An Overview of Schistosomiasis in Africa:
Evolutional Origin and Diagnostic Challenges

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Authors’ contributions
This work was carried out in collaboration between both authors. Author SAH designed the study, wrote the protocol and wrote the first draft of the manuscript. Author SU managed the literature searches. Both authors read and approved the final manuscript.

Article Information
DOI: 10.9734/AJMAH/2019/v17i230160
(1) Prof. Darko Nozic, University of Belgrade, Serbia.
(2) Rowel Candia, Uganda.
Complete Peer review History: http://www.sdiarticle4.com/review-history/52646
Received 20 September 2019
Accepted 25 November 2019
Published 09 December 2019

ABSTRACT

Schistosomiasis is an overlooked tropical illness that is caused by the blood fluke that dwells human blood vessels. It manifest as an acute yet chronic sickness and is normally found in all developing nations with poor health services frameworks. Strong phylogenies dependent on molecular data of species in the family of Schistosoma have been produced lately. The impressive advancement made in understanding the connections between considerable perceived types of Schistosoma is explored with specific consideration being given to the discovery and investigation of parasite variety as reported by studies on ribosomal RNA qualities, mitochondrial DNA and Random Amplified Polymorphic DNA. Generally, molecular phylogenies concur with perceptions depending on morphological or life-history attributes. Basically the parasites of man don't form a monophyletic group and that cozy connections exist between parasites inside species groups, particularly in the S. haematobium group of species. The S. japonicum group has all the hallmarks of being the most dissimilar of the species groups but then little DNA succession variety has been seen between different isolates of S. japonicum. Diagnosis is a key to all aspects of schistosomiasis and decision on individual and community treatment, estimations on prognosis and

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1. INTRODUCTION

Schistosomiasis is infectious parasitic disease caused by *trematode* (flukes) of the genus schistosome. The parasite can be classified on the basis of their egg/ova morphology; they are characterized as the second most devastating parasite commonly found in the tropical region of the world. They cause mortality and morbidity in developing countries especially in Sub-Saharan Africa, although places like the Caribbean, the South America, Middle East and Asia has reported a significant prevalence [1]. Human become infected through skin penetration when it comes in contact with an infective stage of the parasite/cercariae often during various water contact activities in infested water [2]. Five major species are known to infect humans; *S. haematobium*, *S. mansoni*, *S. japonicum* *S. mekongi* and *S. intercalatum* (the first three are of most significant public health problem) [3]. Other schistosome that occasionally infects human causing dermatitis or insignificant infection includes *S. bovis* and *S. mattheei* together with some avian schistosome. These species differ biologically from one another and to their geographic distribution and the type of disease they cause [3].

*S. haematobium* a causative agent of urinary schistosomiasis causes complication like fibrosis, stricture and urinary tract calcification similarly; advanced intestinal schistosomiasis may result in hepatosplenomegaly, liver fibrosis and portal hypertension [2]. However, at the time of infection most of the eggs become trapped in some tissues. Organ in the body like liver become infected with *S. mansoni* and *S. japonicum* worm, on the other hand bladder and urethras are the major organs affected by the species of *S. haematobium* worms [1]. The pathology is not due to the number of worms in the body but instead due to immunologic response to schistosome egg that is being trapped in the host tissues. Antigen which is secreted by the egg stimulates a granulomatous reaction involving T-lymphocyte, macrophages and eosinophils, which initiate the clinical signs and symptoms [1].

Urinary schistosomiasis is endemic in Sub-Saharan Africa including Nigeria with substantial transmission in most of the state of the federation and high prevalence among school age children. Inadequate basic social amenities, weak primary health-care and poor infrastructure, facilitate transmission of schistosomiasis in the country and up till now control program is not available due to the absence of detail distribution of the disease. Intestinal schistosomiasis represents another form of *Schistosoma* infection. Among spectrum of intestinal lesions, polyps are the commonest. Intestinal schistosomiasis is essentially due to *Schistosoma mansoni*, *S. japonicum* and *S. mekongi* infection are distributed primarily across sub-Saharan Africa and South America. Intestinal schistosomiasis are responsible the major public health burden and health problem, concentrated in sub-Saharan Africa.

2. LITERATURES ACQUISITION

Peer-reviewed journal articles search were conducted via Medline, SCOPUS and Google Scholar search engines. All searches were limited to, publication from 1999 to 2019 except where necessary an older publication was used. All publications were in English, and duplicates, conference abstract, comments and short communications were excluded. The initial search result gave us 2584 articles which were screened based on the title relevance, retaining 196 full text review articles out of which we cited 71 articles in this review based on relevance.

3. EPIDEMIOLOGY

Schistosomiasis is one of the most common in the world found in approximately 76 countries more located in Africa, Middle East, South America and the Southern Asian continent [4]. An estimation of 779 million people are at risk of infection by schistosome with 207 million reported to have been infected, most of which are in Africa (97%) [5]. It was estimated that at least 20,000 deaths were reported as a result of schistosomiasis infection yearly [6].

Many African countries provide epidemiological data regularly regarding the prevalence and...
extents of infection cause to humans, and associated with malacological surveys and schistosome prevalence in intermediate host snails. West African epidemiological data related to schistosomiasis prevalence are available from Burkina Faso [7], Ivory Coast [8], and Niger [9]. Most of the schistosome found in Africa is S. haematobium and S. mansoni as shown in Fig. 1. Previous evolutionary history, climate and environmental changes shows a significant selection and pattern of schistosome with increase in fluctuation and pattern of the population size, definitive mammalian hosts, including human and the snail distribution has tremendous impact on the infection change in human social development, in addition has immensely favoured the distribution and survival of the parasite and help in changing the prevalence of the infection in recent years [10]. However, some researchers highlighted that establishing freshwater sites from dam irrigation system on constructed dams, site of human settlements close to freshwater sites has relationship with potential host, with high increase in size and number of human [5,11,12].

It was equally pointed out that, large rivers may not be the main source of schistosomiasis, but waters retained by them through seasonal flooding, and extraction for irrigation schemes and impoundment are likely to be the major transmission sites [13]. The disease is distributed over rural and agricultural areas where the life style of people increases contamination of inland water with human excreta. In Nigeria more than 62% of the constructed dams were built in ten most endemic states, the remaining ones in twenty six less endemic states [14]. An example of such dams located in South-West Nigeria is that of Oyam dam found in Ogun State which were constructed in was 1984. However, there was an outbreak of urinary schistosomiasis in some couples of years after its construction in two resettlement communities around the dam [15].

The study of the prevalence and severity of Schistosoma infection based on age and water contact activities has revealed that adolescents and children are most infected. The severity of the infection is gender and age specific with globally, rate of infections peak at age 10 to 19 years. In some areas, the prevalence in this group may approach up to 100% [16,17]. In relation to gender, more males than females were often been reported to had higher infection in most of the studies reviewed. In the study conducted by Houmsou et al. [18] lower prevalence rate of 37.2% was observed among the females when compared to their counterpart.

![Fig. 1. Distribution of Schistosoma species in Africa](image-url)
males having prevalence of 45.5%. This is attributed to the fact that males are more in contact with the breeding site through swimming, fishing and related activities than their female counterparts who may just visit the stream to fetch water for primary domestic activities [16]. However, other studies revealed that gender distribution of urinary schistosomiasis could change due to some variation in behaviour and cultural practices regarding water activities [19]. Therefore, infection pattern may be attributed to differences in geographic and environmental setting, their cultural and religious beliefs.

4. EVOLUTIONARY ORIGIN OF SCHISTOSOMA

Schistosoma species are found throughout the tropical and developing region of the world, namely Africa, Asia and South America (Fig. 2). The distribution of schistosome parasite is related with the distribution of Schistosoma parasite is related with water contact and geography of the intermediate hosts that each species has adapted to exploit within given regions [20,21]. S. japonicum in Asia has an egg with minute spine and develop in operculate snails of the family Pomatiopsidae. On the other hand, species of the S. mansoni and S. haematobium groups in Africa (and adjacent regions) and South America (S. mansoni only) utilize pulmunate snails. Members of the former group have eggs with a lateral spine, while the S. haematobium group has eggs with a terminal spine, with species from both groups often sharing overlapping geographical ranges [22,23]. Therefore, the size and shape of the egg, intermediate host specificity and the geographical origin of isolates are sufficient to identify species infecting man.

A considerable amount of literature has been published on the origin of schistosome which had earlier developed and reviewed, following argument for both an African and Asian origin [23,24]. However, it was suggested that the genus Schistosoma originated before the separation of the super continent Gondwana land (that was made from today’s Africa, Antarctica, South America, and Australia) for the past 150 million years [24,25]. Moreover, are said to begun in order to exploit pomatiopsid and pulmonate snails, to which extensive fossils records suggested that, is from Gondwanan origin. This shows that spread of parasite were due to continental drift. Similarly suggested that ancestors of the Asian schistosome were translated to Asian when India separated from Africa and merge to Asia, this gives rise to the S. japonicum and S. indicum [24]. This theory also suggest that the remaining other African stock began to radiate over, given rise to the ancestral lineage of the S. haematobium and S. mansoni groups [23].

Fig. 2. Map showing the African origin Schistosoma moving over to Asia on the Indian giving rise to the S. indicum group and diversifying into the S. japonicum suggest the Schistosoma ancestor remaining in Africa diverged giving rise to the S. mansoni and S. haematobium groups
5. SCHISTOSOMA PHYLOGENY AND SPECIATION

Recent years, progress in advances DNA sequencing, molecular tools also have promoted greater records and exploration of genetic diversity of schistosome species and their hosts [23]. A range of genetic markers have been used to construct phylogenetic trees, the nuclear marker ribosomal RNA gene unit (18S, 5.8S, 28S, internal transcribed spacer region (ITS)). Variations in the sequences of mitochondrial cytochrome oxidase sub-unit 1 (cox1) gene were compared among some samples specimens in order to identify the evolutionary differences, and possible similarities. Those studies show a benefit on the knowledge of the complete mitochondrial genome of S. haematobium, and thereby enhancing population focus-studies [27]. Previous study reported that, the genus schistosome was categorized in to 6 clades that correlate to different geographical distributions of the parasites [20,23,28]. In addition, species forming S. japonicum complex (S. ovuncatium, S. sinensium, S. japonicum, S. mekongi, and S. malayensis) are seen on the phylogenetic tree, the position of the clade gives an evidence for the Asiatic origins of this parasite, it suggest an oriental origin and its colonization for both south eastern and the eastern provinces of Asia [20,29,30] as shown in Fig. 3.

On the other hand, the second major split in the phylogeny encompasses the species found throughout Africa. However, there appears to be several distinct clades within this major split, most basal lineage is S. hippopotami clade, containing both S. hippopotami and S. edwardiense, which were both described from the western Uganda [21]. The third category within the phylogeny was described as proto-S. mansoni clade, proven to be basal to the true African species which is represented by two

Fig. 3. Schematic phylogeny of the interrelationships of members of the species within the Schistosoma genus estimated with a Bayesian analysis of combined partial IsrDNA, complete ssrDNA and partial cox1. The tree also illustrates the basal nature of Asian schistosomes, being ancestral to the African stock due to the relative positions of S. hippopotami, S. edwardiense (Inferred from partial cox1), Oriento-bilharzia and S. incognitum (adopted with modification [26])
species, *Orientobilhazia turkestanicum*, now known to belong to the genus *Schistosoma* and *S. incognitum* [23]. Two species are known to be found in Southern and Western Asia, throughout the Middle East, adding more emphasis on the argument of an Asian origin for the African schistosome and may even give an insight into different species that occurred before the colonization of the African continent [20,29,31]. Moreover, African schistosomes form two distinct clades, the *S. mansoni* clade (*S. mansoni* and *S. rodhaini*) and the *S. haematobium* clade (*S. margrebowiei*, *S. guineesis*, *S. leipri*, *S. mattheei*, *S. intercalatum*, *S. haematobium*, *S. curassoni*, *S. bovis* and *S. kisumuensis*). However, between the two major African clades is the *S. indicum* group (*S. indicum*, *S. spindale* and *S. nasale*) which are found throughout Southern and Western Asia suggesting a reinvasion of the Asiatic subcontinent [20,32,33].

6. LABORATORY DIAGNOSIS OF SCHISTOSOMIASIS

Despite the fact that recent information is available among places of genitourinary schistosomiasis like that of global burden of the disease, control and prevention, and other associated morbidity, some practical issues are yet to be clarified as regard to basic diagnostic tools associated with this public health important parasitic disease [34,35]. The direct microscopic sedimentation technique of urine, and examination of urine using filtration method for the detection of schistosome ova including the indirect methods i.e. detection of haematuria, circulating egg antigens, schistosome specific antibodies and ultrasound scans are among others of the genitourinary system of *Schistosoma* infection. Therefore, many reasons are related to the inability in the detection of genitourinary schistosomiasis, citing example with differences in egg deposition levels and/or duration of infection which may influence the potentiality and sensitivity in achieving the most exact and recent infection status [36]. Conventional and new tools to diagnosis, determination of response to therapy, and morbidity assessment are indicated under community and institutional settings in hierarchical order (Fig. 4).

6.1 Filtration Technique

Membrane used in urine filtration remain a common egg concentration technique, although the actual cases of schistosomiasis are high than what is documented after first screening [37,38]. This is especially among individuals who excrete few numbers of eggs [39]. Although the sensitivity of the method can be improved by either increasing the number of samples tested and/or sampling of a larger volume of urine, but such methods are not easily performed. In a testing of monoclonal antibody assay [40] evidence proved that more than eight samples are needed for the examinations of urine through centrifugation and filtration and which are necessary before an infected individual can be identified. However, several studies that assess the accuracy with regards to its sensitivity and specificity of urine filtration method were not seen established. During recent time as mass praziquantel were distributed and well informed mass education, a significant light infection were reported, therefore, it is important that sensitivity of this diagnostic methods should perfectly be characterized [41,42].

6.2 Sedimentation Method

Sedimentation is an important substitute to other techniques especially filtration methods. The sensitivity and specificity of sedimentation method was compared to filtration of 10 mls in low intensity infection, must probably because the intrinsic loss rate is lower and the sample size larger. The presence of epithelial cells for example might likely disguise egg of *S. haematobium* in the sediment. In addition menstrual blood also contaminates urine and lysis of the red blood cells discloses ova in sediment or on filter membrane. This can record success as few drops of 10% hydrochloric acid are added to the urine sample.

6.3 Haematuria (dipstick) Method

Haematuria is a common symptoms of urogenital schistosomiasis, usually appears when *S. haematobium* eggs induced inflammation and blood vessels ruptures [43]. Reagent dipsticks are readily available for testing presence of blood in urine (haematuria), though it often indicates infection; however it is a non-specific marker since haematuria is not specifically caused by *S. haematobium* infection [44]. Several studies carried out were able to estimate the sensitivity, specificity, and false positive rates of the dipstick test [45,46]. Further studies has shown that number of screening tests are possible and could be considered in other to facilitate and improved detection rate, accurately diagnose as well as proper estimates of incidence and prevalence of urinary schistosomiasis within sub-samples of many populations [47,48]. The sensitivity and
specificity of two screening methods; reagent dipstick for haematuria and urine filtration were evaluated among Ghanaian children with light S. haematobium infection. It was highlighted that strips reagent for urine analysis yielded a significant positive result for S. haematobium after diagnosis, with sensitivities of 70% and 97% and specificities of 59 – 80% [48,49], showing concerns as to the possibility of this as the overall technique for screening populations in an endemic setting.

6.4 Immunological Techniques for Diagnosis of Schistosomiasis

To date various methods have been developed and introduced to measure immunological response using crude or purified egg and adult worm antigens in other to detect anti-S. haematobium antibodies [50,51]. Immunodiagnostic technique for detection of light infections is developed by aiming to detect the presence of schistosome circulating Antigens (SCA) in the urine, serum or antibodies specific to schistosome antigens [52]. The commonly reported assays are Enzyme Linked Immunosorbent Assay (ELISA) [53,54], Immunofluorescence or Western Blotting (WB) [55]. However, ELISA is a laboratory based test that is useful for larger-scale operations but its importance in the field is limited [56]. The detection of Soluble Egg Antigen (SEA) in serum and urine could be more valuable in diagnosis; hence early treatment would be applied before irreparable damage occurs [1]. However, an antibody detection method often has its own limitations as it cannot distinguish between current and past infection. Furthermore, it may also present a high level of cross reactivity, molecular methods should be considered for sensitive and specific assay [57]. Kato Katz technique is used for qualitative and semi-quantitative diagnosis of intestinal helminthic infestations; caused by Schistosoma spp. World health organization has recommended Kato Katz technique in areas with moderate to high transmission rates of soil transmitted helminths. People infected with intestinal schistosome pass the eggs of the worms through their faeces. In the Kato-Katz technique faeces are pressed
through a mesh screen to remove large particles. A portion of sieved sample is then transferred to the hole of a template on a slide. After filling the hole, the template is removed and the remaining sample is covered with a piece of cellophane soaked in glycerol. The glycerol clears the faecal material from around the eggs. The eggs are then counted and the number calculated per gram of faeces. The Kato-Katz technique is the diagnostic method recommended for a known species of parasite that is associated with intestinal schistosomiasis and part of the plan for 2020 that focuses on "elimination of disease as a public important". Similarly, direct stool smear, formol-ether concentration technique can detect up to 65.26% of positive intestinal Schistosoma specimens.

6.5 Molecular Based Technique

Recently, many investigators launched molecular approaches for detecting schistosomiasis: Detection of parasite (schistosome) DNA in clinical human samples such as faeces [58], sera [59] and urine [60], confirmed the existence of parasite DNA in the host and provide evidence of the infection. Parasite DNA may be a helpful guideline for selecting the appropriate treatment for schistosomiasis. In another study conducted by [61] reported that data generated obtained from genetic techniques do not only give an insight of the history of these organisms, also into the biology of these parasites that has direct impact on species identification and disease epidemiology. In addition [62] demonstrated new polymerase chain reaction (PCR) primers for the identification and differentiation of major human Schistosoma: S. haematobium, S. mansoni, S. mekongi and S. japonicum. A considerable amount of literature has been published on PCR methods and is proven to have improved the direct detection of Schistosoma. DNA seems to be prepared from egg prior to PCR amplification from urine, stool, or organ biopsy as samples [58,60]. However, DNA is extracted from a small volume of sample success is dependent whether sample processed contains eggs or not [63].

Several studies have revealed that the random amplified polymorphic DNA markers (RAPD) technique can be applied successfully to study helminths [64]. Several DNA-based methods have been developed to investigate genetic diversity, differentiate strains and species and an analyse phylogenies of schistosome [62]. It has been demonstrated that RAPD can be employed to differentiate between strains of S. mansoni and species of Schistosoma.

In order to improve understanding of the epidemiology of the S. haematobium, genetic variation within this species has to be studied extensively [65]. Genomic studies are often carried out on DNA extracted from miracidia, from hatched egg, cercariae released from the snail or adult worms, most often collected from the laboratory animals. In addition, facilities were so limited making the practice so difficult for the isolation and storage of this parasite. Although several attempts has been made to simplify isolation of eggs on the filter paper since it does not require laboratory facilities, DNA can still be successfully extracted at latter stage [66]. Mitochondrial genes have been used in molecular analysis to demonstrate the relationship between different species, because they showed high mutation rates compared to nuclear markers and exist in high number. Moreover, this led to their exploitation for studying phylogenetic variation in these parasites [67,68]. Similarly Schistosoma phylogeny has primarily been based on alignment of multiple gene sequences from a handful of genes such as cox1, cox2, ITS1, ITS2, nad4, rml, and rrnS [21,23]. This might be very vital because it allows the exploration of genetic heterogeneity of parasites among a sample and also between children coming from the same area, the methods could also provide an insight into the knowledge of hybridization among different species after a long period of time as it has been observed that in west Africa, S. haematobium can hybridize S. bovis [69]. However, many children could be infected with one or more schistosome haplotype or even species at a time [70].

Previous studies have reported that presently there were complete mitochondrial genomes available for 6 species of schistosome including S. japonicum, S. malayensis, S. mekongi, S. mansoni, S. spindale and S. haematobium [68,71]. Moreover, in 2001, Le et al. demonstrated that the mitochondrial genomes of S. japonicum and S. mekongi displayed the same gene order as each other and that of other digenea and Cestodes [68]. However, S. mansoni shows several differences in gene order when compared to the species from the S. japonicum clade. The S. mansoni type rearrangement was also seen in S. haematobium and S. spindale [68]. The rearrangement of the gene order among African schistosome and their descendants is of particular interest putting in to consideration the phylogenetic position of S. incognitum and O. turkestanicum and their
geographical differences. These species might provide a clue into the exact time on when these genomic changes happened during the inversion of Africa and the radiation of the schistosome.

7. CONCLUSION

The greatness of molecular methodologies for the examination of schistosome at the individual and community level is presently obvious, particularly for *S. mansoni*. Significant inquiries remain concerning the upkeep of parasite diversity and how schistosome react to selection pressures either during characteristic movement through the life-push or through medication treatment or immunization. Gene revelation and gene mapping activities are prompting a superior comprehension of the schistosome genome and can be relied upon to contribute fundamentally to future relative transformative examinations. Choice and utilization of strategies ought to subsequently compare to the sort of data looked for by the general wellbeing of the epidemiologist or the medical officer and test result interpretation must mull over the downsides and limitations characteristic to every one of the different diagnostic procedures.

CONSENT AND ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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