

## Sandfly and Leishmaniasis: A Review

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### Abstract

Sandfly has a long history of association with humans, which still are suffering from its harmful impacts. It parasites humans and other animals and acts as a source of nuisance and annoyance to them. The present bibliographical study explains the role of Sandfly as a disease vector of Leishmaniasis. This article highlights various aspects of the life of the vector which includes its general description of morphology, biology, life cycle, and major control strategies. Regarding to control strategies of the vector, it was reviewed that chemical control is the most common and efficient technique but sensitivity to is decreasing due to increased insecticide resistance and environmental constraints. This article also highlights the different types of Leishmaniasis and symptoms, treatment, mortality rate and species diversity of Sandfly in Pakistan.

**Keywords:** Sandfly; Phlebotomine flies; Leishmaniasis; Sandfly diversity in Pakistan

### Introduction (Leishmaniasis and Sandfly as Vector)

Leishmaniasis is found every continent except for Antarctica and Australia. Sandflies are the carrier of *leishmaniasis*, affecting human health in more than 90 countries in the subtropics, tropics, and southern Europe [1]. It is more common in rural areas as compare to urban ones. This disease is more common in male adults [2]. Leishmaniasis disease is transmitted by the *Phlebotomine* flies, genus *phlebotomus* in the new world and *Lutzomyia* in the old world. The vector of this disease belongs to the order *Diptera*, class *insecta*, Family: *Psychodidae* [3]. Sandfly is about 3 mm in length, characterized as “hopping” flight. They have dark and large eyes, long antennae the mouthparts are oriented downward, dagger shaped and short, the legs are delicate and long [4].

This disease is caused by 20 *Leishmania* species by the bite of female sandflies. Thirty species of sandflies are the vector of disease. Their reservoir hosts are humans and wild or domestic animals. Female Sandflies feed on reservoir hosts and get infected [5]. This parasite is transmitted through the use of infected syringes from infected individuals. The common species of *Leishmania* are *L. chagasi*, *L. infantum* and *L. donovani* [6]. There has been chance of increase in Leishmaniasis in the last two decades due to the migration of people towards urban area [7]. Through travelling, Leishmaniasis spread in people which are living in non-endemic areas.

### Habitat and flight range

Sand flies are nocturnal and sensitive to dehydration. They shelter in caves, rocks, animal burrows, tree holes and human rooms or accommodation. They fly close to the ground in short hops (jumps) therefore they are called weak flyers. Their flying range is 300 m but in deserted environments some species can travel up to 2300 m [8]. Due to short range of flight, adult stay near to the larval developmental site. Flies in the New World are found near tree holes and caves [9]. *Lutzomyia shannoni* in the United States was found in hardwood forest

and meadow. In the Old World sand flies are found associated with contaminated soils of animal shelters, rodent burrows and termite mounds, also in the earthen floors of human habitations [10].

### Feeding behavior

Male and female sandfly feed on nectar from fruits, flowers and plant juices. Carbohydrates are the source of energy. Female flies suck a blood meal to complete the development of egg batches. Some species of sandfly are autogenous, these species lay butch of eggs without first feeding on blood, female species of sandfly are the disease causing agents [11]. Most anthropophilic (blood sucking arthropods) sand flies bite people outside their tents, houses and accommodation [12].

### Growth and development of Sandfly

Sandfly requires 28 degree centigrade temperature and 40% humidity for growth and development. In laboratory, these flies take 20-40 days to complete the life cycle. The sandfly shows holometabolous metamorphosis (egg, larva, pupa and adult). They can laid between 30 to 70 eggs. Within 1-2 weeks, they hatch. Larvae feed on dead organic matter. Pupal development is completed in 5 to 10 days. Before the sun rising adult emerge from pupa, during day time sandfly cannot survive in dry environment so, resting at humid sites. At night this fly drops the ambient temperature and become active by increasing the humidity [13].

### Diagnosis

Due to high specificity, parasitological diagnosis remains the gold standard in leishmaniasis diagnosis. Conventional techniques like microscopy and culture still the standard approach for the diagnosis of leishmaniasis at primary health levels in areas of endemicity, because more sophisticated techniques like molecular diagnosis are currently expansive and rarely available [14].

Visceral leishmaniasis are commonly diagnose through serological approaches; identification of antibodies in the serum. Immunochromatographic dipstick tests and freeze dried antigen based

direct agglutination test have high specificity, sensitivity and easy to use, require minimal laboratory setup [15].

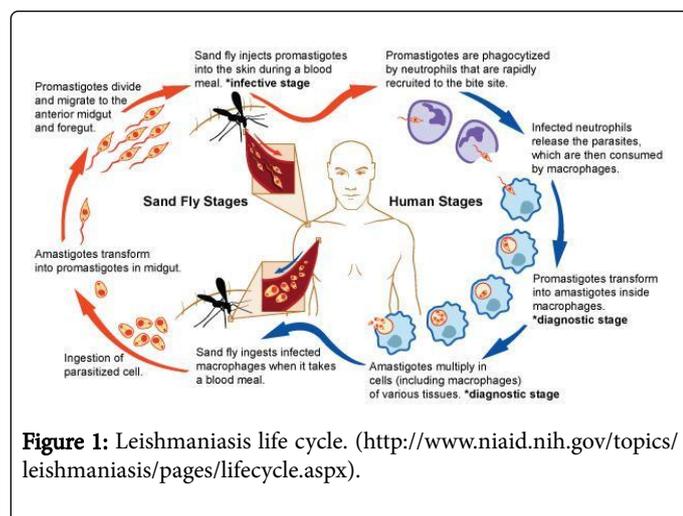
Morphologically identical species can be differentiated with molecular techniques such as PCR, isoenzymes, etc. Sibling species of sandflies can be identified by polymerase restriction fragment length polymorphism (PCR-RFLP) of the 18S rRNA gene. PCR is a powerful tool for research on sandfly species and the relationship between *Leishmania* species and their vectors [16].

The infection of sandflies with *Leishmania* can also be examined by dissecting an individual under a microscope. Fresh specimen should be used. The expertise and skills are needed for the study of small size individuals. This process takes long time, large numbers of individuals have to be examined to obtain data for each area, and the rate of infection of sand flies with *Leishmania* is generally low 0.01-1% [17].

To identify *Leishmaniasis* infections in experimentally infected and field captured phlebotomine sandflies molecular techniques are used such as PCR-RFLP, KDNA-PCR, flurescent quantitative PCR and mini exon PCR. Real time PCR also used to detect the sandfly infection of *Leishmania* [18-20].

### Sandfly and Leshmania Life Cycle

In infected host, the ingested amastigotes (protist cell, non-motile) transform into promastigotes (external flagellum) in the blood meal of different hematophagus arthropods. If the host is unsuitable, than parasites passed out with the feces [21] *Leishmania* species and other trypanosomatids attached to the parasites on the midgut epithelium [16] during digestion of the blood meal, the parasite is retained in the gut and starts a stage differentiation [22]. In next phase promastigotes move to the cuticle lined foregut of the sandfly. In the foregut some attach and some remain free for next transmission by bite [23] into the vertebrate host.



In the alimentary tract of the sandfly, promastigotes multiply by binary fission. After 7 days promastigotes undergo metacyclogenesis and become infectious (metacyclic promastigotes). When the sandfly pokes the skin with its proboscis during feeding, metacyclic promastigotes are released into the host together with saliva (Figure 1). In host macrophage, metamorphose into the amastigote form. By binary fission, they increase in number within the phagolysosome until the cell bursts and infects other phagocytic cells. In this way the cycle continues.

### Forms of Leishmaniasis Disease

There are four major forms of *Leishmaniasis*

*Cutaneous leishmaniasis*

*Visceral leishmaniasis*

*Diffuse cutaneous leishmaniasis*

*Mucocutaneous leishmaniasis*

**Cutaneous Leishmaniasis:** It is characterized by skin ulcers, lesions may be associated with *sporotrichotic* (fungus infection). After being bitten, people develop skin sores a few weeks to months. In the old world this disease is caused by *L. tropica*, *L. aethiopica* and *L. infantum* and in New world by *L. guyanensis*, *L. Mexicana* and *L. braziliensis* [16,23]. There are two epidemiological forms of cutaneous *Leishmaniasis* which are arthropod-borne: cutaneous *Leishmaniasis* and zoonotic cutaneous *Leishmaniasis*. The main reservoir hosts are humans and rodents. The main vector in arthropod-borne cutaneous *leishmaniasis* and zoonotic cutaneous *leishmaniasis* are *P. sergenti* and *P. papatasi* [23-25]. The infection manifests mainly on adults and young children [26].

**Visceral Leishmaniasis:** This is the most severe form of the disease also known as Kala azar (black fever) caused by *L. donovani* and if left untreated are always fatal and mortality rate is 100%. Symptoms of the disease are weight loss, fever (within several weeks or months), hepatomegaly, splenomegaly, pancytopenia, anaemia and lymphadenopathy. This disease is most common in developing countries [6,27]. It is distributed worldwide in both the old and new world. The occurrence of visceral *leishmaniasis* can also be influenced by environmental changes such as incursion of agricultural farms, urbanization and settlements into the forest areas. The distribution of parasites also is affected by humidity, atmospheric temperature, land degradation, global warming and rainfall [28]. Clinical diagnosis of visceral *leishmaniasis* is complex because its features are shared by host of other diseases such as *asthphoid*, malaria and tuberculosis. Diagnosis of this disease is *in vitro* culture and DNA of parasites can be detected in tissue samples (blood or urine). Visceral *leishmania* caused by *L. donovani* complex and *L. infantum* especially [29]. Visceral *leishmaniasis* is anthroponotic (disease causing agents carried by humans to other animals) in transmission [30].

**Diffuse Cutaneous Leishmaniasis:** This disease is characterized by many non-ulcerative skin lesions on the entire body. These lesions contain vacuolated, infected macrophages with only few lymphocytes present. It is wide spread, similar to lepromatous leprosy lesions [31]. The uncontrolled parasite growth results from lack of cell mediated immunity to *leishmanial* antigen [32].

**Mucocutaneous Leishmaniasis:** After the onset of cutaneous *leishmaniasis*, mucocutaneous *leishmaniasis* occurs and it is characterized by the destruction of pharyngeal and oral- nasal cavities. Genetic factors are also important in the incidence of this disease. The initial symptoms of mucocutaneous *leishmaniasis* are not severe with stuffiness and nasal inflammation, but slowly perforation of the septum and ulceration may occur. The lesion expands to the larynx, gums, pharynx, soft palate and face [33]. Bones are not affected due to bacterial infections untreated disease may lead to diarrhea, pneumonia and tuberculosis [34]. Death may also occur due to malnutrition (difficulty in swallowing), lung infections, and suffocation (due to closure of laryngeal aperture) [35].

## Mortality Rate

Leishmaniasis currently is threatening about 350 million people in endemic areas. Global there are 10 to 12 million cases [36-38]. A most recent study indicates that 50,000 deaths occur each year [36].

## Sandfly in Pakistan

According to Durrani et al. [13] conducted a study from May 2007 to June 2008 and recorded the sandfly fauna in four regions of Pakistan. They collected 20,683 sandflies which belong to the genus *Phlebotomus*. No sandflies were recorded in Eastern Pakistan.

Durrani et al. [13] collected the following species:

*P. papatasi*

*P. major*

*P. bergeroti*

*P. alexandri*

*P. orientalis*

*P. longipes*

*P. sergenti*

*P. pedifer*

Endemic areas for the disease vector, described by Bhattacharia et al. [39] were Swat and Gilgit, Lasbela, Rawalpindi, Abbottabad, Chilas, Skardu, Mansehra, Chitral, Dir, D.G. Khan, Rajanpur, Quetta, QilaSaifullah, Qila Abdullah, Pishin, Larkana, AzadKashmir and Dadu. These areas are the North, West and South Western Pakistan. Ali et al. [40] confirmed that south-eastern areas of Pakistan are non-endemic.

## Sandfly Control Techniques

Sandfly control is similar to that used for mosquito because both vector share similar characteristics. *Bacillus sphaericus* is used to control sandfly larvae. In this innovative technique, bait-fed adults were used to carry the bacterial control agent to larval habitats, resulting in larval mortality in burrows up to 10-30 m away from the baited solution [41].

## Insecticides

Spray of insecticides on burrows, walls and humans accommodation. The first insecticide used to control sandfly was DDT (dichlorodiphenyltrichloroethane) and it was sprayed in India, Brazil, China, Soviet Union. It caused reduction in population of sandfly, but there was no direct impact on disease control [42]. From the Middle East, study of four species of sandflies indicates that malathion and DDT less toxic than the newer pyrethroid insecticides [43].

In 1958 and 1970 there was a program to control visceral leishmaniasis in Bihar (India). No visceral leishmania disease was reported during these years but at the end of this control program, disease resurfaced. Antimalarial programs reduce the population of sandfly in Iran, Bangladesh, Peru and Italy. But this did not reduce leishmaniasis [42].

Following insecticides are currently used to control sandfly [44]

Talstar P<sup>®</sup> (bifenthrin, NSN 6840-01-525-6888)

Aqualure<sup>®</sup> 20+20 (permethrin, PBO; NSN 6840-01-606-8581)

Demon<sup>®</sup> WP (cypermethrin, NSN 6840-01-390-4822)

Demand<sup>®</sup> CS (λ-cyhalothrin, NSN 6840-01-428-6646)

Pestabs<sup>®</sup> (λ-cyhalothrin, NSN 6840-01-431-3357)

## Habitat modification and pesticides

Habitat modification such as ploughing or flooding in the host ecosystem reduces the sandfly density. Pesticide application is effective to control the density of sandflies if target sites are known. But this application is not sustainable, limited by resources and time [21].

## Treatment of Leishmaniasis

For all four major forms of Leishmaniasis chemotherapy is effective. Certain factors compromise chemotherapeutic options such as toxicity, costs, complicated and long term regimens "establishment" [45,46]. Many drugs are also used to get recovery from Leishmaniasis.

The eradication of this disease will be achieved through vaccination [47]. Bacon [48] describes that if population of Venezuela, Peru, Colombia, Ecuador, Mexico, Brazil and Bolivia were vaccinated through vaccine than it would provide ten years of protection, 41,000-144,000 cases of cutaneous leishmaniasis could be averted, and this would be lower cost than other treatments. Vaccine provides 5 years of protection and 50% efficacy [48]. In Bihar State similar study for visceral leishmaniasis also confirmed the cost effectiveness of vaccine [48]. The development of vaccine is under processing.

## Conclusion

Leishmaniasis occurs due to sandfly species that feed on human and other reservoir hosts. Young children and male adults are mostly affected. Each year millions of deaths occur due to this disease. Waste openly disposed is the major factor for attracting the vector [49-52]. It is concluded that vector and parasites have adaptability to spread in new environments such as urban and suburban areas. Migration of people from rural to urban areas increases the risk of leishmaniasis. Slow progress has been made on the development of a vaccine. The development of vaccine has proven a challenging and difficult task due to inadequate knowledge of the parasite pathogenesis and complexity of the immune response. Sandfly population should be controlled to prevent Leishmaniasis. There is lack of information and interest to evaluate the significance of Sandfly control in disease control. This is a neglected disease due to limited resources for diagnosis, treatment and control in developing countries.

## Recommendations

Following are some suggestions to control the vector and Leishmaniasis:

- To get protection from sandflies keep the environment clean.
- Use bed net to protect yourself and use pyrethroid-containing insecticide to spray the bed net
- Use insecticides to control sandfly.
- Do not expose the skin, wear long sleeved shirts.
- Use insect repellent on exposed skin.
- Waste should not be openly disposed.
- Drug treatments are expensive, poor people cannot afford. Drugs are also toxic and have numerous side effects.

- To control this disease efficacious vaccine is needed. Vaccines are well tolerated, safe and immunogenic. Vaccines effective in treatment and prevention of different form of leishmaniasis.
- If Leishmaniasis symptoms appear it is important to consult a physician immediately without wasting time.
- There should be need to develop public awareness about this vector.
- There should be need to aware people about this vector.

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