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CUTANEOUS LEISHMANIASIS IN DADU, HYDERABAD AND JAMSHORO DISTRICTS OF SINDH,PAKISTAN AGE WISE INTENSITY CORRELATION WITH PRIMARY HOST DOGS,CAUSES, TREATMENT, PREVENTION AND ERADICATION OF DISEASE

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ABSTRACT

The Cutaneous Leishmaniasis(CL) in Hyderabad, Dadu and Jamshoro districts of Sindh province, Pakistan, age wise correlated with the primary host i.e. dogs with reference to intensity of skin disease in humans was found statistically positively correlated. Causes, treatments and cure in the light of prevention and eradication of the disease is also briefly discussed.

KEYWORDS: Cutaneous Leishmaniasis.

INTRODUCTION

The purpose of this study was to see the situation of cutaneous Leishmaniasis (CL)in the above three districts of Sindh and possible threats of epidemics and spread to neighboring districts. In the far Eastern Mediterranean Region (EMR) of the World Health Organization (WHO) Leishmaniasis is considered as a principal health problem. The disease most commonly affects the children due to less developed immunity. In most of the cases disease also affects the other family members indicating limited flight range of sand fly, person to person transmission or genetic predisposition.(Reithinger et al.2010; Oliveira et al.2015)Cutaneous leishmaniasis is a vector born, usually a self-healing disease which spread through the bite of a female sandfly of dipteran, family Psychodidae, subfamily Phlebotominae, Phlebotomus papatasi. The vector Phlebotomus papatasi is chiefly distributed in Asia Minor regions, Southern Asia (including Pakistan), Northern Africa and Southern Europe (WHO, 1998). Cutaneous leishmaniasis(CL) in India is called Delhi Boil, in Iraq Baghdad, Boil, in Bangladesh, Bangladeshi Boil and in Afghanistan, Saldana (Abdul Ghani, et al. 2008). About 1.5 million new cases of this disease are reported every year from world mainlyfrom across the Algeria, Afghanistan, Sudan, Bangladesh, India, Pakistan, Nepal and Brazil (WHO, 1996) leading to about one hundred thousanddeaths due to visceral Leishmaniasis (VL) annually (Ashford, 2000).

More research work on prevalence and molecular studies of cutaneous leishmaniasis are reported from Sindh province of Pakistan, as compared to other regions, for example (Nawab et al.1997) noted 90 confirmed cases out of 120 referred to Dr. Ehsanullah's Laboratory, Karachi for diagnosis. But the present study areas i.e. Dadu,Hyderabad and Jamshoro districts appear to be neglected in literature although Dadu especially Juhi taluka has been mentioned often, with reference to immigration of CL patients brought from Baluchistan and Afghanistan as it shares borders with Baluchistan with patients from Afghanistan refugees (Bhutto et al., 2004; Kolachi et al., 2005;Ejaz et al., 2008; Reithinger et al 2010; Sharif et al., 2017).

Dawit et al. (2013) stressed that epidemiological studies on leishmaniasis must not be limited to endemic areas but it should also continue in other areas. Kassi et al. (2008) only observed that CL was reported from all the provinces of Pakistan but Bhutto et al. (2008) deduced that it is endemic in Pakistan. ZCL (Zoonotic Cutaneous Leishmaniasis) spread from animal to animal or from animal to man, (ACL) Anthroponotic Cutaneous Leishmaniasis from human to human or DCL (Diffused Cutaneous Leishmaniasis resulting in chronic skin and disseminated lesion similar to that of lepromatous leprosy was reported by Bari et al. (2011). Although Klaus and Frankenburg (1999) and Ashford (2000) stated that *Leishmania* parasitic spp.(i.e. *L.tropica, L.major* and *L.infantom*) are found intracellularly in monocytes and macrophages and they mayalso be found extracellularly in a giemsa stained smear. The giemsa stained amastigotes appear as blue oval bodies,2-5um in diameter having a blue oval nucleus in vitro culture. The flagellated motile promastigotes are10-15 (um).

MATERIAL AND METHOD

During the present study a survey was made in the city of Hyderabad of Sindh province which embraces three districts i.e. Dadu,Jamshoro and Hyderabad. The city is the second largest (population wise) in Sindh province and there are many skin diseases hospitals, health centersand pathological laboratories which were visited to locate and identify CL patients with special reference to history of leishmaniasis in a survey report and an analysis of diseased(CL) patients district wise and age wise statistically are presented using Minitab recent softwareversion 17 and age groups are made in human beings and in dogs and explanation is given under.

RESULT

Agewise correlation between C1(Human) and C2(Dogs)



Fig 1: shows a regressive analysis between C1 and C2as under C1= 5.019+3.139 C2 Positive Correlation. Intensity of CL corresponds to the age of humans between 10 to 25 years in Dadu districtfollowed by Hyderabadand Jamshoro and for the dogs 1 year to 8 years.

Co-efficient of determination of this relationship between C1 and C2 is about 67% i.e. adjusted to the Coefficient of determination which is 65%.



Fig 2: This histogram also shows the same relation between C1 and C2. For C1 the mean is 21.5 and for C2 mean is 5.25. The Standard deviation for C1 is 10.23 and SD for C2 is 2.67. The Co-efficient of variation for man is 47% whereas the Co-efficient of variation of dog is 51%.

DISCUSSION

For studying the reservoir of *leishmania* parasites in the above districts, the primary animal host i.e. the dogs with reference to districts wise and age wise intensity of Cutaneous Leishmaniasis(CL) were also surveyed and statistically analyzed and it was seen if intensity of the skin disease in the primary animal host i.e. dogs is directly proportional and positively correlated with the end host i.e. human patients(Qureshi et al,2018). Photographs of the lesions of the disease in the primary and end host are also given. Diagnosis, causes, cures and treatment of the human patients with reference to prevention and eradication of leishmaniasis are also briefly discussed.Most cutaneousleishmaniasis lesion heals in 8 to 12 months, however it is guite painful to live for one year with a facial lesion (on cheek, nose, lips and chin) which could disfigure face and often cause serious complications because of secondary bacterial and fungal infections. Appropriate treatment is the only choice for these patients for the early healing of their lesions. Dowlati (1996), Al-Majali et al. (1997) and Reithringer et al. (2005) have discussed (a) thermotherapy using hot objectsoracids but these have side effects, sometimes leaving permanent scars of lesions.(b) Cryotherapy using liquid nitrogen and surgery of the lesions and (c) the intralesional injections of pentavalent antimonials i.e meglumine antimonite and sodium stibogluconate commercially called glucantime with generic name "Avantis and pentostam (Glaxo/ Wellcome) are practiced around the world (WHO, 1990). Also plant alkaloid Berberine was found seriouslyaffectedby cutaneous leishmaniasis in rats. Harmaline extracted from Peganum harmalashowed antiprotozoan activity (Evans et al.,1987,Wright et al. 1990). Steroidal alkaloid holamineand Hydroxyl holamine extracted from the Hoarthena and Curtisii also showed plant Leishmanicidal against activity Leishmania donovani(Kam et al., 1998).

Cure and eradication of Leishmaniasis.

Early diagnosis and treatment of patients along with curing of skin lesion are important to prevent spread of disease. Breaking the chain of the life cycle of the Leishmania parasite (i.e. elimination of sandfly vector) or destruction of animal reservoir i.e. killing Rodents and Dogs is remarkably effective in leishmaniasis eradication. Ashford et al. (1995) and Klaus and Frankenburg (1999) have pointed out that by destroying animalfood places and by eliminating burrows of rodentsis quite effective. Successful control of leishmaniasis is also achieved bythe confinement of sandfly including the elimination and annihilation of itsbreeding sites by cleaning garbage, litters and debris near human residence and by covering cracks, Crevices in walls and roofs and spraying with effective synthetic insecticides i.e. deltamethrin (pyrethroid group) with generic name Cyhalothrinunder windows, could be very effective. Mosquito nets are not helpful in preventing bite of sandfly because of larger mesh size. Smaller mesh size is required. Nets if impregnated with insecticide

such as permethrin or deltamethrin are more helpful.(Ejaz et al.2008). By plantation Bouginvilla Glibra as stated by Schlein et al.(2001) and Alten et al. (2003) could significantly reduce the occurrence of cutaneous leishmaniasis.

REFERENCES

- 1. Abdulghani, M. A., Saher, H.and Obaidi, Al. 2008. Cutaneous leishmaniasis in Iraq, *J. Infect Developing Countries*, *3*(2): 123-129.
- Al-Majali, O., Routh, Hirak. B., Abuloham, O., Bhowinic, K. R., Mushsen, M., Hebeheba, H. 1997.
 A- 2-year study of liquid nitrogen theropy in cutaneous leishmaneisis. *Intl. J. Dermatol, 36*(6): 460-462.
- 3. Alten, B., Caglar, S. S., Kaynas, S. and Simset, F. M. 2003. Evaluation of protective efficacy of K-OTAB impregnated bendnets for cutaneous leishmaniasis control in Southeast Anatolia- Turkey. *J. Vec. Ecol*, 53-64.
- Ashford, D. A., Bozza, M., Frerire, M., Miranda, J. C., Shelock, I., Eulaio, C., Lopes, U., Fernandes, O. and David, J. R. 1995. Comparison of the polymerase chain reaction and serology for the detection of canine visceral leishmaniasis. *Am. J. trop. Med. Hyg*, 53: 251-255.
- 5. Ashford, R. W. 2000. The leishmaniasis as emerging and re-emerging zoonoses. *Int. J. Parasital, 30*: 1269-1281.
- 6. Bari, A. U., Rizwan, H., Mahmood, K., Iqbal, M., Shahbaz, N., Tariq, K. M. 2011. Clinicoepidemiological pattern of cutaneous leishmaniasis in armed forces personnel fighting war against terrorism in Khyber Pakhtonkhuwa Province and Fata regions. J. Pak, Assoc. Dermatol, 21: 10-15.
- 7. Bhutto, A. M., Soomro, F. R. and Katakura, K. 2008. Leishmaniasis in Sindh, Pakistan: outbreak and review of the literature. *Journal of Pakistan Association of Dermatologists*, 18: 212-219.
- Bhutto, A. M., Soomro, F. R., Shah, S. S., Solangi, A., Ahmed, A., Uezato, H., Kato, H., Katakura, H., Nonaka, S. and Hashiguchi, Y. 2004. Epidemiology of Leishmaniaes in Pakistan and a literature review. In: Hashiguchi, Y. (ed.), Studies on the New World Leishmaniases and its Transmission with Particular Reference to Ecuador, Argentina and Pakistan, 119-130. *Res. Rept. Series* No. 7, Kyowa Printing and Co. Ltd. Kochi, Japan.
- 9. Dawit, G., Girma, Z. and Simenew, K. 2013. A Review on Biology, Epidemiology and Public Health Significance of Leishmaniasis. *J Bacteriol Parasitol*, 4:166.
- 10. Dowlati, Y. 1996. Cutaneous leishmaniasis: clinical aspect. *Clin Dermatol*, 14: 425-431.
- Ejaz A, Raza N, Din QU, Bux H. Outbreak of cutaneous leishmaniasis in Somniani, Balochistan – implementation of preventive measures for deployed personnel of armed forces. J Pak Assoc Dermatol, 2008; 18: 220-5.

- Evans, D. A., Kenndy, W. P., Elbiharis, S., Chapman, C. J., Smith, V. and Peters, W. 1987. Hybrid formation within the genus Leishmaneiasis? *Parasitologia*, 29(2-3): 65-73.
- Kam, T. S., Sim, K. M., Koyane, T., Toyoshina, M., Hayashi, M. and Kamiyama, K. 1998. Cytotoxic and leishmanicidal aminoglycosteroids and aminosteroids from Holorrhena curtisii. *J. Nat. Prod*, *61*(11): 6-1332.
- Kassi, M., Kassi, A. K., Afghan, R., Rehman, P. and Kasi, M. 2008. Marring leishmaniasis the stigematization and the impact of cutaneous Nawab H., Hafiz A., Ehsanullah, Haider W, Khanani R. 1997. Visceral leishmaniasis in Karachi, *Pakistan. J. med Sci, 13*: 383-388.
- 15. Klaus, S. and Frankenburg, S. 1999. Cutneous leishmaniasis in the Middle East. *Clin. Dermatol*, *17*: 137141.
- Kolachi, H. B., Dahar, M. Y., Rathi, S. L. and Khaskheli, A. 2005. Epidemic of cutaneous leishmaniasis in Taluka johi, Dist, Dadu, Sindh. *Infect. Dis. J. Pak*, 37-40.
- Nawab, H., Hafiz, A., Ehsanullah, S., Haider, W. and Khanani, R. 1997. Visceral leishmaniasis in Karachi, *Pakistan. J. Med. Sci, 13*: 383-388.
- Oliveira PR, Dessein H, Romano A, Cabantous S, de Brito ME, Santoro F, et al. Il2ra genetic variants reduce il-2–dependent responses and aggravate human cutaneous leishmaniasis. J Immunol 2015; 194(6): 2664-72.
- 19. Reithinger R, Mohsen M, Leslie T. Risk factors for anthroponotic cutaneous leishmaniasis at the household level in Kabul, Afghanistan. PLoS Negl Trop Dis, 2010; 4(3): e639.
- Reithinger, R., Mohsen, M., Wahid, M., Bismullah, M., Quinnell, R. J., Davies, C. R., Kolaczinski, J., David, J. R. 2005. Efficacy of thermotherapy to treat cutaneous leishmaniasis caused by Leishmania tropica in Kabul, Afghanistan: a randomized, controlled trial. *Clin. Infect. Dis, 40*: 1148-1155.
- Schlein, Y., Jacobson, R. L. and Muller, G. C. 2001. Sandfly feeding on noxious plants: a potential method for the control of leishmaniasis. *Am. J. Trop. Med. Hyg*, 65(4): 300-3.
- 22. WHO. 1990. Control of the leishmaniasis. Technical report series 793. Geneva: WHO; 1990. EE.
- WHO. 1996. In: Manual on visceral leishmaniasiscontrol. WHO/LEISH/96.40. 4th edn. World Health Organization, Geneva, 1996.
- 24. WHO. 1998. *Leishmania* and HIV in gridlock. Geneva: World Health Organization. WHO/UNAIDS report, 15-25.
- 25. Wright. S. D., Detmers, P. A., Aida, Y., Adamowski, R., Anderson, D. C. 2. Chad, L. Kabbash. G. and Pabst, M. J. 1990. CD 18 deficient cells respond to lipopolysaccharlde in vitro. J. Immunol, 144: 2566.