1	Original Research Article
2	Environmental parameters and Biomphalariasnail distribution along River Kochi, West Nile
3	region, Uganda
4	
5	
6 7	

8 ABSTRACT

Aims: To explore the abundance and distribution of the common fresh water mollusks in River Kochi, with a special focus on *Biomphalaria* species, a vector responsible for transmitting *Schistosomamansoni* to humans.

Study design:

Place and Duration of Study:This study was conducted between October 2007 and March 2008 as a master's project along Kochi River in Koboko, Yumbe and Moyo in the West Nile region of Uganda.

Methodology: Five sites along the river approximately 20 km apart, were selected and data on snail abundance and various environmental variables thought to be influencing the distribution of snails along this river were collected. These variables included: altitude, season of the year, water flow velocity, water pH, water temperature and concentrations of total dissolved solids in the water.

Results:Findings indicate that numbers of *Biomphalaria* species of snails increased with decreasing altitude (mean numbers 0, 15.33, 19, 50 & 73.33 from highest to lowest altitude points) and no snails of this species were recorded during the wet season. Water flow velocity and pH were the main predictors of the presence of *Biomphalaria* snails (Pearson correlations -0.749 and 0.614 for flow velocity and pH respectively). *Biomphalaria* snail numbers increased when water velocity decreased and the reverse was true for pH.

Conclusion: Altitude influences the distribution Biomphalarias nails and hence potential

prevalence of schistosomiasis. Water users of Kochi River should therefore try to minimize contact with water in this river especially during the dry season. Local leaders should lobby to government for alternative sources of water during the dry season.

9

10 Keywords: Biomphalaria, Schistosomamansoni, River Kochi, West Nile

11

12 1. INTRODUCTION

13

14 Approximately 30 species of *Biomphalaria* are recognized and the genus is widely distributed 15 in South America and on the African continent [1]. Biomphalaria is an aquatic snail that acts as 16 a host for a human blood fluke Schistosomamansonithat cause the disease intestinal 17 schistosomiasis (bilharzia) in humans [2]. With its many lakes, rivers, streams, swamps and 18 ponds. Uganda has a diverse fresh water environment that offers numerous and suitable 19 habitats for the Biomphalariaspecies. Currently two species of Biomphalarianamely: B. 20 stanleyiand B. sudanica(hereby known as Biomphalaria) are the most common in the west Nile 21 region of Uganda [3].

22 Whereas Nelson [4] in 1950s pioneered research works on schistosomiasis in West Nile region 23 approaching the infections from ecological and geographical points of view and from both 24 human populations and snail vectors in water bodies, research that followed his works mainly 25 concentrated either in human communities that live close to the shores of lake Albert/ Albert 26 Nile [5, 6, 7, 8, 9&10] or from hospital records [5, 11], with the exception of findings of Kazibwe 27 et al [3].Contrary to the aforementioned studies, Kazibwe [3] looked at the effect of 28 environmental factors on the distribution of Biomphalaria in Lake Albert, Western Uganda. 29 Findings from this study revealed that climatic conditions primarily air temperature, rainfall, lake 30 depth level, water temperature; water conductivity and water pH influence the distribution and 31 abundance of snails in Lake Albert. Similarly Appleton [11] and Thieltges et al. [13] showed that

climatic conditions primarily rainfall and temperature influence the distribution and abundance
 of snails because they have an effect on their breeding and the rate of schistosomal
 development.

35 It is clearly evident from the aforementioned studies that the studies on Biomphalaria snail 36 species ecology was restricted to large water bodies, with little or no attention given to small 37 ones, which are also a source of water and fish for the local communities. However, it is 38 important to mention that Odongo-Aginya and others [14] conducted a research on urban 39 Schistosomamansoninear Enyau River in Arua town, a small river in the highland areas of the 40 region further away from the Nile River but his focus was on infections in humans. In addition, 41 human populations in the township comprise of people from different origins and locations, and 42 may therefore not have given dependable results since all the S. mansonicases registered may 43 not have been contracted from Enyau River. We therefore strongly believe that conducting 44 research on vector dynamics in smaller water bodies will result into better understanding of the 45 disease prevalence and its distribution in the region considering that a lower number of snails 46 mean a lower number of cercariae and therefore a diminished risk of infection. Studies on 47 diseases vectors are very important for evidence based mitigation and control measure. The 48 main objective of the study was therefore to investigate effects of altitude, season and water environmental variables on the distribution of *Biomphalariasnail* species along the Kochi River, 49 50 West Nile region. Findings from this study will generate some current information on the 51 distribution of the disease vectorin the different infection zones and altitudes in the study area.

52

53 2. MATERIAL AND METHODS

54 2.1 Study sites

The study took place in Kochi River located in Koboko, Yumbe and Moyo districts of West Nile region. Generally, the study area was divided into three altitude zones i.e. Koboko with altitude of above 1000 m; Yumbe (two sites: Yumbe 1 and Yumbe 2) with altitude range of 700 m to

58 1000 m and Moyo (two sites: Moyo 1 and Moyo 2) with altitude range of 600 m to 700 m above 59 sea level. Kochi River has its origin in Koboko district near Uganda-DR Congo boarder at an 60 altitude of above 1000 m where it starts as a small stream and gradually widens downstream as it passes through Yumbe district and finally joins the Albert Nile in Moyo at an altitude of 61 62 about 600 m above sea level (Fig 1). This river stretches all the schistosomiasis infection belts 63 of the region that Nelson [4] hadestablished in 1958. The rainfall pattern in this region is bimodal peaking in late March to May (about 900 mm), and August to December (above 900 64 mm) each year. The rest of the months experience dry spells with sporadic rainfall which 65 66 fluctuate the water levels of the rivers and its stream tributaries where some seasonal ones dry 67 up completely.

68

69

70 **Fig.1**. Map showing the study sites



71

72 **2.2 Estimating snail abundance and water parameters**

73 Biomphalaria snail abundance was estimated from well-defined areas along the river. These 74 areas measured 30 m along the bank and 3 m into the main body of the water. The corners of 75 these rectangular sampling areas were marked by pegs so that successive samplings could be 76 performed across the same area. These areas were searched for a period of 30mins and all 77 snails found floating or attached to vegetation were collected using a scooping net with a long 78 handle and placed on white plastic trays in order to be able to rapidly identify the different 79 species based on the standard field identification key guide of the Danish bilharziasis laboratory [15]. Each site was visited weekly and snail samples collected over a period of six 80 81 months between October 2007 and March 2008. Three of the months (October, November and

December) experienced heavy rains of above 900 mm and have been recorded as wet, whilst the other three months (January, February and March) experienced little or no rains and have been recorded as dry. Although our main focus was *Biomphalaria*, snail types like *Lymnae*, *Bulinus* and *Pila* species were collected because they coexist with the *Biomphalaria* snails and are intermediate hosts to other human and animal diseases. Snails were collected from 5 altitude belts spread across the study area at intervals of about 20 km apart.

We took measurements on water flow velocity, water pH, water temperature and concentration of total dissolved solids (TDS) in the water shortly before collecting the snail samples. Water flow velocity was obtained by sprinkling methyl orange dye from the upstream mark of the sampling area and recording the time taken for the dye colour to cover the 30 m distance to the downstream mark and velocities computed. Values for pH and temperature were obtained by using a pH meter integrated with a temperature probe (Model 3150/REV A/04-95). TDS concentration was determined using a conductivity meter (Model 4200/REV A/05-95).

95 2.3 Data analysis

96 Data were analysed using Genstat version 3. Firstly, we made a descriptive summary of the 97 abundance of all snails in total, and then secondly we singled out *Biomphalaria*species and 98 explored how its distribution is affected by the environmental factors considered in this study. 99 Pearson-r Correlation Coefficient tests were done to establish associations between the 100 different variables. Environmental variables that had strong associations with the abundance of 101 *Biomphalaria*species were then used in Simple Regression models.

102

103 3. RESULTS AND DISCUSSION

Higher numbers of *Biomphalaria* and *Pila* were recorded in Yumbe and Moyo while that of *Lymnae* and *Bulinus* snail species were registered in Koboko (Fig 2). Considering that these locations are positioned at different altitudinal zones, these results already indicate an effect of

107 altitude on the distribution of these snail species. The results further show that no *Biomphalaria*

108 species of snails were recorded during the wet season, and the number of snails increased

- 109 with decreasing altitude during the dry season from none recorded at an altitude of 1189 m to a
- 110 mean of 62 snails recorded per month at an altitude of 638 m or 639 m (Table 1).

111 Fig. 2. Abundance and distribution of common snail species at sites along River Kochi



	Mean			
Season	Dry	Wet		
Altitude				
638.0	73.33	0.00		
639.0	50.00	0.00		
898.0	19.00	0.00		
933.0	15.33	0.00		
1189.0	0.00	0.00		

115 Table 1: Altitudinal and seasonal variation in the number of *Biomphalaria* species

116

117 A correlation analysis indicated that there was an association between the number of 118 Biomphalaria snails and water flow velocity (-0.749) and between Biomphalaria snails and pH 119 (0.614) (Table 2). Water flow velocity and pH were in turn highly correlated (-0.899) indicating a 120 strong association between them (Table 2 and Fig 3). Weak associations existed between 121 number of Biomphalaria snails and Temperature and Total dissolved solids (Table 2). In view 122 of the result shown in table 1 that show that snail incidence varied with altitude and season we 123 found it important to summarize mean values for water flow velocity and pH levels in the same 124 way. Results for water flow velocity show that snails were found only in the dry season at the 125 five sites where the velocity was in a range of 0.19 to 0.31 m/s. No snails were recorded at 126 Koboko, which is the site at the highest altitude. The water velocity in the dry season was 0.48 127 m/s, which is at the lower end of the range of values shown for the wet season and above the 128 value of 0.4 m/s (the upper limit for presence of snails shown in Table 1). A similar trend can 129 be seen for pH. Snails were found only when pH values were 7.1 or above (Table 3).

130 Table 2: A correlation matrix between numbers of *Biomphalaria*and environmental

131 variables

	*** Correlation NBiom	matrix 1.000	***			
	TDS	-0.264	1.000			
	Temp	0.012		1.000	1000000	
	Velocity	-0.749	0.182	0.027	1.000	
	pH	0.614	0.099	0.084	-0.899	1.000
132		NBiom	TDS	Temp	Velocity	pH
1 <u>32</u> 133	*NBiom = number of Biomph	alaria snails;	TDS = Total Dissol			

134 Fig. 3: Relationship between water flow velocity and water pH



135



	Mean (Water flow velocity)		Mean (water p		(water pH)
Season	Dry	Wet	Season	Wet	Wet
Altitude			Altitude		
638.0	0.1867	0.4733	638.0	7.200	6.947
639.0	0.2267	0.5400	639.0	7.187	6.837
898.0	0.2917	0.5683	898.0	7.183	6.893
933.0	0.3083	0.5600	933.0	7.080	6.900
1189.0	0.4450	0.7283	1189.0	6.813	6.533

138 * Altitude decreases from Koboko to Moyo

139 Considering that pH and flow velocity were strongly correlated with each other, we decided to 140 use Simple linear regression models for each variable to predict Biomphalaria snail 141 abundance. The results showed highly significant relationships between water pH and flow 142 velocity (P<.001) and that pH accounted for 38% of the variation while water flow velocity 143 accounted for 56% of the variation in Biomphalaria numbers. There was a positive association 144 between Biomphalaria snails and pH when pH is 7.1 or above and no snails were found when 145 pH was below 7.1 (Fig 4). In addition, the number of Biomphalaria snails increased when 146 water flow velocity decreased below about 0.4 m/s and no snails were found when velocity was 147 0.4 m/s or above (Fig 5).

148



150 Fig. 4: Relationship between numbers of *Biomphalaria* snails and water pH



152

153 Fig. 5: Relationship between numbers of *Biomphalaria* snails and water flow velocity



155

156 **DISCUSSION**

157 Bulinus and Lymnae snail species were distributed throughout the river length. On the other 158 hand, Biomphalaria and Pila species were not however found at all in some of the sites 159 especially those towards the source of the river in Koboko above 1000 m. They were only 160 found in sites towards the Nile in Yumbe and Moyo at fairly lower altitudes. The presence of 161 Lymnaespecies in the river poses a threat of Fasciola hepatica (liver fluke) transmissions in 162 domestic animals that graze along the river in case some of the animals happen to be infected 163 with the disease. In the same token Schistosomahaematobium (urinary schistosomiasis) could 164 easily spread in this area as their intermediate snail vectors (Bulinus species) are readily 165 available in the river and the fact that major roads that cross into South Sudan where S. 166 haematobium exists also cross this river. With high human mobility across these two countries, 167 existence of S. haematobium in the area if not the region is highly likely and therefore needs 168 investigation.

169 Biomphalaria numbers in the dry season were associated with water flow velocity and pH 170 levels. A possible explanation for the association is that the river becomes wider and so the 171 flow speed of the water reduces further downstream. Stable water conditions downstream 172 would be particularly prevalent during the dry season. Such conditions would enable the snails 173 to anchor more easily on the water vegetation. Also, as the debris carried down the river settles 174 and rots down, so the pH of the water gradually increases. This would explain why higher 175 numbers of Biomphalaria species of snails are associated with lower water flow rates and 176 higher pH levels (See also [3]). It is however important to note that not all the variation in 177 Biomphalaria species numbers was explained by water velocity and pH. It is possible that this 178 unexplained variation could be due to other factors such as amount of vegetation and snail 179 prey present at the study sites, which were out of the scope of this study.

Numbers of *Biomphalaria* species of snails increased with decreasing altitude and no snails of this species were found during the wet season. The complete absence of snails in the wet season is very difficult to explain. However, we think that this is attributed to the fact that during this time of the year, the water flow velocity in river Kochi was very high thereby drifting the snails away. In addition the water table was also very high therefore submerging the vegetation onto which the snails attach. This seems to suggest that there is need for lowering the sweep net further deeper into the water to search the snails.

187 Furthermore Kabetereine, [16] recorded bigger numbers of Biomphalariastanleyi in shallow 188 waters along Lake Albert during dry season and this was mainly attributed to the effects of light 189 penetration on the growth of Vallisneria weeds which serve as food for the snails in the river. 190 Fewer numbers of snails were recorded when lake levels increased and light penetration to 191 support growth of the weeds reduced. In addition, warmer and wetter conditions encouraged 192 snails to lay more eggs thereby increasing the densities of young snails several weeks later. In 193 this current study we attribute the big numbers of Biomphalaria snails collected during dry 194 season to the preceding wetter and warmer months of wet season that resulted in mass eqg 195 laying and subsequent development of these eggs into the large number of adult snails 196 registered later in the drier months of dry season.

197 There was no significant effect of temperature on the distribution and abundance of 198 *Biomphalaria* snail species along Kochi river. This finding is rather contrary to studies 199 conducted elsewhere in the world [16, 11, 18, 19 & 20]. These findings could be attributed to 200 the fact that there was/is no severe fluctuations in temperature as compared to the extreme 201 cold and hot temperatures experienced in the study sites considered in the above mentioned 202 studies.

204 4. CONCLUSION

205 In conclusion, we want to acknowledge that although this study was limited in time scope 206 compared to earlier studies, our findings are consistent as they seem to indicate that despite 207 national schistosomiasis control efforts, Biomphalaria snail species are still present within the 208 West Nile region. The continued presence of these snail species and other associated snail 209 species in smaller water bodies could thwart the efforts to contain schistosomiasis in this 210 region and pose an unforeseen threat to a number of snail transmitted diseases to humans, 211 and domestic animals in communities along the rivers most especially in the lower altitudes. 212 We recommend regular community sensitisation by the Ministry of health about the risks of 213 getting into contact with the river water during the dry season and that the concerned local 214 governments lobby to government to provide alternative sources of water e.g. boreholes that 215 can be used in the dry season to minimise peoples' contact with the river water. Mass control 216 interventions by the government to the schistosomiasis pandemic in this region following quick 217 diseases surveys in human communities will yield little results if no focus is paid to the water 218 sources where the disease is contracted. Further research may be directed towards scaling up 219 the study along other rivers in the west Nile region and also incorporating other variables like 220 the amount of vegetation and snail prey present at the sites.

221

222

223 **REFERENCES**

- Brown, D. S. Freshwater snails of Africa and their medical importance. Taylor &
 Francis, London; 1994.
- Jordan, P. & G. Webbe. Epidemiology. In: Human schistosomiasis(Eds P Jordan, G.
 Webbe, & R. F. Sturrock). Commonwealth Agricultural Bureau International,
 Wallingford;1993:87-158.
- Kazibwe, F., B. Makanga, C. Rubaire-Akiiki, J. Ouma, C. Kariuki, N. B. Kabatereine, M.
 Booth, B. J. Vennervald, R. F. Sturrock, J. R. Stothard. Ecology of Biomphalaria

231 (Gastropoda: Planorbidae) in Lake Albert, Western Uganda: Snail distribution,

- 232 infections with schistosomes and temporal associations with environmental dynamics.
- 233 Hydrobiologia. 2006;568(1):433-444.
- 4. Nelson, G.S.Schistosomamansoni infection in West Nile District of Uganda. Part II. The
 distribution of S. mansoni with a note on the probable vectors. East African Medical
 Journal. 1958;35:335-344.
- 5. Ongom V.L. & Bradley D.J. 1972. The epidemiology and consequences of
 Schistosomamansoni infection in West Nile. Part I. Field studies of a community at
 Panyagoro. Transaction of Royal Society of Tropical Medicine and Hygiene.
 1972;66:835-851
- 241 6. Ongom, V.L. The epidemiology of et al. and Consequences 242 SchistosomamansoniInfection in West Nile, Uganda. II. Hospital investigation of a 243 sample from the Panyagoro Community. Transaction of Royal Society of Tropical 244 Medicine and Hygiene. 1972;66:851-863.
- 7. Bukenya, G. & S. Andama. Circumstantial epidemiology of Schistosomamansoniin the
 West Nile District of Uganda: Results of a cross-sectional study in the Rhino Camp
 Area. Journal of Tropical Medicine and Hygiene. 1986;89:243-248.
- Kabatereine N.B., C. Ariho, &N.O.Christensen.Schistosomamansoniin Pakwach, Nebbi
 District, Uganda, 40 years after Nelson. Tropical Medicine and Parasitology.
 1992;43:162-166.
- 9. Laroni-Lakwo T, Odongo-Aginya EI, Schweigmann U, Schickerling S, Linder D,
 Doehring-Schwerdtfeger E. Transmission of Schistosoma mansoni in Rhino Camp,
 Uganda. E. Afr. Med. J. 1994;71:165-166.
- 10. Kabatereine, N.B., F. Kazibwe, & J. Kemijumbi. Epidemiology of schistosomiasis in
 Kampala, Uganda. East African Medical Journal. 1996;73:795-800.
- 256 11. Williams, E.H., R.J. Hayes, & P.G. Smith. Admissions to rural hospital in the West Nile
 257 District of Uganda over a 27 year period. Journal of Tropical Medicine and Hygiene.

258	1986;89:193-211.
259	12. Appleton, C.C. Review of literature on abiotic factors influencing the distribution and
260	life cycle of bilharziasis intermediate snail. Malacological Review. 1978;11:1-25.
261	13. Thieltges, D., K.Jensen& R. Poulin. 2008. The role of biotic factors in the transmission
262	of free living endoheliminth stages. Parasitology. 2008;135:407-426
263	14. Odongo-Aginya, E.I. Lakwo TL, Schweigmann U, Schickerling S, Lindner D, Mueller
264	A, Doehring-Schwerdtfeger E.Urban Schistosomamansoninear Enyau River in Arua
265	Town, Uganda. East African Medical Journal. 1994;71:604-606.
266	15. Danish Bilharziasis Laboratory. A Field Guide to African Freshwater Snails, (2nd edn.)
267	East African species. World Health Organization Collaborating Centre,
268	Copenhagen.1987:29–32
269	16. Kabatereine, N.B., F.M. Flemming, U. Nyandindi, J.C.L. Mwanza, and B. Lynsey. The
270	control of schistosomiasis and soil transmitted helminths in East Africa. Trends in
271	Parasitology. 2006;7:332-339.
272	17. Shati, A. A. Factors affecting the prevalence of human schistosomiasis in Aseer
273	regions, Saudi Arabia. Journal of Biological Sciences. 2009;9:815-819.
274	18. Martens, W.J.M.Modelling the effect of global warming on schistosomiasis on the
275	prevalence of schistosomiasis. GLOBO Report Service. 1995;10:1-31.
276	19. Gryseels, B., K. Polman., J. Clerinx& L. Kestens. Human schistosomiasis. Lancet.
277	2006;368:1106-1118.
278	20. Brooker, S. Spatial epidemiology of human schistosomiasis in Africa risk models
279	transmission dynamics and control. Transaction of Royal Society of Tropical Medicine
280	and Hygiene. 2007;101:1-8.
281	
282	