Mapping freshwater snails in north-western Angola: distribution, identity and molecular diversity of medically important taxa

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Abstract

Background: This study was designed to determine the distribution and identity of potential intermediate snail hosts of Schistosoma spp. in Bengo, Luanda, Kwanza Norte and Malanje Provinces in north-western Angola. This is an area where infection with Schistosoma haematobium, causing urogenital schistosomiasis, is common but little is yet known about transmission of the disease. Angola has had a varied past with regard to disease control and is revitalising efforts to combat neglected tropical diseases.

Methods: Snails were sampled from 60 water-contact points. Specimens of the genera Bulinus, Biomphalaria or Lymnaea were screened for trematode infections by inducing cercarial shedding. Snails were initially identified using shell morphology; subsequently a cytochrome c oxidase subunit 1 (cox1) gene fragment was amplified from a subset of snails from each site, for molecular identification. Cercariae were captured onto FTA cards for molecular analysis. Specimens of Bulinus angolensis collected from the original locality of the type specimen have been characterised and comparisons made with snails collected in 1957 held at the Natural History Museum, London, UK.

Results: In total snails of nine genera were identified using morphological characteristics: Biomphalaria, Bulinus, Gyraulus, Lanistes, Lentorbis, Lymnaea, Melanoides, Physa and Succinea. Significant for schistosomiasis transmission, was the discovery of Bulinus globosus, B. canescens, B. angolensis, B. crystallinus and Biomphalaria salinarum in their type-localities and elsewhere. Bulinus globosus and B. angolensis occurred in two distinct geographical areas. The cox1 sequence for B. globosus differed markedly from those from specimens of this species collected from other countries. Bulinus angolensis is more closely related to B. globosus than originally documented and should be included in the B. africanus group. Schistosoma haematobium cercariae were recovered from B. globosus from two locations: Cabungo, Bengo (20 snails) and Calandula, Malanje (5 snails). Schistosoma haematobium cercariae were identified as group 1 cox1 corresponding to the type common throughout the African mainland.

Conclusions: 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 to identify areas where transmission can occur and where control efforts should be targeted. The molecular phylogeny generated from the samples confirmed that considerable variation exists in B. globosus, which is the primary snail host for S. haematobium in many regions of Africa.

Keywords: Angola, Schistosomiasis transmission, Bulinus globosus, Biomphalaria spp., Schistosoma haematobium

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Background
Intestinal and urogenital schistosomiasis, caused by the trematode worms *Schistosoma mansoni* and *S. haematobium*, respectively, are endemic in Angola. In 2010, the World Health Organization (WHO) estimated that 60% of Angola’s population were living at significant risk of schistosomiasis infection and required preventive chemotherapy [1]. This prevalence equates to 14.6 million people according to the most recent census [2]. However, these estimates are based on surprisingly few epidemiological surveys and non-systematically gathered data. Due to this lack of up-to-date information, there is an urgent need for further investigations relating to schistosomiasis and other neglected tropical diseases (NTDs) in Angola encompassing epidemiology, transmission studies and operational research.

While much can be done to control morbidity of schistosomiasis using large-scale community or school mass drug administration of praziquantel [3, 4], it is clear that sustainable control and elimination will need a better understanding of transmission and water contact, as highlighted in the WHO roadmap for control of NTDs [5]. Like many other sub-Saharan African countries, Angola is now mapping schistosomiasis, among other NTDs, in humans. However, very little is known about the diversity of freshwater snails responsible for transmission; in fact, the last large-scale survey of freshwater snails was published more than 50 years ago by Wright in 1963 [6].

Looking back further, to the work of Morelet in 1866 and 1868 [7, 8], it is clear that Angola is an important country in the history of medical malacology, from which many important snails were first described. In fact, the type-localities of many species of *Bulinus* and one species of *Biomphalaria* are in Angola: *Bulinus globosus* (Morelet, 1866), River Dande, Bengo Province; *B. angolensis* (Morelet, 1866), district of Duque de Braganza (now Calandula); *B. canescens* (Morelet, 1868), marshes near River Bengo near Quicuxi; *B. crystallinus* (Morelet, 1868); river close to Golungo Alto and *Biomphalaria salinarum* (Morelet, 1868), tributaries to the River Cuie near Malange (see Brown [9]). Snails from Angola have not previously been collected and examined by modern methods of molecular characterization; hence, there is a need to carry out a comprehensive study to learn more about schistosomiasis and the snails responsible for the transmission of this debilitating disease in Angola and their relationships with other African taxa.

This investigation aimed to determine freshwater snail biodiversity and to identify potential intermediate snail hosts of schistosomes in four provinces in north-western Angola. By collecting fresh samples, comparing them to samples collected in the same locations by C. A. Wright nearly 60 years before [6], and subjecting them to modern molecular analysis we attempted to better characterise the species of the genera *Bulinus* and *Biomphalaria* currently found in Angola and to provide new molecular markers for species identification.

Methods

Locations
Fieldwork was conducted during November and December 2013, by JCSF, AL, AMS, CM and DR. The Dande, Bengo, Kwanza, Lucala and Cuie river basins were visited. Accordingly, a total of four provinces were visited and extensively surveyed: Bengo (25 sites), Luanda (10 sites), Kwanza Norte (12 sites) and Malanje (13 sites), see Fig. 1a for graphical representation and Table 1 for site details.

Malacology procedures
Water bodies were examined for freshwater snails using a standard protocol [9]. Information was recorded on pre-designed forms and included: general locality information (water body name, type, GPS coordinates, nearby village); water properties data (temperature, pH, salinity, dissolved solids, flow rate, water level, conductivity, depth); collector information (number of collectors, time/length of time of collection), ecological data (substrate, all species of snail present, vegetation, animal contact, human contact), and snail data (snails species, abundance, number infected).

Snails were collected using large scoops, hand sieves and forceps for approximately 22 min per site usually with 3–4 persons. Snails were carefully picked out with forceps taking care not to damage the shells. The collection at each site was separated into genera and placed in screw cap pots with clean mineral water before securing in a cooler with ice packs when available) for transportation. Photographs were taken of each site visited. Upon return to Centro de Investigação em Saúde de Angola (CISA) (Angolan Health Research Center, in Caxito) *Bulinus* and *Biomphalaria* specimens were placed in fresh water and checked for the shedding of cercariae by exposing them to a lamp or sunlight, any emerging cercariae were examined under the microscope and identified by morphology. Snails were screened, individually, for a period of 24 h and then fixed in ethanol. Individual cercariae in 4 μl of water, collected by pipetting, were placed onto Whatman FTA Classic cards (GE Healthcare Life Sciences, Amersham, UK) for future molecular analyses [10]. Snails were relaxed by briefly placing in a -20 °C freezer and then fixed in ethanol (95%). Screw top 25 ml universal tubes were labelled both internally using pencil and parchment paper and externally with permanent marker.
Molecular characterization studies

Sample preparation and DNA extraction

The snail samples selected for the molecular analyses represented individuals from each collection site. Examples of all species of *Bulinus* and *Biomphalaria* found in this collection trip were included. Additionally, a single sample from C. A. Wright’s 1957 collection held at the Natural History Museum was included in these analyses. All specimens were stored in 95% ethanol in the field and the number of snails recounted, identified by morphological characters and re-spirited (absolute ethanol) on arrival at the Natural History Museum, London (NHM) for incorporation into the Schistosomiasis Collection at the Natural History Museum (SCAN) [11]. Photographic images were taken of the snail shells prior to DNA extraction. Specimens were placed in TE buffer (10 mM Tris, 0.1 mM EDTA) pH 7.4 for 1 h in order to remove any remaining alcohol from within the tissue, which might interfere with subsequent extraction techniques. Total genomic DNA was isolated from head-foot snail tissue using the DNeasy Blood and Tissue kit (Qiagen, Crawley, UK) according to the manufacturer’s instructions except that the volume of buffers was doubled. DNA was eluted into 200 μl sterile water.

Amplification of cox1 fragments of snail DNA

A polymerase chain reaction (PCR) amplification of a partial cytochrome c oxidase subunit 1 (cox1) sequence was performed using primers LCO1490 (5'-GGT CAA CAA ATC ATA AAG ATA TTG G-3' forward) and HCO2198 (5'-TAA ACT TCA GGG TGA CCA AAA AAT CA-3' reverse) [12]. PCR investigations and sequencing conditions were chosen as previously outlined [13]. A second set of primers was required for the PCR and sequencing of *B. globosus* and *B. angolensis* specimens; these were designed using whole mitochondrial genome data for these species (novel): BulAng61F (5'-GTA TGA TGC GGC CTG GTA GG-3') and BulAng895R (5'-AAG CCC GAG TAT CCA CAT CT-3').
## Table 1: Sites visited during field-work

| Code | Name                                                                 | Province   | Type                   | Closest village       | Latitude (E) | Longitude (S) | Altitude (m) |
|------|----------------------------------------------------------------------|------------|------------------------|-----------------------|--------------|---------------|--------------|
| 1    | Mabubas Dam                                                          | Bengo      | Dam                    | Mabubas               | 13.6999      | 8.5343        | 70           |
| 2    | River Dande and marshes                                              | Bengo      | Marshes                | Sogramo Farm          | 13.74204     | 8.52106       | 65           |
| 3    | Caxito canal (North)                                                 | Bengo      | Canal                  | Caxito                | 13.68946     | 8.55546       | 33           |
| 4    | Caxito canal (Centre)                                                | Bengo      | Canal                  | Caxito                | 13.65779     | 8.57986       | 19           |
| 5    | Cabungo stream (type-locality for *Bulinus globosus*)                | Bengo      | Stream                 | Cabungo               | 13.53737     | 8.57579       | 22           |
| 6    | Lake Cabundo                                                         | Bengo      | Lake                   | Cabungo               | 13.53983     | 8.57282       | 19           |
| 7    | Natural dam                                                          | Bengo      | Dam                    | Cabungo               | 13.53779     | 8.58734       | 12           |
| 8    | Irrigation canal                                                     | Bengo      | Canal                  | Cabungo               | 13.53824     | 8.58765       | 22           |
| 9    | River Dande flood plain                                              | Bengo      | River                  | Talelo                | 13.731       | 8.52747       | 69           |
| 10   | Decommissioned irrigation canal                                       | Bengo      | Canal                  | Icau Centro           | 13.77906     | 8.64375       | 42           |
| 11   | Lake formed by the River Úcua floodplain                             | Bengo      | Lake                   | Icau Wando            | 13.79642     | 8.6451        | 49           |
| 12   | River Lifune                                                         | Bengo      | River                  | Libongo               | 13.46949     | 8.39041       | 56           |
| 13   | Irrigation canal close to Libongo                                    | Bengo      | Canal                  | Libongo               | 13.49718     | 8.38692       | 56           |
| 14   | River Tanda (downstream of dam)                                      | Bengo      | River                  | Quibaxe               | 14.59702     | 8.51995       | 758          |
| 15   | River Tanda (upstream of dam)                                        | Bengo      | River                  | Quibaxe               | 14.59674     | 8.52083       | 762          |
| 16   | Stream Quizende                                                      | Bengo      | Stream                 | Quibaxe               | 14.59362     | 8.50073       | 700          |
| 17   | Merge point between the rivers Tanda and Quizende                    | Bengo      | River                  | Quibaxe               | 14.60326     | 8.51774       | 742          |
| 18   | River Calua                                                          | Bengo      | River                  | Pango Aluquem         | 14.4808      | 8.66434       | 580          |
| 19   | River Úcua (1)                                                       | Bengo      | River                  | Cacamba               | 14.64988     | 8.64988       | 329          |
| 20   | River Úcua (2)                                                       | Bengo      | River                  | Ucau                  | 14.16441     | 8.65763       | 240          |
| 21   | Lake Panguila (1) (potential type-locality for *Bulinus canescens*)  | Bengo      | Lake                   | Porto mangueiras      | 13.45476     | 8.71929       | 0            |
| 22   | Lake Panguila (2)                                                    | Bengo      | Lake                   | Porto mangueiras      | 13.45443     | 8.72003       | 9            |
| 23   | Canal from Lake Panguila (1)                                         | Bengo      | Canal                  | Burgalheira escola    | 13.55499     | 8.79666       | 18           |
| 24   | Canal from Lake Panguila (2)                                         | Bengo      | Canal                  | Burgalheira escola    | 13.55088     | 8.80342       | 15           |
| 25   | Canal from Lake Panguila (3)                                         | Bengo      | Canal                  | Burgalheira escola    | 13.54862     | 8.80783       | 16           |
| 26   | Bridge over the Lake Panguila                                        | Luanda     | Lake                   | Panguila              | 13.43812     | 8.73092       | 18           |
| 27   | River Bengo                                                          | Luanda     | River                  | Funda                 | 13.55313     | 8.84427       | 19           |
| 28   | Lake Quilunda (1) (potential type-locality for *Bulinus canescens*) | Luanda     | Lake                   | Mucoilo               | 13.58681     | 8.86712       | 25           |
| 29   | Lake Quilunda (2) (potential type-locality for *Bulinus canescens*) | Luanda     | Lake                   | Mucoilo               | 13.58789     | 8.86708       | 23           |
| 30   | Lake Quilunda (3)                                                    | Luanda     | Lake                   | Cadianzala            | 13.62339     | 8.90514       | 20           |
| 31   | Artificial lagoon on Aurora farm                                     | Luanda     | Lagoon                 | Onga Zanga            | 13.67903     | 9.0191        | 31           |
| 32   | Pool formed by River Kwanza                                          | Luanda     | Pool                   | Candimba/Muxima       | 13.96911     | 9.53437       | 21           |
| 33   | Cambemba Lagoon (1)                                                  | Luanda     | Lagoon                 | Cacefo                | 13.77959     | 9.27866       | 18           |
| 34   | Cambemba Lagoon (2)                                                  | Luanda     | Lagoon                 | Cacefo                | 13.78779     | 9.26822       | 16           |
| 35   | Canal from the River Kwanza                                          | Luanda     | Canal                  | Caquila               | 13.468       | 9.1594        | 4            |
| 36   | Pool near Aldeia Nova (1) (type-locality for *Bulinus crystallinus*) | Kwanza Norte| Pool                   | Aldeia Nova           | 14.24768     | 9.21573       | 201          |
| 37   | Pool near Aldeia Nova (2)                                             | Kwanza Norte| Pool                   | Aldeia Nova           | 14.36089     | 9.1771        | 239          |
| 38   | Unknown stream near Golungo Alto                                     | Kwanza Norte| River                  | Golungo Alto          | 14.59253     | 9.17718       | 308          |
| 39   | River Luinha                                                         | Kwanza Norte| River                  | Luinha                | 14.66055     | 9.21806       | 333          |
| 40   | River Lussué                                                        | Kwanza Norte| River                  | Lussue                | 14.83917     | 9.2864        | 591          |
The PCR conditions were as outlined by [13] except an annealing temperature of 60 °C instead of 40 °C was used. Sequencing was performed on an Applied Biosystems 3730XL analyser (Life Technologies, UK).

Extraction and amplification of cox1 and ITS2 fragments of Schistosoma haematobium

Punches (2.0 mm) were taken from the centre of each cercarial spot on FTA cards and pH elution performed to remove the DNA from the card matrix [14]. A 1500 bp fragment was amplified from 3 μl of genomic DNA using the Cox1_Schist_5′ and Cox1_Schist_3′ [15] primers and illustra™ puReTaq Ready-To-Go PCR Beads (GE Healthcare Life Sciences, UK). The following cycling conditions were used: 95 °C for 1 min, 40 cycles of 95 °C for 30 s, 40 °C for 30 s and 72 °C for 2 min, with a 7 min extension of 72 °C. In addition, ITS2 was amplified as it would identify schistosome hybrids, the PCR used ITTS1 and ITTS2 primers [15] with the above conditions except the annealing temperature of 56 °C instead of 40 °C. PCR products were cleaned using the QIAquick PCR purification kit (Qiagen, UK). Samples were sequenced in both orientations on an Applied Biosystems 3730XL analyser running BigDye v3.1 sequencing chemistry.

Phylogenetic analysis of sequence data

The electropherograms produced were checked and cox1 sequences edited using Geneious, version 6.1.8 (http://www.geneious.com [16]). Sequences were compared to database entries by performing BLAST searches via the National Center for Biotechnology Information against GenBank and EMBL sequence databases; and aligned with reference material [13] using Geneious version 5.6. The cox1 data for all taxa were analysed solely as nucleotides and phylograms were produced from the alignments using PhyML with automatic model selection by SMS [17, 18] using the Akaike Information Criterion. Branch support was estimated using 1000 bootstrap replications. Additionally, B. truncatus (Niger [13]) was used as an outgroup for the Bulinus forskalii species group, and B. forskalii (Niger [13]) was used for the Bulinus africanus species group. Different outgroups were used for the different analyses based on suitability (geographical and sister taxa). DNA sequences have been submitted to the European Nucleotide Archive with accession numbers LT671915–LT671982.
Results

Locations
A total of 60 sites were visited and included rivers, streams and canals. Sites on the Bengo, Kwanza and Cuije River Basins displayed the highest mean snail biodiversity, with more than 3 genera identified (Table 2). In total, 73% of sites had a mud substrate (usually associated with roots of aquatic and peripheral vegetation), with little domestic animal water contact observed (of the 60 sites 3% had cows, 8% goats, 13% pigs, 7% dogs, 3% horses and 7% had chickens). Wildlife was present in some sites: in 18% of the sites aquatic reptiles were observed or believed to be present, in 3% of the sites hippopotamuses were observed or believed to be present, and in 27% of the sites aquatic birds were observed. Finally, the most prevalent vegetation identified in all 60 sites was grass (57%), macrophyte plants (52%), lilies (37%), rushes (35%) and water hyacinths (13%), in descending order. The water chemistry and other factors did not significantly impact on the presence or absence of snail species (Table 2; Additional file 1: Table S1).

Malacology
In total 1265 snail samples were accessioned into SCAN. According to morphological evaluations, species of a total of 41 species were used for molecular analyses (Table 2; Additional file 1: Table S1). In total 1265 snail samples were accessioned into SCAN.

Analysis of sequence data
The mitochondrial cox1 gene sequence generated for all the Bulinus species in this study was found to be variable, in agreement with previous work [13]. Initially the universal “Folmer” primers, effective across a wide range of taxa, were used that cover approximately the first 600 bp of cox1; however for B. globosus and B. angolensis these standard primers did not amplify the target region. By examining whole mitochondrial DNA sequence (unpublished) new primers were designed for these species, which allowed the amplification of a comparable region of DNA sequence. Phylogenetic analysis of the sequence data for snails identified as B. globosus and B. angolensis revealed that the B. globosus snails collected for this study formed a monophyletic group distinct from B. globosus found elsewhere (Fig. 2). Additionally, the snails identified as B. angolensis formed another distinct clade with the specimen from Wright’s 1957 study, previously identified as B. globosus. Snails from Angola identified as B. forskalii were not monophyletic with respect to B. canescens or B. crystallinus, whereas B. forskalii (and B. camerunensis) from elsewhere formed a separate clade (Fig. 3). Phylogenetic analysis of Biomphalaria salinarum suggested a close affinity with Biomphalaria pfeifferi (not shown); DNA sequence was highly similar to B. pfeifferi for the cox1 barcoding region (up to 99% similarity).

Parasitology
Of the Bulinus spp. and Biomphalaria spp. specimens collected, only Bulinus globosus (25 in total) was found to be shedding Schistosoma cercariae. These were found in 2 sites: Site 5 [20 out of 105 snails shedding (19% prevalence)] and site 60, the source of the River Cata [5 out of 68 (0.07% prevalence)]. The cox1 and ITS2 sequences of the cercariae sampled corresponded to haplotype H1 of S. haematobium, by far the most widespread across Africa [19]. Apart from human parasites, B. globosus specimens were also found to be shedding Trichobilharzia, echinostome, amphistome and strigeid cercariae. Biomphalaria crystallinus specimens were found to be shedding cercariae of Trichobilharzia

Table 2 Summary of water chemistry parameters and biodiversity of each river basin

| River basin | Dande/Úcua | Bengo | Kwanza | Luella | Cuije |
|-------------|------------|-------|--------|--------|------|
| Number of sites | 20 | 11 | 4 | 22 | 3 |
| Altitude (m) | 232 (12–762) | 17 (0–31) | 14 (4–21) | 220 (201–1111) | 1045 (1038–1051) |
| Temperature (°C) | 28.3 (25.4–32.8) | 30.4 (27.8–33.1) | 32.7 (30.9–33.6) | 31.5 (22.5–32.1) | 24.2 (22.1–27.2) |
| TDS (ppm) | 205 (42–736) | 398 (138–830) | 69 (59–80) | 119 (26–612) | 204 (156–276) |
| pH | 7.75 (6.96–8.79) | 7.71 (6.88–8.70) | 7.02 (6.54–7.82) | 9.26 (6.85–9.60) | 7.76 (7.07–8.31) |
| Conductivity (m/s) | 245 (59–795) | 562 (194–1171) | 96 (83–112) | 159 (65–860) | 288 (220–397) |
| Salinity (ppm) | 125 (33–389) | 276 (95–581) | 52 (46–58) | 84 (35–415) | 139 (106–190) |
| Average biodiversity (no. of genera) | 1.35 (0–4) | 3.27 (2–5) | 3.25 (3–4) | 1.5 (0–2) | 3.3 (1–5) |

Note: Data are given as mean (range)
and echinostomes. The non-schistosome cercariae were identified using morphology and swimming characteristics.

**Discussion**

Angola is a particularly rich area for type-localities of both *Bulinus* and *Biomphalaria* species and it is of interest that similar diversity of snails still exists in locations sampled 56 years previously. The fact that the survey was conducted in November/December had its advantages and disadvantages. Importantly, C. A. Wright visited these same areas in November/December of 1957 [6], giving concordance in sampling season between the two studies. However, the survey was conducted at the start of the rainy season, when many of the rivers and tributaries were running low or even dry (namely in the Úcua River basin). Nevertheless, the level of snail biodiversity in some sites was high, as well as the abundance. For example, in the Caxito canal (sites 3 and 4), the number of *Melanoides tuberculata* and *Physa acuta* was large and may in part account for the absence of *Bulinus* spp.

The 37 recognised species of *Bulinus* have been divided for convenience into four groups: the *B. africanus* group, the *B. reticulatus* group, the *B. forskalli* group and the *B. truncatus/tropicus* complex [9], while others studies have argued for subdivision of the genus into three genera [20]. The lack of clear morphological characters for species identification provided the impetus to search for molecular solutions and *cox1* barcoding was shown to provide good species discrimination and agreed for the most part with the taxonomy based on morphological criteria [9]. The dataset generated here can be added to and compared with the earlier *cox1* sequences considered by Kane et al. [13].

One of the most striking findings is the close association observed between *B. globosus* and *B. angolensis*. The latter species had previously been placed in a different species groups: *B. globosus* within the *B. africanus* group while *B. angolensis* had been included in the *B. tropicus/truncatus* complex. Brown [9] made the point that treatment of this species is unclear partly because the chromosome number and molecular properties were unknown. Wright [6] noted that he did not have sufficient material to assess properly the relationships of this species or to fully describe the range of variation. Our own material has so far been limited but based on current data it is safe to consider *B. angolensis* as a member of the *B. africanus* group and not a member of the *B. tropicus/truncatus* complex.

Another observation that deserves further, detailed studies is the relationship of the samples included as *B.
globosus. Kane et al. [13] drew attention to the fact that there was a clear division between samples originating from East and West Africa. Bulinus globosus has almost a pan-African distribution. The trees generated here on cox1 sequence data suggest that B. globosus from Angola are quite distinct from other geographical areas. This observation might cause a taxonomic dilemma as the samples analysed here are in fact from the type-locality and hence represent the "true" B. globosus. It is clear that this snail species, which acts as an important host for S. haematobium and other schistosome species throughout its range, needs a more thorough investigation. While changes in nomenclature might eventually be warranted we suggest that further sampling is required and more comprehensive molecular data are needed before such changes are made.

Additionally, a single sample from C. A. Wright’s 1957 collection held at the Natural History Museum as part of SCAN was included in these analyses and can be seen to be most similar to B. angolensis from the 2013 collection. The cox1 sequence gained from this snail is particularly interesting as the specimen had been held in industrial methylated spirit for the past 60 years; samples held in this way are generally fragmented as the DNA can be damaged by this storage medium. Wright’s original identification on morphological grounds alone was that this specimen was best considered as B. globosus but the molecular data strongly supports the grouping with B. angolensis.

Other specimens collected at type-localities were from the B. forskali group. Again, analysis of the cox1 regions shows that the samples from Angola are diverse and not necessarily falling into the groups that were previously expected. It is clear that there are at least two different groupings (Fig. 3); first “B. forskali” types with the inclusion of B. canescens potentially as a subgroup of B. forskali, and the B. crystallinus types which are distinct from the main clade. It is of interest to note here that Brown [9] discussed the lumping of these species and thought that there may be up to 12 separate species within Angola; these points were also made by Jones et al. [20] and Kane et al. [13]. Species assignment based on locality and morphology looks somewhat arbitrary when compared with phylogenetic analysis using sequence data, especially for the B. forskali group. While cox1 barcoding has been used extensively for snail vectors (e.g. [13]), it is clear that further investigation at the molecular level is needed to elucidate and identify and relationship of species within this species group prior to any changes in nomenclature.

Biomphalaria species were collected from only a couple of locations in this study, ultimately only two
specimens had unique cox1 sequence and therefore were included in this analysis. From the analysis B. salinarum is found at two sites and is most closely related to B. pfeifferi.

With reference to the schistosome species found in this study, to our knowledge this is the first time that S. haematobium has been genetically typed from Angola. Schistosoma haematobium cercariae recovered from B. globosus showed to be standard H1 cox1 haplotype and the three samples examined had a low diversity (only 3 single nucleotide changes). This is what we would expect from the African mainland from previous studies [19]. No other animal schistosomes were encountered, but a range of species of cercariae were produced by a number of species of snail; this is usual in water-contact sites in use by humans, animals (wild and domestic) and birds. Interestingly, Biomphalaria species were not common suggesting that intestinal schistosomiasis caused by S. mansoni is unlikely to be a major public health problem in the region [21].

As schistosomiasis control programmes gather pace it is important that malacological studies are carried out to help guide mapping of disease transmission and the planning of control interventions. Focus is moving towards the elimination of schistosomiasis [4, 5] and a recent analysis has shown the value of interventions that control snail populations [22]. The increase in data collection from sites, with geo-referencing, and with molecular techniques gives a more detailed view on transmission and where to target control efforts. This study has initiated mapping of snail distributions in four provinces of Angola, important site localities have been geo-referenced and photographic records have been made at each of the collection points. There is now a need to extend the survey into other regions of Angola.

Conclusions
The findings reported here provide new insights into the molecular diversity within and between the Bulinus species examined. The samples of medically important snails collected and identified by the barcoding approach will be subjected to further more in depth molecular sequencing to help elucidate the relationships with Bulinus and Biomphalaria species from other African regions.

Additional file

Additional file 1: Table S1. Details of snails at each site. (XLS 48 kb)

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Availability of data and materials
The datasets generated during and/or analysed during the current study that are not included in this published article are available from the Schistosomiasis Collection at the Natural History Museum (SCAN) http://scan.myspecies.info. DNA sequences have been submitted to the European Nucleotide Archive with accession numbers LT671915–LT671982.

Authors’ contributions
Conception and design of the study: JCSF, DR, FA and MB. Acquisition of data: FA, JCSF, DR, AE, RP, CM, AS and MB. Analysis and interpretation of data: FA, JCSF and AE. Drafting the article: FA, JCSF and DR. Revising the article critically for important intellectual content: FA, AE, JCSF, MB and DR. All authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

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References
1. WHO. Country Profile: Preventive chemotherapy and transmission control. Geneva: World Health Organization; 2010. http://www.who.int/neglected_diseases/preventive_chemistry/databank/CP_Angola.pdf?ua=1
2. INE. 2014 National Census, vol. 2014: Instituto Nacional de Estatística. http://www.ine.gov.ao
3. WHO. Preventive Chemotherapy in Human Helminthiasis. Coordinated use of Anthelminthic Drugs in Human Interventions: a Manual for Health Professionals and Programme Managers. Geneva: World Health Organization; 2006.
4. Rollinson D, Knopp S, Levitz S, Stothard JR, Tchueni Tchuenté L-A, Garba A, et al. Time to set the agenda for schistosomiasis elimination. Acta Trop. 2013;128:423–40.
5. WHO. Accelerating work to overcome the global impact of neglected tropical diseases: a roadmap for implementation. Geneva: World Health Organization; 2011.
6. Wright CA. The freshwater gastropod mollusca of Angola. Bull Brit Mus (Nat Hist). 1963;10(8):447–528.
7. Morelet A. Coquilles nouvelles recueillies par le Dr. Fr. Welwitch dans l’Afrique équatoriale, et particulièrement dans les provinces portugaises d’Angola et de Benguela. Conchylol. 1866;6(3):153–63.

8. Morelet A. Molliusques terrestres et fluviatiles. Voyage du Dr. Friederich Welwitch. Bailliere, Paris: Bailliere; 1868.

9. Brown D. Freshwater snails of Africa and their medical importance. Oxford: Taylor & Francis; 1994.

10. Gower CM, Gabrielli AF, Sacko M, Dembélé R, Golan R, Emery AM, et al. Population genetics of Schistosoma haematobium: development of novel microsatellite markers and their application to schistosomiasis control in Mali. Parasitology. 2011;138:978–94.

11. Emery AM, Allan FE, Rabone ME, Rollinson D. Schistosomiasis collection at NHM (SCAN). Parasit Vectors. 2012;5:185.

12. Folmer O, Black M, Hoeh W, Lutz R. DNA primers for amplification of mitochondrial cytochrome c oxidase subunit I from diverse metazoan invertebrates. Mol Mar Biol Biotechnol. 1994;5:294–9.

13. Kane RA, Stothard JR, Emery AM, Rollinson D. Molecular characterization of freshwater snails in the genus Bulinus: a role for barcodes? Parasit Vectors. 2008;1:15.

14. Webster BL, Rabone M, Pennance T, Emery AM, Allan F, Gouvras AN, et al. Development of novel multiplex microsatellite polymerase chain reactions to enable high-throughput population genetic studies of Schistosoma haematobium. Parasit Vectors. 2015;8:432.

15. Lockyer AE, Olson PD, Østergaard P, Rollinson D, Johnston DA, Attwood SW, et al. The phylogeny of the Schistosomatidae based on three genes with emphasis on the interrelationships of Schistosoma Weinland, 1858. Parasitology. 2003;126(03):203–24.

16. Keanse M, Moir R, Wilson A, Stones-Havas S, Cheung M, Sturrock S, et al. Geneious Basic: an integrated and extendable desktop software platform for the organization and analysis of sequence data. Bioinformatics. 2012;28(12):1647–9.

17. Guindon S, Gascuel O. A Simple, fast, and accurate algorithm to estimate large phylogenies by maximum likelihood. Syst Biol. 2003;52:696–704.

18. Lefort V, Longueville J-E, Gascuel O. ATGC: PhyML with automatic model selection by SMS. http://www.atgc-montpellier.fr/phyml_sms. Accessed 12 Oct 2016.

19. Webster BL, Emery AM, Webster JP, Gouvras A, Garba A, Diaw O, et al. Genetic diversity within Schistosoma haematobium: DNA barcoding reveals two distinct groups. PLoS Negl Trop Dis. 2012;7(2):10.

20. Jones CS, Rollinson D, Mimpfoundi J, Ouma J, Kariuki HC, Noble LR. Molecular evolution of freshwater snail intermediate hosts within the Bulinus forskali group. Parasitology. 2001;123:5277–92.

21. Sousa-Figueiredo JC, Gamboa D, Pedro JM, Fançony C, Langa AJ, Soares Magalhães RJ, et al. (2012) Epidemiology of malaria, schistosomiasis, geohelminths, anemia and malnutrition in the context of a demographic surveillance system in Northern Angola. PLoS One. 2012;7(4):e33189.

22. Sokolow SH, Wood CL, Jones LI, Swartz SJ, Lopez M, Hiesh MH, et al. Global assessment of schistosomiasis control over the past century shows targeting the snail intermediate host works best. PLoS Negl Trop Dis. 2016; https://doi.org/10.1371/journal.pntd.0004794.