

Dermatology Reports

https://www.pagepress.org/journals/index.php/dr/index

eISSN 2036-7406







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The final version of the manuscript will then appear on a regular issue of the journal. E-publishing of this PDF file has been approved by the authors.

Please cite this article as: Gashaw B, Yizengaw E, Nibret E, et al. Epidemiological and clinical profiles of cutaneous leishmaniasis cases in Amhara National Regional State, Northwest Ethiopia: a multicenter retrospective study.

Dermatol Rep 2024 [Epub Ahead of Print] doi: 10.4081/dr.2024.10089

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Submitted 12/07/24 - Accepted 31/08/24

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Epidemiological and clinical profiles of cutaneous leishmaniasis cases in Amhara

National Regional State, Northwest Ethiopia: a multicenter retrospective study

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Acknowledgment

We would like to thank all the research and administrative staffs of Amhara Public Health

Institute for arranging transportation and support letter for study hospitals. We are also

thankful to Amhara Regional Health Bureau and all staff of study hospitals for their

unreserved support in all the steps of this study.

Key words: cutaneous leishmaniasis, Amhara Regional State, Ethiopia.

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Contributions: BG: conceptualization, data collection, analysis, visualization, writing,

review and editing; EY: data collection, analysis, visualization, review, and editing; EN:

writing, review, visualization, analysis and editing; AW: data collection, visualization,

analysis; AA: analysis, review, and editing.

Conflict of interest: The authors declared that no conflict of interest.

Ethics approval and consent to participate: This study was granted by Amhara Public

Health Institute (APHI) (NoH/R/T/T/D/07/83). And support letter was written from APHI to

all study hospitals (03/1691). The study participants were kept anonymous to maintain their

medical confidentiality rights. Personal identifier variables like names were not included in

the data collection checklist. All information was anonymized with an anonymized link to the

individual patient.

Availability of data and materials: All the data reported in this manuscript can be available

by requesting the corresponding author.

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Abstract

Cutaneous leishmaniasis (CL) is caused by *Leishmania* parasites. Ethiopia is one of the top ten countries with high CL load. Amhara National Regional State (ANRS) is one of the CL hotspot areas in Ethiopia. This study determined the epidemiology and clinical profiles of CL in ANRS. This study was conducted from April to October 2023 in eight Leishmaniasis Treatment Centres (LTCs). A data review was done from patients presenting to these centres between June 2018 and July 2023. Chi-square test and logistic regression were performed using SPSS-23. A total of 1729 CL patients were recorded. The overall burden of CL per 10,000 outpatients was 900. Most of the patients (71.1%) presented with localised cutaneous leishmaniasis (LCL). The patients were from 112 districts. About 12% of the patients lived with the disease for over a year without treatment. Multiple-time comer patients accounted for 13.2% of the patients. Cutaneous leishmaniasis is still a major public health problem in ANRS. One-third of CL patients presented with the MCL clinical form. There was longer delay among CL patients for seeking diagnosis and treatment. Large scale community based study and traditional and modern treatment centers focused studies should be included to estimate the actual number of CL in the region. Follow-up and molecular studies are important to better understand the clinical features of the disease. Moreover, awareness of the community about the CL prevention and control help the patients to get early diagnosis and treatment.

Introduction

Cutaneous leishmaniasis (CL) is one of the major public health problems worldwide, especially in low- and middle-income countries.¹ It is more common than the fatal visceral leishmaniasis (VL) form with more than 1 billion people live in CL endemic areas worldwide and annually, an estimated of more than one million new cases of CL occur.²⁻⁴ The CL is

caused by *Leishmania* parasites and spread by the bite of a Phlebotomine sand fly. Leishmaniasis (Both CL and VL) is the second cause of deaths attributed to vector-borne parasitic disease next to malaria.⁵ It mainly presented in three different clinical forms:^{6, 7} localised CL (LCL), mucocutaneous leishmaniasis (MCL), and diffused CL (DCL). LCL is usually self-healing over time while the latter two are non-healing forms causing deformity of affected areas and associated with high social stigma.^{8, 9}

Ethiopia is among east African countries with high burden of CL.¹⁰⁻¹² There are 30 million people at risk.¹³⁻¹⁵ The CL has been known in the country since 1913 in Kutaber (10),¹⁰ one of CL endemic sites. However, CL is the one of the neglected tropical diseases (NTDs) in the country. Our recent work in Lay Gayint district, Northwest Ethiopia, showed that significant proportions of the CL cases have low knowledge about the disease and use traditional drugs (Yizengaw 2024, unpublished data). Low awareness and knowledge of the community regarding CL has also been shown in Ethiopia ^{4, 10, 11, 16, 17}.

Nowadays, in ANRS it is spreading and covering new areas that were not endemic to CL before.¹² Different environmental factors such as change in temperature, expansion of irrigation, deforestation, climate changes and development of drug resistance ^{12, 17} could be the factors for the spreading of CL. Local war, poor socioeconomic status, and poor access for health facilities that can diagnose and treat CL are also major contributors to the spread of this disease.¹²

According to the Amhara Regional Health Bureau (ARHB) 2018 report, the common CL endemic sites include Gayint, Addis Zemen, Finote Selam, Ankesha, Boru Meda, Sekota, and Kutaber. However, there is a scarcity of research dealing with the overall burden and clinical characteristics of CL in ANRS. This study aimed to determine the epidemiology and clinical profiles of CL in eight leishmaniasis treatment centres in ANRS.

Materials and Methods

Study design and period

A multicenter retrospective study was done from April to October 2023.

Study area

This study was conducted at Leishmaniasis Treatment Centres (LTCs) of: Addis Zemen Hospital, located in South Gondar Zone; Addis Alem Hospital, a primary hospital in Bahir Dar City; Fenote Selam Hospital, in West Gojjam Zone; Agewgemjabet Hospital, in Awi Zone; the University of Gondar Hospital, in Central Gondar Zone; Boru Meda Hospital, in South Wollo Zone; Nefas Mewcha Hospital, in South Gondar Zone; and Tefera Hailu Hospital, in Wag Hemra Zone (Figure 1).

The LTCs are established at different times. Thus, the data incorporated in this study from each centre included different periods. Data were collected from CL patients registered in Addis Zemen Hospital LTC from June 2018 to August 2022, Addis Alem Hospital LTC from June 2018 to August 2022, Fenote Selam Hospital LTC from July 2018 to June 2022, Agewgemjabet Hospital LTC from August 2021 to July 2022, University of Gondar Hospital LTC from January 2022 to July 2023, Boru Meda Hospital LTC from October 2021 to February 2023, Nefas Mewcha Hospital LTC from January to March 2023, and Tefera Hailu Hospital LTC from August 2018 to April 2022.

The ANRS has a population of 22.5 million, according to the ANRS Plan Commission. The majority (79.9%) of the population live in rural areas, while 20.1% are urban dwellers. The health care system in the region consists of 100 government hospitals and 900 health centers. According to the ANRS Plan Commission 2019/20 report (unpublished), the altitude distribution of the region was as follows: between 500 and 1500 (29.3%), 1500 and 2300 (43.5%), 2300 and 3200 (24.2%), and >3200 (3.1%) metres above sea level.

Data collection

We retrieved the records of CL patients from each of LTC. We have retrieved the records of 492 CL patients from Boru Meda Hospital, 354 from Addis Zemen Hospital, 347 from University of Gondar Hospital, 147 from Nefas Mewcha Hospital, 126 from Addis Alem Hospital, 116 from Tefera Hailu Hospital, 94 from Fenote Selam Hospital and 37 from Agewgemjabet Hospital. The data was retrieved using a structured data collection questionnaire developed for this purpose. The data collection questionnaire was comprised of: Age, Sex, CL type, treatment history, lesion size, duration of illness, microscopy result, and parasite load. Moreover, the total outpatient cases were retrieved from the triage registration logbook in each hospital. Inconsistent or incomplete patient data were excluded from the study.

Patients were Diagnosed and treated based on the Guidelines for Diagnosis, Treatment, and Prevention of Leishmaniasis in Ethiopia. Patient diagnosis was made both by parasitological parasite detection and clinical decision. Parasitological diagnosis was done by parasite identification and quantification of the load from a skin slit smear under a microscope. A clinical decision was made by characterising the skin lesion.

Inclusion and exclusion criteria

In the context of this study, all CL confirmed patient records were included in each LTC since the establishment of the respective centre. However, patient records with data previously published were excluded from the study.

Data Quality Management

Patient records with incomplete or ambiguous information in the registration logbook were meticulously cross-referenced with corresponding patient charts for further clarification. Health professionals working in the LTC of each hospital were consulted for unreadable information in the registration logbook. The completeness and consistency of the data was checked before analysis of the data.

Diagnosis and treatment of cutaneous leishmaniasis

The Diagnosis and treatment of CL in the study hospitals were done based on the Guidelines for Diagnosis, Treatment, and Prevention of Leishmaniasis in Ethiopia. Diagnosis of CL was by parasitological detection of the parasite from a skin scraping smear and by clinical decisions. A skin scraping was taken from the active lesion and examined under a light microscope to detect and quantify the parasitic load of amastigotes. A clinical decision was made by characterising the skin lesion.

Parasite load was determined for microscopy positive CL cases. The parasite load was reported as follow: Parasite load 6+=100 parasites per field, 5+=10-100 parasites per field, 4+=1-10 parasites per field, 3+=1-10 parasites per 10 field, 2+=1-10 parasites per 100 field, 1+=1-10 parasites per 1000 field.

Confirmed CL patients were treated commonly with intramuscular injection of Sodium Stibogluconate (SSG) 20 mg/kg/day for 28 days.

Data analysis

The data were entered and analysed using Statistical Package for Social Science 23 (SPSS-23). The prevalence of CL was expressed per 10, 000 total outpatient cases. The total outpatient cases were also retrieved from each study hospital during the same period. The percentage of different clinical forms of CL was calculated from the total CL cases. Both univariate and multivariate logistic regressions were used to measure the strength of the association. Variables with p<0.25 in the univariate analysis were entered to multivariate logistic regressions analysis. A chi-square test was employed to assess the relationship between dependent and independent variables. Statistical significance was declared at p < 0.05.

Results

Demographic variables

Of the total 2,019,217 outpatients, 1729 CL cases were diagnosed and treated in eight LTCs in the region. Of 1729 CL cases, 585 (33.8%) were females. The mean age of CL patients was 25.88 ± 17.5 years. The highest numbers of CL patients presenting at the LTCs were in the 15–29 (44.7%) and <15 (25.6%) age groups. The age was not recorded for one patient.

Prevalence of cutaneous leishmaniasis

The highest number of reported CL cases was from Boru Meda Hospital (492 cases). The pooled average prevalence of CL was 8.6 per 10,000 total outpatients registered in each hospital during the same period (Table 1).

Clinical characteristics of cutaneous leishmaniasis patients

The proportions of LCL, MCL, and DCL clinical forms were 71.1%, 28.1%, and 0.8%, respectively. Significant difference was seen in distribution of CL clinical forms (χ^2 :1296; p < 0.001) (Table 2).

Clinical data was not recorded for seven patients. Majority of CL patients were diagnosed and treated late, after six month of having the disease (Table 2).

The longest period of time without treatment for CL patients was 158 months, with a mean duration of 11 months (SD: 14.1 months). The median duration of illness was 7.5 months.

A total of 228 (13.2%) CL patients had previous treatment history, and were considered as multiple-time comers. Out of these repeat comers, 155 (68%), 65 (28.5%), and 8 (3.5%) had LCL, MCL, and DCL clinical forms, respectively (Table 2).

Skin scraping microscopy and duration of illness

From the registration logbook, we found that 954 (83.9%) CL cases were microscopically positive for leishmania parasites. The slide positivity rate was higher in patients with a short duration of illness compared to those with a long duration of illness.

Regarding the parasite density, only 208 slides were graded for parasitic load. Among these, the majorities (54; 26%) were in grade two. The remaining 592 CL patients and microscopically negative CL patients (183) were diagnosed and treated clinically.

Lesion size of cutaneous leishmaniasis and associated factors

Duration of illness was found to be an independent explanatory risk factor for bigger lesion size. A multivariate logistic regression showed that duration of illness <6 months and 6-12 months showed a strong association (p = 0.01 and 0.03, respectively) with lesion size (Table 3).

Discussion

This study describes the overall prevalence and clinical characteristics of CL in ANRS, northwest Ethiopia. Unlike previous studies, we enrolled a large dataset collected from all LTCs in the region. It showed that the overall prevalence of CL was 8.6 per 10,000 outpatients. This, however, does not represent the true burden of the disease, as there is under-reporting of patients and a limited number of diagnosis and treatment centres for CL. Significant proportion of CL patients used traditional treatment in Ethiopia. 12, 18 A similar study conducted at ALERT Hospital in Addis Ababa, reported that 33% of outpatients were diagnosed and confirmed to have CL. 19 This higher prevalence might be because ALERT Hospital is well-known for treating skin diseases and serves as a referral centre for CL from all over the country.

The prevalence of CL per 10,000 total outpatient cases was higher in Boru Meda Hospital (49.0) followed by Nefas Mewcha Hospital (27.5). This might be because the awareness of the CL cases and the community in the catchment area of Boru Meda Hospital is better as it is the oldest CL treatment center in the region. Our previous work in Nefas Mewcha Hospital showed that the treatment seeking behaviour of CL cases was improved by awareness training we did.¹¹ The lowest prevalence of CL cases per 10,000 total outpatient cases (2.3)

was in Fenote Selam Hospital. This is due to the fact that the number of CL cases decreased because of the establishment of a new LTC, Agewgemjabet Hospital LTC, which is very close. Even though, the LTC in the University of Gondar Hospital is among the oldest LTC in the country, it is basically established for VL cases.

Our study showed that CL was predominantly affecting children and the youngest population. This is similar with the trend in Ethiopia where CL is more common in children, with the highest prevalence occurring between 10 and 15 years of age.²⁰⁻²² This might be because children have immature immune response as compared to adults.²³ An alternative explanation may be the development of immunity to CL with previous exposure in adults. Prospective data showed this to occur with *L. infantum* in Iran,²⁴ suggesting that protective anti-leishmanial vaccines are achievable.²⁵ It also showed that more males (66.2%) were diagnosed and treated for CL. This might be attributed to gender-based differences in activities and roles, which could potentially lead to increased exposure to sandfly bites. A similar finding has been reported from a retrospective study at the University of Gondar Hospital Leishmaniasis Research and Treatment Centre, Northwest Ethiopia,²¹ and at Boru Meda Hospital.²⁶ However, in Silte Zone, Southern Ethiopia, the frequency of CL infection between males and females was nearly the same.²⁷ Another study conducted in Tigray, northern Ethiopia, indicated that the odds of CL occurrence were 2.1 times higher in males compared to females.²⁸

The age group between 15 and 29 years old was the most affected. Similar to other studies in Ethiopia, it is evident that CL affects a population with a wide age range.¹⁰ In our study, it was found out that one year old to 72 years old were infected. However, it was indicated that the most affected age group was between 16 and 45 years old, accounting for 63.4%,²⁸ and 70.2%,²⁹ of the total cases in Ayder referral hospital, northern Ethiopia, and Boru Meda

Hospital, northwest Ethiopia, respectively. The difference might be explained by the population pyramid of the ANRS.

Most CL patients presented with LCL forms. It is the most common form of CL both worldwide and in Ethiopia. 12, 13, 30 We also observed a higher number of CL patients with MCL clinical forms compared to other World Old reports. We had reported similar findings from Lay Gayint district. 11 This finding might not be entirely attributed to *L. aethiopica*. Rather, MCL cases seek diagnosis and treatment more readily as MCL disfigures the mucosal tissues. Another potential reason could be co-infection with the *Leishmania* RNA virus, which induces severe MCL forms. 31 In our study, a few CL patients presented with DCL. It might indicate that DCL is rare in ANRS. Similar findings have been reported in other studies. 11, 26 However, more DCL (22.9%) has been reported from Northern Ethiopia. 28 Furthermore, the exchange of genomic materials between *L. aethiopica* and other *Leishmania* species like *L. tropica* and *L. major* could contribute. 20

The majority (56.1%) of CL patients lived with the skin lesion for a significant period of time (6 to 12 months) without being diagnosed and treated. This finding is supported by a study conducted in.^{29, 32} In our study, the average time that patients lived with the lesions was nearly 11 months. Other previous studies reported 12.9 months of illness.³² There is a significant delay in seeking diagnosis and treatment for CL patients. This might have implications for disease progression and complicating case management. This could be attributed to the lack of awareness and low of knowledge of the community regarding CL.^{4, 10, 11, 17, 22, 33, 34} This will contribute to the continuous transmission of CL.

Microscopic identification of the parasite was higher than previous reports by most studies.^{19,}
²² This might be associated with experience of laboratory technologists working in LTC.

Health professionals and others working in hospitals with LTC are taking continuous training

on CL, including its differential diagnosis. Skin slit smears from patients with MCL and DCL were more likely to test negative compared to LCL cases. About 21.5% of DCL and 21% of MCL patients had negative skin slit smear microscopic results. The number of LCL patients with negative smear results was only 13.9%. One possible explanation for this finding is that the MCL form of the disease primarily affects soft tissues, making it difficult to obtain an adequate sample for microscopic evaluation. It is also difficult to remove blood from the slit in soft tissues. Additionally, the MCL form had a lower parasitic load compared to the LCL and DCL cases. Moreover, DCL patients might have superinfections as there is a delay in diagnosis. It makes sense as the majority (57.1%) of DCL patients were multiple-time comers. Interestingly, our findings also revealed an association between the duration of the disease and slide negativity. As the disease duration increased, the proportion of microscopically negative cases also increased linearly.

Our findings align with a study in ²⁹ that reported a higher prevalence of CL in patients with lesions less than 12 months old compared to lesions older than 12 months. Similar findings were reported in a study conducted in Sri Lanka, where 54.5% of positive cases were found to have lesions for less than 6 months. There was a higher number of multiple-time comer DCL patients compared to those with other clinical forms. This might indicate a relatively lower response of DCL patients to the current treatment. Our finding is in agreement with existing previous findings that indicated DCL cases posing a great challenge to treatment response, in which a potential treatment failure rate of 75% was, recorded. In our study, it was also worth noting that the highest number (32.5 %) of multiple-time comer CL patients was recorded at Boru Meda Hospital (Figure 2). It was not, however, possible to confirm the reason (e.g. re-infection, treatment failure or drug resistance associated with the parasite) for repeated visiting of LTCs by CL cases in the region. A previous hospital-based study in

northcentral Ethiopia also reported that 367 (41.3%) patients were previously treated and came back to the treatment center.¹⁷

Conclusions

This study showed that CL is a major public health problem in Amhara National Regional State. One-third of CL patients presented with the MCL clinical form. A relatively longer delay was observed among CL patients for seeking diagnosis and treatment. This is not only increases the severity and burden of the disease to the population but it also contributes to the ongoing transmission cycle of CL in the region.

Recommendations

Large scale community based and in both traditional and modern treatments center studies should be included to estimate the actual number of CL in the region. Follow-up and molecular studies are important to better understand the clinical features of the disease. Moreover, awareness of the community about the CL prevention and control help the patients to get early diagnosis and treatment.

References

- 1. Ruiz-Postigo JA, Grout L, Jain S. Global leishmaniasis surveillance, 2017–2018, and first report on 5 additional indicators Weekly epidemiological record 2020;25:165-280.
- 2. Singh OP, Sundar S. Immunotherapy and targeted therapies in treatment of visceral leishmaniasis: current status and future prospects. Front Immunol. 2014;5:296.
- 3. Burza S, Croft SL, Boelaert M. Leishmaniasis. Lancet. 2018;392:951-70.
- 4. World Health Organization (WHO). Leishmaniasis, 2022.
- 5. den Boer M, Argaw D, Jannin J, et al. Leishmaniasis impact and treatment access. Clin Microbiol Infect. 2011;17:1471-7.
- 6. Reithinger R, Dujardin JC, Louzir H, et al. Cutaneous leishmaniasis. Lancet Infect Dis. 2007;7:581-96.

- 7. Neitzke-Abreu HC, Venazzi MS, de Lima Scodro RB, et al. Cutaneous leishmaniasis with atypical clinical manifestations: Case report. IDCases. 2014;1:60-2.
- 8. Libardo J. Gómez RvW, Lena van Selm, Alberto Rivera, et al. Stigma, participation restriction and mental distress in patients affected by leprosy, cutaneous leishmaniasis and Chagas disease: a pilot study in two co-endemic regions of eastern Colombia. Transactions of the Royal Society of Tropical Medicine and Hygiene. 2019:1-7.
- 9. Bennis I, Thys S, Filali H, et al. Psychosocial impact of scars due to cutaneous leishmaniasis on high school students in Errachidia province, Morocco. Infect Dis Poverty. 2017;6:46.
- 10. Mengistu G, Laskay T, Gemetchu T, et al. Cutaneous leishmaniasis in south-western Ethiopia: Ocholo revisited. Trans R Soc Trop Med Hyg. 1992;86:149-53.
- 11. Yizengaw E, Nibret E, Yismaw G, et al. Cutaneous leishmaniasis in a newly established treatment centre in the Lay Gayint district, Northwest Ethiopia. Skin health and Disease. 2023:1-8.
- 12. Shita EY, Nibret E, Munshea A, et al. Burden and risk factors of cutaneous leishmaniasis in Ethiopia: a systematic review and meta-analysis. Int J Dermatol. 2022;61:1336-45.
- 13. Alvar J, Canavate C, Gutierrez-Solar B, et al. Leishmania and human immunodeficiency virus coinfection: the first 10 years. Clin Microbiol Rev. 1997;10:298-319.
- 14. Alvar J, Velez ID, Bern C, et al. Leishmaniasis worldwide and global estimates of its incidence. PLoS One. 2012;7:e35671.
- 15. Guideline for Diagnosis, Treatment and Prevention of Leishmaniasis in Ethiopia 2013.
- 16. Gashaw G. Yizengaw E, Sebsibe B, et al. Clinical Manifestations of Cutaneous Leishmaniasis (CL): Does Elevated Blood Sugar Level have Implications for Clinical Management of CL? Ethiopian Journal of Health Development. 2023;37.

- 17. Eshetu B, Mamo H. Cutaneous leishmaniasis in north-central Ethiopia: trend, clinical forms, geographic distribution, and determinants. Trop Med Health. 2020;48:39.
- 18. van Henten S, Bialfew F, Hassen S, et al.Treatment of cutaneous leishmaniasis with sodium stibogluconate and allopurinol in a routine setting in Ethiopia: Clinical and Patient-Reported Outcomes and Operational Challenges. Trop Med Infect Dis. 2023;8.
- 19. Neway B.Y.S, Mebrat B, Ayalew G. Prevalence of cutaneous leishmaniasis in alert center, retrospective analysis, Addis Ababa. Journal of Health Systems and Policies. 2021;3:110-21.
- 20. van Henten S, Adriaensen W, Fikre H, et al. Cutaneous Leishmaniasis Due to Leishmania aethiopica. EClinicalMedicine. 2018;6:69-81.
- 21. Fikre H, Mohammed R, Atinafu S, et al. Clinical features and treatment response of cutaneous leishmaniasis in North-West Ethiopia. Trop Med Int Health. 2017;22:1293-301.
- 22. Yohannes M, Abebe Z, Boelee E. Prevalence and environmental determinants of cutaneous leishmaniasis in rural communities in Tigray, northern Ethiopia. PLoS Negl Trop Dis. 2019;13:e0007722.
- 23. Simon AK, Hollander GA, McMichael A. Evolution of the immune system in humans from infancy to old age. Proc Biol Sci. 2015;282:20143085.
- 24. Davies CR, Gavgani ASM. Age, acquired immunity and the risk of visceral leishmaniasis: a prospective study in Iran. Parasitology. 1999;119:247-57.
- 25. Okwor I, Mou Z, Liu D, et al. Protective immunity and vaccination against cutaneous leishmaniasis. Front Immunol. 2012;3:128.
- 26. Seife AKBT, Zewdu TF, Ayal A, et al. Treatment Patterns and Effectivness of Anti-Leishmaniasis Agents for Patients with Cutaneous Leishmaniasis at Boru Meda Hospital, South Wollo, North East Ethiopia, 2017/18. Journal of Clinical & Experimental Dermatology Research. 2018;9.

- 27. Negera E, Gadisa E, Yamuah L, et al. Outbreak of cutaneous leishmaniasis in Silti woreda, Ethiopia: risk factor assessment and causative agent identification. Trans R Soc Trop Med Hyg. 2008;102:883-90.
- 28. Tilahun WAT, Mulatu G. Magnitude and associated factors of cutaneous leishmaniasis; in Mekelle city, Ayder referral hospital, Tigray, Northern Ethiopia, 2014 Clinical Medicine Research. 2014;3:.
- 29. Bisetegn H, Zeleke AJ, Gadisa E, et al. Clinical, parasitological and molecular profiles of Cutaneous Leishmaniasis and its associated factors among clinically suspected patients attending Borumeda Hospital, North-East Ethiopia. PLoS Negl Trop Dis. 2020;14:e0008507.
- 30. van Henten S, Tesfaye AB, Abdela SG, et al. Miltefosine for the treatment of cutaneous leishmaniasis-A pilot study from Ethiopia. PLoS Negl Trop Dis. 2021;15:e0009460.
- 31. Cantanhede LM, da Silva Junior CF, Ito MM, et al. Further Evidence of an Association between the Presence of Leishmania RNA Virus 1 and the Mucosal Manifestations in Tegumentary Leishmaniasis Patients. PLoS Negl Trop Dis. 2015;9:e0004079.
- 32. Aberra L, Abera A, Belay T, et al. Evaluation of microcapillary culture method for the isolation of Leishmania aethiopica parasites from patients with cutaneous lesions in Ethiopia. Diagn Progn Res. 2019;3:4.
- 33. A Lemma WAF, T Gemetchu, P M Preston, et al. A Studies on leishmaniasis in Ethiopia.
- I. Preliminary investigations into the epidemiology of cutaneous leishmaniasis in the highlands. Ann Trop Med Parasitol. 1969;63:455-72.
- 34. Padovese V, Terranova M, Toma L, et al. Cutaneous and mucocutaneous leishmaniasis in Tigray, northern Ethiopia: clinical aspects and therapeutic concerns. Trans R Soc Trop Med Hyg. 2009;103:707-11.

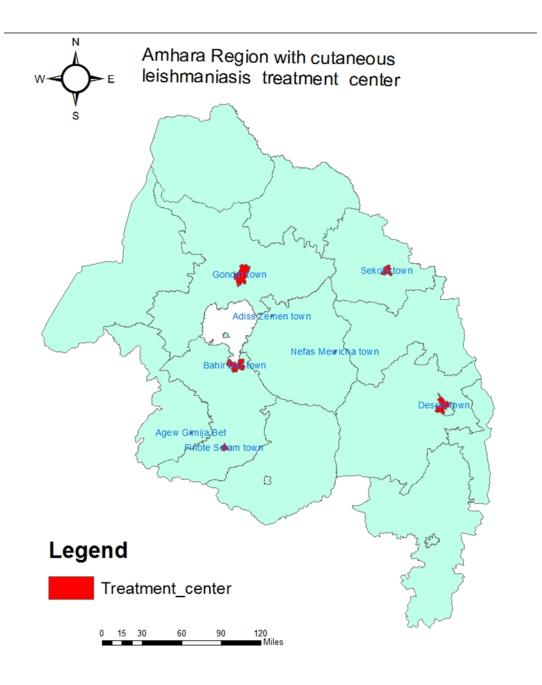


Figure 1. Leishmaniasis Treatment Centres in Amhara Region, September 2023 (Google map).

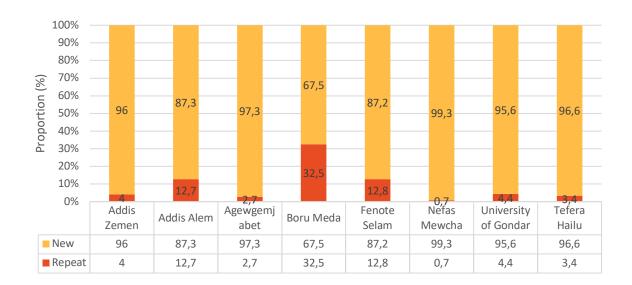


Figure 2. Proportions of both new and repeat (multiple-time comer) CL patients in ANRS, 2023.

Table 1. Prevalence of CL among the total outpatients in ANRS, 2023

Hospitals with LTC	Distance from Total		CL	Prevalence (per	
	Bahir Dar (Km)	outpatients	cases	10,000 outpatients)	
Addis Zemen Hospital	88	235,941	354	15.0	
Addis Alem Hospital	10	510,142	126	2.4	
Agewgemjabet Hospital	131	56,761	37	6.5	
Boru Meda Hospital	490	100,373	492	49.0	
Fenote Selam Hospital	172	403,933	94	2.3	
Nefas Mewcha Hospital	180	53,492	147	27.5	
University of Gondar Hospital	172	417,580	347	8.3	
Tefera Hailu Hospital	436	240,995	116	4.8	
Total		2,019,217	1729	8.6	

Table 2. Clinical characteristics of CL patients in ANRS, 2023

Variable		Frequency	Percent	X ² (p-value)
CL types				
LCL	1224	71.1		
MCL		484	28.1	1296.5(p < 0.001)
DCL	14	0.8		
Treatment History				
New	New		86.8	
Multiple-time comer	LCL	155	68%	932.6(p < 0.001)
(228, 13.2%)	MCL	65	28.55	
	DCL	8	3.5%	
Lesion size (millimeter)				
<u>≥4</u>	570	(60.4)		
<4	374	(39.6)	40.6 (p <0.001)	
Duration of illness (month)				
<6	309	32.1		
6-12		540	56.1	
13-24		54	5.6	674 (p <0.001)
>24	59	6.1		
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DCL, Diffuse Cutaneous Leishmaniasis, LCL, Localised Cutaneous Leishmaniasis, MCL, Mucocutaneous Leishmaniasis

Table 3. Lesion size and associated factors for CL in ANRS, 2023

Variable		Lesion	size						
		(centimeter)		Univariate analysis		Multivariate Analysis			
		≥4	<4	COR	95% CI	P-value	AOR	95%CI	P-value
CL type	LCL	380	311	1.6	0.48-5.48	0.42	NA	NA	NA
	MCL	180	57	0.63	0.18-2.18	0.46	NA	NA	NA
	DCL	8	4	1	1	1	NA	NA	NA
Parasite	1+-3+	89	33	0.58	0.3-1.0	0.09	NA	NA	NA
load	4+-6+	46	29	1	1	1	NA	NA	NA
Sex	Female	154	116		1	1	NA	NA	NA
	Male	416	258	1.21	0.91-1.6	0.18	NA	NA	NA
Treatment	New	526	364	2.9	1.44-5.86	0.01	0.6	0.29-1.4	0.27
history	Multiple	42	10	1	1	1	1	1	1
Duration	<6	126	135	4.5	2.1-9.7	0.01	2.7	0.12-0.63	0.01
of illness	6-12	258	197	3.2	1.5-6.8	0.01	4.1	1.18-0.93	0.03
(month)	13-24	32	7	0.92	0.3-2.7	0.89	1.6	0.47-5.56	0.44
	≥24	39	9	1	1	1	1	1	1
	<u> </u>		l					. 11 DCI	

AOR, Adjusted Odds Ratio, COR, Crude Odds Ratio, NA, Not Applicable, DCL, Diffuse Cutaneous Leishmaniasis, LCL, Localised Cutaneous Leishmaniasis, MCL, Mucocutaneous Leishmaniasis