Species Identification of Cutaneous Leishmaniasis in Quchan, Northeast of Iran

Elham Moghaddas¹, Abdolmajid Fata¹, Ameneh Gholampour¹, Lida Jarahi², Saman Soleimanpour³ and Seyed Aliakbar Shamsian¹,*

¹Department of Parasitology, Faculty of Medicine, University of Medical Sciences, Mashhad, Iran
²Department of Community Medicine, Faculty of Medicine, Mashhad University of Medical Sciences, Mashhad, Iran
³Department of Microbiology and Virology, School of Medicine, Mashhad University of Medical Sciences, Mashhad, Iran

*Corresponding author: Department of Parasitology, Faculty of Medicine, University of Medical Sciences, Mashhad, Iran. Fax: +98-5138002401-3, Email: shamsianaa@mums.ac.ir

Received 2019 February 20; Revised 2019 May 28; Accepted 2019 June 11.

Abstract

Background: Khorasan Province is an endemic region for cutaneous leishmaniasis. The species identification of Leishmania protozoa is useful for the control and prevention of leishmaniasis.

Objectives: The present study is the first to identify Leishmania species by evaluating their risk factors in Quchan, Northern Khorasan, Northeast of Iran.

Methods: A questionnaire and slide smears were collected from 103 individuals suspected of having leishmaniasis. Optimized PCR was performed using specific kDNA primers on all the slides. The data obtained were analyzed in SPSS-20 software.

Results: Among the 103 subjects with skin ulcers suspected of having CL, 77 (74.8%) showed positive results in their direct microscopic smear. Specific Leishmania PCR bands, however, were observed in 86 (83.4%), including 57 subjects with L. tropica and 29 with L. major. The most frequent age range involved was 20 - 30 years and the most common site of the lesions was the hands. From the 57 cases of L. tropica, 43.9% and 56.1% lived in urban and rural districts, respectively. The sensitivity of microscopy for the diagnosis of Leishmania spp. was calculated as 89.5% in this study.

Conclusions: L. tropica is the dominant causative species for cutaneous leishmaniasis in Quchan. This study identified a new rural focus for cutaneous leishmaniasis caused by L. tropica in Quchan suburbs.

Keywords: Leishmania, PCR, Quchan, Species, Iran

1. Background

Cutaneous Leishmaniasis (CL) is among the neglected endemic diseases of America, Africa, India, southwest Asia and the Mediterranean (1). Nowadays, 12 million people in the world suffer from this disease. The annual incidence of leishmaniasis in the world is estimated at 1 to 1.5 million. Iran is one of the ten countries in which cutaneous leishmaniasis has constantly been reported to have a high prevalence (2). About 20,000 cases of leishmaniasis are reported annually in this country, but the actual prevalence of the disease appears to be four to five times higher. The prevalence of infection has been reported as 1.8% to 37.9% in different provinces of Iran (3). Cutaneous leishmaniasis has two forms, including Anthroponotic Cutaneous Leishmaniasis (ACL), which is caused by Leishmania tropica (L. tropica), and Zoonotic Cutaneous Leishmaniasis (ZCL), which is caused by Leishmania major (L. major). L. tropica is transmitted to human beings by the bite of infected female Phlebotomus sergenti. Phlebotomus papatasi, the most prevalent species of the Phlebotomus genus, is the only known vector of L. major (4). The identification and characterization of different species of leishmaniasis is one of the main factors involved in the prevention and control of this disease. Determining the dominant strain in each region has applications in the pharmaceutical industry, vaccination and the production of antigens for disease diagnosis (5).
3. Methods

3.1. Study Area

This cross-sectional study was carried out in Quchan (37.1293° N, 58.4744° E) with a population of 96,953, consisting of 25,066 families (Figure 1).

3.2. Patients

This study was conducted in collaboration between Mashhad University of Medical Sciences and the health centers of Quchan in 2017. One-hundred three individuals who had at least one skin ulcer and presented to medical centers were included in this study. Scraping was performed for the direct examination of the edge of each skin lesion using a sterile scalpel after obtaining informed consent and completing a questionnaire.

3.3. PCR

After preparing direct Giemsa stained smears, all the slides were scrapped for DNA extraction (GeNet Bio, Korea). Polymerase chain reaction was performed with specific primers of kDNA pattern as F: 5’-TCGCAGAACGCCCTACC-3’ and R: 5’-AGGGGTTGGTGTAAAATAGG-3’ (Tuba-Negin, Iran), which produced a 615-bp fragment for *L. major* and 744-bp fragment for *L. tropica* (6). PCR was performed as initial denaturation for 5 min at 95°C, followed by 38 cycles of 94°C for 30 s, 60°C for 45 s and 72°C for 60 s, ending with a final extension at 72°C for 7 min. In each round of PCR, two standard *L. major* samples (strain: MRHO/IR/75/ER) and *L. tropica* (strain: MHOM/01/IR/YAZA) were used as a positive control and distilled water as the negative control. Data were analyzed in SPSS-20 (SPSS Inc., Chicago, IL, USA) using the chi-square test.

4. Results

Out of the 103 suspected individuals, 77 (74.8%) and 86 (83.4%) patients were diagnosed with CL by microscopic examination and PCR, respectively. Figure 2 presents the PCR results. The sensitivity of the microscopic method was measured as 89.5%. No significant correlations were observed between habitat (rural and urban) and the *Leishmania* species (Table 1).

Overall, 46 (53.5%) out of the 86 cases were male and 40 (46.5%) were female (P = 0.2). The patients’ age ranged from ten months to 78 years and the highest rate of infection was recorded in the 20-30-year-old age range (33%).

The most common lesion site was observed on the patients’ hand (Table 2). Papule was the most commonly-observed clinical feature (Table 3). There were no statistically significant differences between the type of clinical presentation and *Leishmania* species (P = 0.2). Also, most of the patients had one lesion (Table 4). Most lesion measurements (n = 55, 64.7%) were under ten millimeters (Table 5). There were no correlations between the size of the ulcer and the species of *Leishmania* (P = 0.5).

5. Discussion

To date, no research has been conducted in Quchan as an endemic focus of CL. According to the present findings,
there is a predominant species of *L. tropica* in Quchan, a city neighboring Mashhad, Sabzevar, Bojnord, Esfarayen and Neyshabur. *L. major* is responsible for leishmaniasis in Esfarayen (based on an unpublished study) and Sabzevar (7), as ZCL was reported in all the 153 patients in the latter city. Mashhad and most of the cities in Khorasan-e Razavi are an endemic region for ACL (7, 8). In Iran, ZCL is endemic in the rural regions of the west, the northeast, the center and the southwest, especially in Khuzestan Province. ACL mostly occurs in large urban areas, such as Yazd, Tehran, Shiraz, Mashhad and Kerman (9).

Table 4. The Frequency Distribution of Lesion Number in Patients with Cutaneous Leishmaniasis in Quchan in 2017

| Lesion Number | Species      | *L. major*, Frequency (%) | *L. tropica*, Frequency (%) | Total, Frequency (%) |
|---------------|--------------|---------------------------|----------------------------|---------------------|
| One           | *L. major*   | 14 (48.3)                 | 35 (61.4)                  | 49 (57)             |
| Two           | *L. major*   | 5 (17.2)                  | 15 (26.3)                 | 20 (23.2)           |
| Three or more | *L. major*   | 10 (34.5)                 | 7 (12.3)                  | 17 (19.8)           |
| Total         | *L. major*   | 29 (100)                  | 57 (100)                  | 86 (100)            |
Table 5. The Frequency Distribution of Lesion Measurement (mm) in Patients with Cutaneous Leishmaniasis in Quchan in 2017

| Lesion Measurement, mm | L. major, Frequency | L. tropica, Frequency | Total, Frequency |
|------------------------|---------------------|-----------------------|-----------------|
| 0 - 9                  | 40                  | 15                    | 55 (62.7)       |
| 10 - 19                | 12                  | 5                     | 17 (20)         |
| 20 - 29                | 9                   | 1                     | 10 (11.8)       |
| 30 - 39                | 2                   | 1                     | 3 (3.5)         |
| Total                  | 29 (100)            | 57 (100)              | 86 (100)        |

Table 6. The Relationship Between Season and Disease Incidence in Patients with Cutaneous Leishmaniasis in Quchan in 2017

| Season | L. major, Frequency (%) | L. tropica, Frequency (%) | Total, Frequency (%) |
|--------|-------------------------|---------------------------|----------------------|
| Spring | 1 (3.4)                 | 8 (34.1)                  | 9 (10)               |
| Summer | 9 (31.2)                | 20 (35.1)                 | 29 (34)              |
| Autumn | 14 (48.2)               | 23 (40.3)                 | 37 (43)              |
| Winter | 5 (17.2)                | 6 (10.5)                  | 11 (13)              |
| Total  | 29 (100)                | 57 (100)                  | 86 (100)             |

The most common number of ulcers in CL patients is one and usually occurs in the hands and face, which are un-woven. Among all the risk factors evaluated in the present study, including age, gender, clinical features, living place, lesion size, lesion number, engaged organ, travel history and season, only the relationship between season and leishmaniasis prevalence was statistically significant. All the studies reported a high infection rate in autumn (7). Due to the activity of sand flies in late spring and summer, the incidence of CL is highest in autumn and winter, which had the most recorded number of leishmaniasis patients in Iran (10).

The articles that reported a significant relationship between gender, age and cutaneous leishmaniasis were the same number as the articles that reported the opposite. Despite the many studies conducted in endemic areas of CL, the issue remains unclear. All age groups were affected in the different reports; that is, age ranges 21 - 30 years, 10 - 15 years and 0 - 9 years, all of which fall under the age of 40 years (II-13).

None false negative cases were reported by microscopic examination in this study. KDNA-PCR has a higher sensitivity than the microscopic method with a 75% - 98% rate for the detection of Leishmania species, even in the patients’ urine (14).

The boundaries of leishmaniasis incidence have changed in recent years, as new foci of L. tropica were detected in rural regions (15, 16). In this study, 56.1% of the patients with L. tropica lived in rural districts. Humans are thought to be the reservoir of L. tropica. In rural areas, it is not important to determine the contaminated rodents, but infected patients should be treated.

5.1. Conclusions

According to the present condition, L. tropica is responsible for cutaneous leishmaniasis in either rural or urban areas of Quchan. In rural areas, better vector/reservoir detection, training and control programs should be set in place.

Acknowledgments

The authors wish to express their gratitude to the health centers of Quchan for their help in collecting the samples.

Footnotes

Authors’ Contribution: Seyed Aliakbar Shamsian designed and coordinated this study. Ameneh Gholampour collected the samples. Elham Moghaddas wrote the manuscript. Lida Jarahi verified the analytical methods. All the authors read and approved the final manuscript.

Conflict of Interests: The authors report no conflicts of interest in this work.

Ethical Approval: The study was approved by the Ethical Committee of the Research Department of Mashhad University of Medical Sciences (IR.MUMS.fm.REC.1394.523).

Funding/Support: This work was funded by Mashhad University of Medical Sciences (project grant No: 940900).

Patient Consent: Informed consent has been obtained from all the patients.

References

1. Alvar J, Velez ID, Bern C, Herrera M, Desjeux P, Cano J, et al. Leishmaniasis worldwide and global estimates of its incidence. PLoS One. 2012;7(5). e35671. doi: 10.1371/journal.pone.0035671. [PubMed: 22693548]. [PubMed Central: PMC3365071].
2. Yaghoobi-Ershadi MR, Hanafi-Bojd AA, Javadian E, Jafari R, Zahraei-Ramazani AR, Mohebali M. A new focus of cutaneous leishmaniasis caused by Leishmania tropica. Saudi Med J. 2002;23(3):291-4. [PubMed: 11938418].
3. Khazaeei S, Mohammadian Hafshejani A, Saatchi M, Salehiniya H, Nematiollahi S. Epidemiological aspects of cutaneous leishmaniasis in Iran. Arch Clin Infect Dis. 2015;10(3). doi: 10.5812/archced.28511.
4. Hepburn NC. Cutaneous leishmaniasis. Clin Exp Dermatol. 2006;25(5):363-70. [PubMed: 16125866].
5. Fakhar M, Keyghobadi M, Akramipour R, Ghadiri K, Limouei M. Characterization of Leishmania isolated from Kala-azar infected patients in Kermanshah using PCR. J Kermanshah Univ Med Sci. 2011;15(2).
6. Motazedian H, Karamian M, Noyes HA, Ardehali S. DNA extraction and amplification of Leishmania from archived, Giemsa-stained slides, for the diagnosis of cutaneous leishmaniasis by PCR. Ann Trop Med Parasitol. 2002;96(1):31-4. doi: 10.1079/00034980225000484. [PubMed: 11999533].
7. Namazi MJ, Dehkordi AB, Haghighi F, Mohammadzadeh M, Zarean M, Hasnab M. Molecular detection of Leishmania species in north-east of Iran. Compr Clin Pathol. 2018;27(3):279-33. doi: 10.1077/j00580-018-2658-9.
8. Rezai A, Moghaddas E, Bagherpor MR, Naseri A, Shamsian SA. Identification of Leishmania species for cutaneous leishmaniasis in Gonabad, Bardaskan and Kashmar, Central Khorasan, 2015. Jundishapur J Microbiol. 2017;In press.[In press]. doi: 10.5812/jjm.44469.
9. Ghaee MA, Mirhendi H, Marashifard M, Kanannejad Z, Taylor WR, Sharifi I. Population structure of Leishmania tropica causing anthroponotic cutaneous leishmaniasis in Southern Iran by PCR-RFLP of kinetoplastid DNA. Biomed Res Int. 2018;2018:604998. doi: 10.1155/2018/604998. [PubMed: 29984240]. [PubMed Central: PMC6011176].
10. Khosravani M, Nasti Z, Keshavarz D, Rafat-Panah A. Epidemiological trend of cutaneous leishmaniasis in two endemic focus of disease, south of Iran. J Parasit Dis. 2016;40(4):3609-13. doi: 10.1007/s12639-015-0740-7. [PubMed: 27876994]. [PubMed Central: PMC518366].
11. Salehi M, Ghasemian A, Mostafavi SKS, Masoumi M, Nojoomi F. Epidemiology of cutaneous leishmaniasis in Neyshabur, Iran from 2010 to 2014. J Coastal Life Med. 2016;4(1):887-9. doi: 10.42980/jclm.4.2016.6-167.
12. Momeni AZ, Aminjavaheri M. Clinical picture of cutaneous leishmaniasis in Isfahan, Iran. Int J Dermatol. 1994;33(4):260-5. doi: 10.1111/j.1365-4362.1994.tb01039.x. [PubMed: 802082].
13. Razmjou S, Hejazy H, Motazedian MH, Baghaei M, Emamy M, Kalantary M. A new focus of zoonotic cutaneous leishmaniasis in Shiraz, Iran. Trans R Soc Trop Med Hyg. 2009;103(7):727-30. doi: 10.1016/j.trstmh.2008.12.013. [PubMed: 19221055].
14. Veland N, Espinosa D, Valencia BM, Ramos AP, Calderon F, Arevalo J, et al. Polymerase chain reaction detection of Leishmania kDNA from the urine of Peruvian patients with cutaneous and mucocutaneous leishmaniasis. Am J Trop Med Hyg. 2011;84(4):556-61. doi: 10.4269/ajtmh.2011.10-0556. [PubMed: 21460009]. [PubMed Central: PMC3062448].
15. Mebrahtu YB, Lawyer PG, Ngumbi PM, Kirigi G, Mbugua J, Gachihi G, et al. A new rural focus of cutaneous leishmaniasis caused by Leishmania tropica in Kenya. Trans R Soc Trop Med Hyg. 1992;86(4):385-7. doi: 10.1016/0035-9203(92)90230-a. [PubMed: 1332221].
16. Sharifi I, Poursmaelian S, Aflatoonian MR, Ardakani RF, Mirzaei M, Fekri AR, et al. Emergence of a new focus of anthropopotic cutaneous leishmaniasis due to Leishmania tropica in rural communities of Bam district after the earthquake, Iran. Trop Med Int Health. 2011;16(4):510-3. doi: 10.1111/j.1365-3156.2011.02729.x. [PubMed: 21255206].