

RESEARCH ARTICLE

Detection of cutaneous leishmaniasis in three communities of Oti Region, Ghana

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Abstract

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Background

Cutaneous leishmaniasis (CL) is the most common type of leishmaniasis, a neglected tropical disease caused by parasites of the genus *Leishmania*. In Ghana, some studies in the Volta region have detected *Leishmania* parasites among persons with skin ulcers.

Methodology/Principal findings

Using a cross-sectional study design, the prevalence of CL in three communities of the Oti Region of Ghana was investigated. Demographic and epidemiological data were obtained by a structured interviewer administered questionnaire. A total of 426 (12.4%) out of 3,440 participants screened had at least one skin ulcer. Of 595 skin ulcers sampled and tested by PCR for *Leishmania* infection, 150 (25.2%) ulcers from 136 individuals tested positive, accounting for an overall CL prevalence of 31.9% among persons with skin ulcers. Individual community CL prevalence of 23.2%, 29.8%, and 36.8% was observed in Ashiabre, Keri, and Sibi Hilltop respectively among persons with skin ulcers.

Conclusions/Significance

Confirmation of CL in the study area suggests an active cycle of transmission of *Leishmania* infection. The observation of skin ulcers which tested negative to *Leishmania* infection suggests a need to test for additional causes of skin ulcers such as *Treponema pallidum pertenuis* and *Mycobacterium ulcerans* in the study area.

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Author summary

Cutaneous leishmaniasis (CL) is a neglected tropical disease caused by parasites of the genus *Leishmania* and is transmitted by various species of female sandflies. CL usually begins as painless nodules and is usually characterized by skin ulcers which may be single or multiple. Although ulcers due to CL are often self-healing, they may become painful and result in scarring after healing. Typically, CL occurs on exposed parts of the body such as the neck, limbs, and face, which may be easily accessible to sandflies. Although cases of CL had previously been reported in some parts of the Volta region of Ghana, no previous case of CL had been reported in the Oti region. This study was initiated following reports of skin ulcers which were suggestive of CL in some communities of the Oti region. This study confirmed *Leishmania* infection in 150(25.5%) out of 595 skin ulcer samples obtained from 426 study participants. Given that 445(74.8%) of the skin ulcers tested negative for *Leishmania* parasite suggests a need for investigation of additional causes of skin ulcers such as yaws and Buruli ulcer in the study area.

Introduction

Cutaneous leishmaniasis (CL) is an important neglected tropical skin disease (skin NTD) of public health importance and is the commonest form of leishmaniasis, characterized by skin lesions which may result in ulcers, scars, disability and stigma [1,2]. Globally, it is estimated that between 0.7 to 1.3 million new cases of CL are reported annually [3].

A localized outbreak of skin ulcers suspected to be cases of CL was first reported in Ghana from the Ho municipality of the Volta Region in 1999 based on the identification of *Leishmania* amastigotes in some skin lesion biopsies [4]. Subsequent studies have identified *L. major*, uncharacterized *Leishmania* species, and recently, members of the *Leishmania enriettii* complex from suspected CL cases in the Ho municipality, suggesting a complex epidemiology of CL in the region [5–7].

Although the Oti region has been part of the Volta region until the year 2019, no previous confirmation of CL cases had been made there. This study was therefore initiated following reports of skin ulcers which were suggestive of CL in some communities of the Oti region, after leishmanin skin test (LST) had been conducted to establish *Leishmania* infection (reported elsewhere).

Materials and methods

Ethics statement

Ethical approval to conduct this study was obtained from the ethics review committee of the Ghana Health Service (GHS-ERC006/08/18). Written informed consent was obtained from all study participants. For participants under 18 years, written consent was obtained from a parent or guardian.

Study design

This study was based on a cross-sectional study design approach. The study was conducted from October to December 2018 in three communities of the Oti region of Ghana having at least three suspected cases of active CL (ACL). A suspected ACL lesion was defined clinically as any open ulcer with diameter bigger than 5mm. Prevalence of CL among study participants

with skin ulcers was investigated. Demographic and epidemiological data were obtained by a structured interviewer administered questionnaire.

Study area

This study was conducted in the following three communities of the Oti region of Ghana: Ashiabre, Keri, and Sibi Hilltop. Ashiabre is in the Tutukpene sub-district of the Nkwanta South municipality while Keri is in the Keri sub-district of the municipality. Sibi Hilltop is in the Sibi sub-district of the Nkwanta North district of the region.

The population of Nkwanta South municipality is estimated to be 117,878 with males constituting 49.6% of the population. Covering a land area of approximately 2733 km², the Nkwanta South municipality is located between latitudes 7° 30' and 8° 45' North and longitude 0° 10' and 0° 45' East [8].

The population of the Nkwanta North district is estimated to be 64,553 with males constituting 50.2% of the population. The district is located between Latitude 7° 30' N and 8° 45' N and Longitude 0° 10' W and 045° E. It shares boundaries with Nkwanta South municipality to the south, Nanumba South to the north, Republic of Togo to the east, and Kpandai District to the west [9].

Inclusion criteria

Eligible study participants were residents in the study community for ≥ 12 months, aged between 2 to 65 years (inclusive).

Sample size considerations

For active case detection, assuming a current CL prevalence (P) of 22.1% [4,10], $Z^2 = (1.96)^2$ for 95% confidence interval $D^2 = \text{maximum } 0.05$, a minimum sample size (N) of 265 individuals was required for screening for active case detection using the formula:

$$N = ((Z)^2 P/D^2) * (1-P)$$

Selection of households for study inclusion

Using a sorted list of households, 200 households (with an average of 5–7 persons per household) were selected for study inclusion in each study community using a systematic sampling approach. For this study, a household was defined as a person or a group of persons, who live together in the same house or compound and share the same house-keeping arrangements. The head of each household was defined as a male or female member of the household recognised as such by the other household members. The head of a particular household is generally the person with economic and social responsibility for the household. As a result, household relationships were defined with reference the household head [11]. The community household list was obtained for each study community based on a household census. The number of households per study community determined by household census was 945, 795, and 1184 in Ashiabre, Keri, and Sibi Hilltop respectively.

A sampling interval I was determined, where $I = N/n$ with N being the sum of individual households in the study community while n was the number of households to be selected. The I was rounded to 2 decimal places.

Using Microsoft excel, the RANDBETWEEN command was used to generate a random decimal integer R between 0 and 1 rounded up to two decimal points. The sequence of households that were selected in each study community were R^*I , $R^*I + I$, $R^*I + 2^*I$, $R^*I + 3^*I$, ... $R^*I + (n^{-1})^*I$, each rounded up to the next highest whole number [12]. With the

assistance of community-based volunteers, the selected households were identified after which all members of the selected households aged 2 to 65 years were invited to participate in the study, using a door-to-door invitation approach. Because the invitation to participate in the study was extended to households, a household was not included in the study if the household head declined to allow his or her household to participate in the study. However, the agreement of the household head did not make it compulsory for every household member of age 2 to 65 years to participate in the study. Each household member was given the opportunity to go through the informed consent process to decide whether they wish to participate or not.

Sampling of suspected active cutaneous leishmaniasis (ACL) lesions (ulcers)

Each study participant was asked to disclose the occurrence of any skin ulcer(s) on their body. Interviewers also examined the exposed parts of participants body such as legs, arms, neck, and face to identify any suspected active CL (ACL) lesion(s). The location, size, and duration of each suspected ACL was documented. For each suspected ACL lesion, a non-invasive diagnostic sampling technique using sequential tape strips with a diameter of 22 mm (D-Squame, CuDerm Corporation, Texas, USA) was used to obtain samples for subsequent DNA isolation [13].

For the non-invasive skin sampling, one tape disc was placed on each suspected skin lesion after which even pressure was applied to the disc on the lesion using a plunger which was gently held on the disc and pressed for approximately 20 seconds. The tape disc was then detached and transferred into a sterile 1.5ml Eppendorf vial and stored at 4°C for transportation to the laboratory for further analysis (Fig 1). Participants received standard wound care after sample collection.

DNA isolation from tape strip disc and PCR amplification of *Leishmania* species

DNA extraction was performed using SpeedTools Tissue DNA Extraction Kit (Biotoools, Inc).

A nested polymerase chain reaction (Ln-PCR) approach was used to amplify DNA of *Leishmania* species from the human skin lesions following an adaptation of the protocol by Cruz et al., 2002 [14], with the target being the small subunit ribosomal ribonucleic acid (SSU rRNA) gene. Positive control used was *Leishmania infantum* (JPC strain) with distilled water as negative control.

Data management

Data was managed using Microsoft Access software version 2013 and analyzed using STATA software version 14. Association between nominal variables was assessed using Pearson's chi square test of association and Fishers exact test. All statistical tests were performed at a 95% confidence level.

Results

Of 600 households (200 in each study community) invited to participate in this study, a total of 587 households comprising 189 (32.2%), 200 (34.1%), and 198 (33.7%) from Ashiabre, Keri and Sibi Hilltop respectively, were included in this study. The study households had a total of 3718 members out of which 3,440 (92.5%) consisting of 1,194, 941, 1305 from Ashiabre, Keri, and Sibi Hilltop respectively were enrolled in the study.



Fig 1. Non-invasive sampling of skin lesions.

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The average household size was 6.3 with a range of 1 to 18 household members. Ashiabre and Sibi Hilltop had an average household size of 7 while Keri had an average household size of 5.

Out of 3440 persons physically examined for ulcers, a total of 595 skin ulcers were observed on 426 (12.4%) (Table 1). Of the 426 persons, 314 (73.7%) were within the age group 5–15 years while those under five constituted 13.6%. The number of skin ulcers observed on the participants ranged from 1 to 7 with those having one ulcer (47.1%) and two ulcers (27.6%) being the majority. Although skin ulcers were observed on various parts of the participants' body, majority occurred on the lower legs (71.3%) and feet (17.1%). In Ashiabre, Keri, and Sibi Hilltop, 65.2%, 70.1%, and 74.3% of persons with skin ulcers had the ulcer on their lower legs respectively (Table 1).

PCR test of the 595 ulcer samples indicated that 150 (25.2%) of them were *Leishmania* positive. In the study communities, 14 (20.3%), 62 (22.1%), and 74 (30.2%) of skin ulcers tested from Ashiabre, Keri, and Sibi Hilltop respectively were positive for *Leishmania* (Table 1).

Of the 595 ulcer samples tested, 365 (61.3%) were obtained from males while 90 (60.0%) of the 150 *Leishmania* positive samples were also obtained from males. Also, 437 (73.4%) of the ulcer samples tested as well as 112 (74.7%) of the *Leishmania* positive ulcer samples were obtained from people within the age group 5–15 years (Table 2).

Table 1. Individuals with skin ulcers, ulcers sampled and result of *Leishmania* PCR test.

| Characteristic | Category | Ashiabre | | Keri | | Sibi Hilltop | | Total | | P value |
|-------------------------------------|------------------------------|----------|------|------|------|--------------|------|-------|------|---------|
| | | n | % | n | % | n | % | n | % | |
| Age of individuals with skin ulcers | | | | | | | | | | |
| | <5 years | 12 | 21.4 | 22 | 11.7 | 24 | 13.2 | 58 | 13.6 | 0.141 |
| | 5–15 years | 35 | 62.5 | 145 | 77.1 | 134 | 73.6 | 314 | 73.7 | |
| | 16–45 years | 9 | 16.1 | 19 | 10.1 | 18 | 9.9 | 46 | 10.8 | |
| | >45 years | 0 | 0 | 2 | 1.1 | 6 | 3.3 | 8 | 1.9 | |
| | Total | 56 | 100 | 188 | 100 | 182 | 100 | 426 | 100 | |
| Sex of individuals with skin ulcers | | | | | | | | | | |
| | Male | 36 | 64.3 | 109 | 58 | 110 | 60.4 | 255 | 59.9 | 0.684 |
| | Female | 20 | 35.7 | 79 | 42 | 72 | 39.6 | 171 | 40.1 | |
| | Total | 56 | 100 | 188 | 100 | 182 | 100 | 426 | 100 | |
| Number of Skin ulcers tested | | | | | | | | | | |
| | 1 | 31 | 44.9 | 116 | 41.3 | 133 | 54.3 | 280 | 47.1 | <0.001 |
| | 2 | 28 | 40.6 | 84 | 29.9 | 52 | 21.2 | 164 | 27.6 | |
| | 3 | 8 | 11.6 | 46 | 16.4 | 37 | 15.1 | 91 | 15.3 | |
| | 4 | 0 | 0 | 30 | 10.7 | 11 | 4.5 | 41 | 6.9 | |
| | 5 | 2 | 2.9 | 5 | 1.8 | 0 | 0 | 7 | 1.2 | |
| | 6 | 0 | 0 | 0 | 0 | 5 | 2 | 5 | 0.8 | |
| | 7 | 0 | 0 | 0 | 0 | 7 | 2.9 | 7 | 1.2 | |
| | Total | 69 | 100 | 281 | 100 | 245 | 100 | 595 | 100 | |
| Skin ulcer locations | | | | | | | | | | |
| | Face/Head | 3 | 4.3 | 5 | 1.8 | 6 | 2.4 | 14 | 2.4 | 0.029 |
| | Upper arm | 0 | 0 | 2 | 0.7 | 0 | 0 | 2 | 0.3 | |
| | Lower arm | 1 | 1.4 | 13 | 4.6 | 11 | 4.5 | 25 | 4.2 | |
| | Palm/Back of palm | 0 | 0 | 2 | 0.7 | 3 | 1.2 | 5 | 0.8 | |
| | Chest | 0 | 0 | 1 | 0.4 | 0 | 0 | 1 | 0.2 | |
| | Back (upper part below neck) | 0 | 0 | 0 | 0 | 2 | 0.8 | 2 | 0.3 | |
| | Stomach | 2 | 2.9 | 0 | 0 | 0 | 0 | 2 | 0.3 | |
| | Buttocks | 1 | 1.4 | 0 | 0 | 2 | 0.8 | 3 | 0.5 | |
| | Thighs | 1 | 1.4 | 8 | 2.8 | 6 | 2.4 | 15 | 2.5 | |
| | Lower legs(crus/cnemis) | 45 | 65.2 | 197 | 70.1 | 182 | 74.3 | 424 | 71.3 | |
| | Feet | 16 | 23.2 | 53 | 18.9 | 33 | 13.5 | 102 | 17.1 | |
| | Total | 69 | 100 | 281 | 100 | 245 | 100 | 595 | 100 | |
| <i>Leishmania</i> pcr result | | | | | | | | | | |
| | Negative | 55 | 79.7 | 219 | 77.9 | 171 | 69.8 | 445 | 74.8 | 0.061 |
| | Positive | 14 | 20.3 | 62 | 22.1 | 74 | 30.2 | 150 | 25.2 | |
| | Total | 69 | 100 | 281 | 100 | 245 | 100 | 595 | 100 | |

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The 150 *Leishmania* positive ulcer samples were obtained from 136 study participants of which 123 (90.4%) had single *Leishmania* positive skin ulcer, 12 (8.8%) had two *Leishmania* positive skin ulcers and 1 person had three *Leishmania* positive skin ulcers (Table 3). Majority of individuals with *Leishmania* positive ulcers were within the age group of 5–15 years (73.5%) followed by children under five (14.0%) and persons aged 16–45 years (10.3%). Across the study sites and among males and females respectively, majority of persons with *Leishmania* positive skin ulcer(s) were within the age group 5–15 years (Table 3).

The overall prevalence of cutaneous leishmaniasis (*Leishmania* infection observed among those with skin ulcers) was 31.9% (136/426) with prevalence of 23.2% (13/56), 29.8% (56/188), and 36.8% (67/182) observed in Ashiabre, Keri and Sibi Hilltop respectively.

Table 2. Skin ulcers tested for *Leishmania* parasite using PCR by age and sex.

| Sex | Age | Number of skin | <i>Leishmania</i> positive ulcers |
|---------|-------------|----------------|-----------------------------------|
| | | ulcers tested | n (%) |
| Males | < 5 years | 48 | 8 (16.7) |
| | 5–15 years | 276 | 70 (25.4) |
| | 16–45 years | 36 | 10 (27.8) |
| | >45 years | 5 | 2 (40.0) |
| | Subtotal | 365 | 90 (24.7) |
| Females | < 5 years | 45 | 13 (28.9) |
| | 5–15 years | 161 | 42 (26.1) |
| | 16–45 years | 19 | 4 (21.1) |
| | >45 years | 5 | 1 (20.0) |
| | Subtotal | 230 | 60 (26.1) |
| Total | < 5 years | 93 | 21 (22.6) |
| | 5–15 years | 437 | 112 (25.6) |
| | 16–45 years | 55 | 14 (25.5) |
| | >45 years | 10 | 3 (30.0) |
| | Total | 595 | 150 (25.2) |

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Table 3. Distribution of individuals with *Leishmania* positive skin ulcers by age, sex, and community of residence.

| Characteristic | Category | Ashiabre | | Keri | | Sibi Hilltop | | Total | | Total |
|---|-------------|-----------|------------|-----------|-----------|--------------|-----------|-----------|-----------|------------|
| | | Male (%) | Female (%) | Male | Female | Male | Female | Male | Female | |
| Individuals with one <i>Leishmania</i> positive skin ulcer | | | | | | | | | | |
| | <5 years | 1 (25.0) | 1 (12.5) | 3 (9.4) | 5 (27.8) | 4 (9.8) | 3 (15.0) | 8 (10.4) | 9 (19.6) | 17 (13.8) |
| | 5–15 years | 3 (75.0) | 6 (75.0) | 23 (71.9) | 11 (61.1) | 31 (75.6) | 15 (75.0) | 57 (74.0) | 32 (69.6) | 89 (72.4) |
| | 16–45 years | 0 | 1 (12.5) | 6 (18.8) | 1 (5.6) | 4 (9.8) | 2 (10.0) | 10 (13.0) | 4 (8.7) | 14 (11.4) |
| | >45 years | 0 | 0 | 0 (0) | 1 (5.6) | 2 (4.9) | 0 | 2 (2.6) | 1 (2.2) | 3 (2.4) |
| | Sub total | 4 (100) | 8 (100) | 32 (100) | 18 (100) | 41 (100) | 20 (100) | 77 (100) | 46 (100) | 123 (100) |
| Individuals with two <i>Leishmania</i> positive skin ulcers | | | | | | | | | | |
| | <5 years | 0 | 0 | 0 | 1 (33.3) | 0 | 1 (25.0) | 0 | 2 (28.6) | 2 (16.7) |
| | 5–15 years | 1 (100) | 0 | 3 (100) | 2 (66.7) | 1 (100) | 3 (75.0) | 5 (100) | 5 (71.4) | 10 (83.3) |
| | 16–45 years | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| | >45 years | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| | Sub total | 1 (100) | 0 | 3 (100) | 3 (100) | 1 (100) | 4 (100) | 5 (100) | 7 (100) | 12 (100) |
| Individuals with three <i>Leishmania</i> positive skin ulcers | | | | | | | | | | |
| | <5 years | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| | 5–15 years | 0 | 0 | 0 | 0 | 1 (100.0) | 0 | 1 (100.0) | 0 | 1 (100.0) |
| | 16–45 years | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| | >45 years | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| | Sub total | 0 | 0 | 0 | 0 | 1 (100) | 0 | 1 (100) | 0 | 1 (100) |
| Individuals with <i>Leishmania</i> positive skin ulcer(s) | | | | | | | | | | |
| | <5 years | 1 (20.0) | 1 (12.5) | 3 (8.6) | 6 (28.6) | 4 (9.3) | 4 (16.7) | 8 (9.6) | 11 (20.8) | 19 (14.0) |
| | 5–15 years | 4 (80.0) | 6 (75.0) | 26 (74.3) | 13 (61.9) | 33 (76.7) | 18 (75.0) | 63 (75.9) | 37 (69.8) | 100 (73.5) |
| | 16–45 years | 0 | 1 (12.5) | 6 (17.1) | 1 (4.8) | 4 (9.3) | 2 (8.3) | 10 (12.0) | 4 (7.5) | 14 (10.3) |
| | >45 years | 0 | 0 | 0 (0) | 1 (4.8) | 2 (4.7) | 0 | 2 (2.4) | 1 (1.9) | 3 (2.2) |
| | Sub total | 5 (100.0) | 8 (100.0) | 35 (100) | 21 (100) | 43 (100) | 24 (100) | 83 (100) | 53 (100) | 136 (100) |

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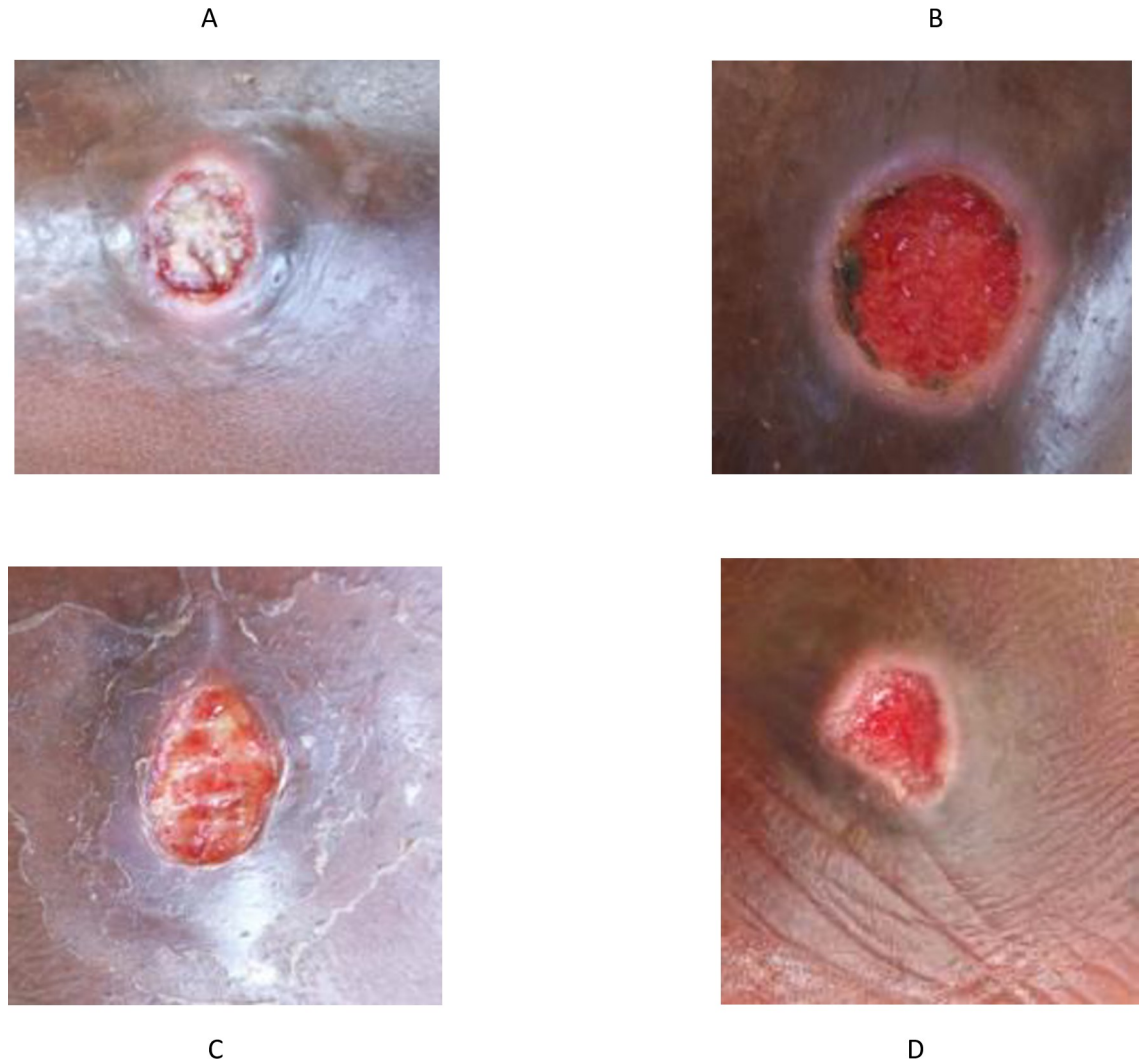


Fig 2. Examples of skin ulcers which tested positive for *Leishmania* parasite. A. Location: Left lower leg; dimension:10.1mm by 5.9mm. B. Location: Left lower arm; dimension:17.0mm by 15.1mm. C. Location: Left lower arm; dimension:17.6mm by 11.0mm. D. Location: Left lower leg; dimension:14.4mm by 5.2mm.

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The average size of the skin ulcers observed was 10.2mm by 10.3mm with 573 (96.3%) of them reported to have started in the year 2018. Among the ulcers which started in the year 2018, 17 (3.0%) started between January to July 2018 while 13 (2.3%), 70 (12.2%), 346 (60.4%), 127 (22.2%) of them started in August, September, October and November of the year 2018 respectively. Examples of *Leishmania* positive skin ulcers observed is captured as Fig 2.

Of the 426 individuals with skin ulcers, 419 (98.4%) indicated that they applied some form of treatment. Majority of them (67.5%) used herbs while 35.3%, and 14.2% of them used hot stone and hot water respectively as treatment of their skin ulcers (Table 4).

Discussion

Cutaneous leishmaniasis among study participants

The control of CL requires an understanding of the disease epidemiology [15]. This study confirmed cutaneous leishmaniasis in the study communities by detecting *Leishmania* infection in

Table 4. Summary of ulcer treatment methods reported by study participants.

| Treatment method | Ashiabre | | Keri | | Sibi Hilltop | | Total | |
|------------------|----------|------|------|------|--------------|------|-------|------|
| | No. | % | No. | % | No. | % | No. | % |
| Herbs | 21 | 40.4 | 117 | 62.6 | 145 | 80.6 | 283 | 67.5 |
| Hot stone | 3 | 5.8 | 72 | 38.5 | 73 | 40.6 | 148 | 35.3 |
| Dermacot | 7 | 13.5 | 31 | 16.6 | 3 | 1.7 | 41 | 9.8 |
| Penicillin | 7 | 13.5 | 14 | 7.5 | 8 | 4.4 | 29 | 8.1 |
| Amoxycillin | 5 | 9.6 | 10 | 5.3 | 2 | 1.1 | 17 | 4.7 |
| Hotwater | 5 | 9.6 | 19 | 10.2 | 27 | 15 | 51 | 14.2 |
| Other treatment | 5 | 9.6 | 4 | 2.1 | 6 | 3.3 | 15 | 4.2 |
| Total | 52 | 100 | 187 | 100 | 180 | 100 | 419 | 100 |

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150 (25.2%) out of 595 ulcer biopsies tested by PCR. The overall prevalence of cutaneous leishmaniasis among persons with skin ulcers was 31.9% (136/426) with prevalence of 23.2% (13/56), 29.8% (56/188), and 36.8% (67/182) observed in Ashiabre, Keri and Sibi Hilltop respectively. In Mali, a systematic review reported a prevalence of 40.3% for cutaneous leishmaniasis among suspected CL cases [10].

Majority of the persons with CL in this study (73.5%) were in the age group of 5–15 years, with males in this age group constituting majority of those infected among persons with skin ulcers. A study in Mali which screened study participants with skin lesions for CL using PCR, confirmed *Leishmania* infection in samples from 8 persons who were all under 18 years [16].

A review of literature on CL suggests that although *Leishmania* infection and subsequent leishmaniasis disease generally tends to be influenced by factors associated with the host, the parasite, as well as the disease vectors, the prevalence of CL usually increases with age till about 15 years [17]. It is assumed that the prevalence of CL levels of at about 15 years because persons exposed early on in life to *Leishmania* infection may have acquired some level of immunity to the infection by then [17]. Observation of the highest prevalence of CL in 5–15 years age group in this study suggest a need to prioritize this group in future CL control planning in the study area.

Treatment of persons with cutaneous leishmaniasis

An important aspect of disease control is treatment of affected people. The data on treatment of skin lesions by study participants indicate that majority of them use herbs (67.5%) followed by those who use hot stone (33.5%) and hot water (14.2%) respectively.

In the case of cutaneous leishmaniasis, the first choice of treatment is pentavalent antimonials with its attendant cost and possible adverse effects [18–22]. However, the evidence for what can be described as optimal treatment for CL has been described as patchy and generally weak. There is therefore a need for the development of improved guidelines for management of CL in addition to the conduct of more robust studies to improve the existing body of evidence for treatment of CL [18,23–25].

Furthermore, although efforts are ongoing to develop a vaccine against leishmaniasis, there is currently no vaccine licensed for use against leishmaniasis [26,27]. Given the gaps in the treatment of leishmaniasis and ongoing global efforts to develop vaccines, there is a need to develop measures in the local Ghanaian context, to protect people who are affected by leishmaniasis while research continues to provide data on critical aspects of the disease such as the vectors and reservoirs.

Need for investigation of skin ulcers which were negative for *Leishmania* infection

Given that not all skin ulcers observed in the study communities were infected with *Leishmania* parasites, there is a need for continuous diagnoses of skin ulcers observed in the study communities in order to identify the ulcers infected by *Leishmania* parasite for the appropriate treatment to be applied [28–30].

Some studies have reported occurrence of other skin ulcers such as buruli ulcer, and yaws in Ghana [31–34]. A pilot study aimed at using azithromycin as treatment for yaws in some communities of the West Akim district of Ghana for instance, used sero-positivity based on a point of care dual treponemal and non-treponemal test as the primary outcome in addition to presentation with clinically active yaws like lesions (as secondary outcome) to select yaws cases [35].

As a result, future studies aimed at screening a larger sample of persons in the study area for yaws and other skin ulcer causing diseases such as buruli ulcer, incorporating more sophisticated laboratory diagnostic approaches may help to better characterize the causes of skin ulcers in the study area.

Conclusions

Out of 426 individuals observed with various numbers of skin ulcers in the study communities, 136 (31.9%) individuals had various numbers of confirmed *Leishmania* positive skin ulcers. The observation of skin ulcers which tested negative to *Leishmania* infection suggests a need to test for additional causes of skin ulcers such as *Treponema pallidum pertenue* and *Mycobacterium ulcerans* in the study area.

Limitations of the study

Molecular characterization of the ulcer samples for agents of other skin ulcer causing diseases reported in Ghana such as yaws, and buruli ulcer would have enriched the data.

Inclusion of a household in the study depended on the consent of the household head. This may have led to the exclusion of a few households, given that 587 households were included out of 600 households invited.

Supporting information

S1 STROBE checklist. Checklist according to The Strengthening the Reporting of Observational studies in Epidemiology (STROBE) guidelines.

(DOCX)

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References

1. Hay R, Asiedu K. Skin-Related Neglected Tropical Diseases (Skin NTDs)—A New Challenge. *Trop Med Infect Dis*. 2018; 4(1):4. <https://doi.org/10.3390/tropicalmed4010004> PMID: 30585179
2. Gabriel Á, Valério-Bolas A, Palma-Marques J, Mourata-Gonçalves P, Ruas P, Dias-Guerreiro T, et al. Cutaneous Leishmaniasis: The Complexity of Host's Effective Immune Response against a Polymorphic Parasitic Disease. *J Immunol Res* [Internet]. 2019; 2019:1–16. Available from: <https://www.hindawi.com/journals/jir/2019/2603730/> <https://doi.org/10.1155/2019/2603730> PMID: 31871953
3. Showler AJ, Boggild AK. Cutaneous Leishmaniasis in Travellers: a Focus on Epidemiology and Treatment in 2015. *Curr Infect Dis Rep* [Internet]. 2015; 17(7):37. Available from: <https://doi.org/10.1007/s11908-015-0489-2> PMID: 26031962
4. Kweku MA, Odoom S, Pupilampu N, Desewu K, Nuako GK, Gyan B, et al. An outbreak of suspected cutaneous leishmaniasis in Ghana: lessons learnt and preparation for future outbreaks. *Glob Health Action*. 2011; 4:1–9. <https://doi.org/10.3402/gha.v4i0.5527> PMID: 21765823
5. Fryauff DJ, Hanafi HA, Klena JD, Hoel DF, Appawu M, Rogers W, et al. Short report: ITS-1 DNA sequence confirmation of *Leishmania major* as a cause of cutaneous leishmaniasis from an outbreak focus in the Ho District, southeastern Ghana. *Am J Trop Med Hyg*. 2006; 75(3):502–4. PMID: 16968929
6. Villinski JT, Klena JD, Abbassy M, Hoel DF, Pupilampu N, Mechta S, et al. Evidence for a new species of *Leishmania* associated with a focal disease outbreak in Ghana. *Diagn Microbiol Infect Dis*. 2008; 60(3):323–7. <https://doi.org/10.1016/j.diagmicrobio.2007.09.013> PMID: 18031968
7. Kwakye-Nuako G, Mosore MT, Duplessis C, Bates MD, Pupilampu N, Mensah-Attipoe I, et al. First isolation of a new species of *Leishmania* responsible for human cutaneous leishmaniasis in Ghana and classification in the *Leishmania enriettii* complex. *Int J Parasitol* [Internet]. 2015; 45(11):679–84. Available from: <https://doi.org/10.1016/j.ijpara.2015.05.001> PMID: 26099650
8. Ghana Statistical Service (GSS). 2010 POPULATION & HOUSING CENSUS; DISTRICT ANALYTICAL REPORT: Nkwanta south district. 2014;

9. Ghana Statistical Service. 2010 POPULATION & HOUSING CENSUS; DISTRICT ANALYTICAL REPORT: Nkwanta north district. 2014;
10. Kone AK, Sa D, Thera MA, Kayentao K, Djimde A, Delaunay P, et al. Epidemiology of the outbreak, vectors and reservoirs of cutaneous leishmaniasis in Mali: A systematic review and meta-analysis. *Asian Pac J Trop Med*. 2016; 9(10):985–90. <https://doi.org/10.1016/j.apjtm.2016.07.025> PMID: 27794393
11. Ghana Statistical Service. Ghana Demographic and Health Survey. *Stud Fam Plann*. 2014; 21(1):1–5.
12. Demographic and Health Surveys Methodology. Sampling and Household Listing Manual. Demographic and Health Surveys Methodology. 2012.
13. Taslimi Y, Sadeghipour P, Habibzadeh S, Mashayekhi V, Mortazavi H, Müller I, et al. A novel non-invasive diagnostic sampling technique for cutaneous leishmaniasis. 2017;1–12.
14. Cruz I, Cañavate C, Rubio JM, Morales MA, Chicharro C, Laguna F, et al. A nested polymerase chain reaction (Ln-PCR) for diagnosing and monitoring *Leishmania infantum* infection in patients co-infected with human immunodeficiency virus. *Trans R Soc Trop Med Hyg*. 2002; 96:S185–9. [https://doi.org/10.1016/s0035-9203\(02\)90074-x](https://doi.org/10.1016/s0035-9203(02)90074-x) PMID: 12055836
15. González U, Pinart M, Sinclair D, Firooz A, Enk C, Id V, et al. Vector and reservoir control for preventing leishmaniasis (Review). *Cochrane Database Syst Rev*. 2015;(8). <https://doi.org/10.1002/14651858.CD008736.pub2> PMID: 26246011
16. Traore B, Oliveira F, Faye O, Dicko A, Coulibaly A, Sissoko IM, et al. Prevalence of Cutaneous Leishmaniasis in Districts of High and Low Endemicity in Mali. *PLoS Negl Trop Dis*. 2016; 10(11):1–12. <https://doi.org/10.1371/journal.pntd.0005141> PMID: 27898671
17. Reithinger Richard; Dujardin Jean-Claude; Louzir Hechmi; Pirmez Claude; Alexander Bruce; Brooker S. Cutaneous leishmaniasis. *Lancet*. 2007; 146(3):581–96.
18. Berbert TRN, Mello TFP De, Wolf Nassif P, Mota CA, Silveira AV, Duarte GC, et al. Pentavalent antimonials combined with other therapeutic alternatives for the treatment of cutaneous and mucocutaneous leishmaniasis: A systematic review. *Dermatol Res Pract*. 2018;2018. <https://doi.org/10.1155/2018/9014726> PMID: 30675152
19. Haldar AK, Sen P, Roy S. Use of Antimony in the Treatment of Leishmaniasis: Current Status and Future Directions. *Mol Biol Int*. 2011; 2011:1–23. <https://doi.org/10.4061/2011/571242> PMID: 22091408
20. Hodiament CJ, Kager PA, Bart A, de Vries HJC, van Thiel PPAM, Leenstra T, et al. Species-Directed Therapy for Leishmaniasis in Returning Travellers: A Comprehensive Guide. *PLoS Negl Trop Dis*. 2014; 8(5). <https://doi.org/10.1371/journal.pntd.0002832> PMID: 24787001
21. Aronson NE, Wortmann GW, Byrne WR, Howard RS, Bernstein WB, Marovich MA, et al. A randomized controlled trial of local heat therapy versus intravenous sodium stibogluconate for the treatment of cutaneous *Leishmania major* infection. *PLoS Negl Trop Dis*. 2010; 4(3). <https://doi.org/10.1371/journal.pntd.0000628> PMID: 20231896
22. Asilian A, Sadeghinia A, Faghihi G, Momeni A. Comparative study of the efficacy of combined cryotherapy and intralesional meglumine antimoniate (GlucantimeR) vs. cryotherapy and intralesional meglumine antimoniate (GlucantimeR) alone for the treatment of cutaneous leishmaniasis. *Int J Dermatol [Internet]*. 2004; 43(4):281–3. Available from: <https://doi.org/10.1111/j.1365-4632.2004.02002.x> PMID: 15090013
23. Palumbo E. Current treatment for cutaneous leishmaniasis: a review. *Am J Ther*. 2009; 16(2):178–82. <https://doi.org/10.1097/MJT.0b013e3181822e90> PMID: 19300044
24. Oliario P, Vaillant M, Arana B, Grogl M, Modabber F, Magill A, et al. Methodology of Clinical Trials Aimed at Assessing Interventions for Cutaneous Leishmaniasis. *PLoS Negl Trop Dis*. 2013; 7(3). <https://doi.org/10.1371/journal.pntd.0002130> PMID: 23556016
25. Uribe-Restrepo A, Cossio A, Desai MM, Dávalos D, Castro M del M. Interventions to treat cutaneous leishmaniasis in children: A systematic review. *PLoS Negl Trop Dis*. 2018; 12(12):1–16. <https://doi.org/10.1371/journal.pntd.0006986> PMID: 30550538
26. Davian C Whyte, Rachel Zufferey. Cutaneous Leishmaniasis: Update on Vaccine Development. *Hum Parasit Dis*. 2017;9.
27. De Luca PM, Macedo ABB. Cutaneous leishmaniasis vaccination: A matter of quality. *Front Immunol*. 2016; 7(APR):1–8. <https://doi.org/10.3389/fimmu.2016.00151> PMID: 27148270
28. Cruz I, Millet A, Carrillo E, Chenik M, Salotra P, Verma S, et al. An approach for interlaboratory comparison of conventional and real-time PCR assays for diagnosis of human leishmaniasis. *Exp Parasitol [Internet]*. 2013; 134(3):281–9. Available from: <https://doi.org/10.1016/j.exppara.2013.03.026> PMID: 23562705
29. Foulet F, Botterel F, Buffet P, Morizot G, Rivollet D, Deniau M, et al. Detection and identification of *Leishmania* species from clinical specimens by using a real-time PCR assay and sequencing of the

cytochrome b gene. *J Clin Microbiol*. 2007; 45(7):2110–5. <https://doi.org/10.1128/JCM.02555-06> PMID: 17475750

30. Goto H, Lindoso JAL. Current diagnosis and treatment of cutaneous and mucocutaneous leishmaniasis. *Expert Rev Anti Infect Ther* [Internet]. 2010; 8(4):419–33. Available from: <http://www.tandfonline.com/doi/full/10.1586/eri.10.19> PMID: 20377337
31. Marks M, Mitjà O, Bottomley C, Kwakye C, Houinei W, Bauri M, et al. Comparative efficacy of low-dose versus standard-dose azithromycin for patients with yaws: a randomised non-inferiority trial in Ghana and Papua New Guinea. *Lancet Glob Heal*. 2018; 6(4):e401–10. [https://doi.org/10.1016/S2214-109X\(18\)30023-8](https://doi.org/10.1016/S2214-109X(18)30023-8) PMID: 29456191
32. Agana-Nsiire P, Kaitoo E, Agongo EEA, Bonsu G, Kyei-Faried S, Amponsa-Achiano K, et al. Yaws Prevalence, Lessons from the Field and the Way Forward towards Yaws Eradication in Ghana. *Int Sch Res Not*. 2014; 2014:1–7. <https://doi.org/10.1155/2014/910937> PMID: 27437507
33. Kenu E, Nyarko KM, Seefeld L, Ganu V, Käser M, Lartey M, et al. Risk Factors for Buruli Ulcer in Ghana—A Case Control Study in the Suhum-Kraboia-Coaltar and Akuapem South Districts of the Eastern Region. *PLoS Negl Trop Dis*. 2014; 8(11):1–8. <https://doi.org/10.1371/journal.pntd.0003279> PMID: 25411974
34. Yeboah-Manu D, Aboagye SY, Asare P, Asante-Poku A, Ampah K, Danso E, et al. Laboratory confirmation of Buruli ulcer cases in Ghana, 2008–2016. *PLoS Negl Trop Dis*. 2018; 12(6):2008–16. <https://doi.org/10.1371/journal.pntd.0006560> PMID: 29870529
35. Abdulai AA, Agana-Nsiire P, Biney F, Kwakye-Maclean C, Kyei-Faried S, Amponsa-Achiano K, et al. Community-based mass treatment with azithromycin for the elimination of yaws in Ghana—Results of a pilot study. *PLoS Negl Trop Dis*. 2018; 12(3):1–16. <https://doi.org/10.1371/journal.pntd.0006303> PMID: 29566044