been reported in Kano State and many parts of Nigeria [10,11,12,13,14]. Human activity of gross contamination of the water body perimeter is a major factor for infection of the snails with human schistosome species. Even though snail infection rates may be low, the presence of infected snails portends potential transmission of schistosomiasis. However, marked seasonal fluctuation in snail infections may occur [14]. In the North-western parts of Nigeria, comprising Sokoto, Katsina and Kebbi States, there are about 16 large and much small-scale formal irrigations and many private ones. Here, the general prevalence of urinary schistosomiasis was shown to be 22.3% [15]. In Kano State, Nigeria, a considerable amount of water

development projects has been carried out and more are being proposed which will enhance transmission [13]. In a prevalence study in Katsina State, Idris *et al*. [16] reported infection rates of 12% and 3.3% for *S. haematobium* and *S. mansoni* among primary school pupils. Tukur and Galadima [17] reported prevalence and intensity of *S. haematobium* infection of 50.9% and 151.0 eggs/10ml urine respectively, in Bakolori irrigation project area of Zamfara State. They also found that persons aged 10-19 years had the highest prevalence rate of 70.3% and the mean intensity of 324.33 eggs/10ml urine, while those aged 40 years and above had the least prevalence of 20.8%. Adamu *et al*. [18] reported 41% prevalence for urinary schistosomiasis in Wurno district of Sokoto State, with intensity of 310 eggs/10ml urine. However, low prevalence and intensity of 5% and 10 eggs/gm stools were recorded for intestinal schistosomiasis. In Kano and Bauchi States, where a number of irrigation schemes and other water projects have been executed and still more are expected, there was a high rate of schistosomiasis recorded from these water projects [19]. *Schistosoma haematobium* infection prevalence rates in some parts of Kano State and its neighbours have been monitored. Umar [20] recorded as high as 28.4% prevalence rate for *S. haematobium* among pupils of 8-10 years in Kura Local Government Area of Kano State which is an area that is extensively irrigated as well as being very rich in ponds and rivers. Betterton *et al.* [10] showed the presence of *S. haematobium* among the 813 school children and adults from Tomas and Rimin Gado dam areas of Kano State, with the prevalence of 26.6% and 36.8% respectively. They observed that the prevalence and intensity of *S. haematobium* were low and similar in both study areas and no cases of *S. mansoni* infection were found. The study area is sandwiched between tow village communities, Dakasoye and Dorawar Sallau, with a reported overall prevalence of 32.8% and 16.8% for *S. haematobium* and *S. mansoni* infections respectively [14]. Ali and Ndams [14] further reported an association between infection prevalence and water contact activities in both communities. This research work reports an

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 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.

MATERIALS AND METHODS

Study Area

The study area is an irrigated area lying about 35km southwest of Kano City (Lat. 11°59'N, Long. 8°30'E) on both sides of Kano-Zaria trunk road. The irrigation water is conveyed from Tiga Dam to the project site through an 18km-long main canal, which splits into East and West branches of canals and earthen field channels from where water is finally abstracted for crops irrigation using plastic syphon tubes. Canals are designed such that the west branch canal and the lateral canal are lined with the side slopes kept at 1:1½ and maximum velocity of 1.8m/s while the earthen distributary canals have side slopes kept at 1:2, with a velocity below 0.3m/s to prevent erosion [21]. The study area is bordered by two villages, Dakasoye (Lat. 11°44'N, Long. 8°25'E) and Dorawar Sallau (Lat. 11°39'N, Long. 8°23'E) within the Kano River Project Phase I (KRP I), which is one of the largest and successful irrigation projects in Nigeria. The study area 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.

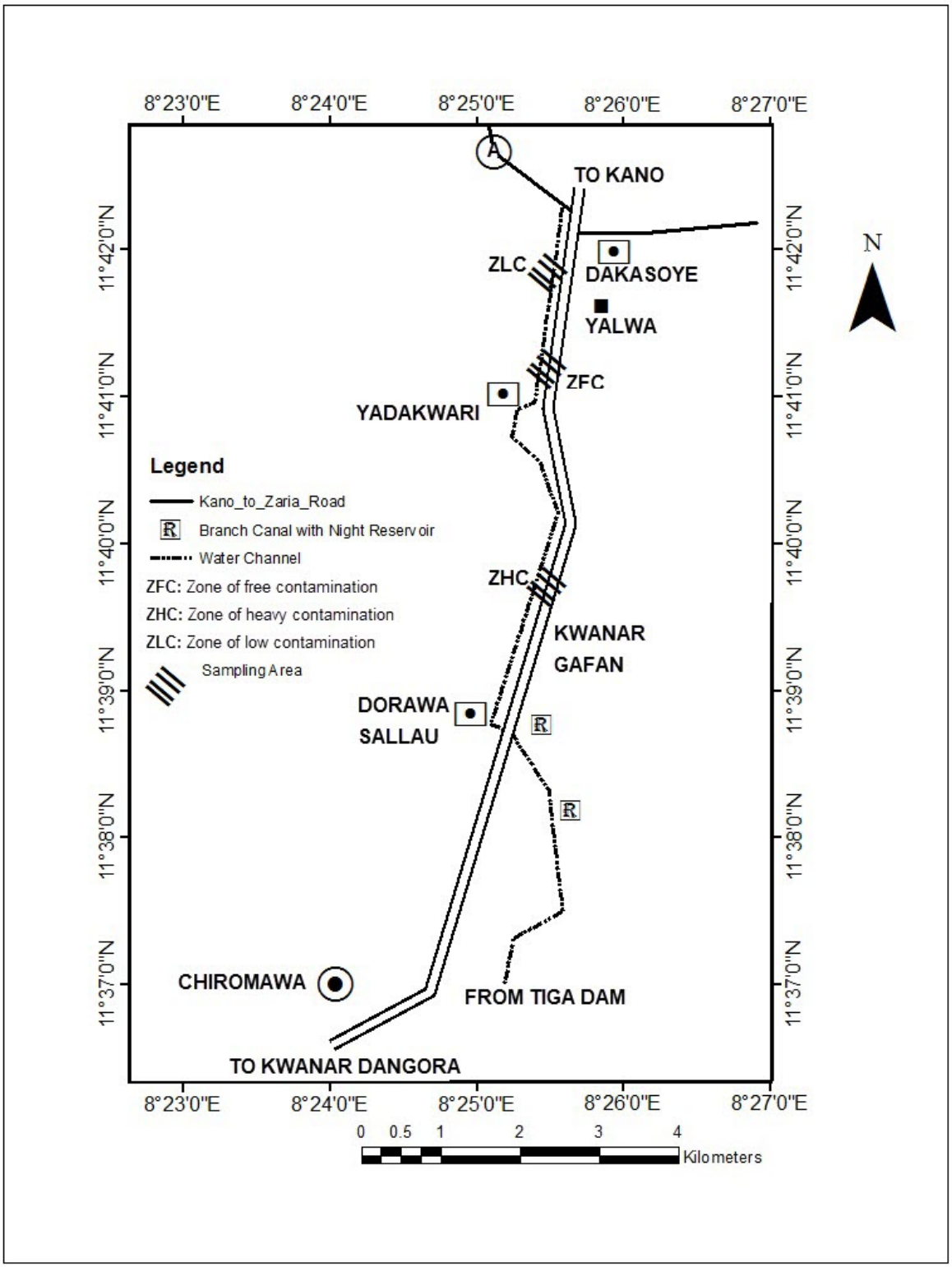

Figure 1: Map of the study area

Study Design

The study area was categorized into three (3) zones: Zone of Heavy Contamination (ZHC), Zone of Light Contamination (ZLC) and Zone of Free Contamination (ZFC). The categorization was on the basis of the observed level of human faecal and urine contamination during pre-sampling 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 density of faecal lumps in each zone, using $1m^2$ quadrat sampling technique. The quadrat was thrown three times at random, on each landing the area covered by it was observed. The number of visible faecal lumps within the quadrat was recorded and the average number of faecal lumps calculated as lumps/m^2. Selection of contamination zones in the study area was made on the basis of faecal density as: ZHC (> 3 lumps/m^2), ZLC (1-3 lumps/m^2) and ZFC (<1 lump/m^2). The distance between ZHC and ZFC was about 2.6km and that between ZFC and ZLC was roughly 1.2km. The distance between points of faecal contamination and the edge of the water canal was also determined using meter rule. The study area covers a distance of about 4km along the water canal and Kano-Zaria Trunk Road, with the direction of the watercourse from ZHC to ZFC to ZLC. The source of water in the irrigation canal was Tiga Dam. The topography of the three zones, in particular, the vegetation covers and the nature of gradient around the perimeter of the water canal, were also noted. All the three zones were measured approximately 8m by 150m to obtain an approximate canal perimeter area of $1200m^2$ along the water canal. ZHC was located proximal to Kwanar Gafan seasonal Vegetable Market. The people attending the market come from various parts of Nigeria transacting in green vegetables which were harvested from 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 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

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

Snail Collection and Identification

Collection of snails was done between January and June 2012 directly from 3 or 4 points adjacent to the respective zones of contamination. The peak period of snail abundance and anthropogenic environmental contamination in the hot dry seasons as well as water contact activities reported earlier by [12] and [14], informed the decision of confining the study to six months. The snails were searched from the aquatic substrates such as macrophytes, plant twigs, rock surfaces and floating objects and collected by hand picking with the aid of tea strainer from the three zones (ZHC, ZLC and ZFC), during the study period, taking into cognizance of the substrates to which the snails were attached. Protective hand gloves were worn during each sampling. The snails were then transferred to labelled plastic beakers containing the canal water 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].

Snail Cercarial Shedding and Counting

Snails were examined for schistosome infection by immersing each snail in 5ml of dechlorinated water in a Petri dish after exposure to light from a lamp-bulb for about 2-3 hours according to [11]. The cercariae observed in the water contained in each Petri dish were counted by adopting the method of [23] as follows. Water sample in each Petri dish was passed through 7cm Whatman No.1 filter paper in a Buchner funnel under partial vacuum. Dechlorinated water was used to 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 power (×10 objective) or dissecting microscope. Only the heads of cercariae were counted since

tails may become detached during sample preparation [23].

RESULTS

Snail Vector Abundance and Temporal Distribution

The results for the abundance and temporal distribution of snail vector species in the three zones of contamination have been presented in Figures 1 and 3. There was a monthly variation in snail abundance in the three zones of contamination. Snail count was generally low in the months of January and February, high between the months of March and May and highest in May in the zones of heavy and light contamination. However, snail count dropped in all the three zones in 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 study area; viz.: *Biomphalaria pfeifferi*, *Bulinus globosus* and *B. rohlfsi*, the former species being predominant in all the three zones.

Snail Infection Prevalence and Intensity

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

178 statistically significant difference in infection prevalence in the three zones (χ^2 = 0.025). Tables
179 2and 3 revealed the results of the mean intensity of schistosome cercarial infection in the snail
180 species. The mean infection intensities for *Bulinus globosus*, *B. rohlfsi* and *Biomphalaria pfeifferi*
181 were 8.6, 5.67 and 3.94, respectively; with a total mean intensity of 4.67. Moreover, infection
182 intensity was significantly different in the three zones and among the three snail species using
183 analysis of variance at P<0.05 (Table 3).

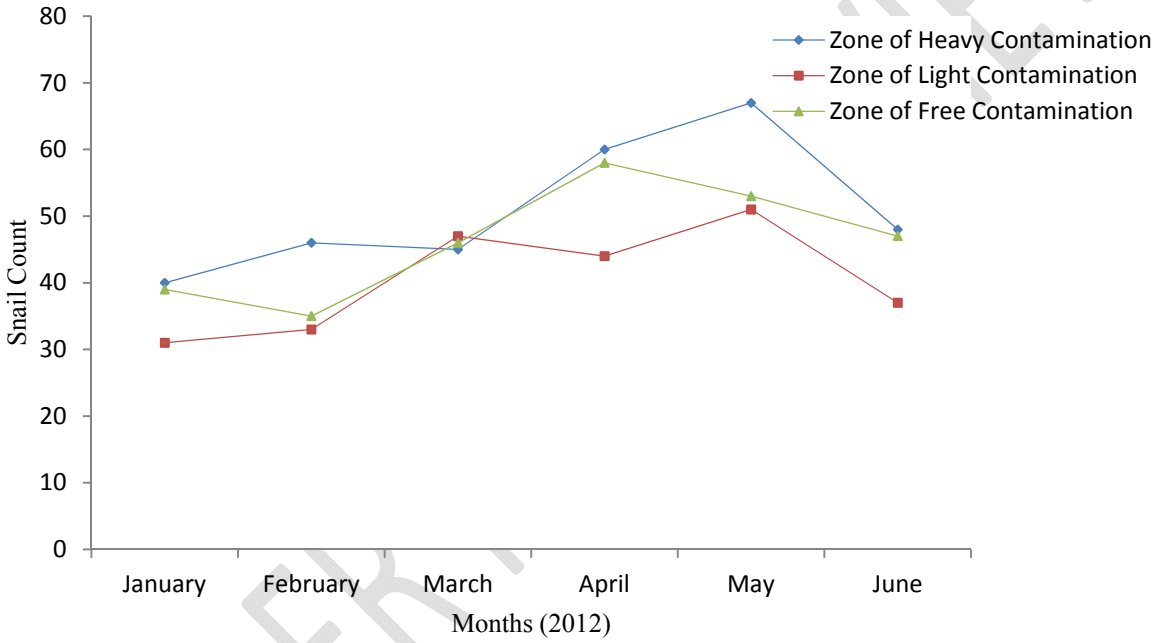

Figure 1: Temporal distribution of snail vectors in the three zones

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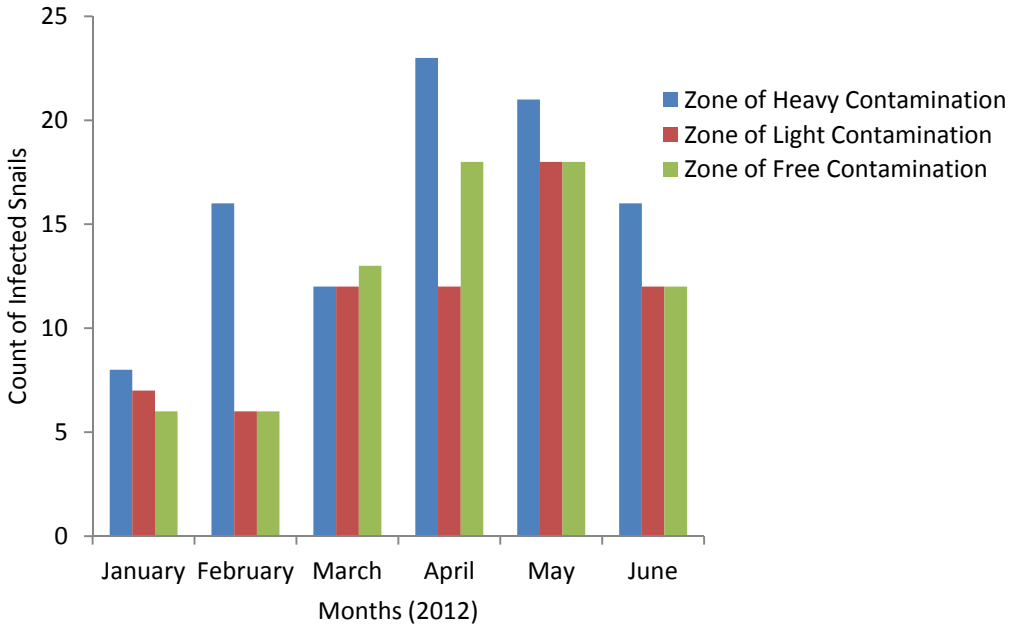

Figure 2: Relative number of infected snails in the three zones

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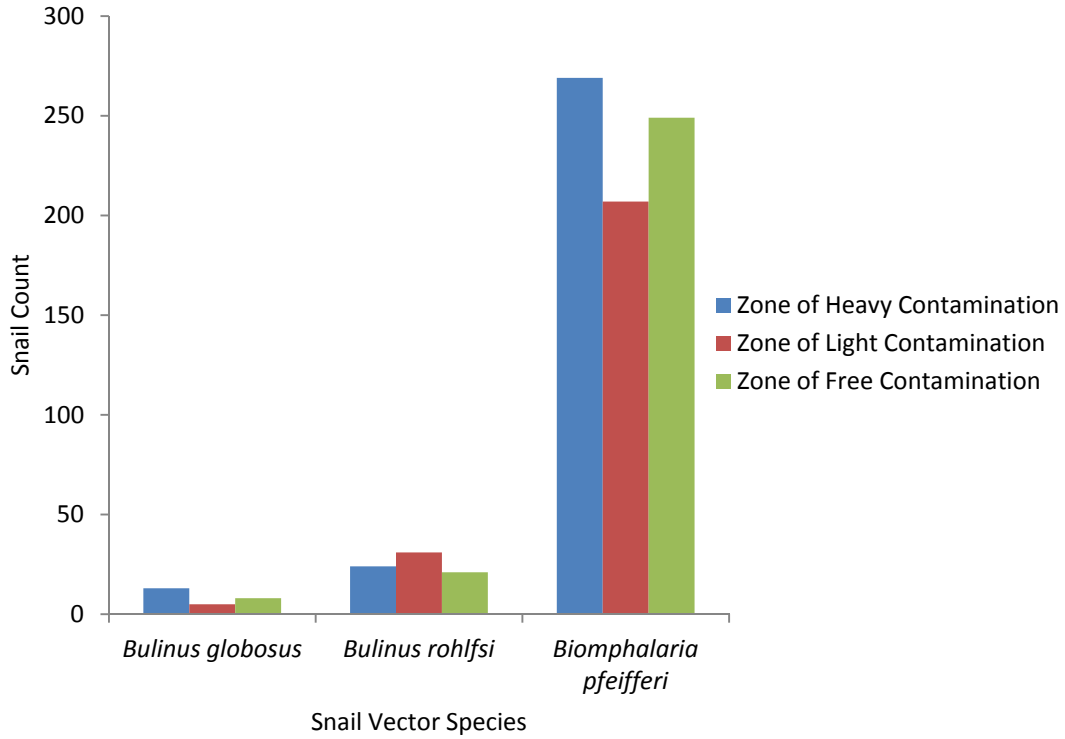

Figure 3: Relative abundance of snail species in the three zones

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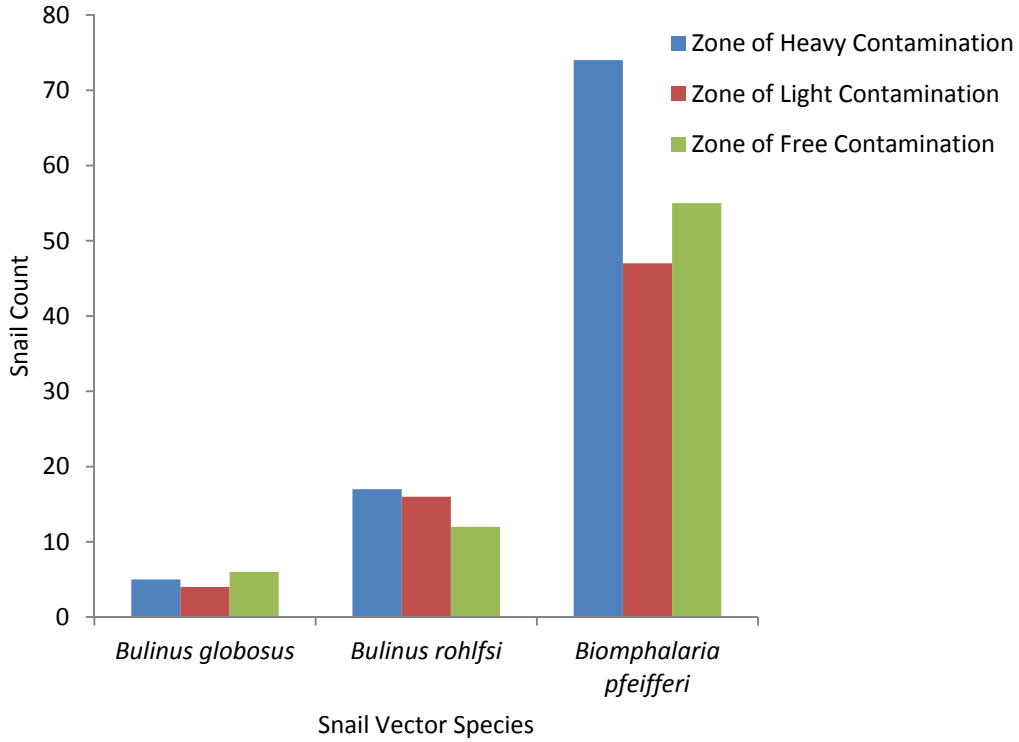

Figure 4: Relative abundance of snail vector species shedding schistosome cercariae.

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190 **Table 1**: **Schistosome Cercarial Infection Prevalence in Snail Vectors**

Zone of Contamination	No. of Snail Vectors	No. of Infected Snails	% Infection Prevalence
ZHC	306	96	31.37
ZLC	242	67	27.69
ZFC	278	73	26.26
Total	827	236	28.54

191 $\chi^2 = 0.025$

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Table 2: **Mean Intensity of Schistosome Cercariae Infection**

	Snail Vector Species			
Zone of Contamination	***Bulinus globosus***	***Bulinus rohlfsi***	***Biomphalaria pfeifferi***	**Mean**
ZHC	12.2	8.24	5.19	6.09
ZLC	6.75	3.94	3.96	4.12
ZFC	6.83	4.33	2.67	3.29
Mean	8.6	5.67	3.94	4.67

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

Source of Variation	**Degree of freedom**	**Sum of Squares**	**Mean Squares**	**$F_{calculated}$**
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

DISCUSSION

The temporal distribution and abundance of snail vectors of schistosomiasis show variation in the three contamination zones. This was similarly observed by [14] who inferred that variation in snail vector population in time and space was due to fluctuating seasonal temperature as water constancy in the canal has a neutral influence on snail abundance throughout the study period. Moreover, the snail vectors were observed to be mainly spatially distributed in the littoral zone of the water canal where water flow velocity was lowest, usually attached to submerged and 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

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

with 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 behaviour and the geographic extent of human travel than environmental factors alone. The elimination of schistosomiasis and other trematode parasite infections will receive a great boost when snail hosts studies and effective snail control programme are prioritized [26]. Based on available data in the P.R. China, Zheng et al. [27] highlight the gaps between control capacity and prevalence levels, and between diagnostic/drug development and population need for treatment at different stages of the national control programme. The study of Allan et al. [28] determines the distribution and identity of potential intermediate snail hosts of Schistosoma spp. in Bengo, Luanda, Kwanza Norte and Malanje Provinces in north-western Angola. Various freshwater bodies in north-western Angola harbour potential intermediate snail hosts for urogenital schistosomiasis, highlighting the need to map the rest of the country. de Assis [29] also probed a pilot version of the microarray with sera from individuals either acutely or chronically infected with *S. mansoni* from endemic areas in Brazil and sera from individuals resident outside the endemic area (USA) to determine if the array is functional and informative.

The significant difference in infection prevalence in the three zones observed in this study indicates the pivotal role of anthropogenic activity of faecal and urine contamination of the edges along water canals in the epidemiology of schistosomiasis; maintenance of infection in snail host reliant upon infected urine and faecal matter that slip into the canal. Similar findings have been reported by [14] who observed that human activity of grossly contaminating the canal periphery contributed to increased infection of snail vectors with human schistosome species. This promiscuous contamination is the sole source of human-to-snail transmission which has an

attendant effect of maintaining schistosome infection in humans in the study area as a result of water contact with cercariae-infested water, hence, the endemicity of schistosomiasis in the study area, as reported by several researchers [9,13,14,20,25]. Moreover, Akullian [7] further observed that in many endemic areas humans contribute heavily to both the parasite's survival and the resulting burden of disease within the human population through continued faecal and urinary contamination of heavily used waterways. This study revealed a significant difference in infection intensity in the three zones and among the three snail species, namely, *Biomphalaria pfeifferi*, *Bulinus globosus* and *B. rohlfsi*. The presence of infected snail vectors in the ZFC might be attributed to the influence of water currents in horizontal transportation of snail infective larval forms, miracidia, thereby seeding the near and distant snail colonies along the watercourse, as well as the greater chance of the surrounding contaminated soil to be blown into the water canal especially by the whirling wind during the hot dry season, precisely the months of 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 susceptible snail vectors and at the advent of the rains due to a slightly slanted topography of the water canal perimeter; a triple tragedy in epidemiological point of view. This finding thus, strengthens the epidemiologic importance of contamination activity in schistosomiasis transmission, in particular, human-to-snail transmission. Moreover, the infectivity of the three snail vector species indicates their competence in hosting and nurturing the developing juveniles of schistosomes, with bulinid species surpassing in vectorial competence, therefore connoting a higher prevalence of urinary schistosomiasis in the study area, as reported by [9,13,20].

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 the anti-schistosomal regimen, Praziquantel, following mass screening should be implemented once a year, targeting children of school age in all schistosomiasis-endemic areas with the intention of providing mass prevention. This exercise should be sole responsibility of the health department under state and local government authorities. However, the WHO [1] criterion for MDA is a primary school prevalence of ≥50% of infection. Moreover, the WHO Control Strategy for urinary schistosomiasis states that the major control plans of urinary schistosomiasis are the provision of Praziquantel to primary school children, provision of safe tap water to the whole community and health education.

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

iii. Periodic community awareness campaign on the health-risk of unprotected water 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.

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 harbours susceptible snail population.

Disclaimer: - This manuscript was presented in a Conference.

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