

EPIDEMIOLOGICAL SURVEY OF BANCROFTIAN FILARIASIS IN TANGEDU (GUNTUR, AP)

V.VIVEKA VARDHANI* AND R. ADINARAYANA

Department of Zoology and Aquaculture

Acharya Nagarjuna University, Nagarjuna Nagar - 522 510, (A.P)

E-mail: vadlamudi-vv@yahoo.co.in

KEY WORDS

Epidemiology
Survey
Bancroftian filariasis

Received on :

28.12.2010

Accepted on :

27.03.2011

*Corresponding author

ABSTRACT

Filariasis is major public health illness in Tangedu (Guntur District) (Andhra Pradesh) (India) and 30-32% of the population is estimated to be at high risk. Inspite of repeated attempts to control filariasis, the disease transmission is perennial and persistent mostly transmitted by *Culex quinquefasciatus*. The transmission of lymphatic filariasis is correlated with Mf rate, disease rate, infection rate and infectivity rate and found that the disease transmission is very high in Tangedu.

INTRODUCTION

Human lymphatic filariasis results from infection with *Wuchereria bancrofti*, *Brugia malayi* and *B. timori*. *W. bancrofti* is prevalent in tropical Africa, South East Asia, Papua New Guinea, Philippines, Candelonia and Thailand. *B. malayi* is spread over South and South East Asia (WHO, 1992). *W. bancrofti* and *B. malayi* co-exist in many places and the mf of these species can be identified based on the measurements of various body characters (Raina et al., 1990; Rajendran et al., 1997); the identification of species (mf) is very important in epidemiological surveys. There are 120 million people harbouring lymphatic filariasis infection world wide (Shenoy, 2003). Lymphatic filariasis (LF) contributes a significant impediment to socioeconomic development in much of Asia, Africa and the western Pacific as well as in certain regions of the central and South America (Singh et al., 2003). LF has been a major public health problem in tropical and sub tropical countries next to malaria (Michael et al., 1996). Bancroftian filariasis is the most predominant infection contributing 99.4 % of the disease problem in India. Human bancroftian lymphatic filariasis (BF) can be differentiated into asymptomatic microfilariae, acute manifestations, and disease manifestations including hydrocoele and occult manifestations (Das et al., 1990; Harinath et al., 2000). The clinical symptoms of BF include filarial fever associated with lymphoedema with pain and lymphadipathy (Ravindran et al., 1994). The susceptibility/survival analysis of *Culex quinquefasciatus* infected with *W. bancrofti* showed that the parasite load in the mosquito is a risk factor for vector survival (Kumar et al., 1996; Kumar, 1997; Krishnamoorthy et al., 2004).

One of the essential requirements (for effective filarial control programme) is a precise and accurate diagnostic test for detection of the infection in individual cases as well as in mass scale in field surveys. The precise diagnosis of filariasis is presently based on the demonstration of microfilariae in night blood and clinical observations for acute or chronic manifestations of the disease. The adult female may produce mf at least for a period of five years and the life span of the mf has some influence on the dynamics of transmission of filariasis (Narasimham et al., 1984; Vanamail et al., 1990; Paily et al., 2009). The occult manifestations can not be diagnosed either by the method of night blood smear examination or by the symptomatology. It can be diagnosed by the method of immunodiagnostics – detection of filarial antibody and antigen by Enzyme Linked Immunosorbent Assay (ELISA). The study on survey of brancroftian filariasis in Tangedu will help in implementing the protection measures against the disease. This may also help in finding out the population of vector species in Tangedu (AP), India.

MATERIALS AND METHODS

Epidemiological survey was carried out in Tangedu which is the rural filarial endemic area of Palanadu (Guntur District) (A.P) (India). Tangedu is surrounded by hillocks and forest, located very nearer to river side belt of Krishna River. Night survey of blood smear collection was conducted during 20.00 to 24.00 hr following standard methods. During the survey, 318 houses were visited, 1636 population surveyed and 942 blood samples were collected. Blood smears were obtained by the finger prick method. A thick blood smear was prepared

on grease free microslide and allowed it to dry over night and stained in the Jaswanth Singh Bhattacharyaji stain - I and buffer solution. The stained blood samples were examined under the compound microscope for the presence of microfilarial parasites. The filariometric parasitic indices were calculated as given below:

$$\text{Mf rate} = \frac{\text{Total no. of persons found to harbour mf}}{\text{Total no. of blood samples examined}} \times 100$$

$$\text{Mean mf Density} = \frac{\text{Total no. of microfilariae}}{\text{Total no. of blood samples } +\text{ve for mf}}$$

$$\text{Disease rate} = \frac{\text{Total no. of persons showing signs and symptoms of filarial disease manifestations}}{\text{No. of persons examined for filarial disease}} \times 100$$

The lymphatic filarial cases were classified into acute clinical filariasis and chronic clinical filariasis (WHO, 1992).

RESULTS AND DISCUSSION

Positive cases: In the filarial night survey conducted in Tangedu, 942 persons were surveyed following the method of Rajendran et al., (1997) (Table 1). Among 942 cases, 29 were found to harbour microfilaria in their peripheral blood. Also, among these 942 cases, 36 and 8 were suffering from disease manifestation and occult filariasis respectively. Filariometric parasitic indices like mf rate, disease rate, endemicity rate and occult cases were found to be 3.07 %, 3.82 %, 6.90 % and 8 respectively.

Table 1: Epidemiological situation in Tangedu

Blood Samples collected	Mf cases detected	Mf rate (%) (positive cases)	Disease cases detected	Disease rate (%)	Endemicity rate(%)	Ocult cases Detected
942	29	3.07	36	3.82	6.90	8
Categories of microfilaria cases (24 males + 12 females)						
Acute cases		Chronic cases		Hydrocoel cases		
Male	Female	Male	Female	Male	Female	
05	08	06	04	13	NIL	
Vector density:						
C. q	Female Anopheline	Other species		Total mosquitoes collected		
2184	417	185		2805		
C.q-Culex quinquefasciatus						

Patho-physiological analysis of the clinical lymphatic filariasis

Out of 36 microfilarial cases detected (24 males, 12 females), 13 (5 males, 8 females) were acute cases, 10 (6 males, 4 females) were chronic cases and 13 (13 males only) were hydrocoel cases. Eight (5 males, 3 females) occult cases were evident. Lymphatic filariasis was found more in males (24 cases) than in females (12 cases) (Table 1).

Entomological evidences: A total of 2805 mosquitoes were collected in 280 households in Tangedu, out of which female *Culex quinquefasciatus* were 2184, female *Anophelines* 417 and other species 184.

Determination of vector man hour dsensity: It is determined by the calculation of vector mosquitoes in a stipulated time (15 minutes) in 10 houses in each multiplied by 10. The Man Hour Density recorded is 312.0 in Tangedu. The calculated

vector infection rate and vector infectivity rate are 1.23% and 0.44%. The identification of vector mosquito blood meal reveals the specification of the vector towards the host which influences the transmission potential of the disease in a community. The human blood index determined is 58.0%. The correlation of transmission potential of lymphatic filariasis with entomological parameters revealed the positive Mf rate (3.07%), disease rate (3.82%), vector 10 MHD (312.0%), infection rate (0.97%), infectivity rate (0.44%) and mosquito blood meal positivity for human blood (58.0%). These observations reveal that the transmission potential was found to be high in Tangedu.

Lymphatic filariasis continues to be a major cause of clinical morbidity with over one third of the world's population at the risk of infection. In India, inspite of National Filarial Control Programme (MFCP) being in operation for about five decades, the disease is showing an upward trend both in urban and rural areas of the countries. The topographical and ecological conditions of the filarial endemic area were favourable for vector breeding. Drainage system and low lying areas are promoting the high densities of vector population. The high vector density, vector infection rate, vector infectivity rate, human blood index, microfilarial rate and disease rate indicate the prevalence of high disease transmission in the communities of Tangedu. *Culex* and *Anopheline* mosquitoes were found as vectors of human lymphatic filariasis in the present study as suggested by WHO (1989). The entomological and parasitological parameters of the present studies confirm the correlation between the host, parasite and the poor sanitary conditions in the communities of Tangedu. The transmission of the disease was so high because of the non-availability of

accurate diagnosis of the disease and imperfect chemotherapeutic applications. The occurrence of 2184 *Culex* mosquitoes (out of 2805 mosquitoes) in the study area confirm the observations of Das (1976), Rajagopalan et al., (1977), Rajagopalan and Das (1987) and Paily et al., (2007) who reported *Culex quinquefasciatus* as vector of *W. bancrofti* in India. Majority of the people in the study area had no knowledge about the disease, its symptoms, and mode of transmission and method of prevention. The present study reveals the poor knowledge on the cause, prevention, control, transmission and treatment of filariasis. Hence, appropriate health education programmes may help to disease control in this rural area.

REFERENCES

Das, M. 1976. Vectors of filariasis with special reference to India. J.

- Commun Dis.* **8:** 101-109.
- Das, P. K., Manoharan, A., Srividya, A., Greenfell, B. T., Bundy, D. A. P. and Vanamail, P.** **1990.** Frequency distribution of *Wuchereria bancrofti* microfilariae in human populations and its relationships with age and sex. *Parasitology*. **101:** 429-434.
- Harinath, B. C., Reddy, M. V. R., Bhunia, B., Bandari, Y. P., Mehta, V. K., Chatthurvedi, P., Prajapati, N. C. and Gupta, R. K. C.** **2000.** *J. Clinical Biochemist*. **15:** 118-126.
- Krishnamoorthy, K., Subramanyan, S., Van Oortmarsen, G. J., Habbema, J. D. F and Das, P. K.** **2004** Vector survival and parasite infection: the effect of *Wuchereria bancrofti* on its vector *Culex quinquefasciatus*. *Parasitology*. **129:** 43-50.
- Kumar, A.** **1997.** Epidemiology of filariasis transmission: estimation of infectivity index in vector population from mF data. *J. Parasit. Dis.* **21:** 182-184.
- Kumar, A., Mansing, G. D. and Jangid, P. K.** **1996.** Correlation between human microfilaremia and vector infection in a bancroftian endemic area, Khurda District. *Indian Tropical Biomedicine*. **13:** 5-7.
- Michael, E., Bundy, D. A. P. and Grenfell, B.** **1996.** Reassessing the global prevalence and distribution of lymphatic filariasis. *Parasitology*. **112:** 409-428.
- Narasimham, M. V., Rao P. K., Venkatanarayana M., Viswanathanam, P and Rao C. K.** **1984.** Longevity of *Wuchereria bancrofti* microfilaria. *J. Commun. Dis.* **16:** 326-327.
- Paily, K. P., Agiesh Kumar, B. and Balaraman, K.** **2007.** Transferrin in the mosquito, *Culex quinquefasciatus* Say (Diptera: Culicidae), up-regulated upon infection and development of the filarial parasite, *Wuchereria bancrofti* (Cobbold) (Spirurida:Onchocercidae). *Parasitol. Res.* **101:** 325-330.
- Paily, K. P., Hoti, S. L. and Das, P. K.** **2009.** A review of the complexity of biology of lymphatic filarial parasites. *J. Parasit. Dis.* **33(1&2):** 3-12.
- Raina, V. K., Joshi, M. C., Singh, S., Joshi, R. D., Bhattacharjee, K., Kumar, A. and Verghese, T.** **1990.** Epidemiology of *Brugia malayi* infection and its co-existence with *Wuchereria bancrofti* in and around Sillaberia PHC, District Midnapur, West Bengal. *J. Commun. Dis.* **22:** 205-208.
- Rajagopalan, P. K., Kazmi, S. J. and Mani, T. R.** **1977.** Some aspects of transmission of *Wuchereria bancrofti* and ecology of *Culex pipiens fatigans* in Pondicherry. *Indian J. Med Res.* **66:** 200-207.
- Rajagopalan, P. K. and Das, P. K.** **1987.** The Pondicherry Project on Integrated Disease vector Control (Filariasis Control Demonstration Project) 1981-1986. Vector Control Research Centre, Pondicherry. p. 164.
- Rajendran, G., Panicker, K. N., Krishnamoorthy, K., Sabesan, S. and Snehalatha, Radhakrishnan R.** **1997.** Current status of filariasis in Chavakkad taluk, Trichur district, Kerala. *J. Commun. Dis.* **29:** 333-343.
- Ravindran, B., Satapathy, A. K. and Sahoo, P. K.** **1994.** Bancroftian filariasis- differential reactivity of antisheath antibodies in microfilariae carriers. *Parasitic Immunol.* **16:** 321-324.
- Shenoy, R.** **2003.** Lymphatic filariasis in children. *The Ind. J. Lymphology*. **1:** 7-13.
- Singh, U., Ruma Gupta., Chatterjee, R. K. and Shailja Mishra-Bhattacharya.** **2003.** Characterization of geographical isolates of *Wuchereria bancrofti* from six states of India. *J. Parasitic Dis.* **27:** 21-28.
- Vanamail, P., Subramanian, S., Das, P. K., Pani, S. P. and Rajagopalan, P. K.** **1990.** Estimation of fecundic life span of *Wuchereria bancrofti* from longitudinal study of human infection in an endemic area of Pondicherry (South India). *Indian J. Med. Res.* **91:** 293-297.
- World Health Organization.** **1989.** Geographical distribution of arthropod borne diseases and their principal vectors. WHO/VBC/89. **967:** 134-139.
- World Health Organization.** **1992.** Lymphatic filariasis. Fifth report of the WHO expert committee on filariasis, Geneva. WHO Technical Report Series No. 821.

