Serum Levels of Interleukin-1β and Interferon-γ in Hansen’s disease

Amul K Butti1, Shakti Kumar Yadav1, Alekh Verma1, V K Khurana2, Sanjay Jain3, Sompal Singh1 and Namrata Sarin1

1Department of Pathology, North Delhi Municipal Corporation Medical College and Hindu Rao Hospital, 110007, India; 2Department of Dermatology, North Delhi Municipal Corporation Medical College and Hindu Rao Hospital, 110007, India; 3Department of Microbiology, North Delhi Municipal Corporation Medical College and Hindu Rao Hospital, 110007, India

Corresponding author: Dr Namrata Sarin, Department of Pathology, North Delhi Municipal Corporation Medical College and Hindu Rao Hospital, 110007, India, Tel: 91123905652, E-mail: drnamratasarin@gmail.com

Abstract

Background: Hansen’s disease is a chronic infectious granulomatous disease caused by Mycobacterium leprae. In leprosy the final immune response depends on predominant effector pathway, M. leprae bacilli are presented by antigen presenting cells to CD4+ T cells with subsequent release of Interleukin -12 leading to TH1 response. Release of interferon gamma (IF-γ) promotes further TH1 response development. IF-γ along with tumor necrosis factor and IL-2 leads to recruitment of monocytes and their transformation into epithelioid cells and granuloma formation. Aims: Our aim of the study was to analyse serum levels of interleukin-1β (IL-1β) and interferon-γ in newly diagnosed untreated patients of leprosy and to co-relate their levels with clinico-histopathological profile. Methods: (including settings and design and statistical analysis used). The study was an analytical cross-sectional study. Fifty newly diagnosed leprosy patients were included in the study. They were categorized into paucibacillary (30) and multibacillary (20) groups. Serum cytokines assay for IL-1β and IFN-γ were performed by ELISA and were compared between the two groups. Mean serum cytokine levels of IL-1β and IFN-γ in both paucibacillary and multibacillary groups were calculated. Significance of mean difference between IL-1β and IFN-γ was assessed using Mann Whitney U test between the two groups. Results: Significantly high serum levels of IFN-γ were seen in paucibacillary group compared to multibacillary group. Although serum levels of IL-1β were raised in multibacillary group compared to paucibacillary group. The mean difference was not statistically significant. Limitations: The limitation of present study is low sample size. Conclusions: Serum interleukin levels (IL-1β and IFN-γ) are different among the paucibacillary and multibacillary cases. The fact is useful not only to understand the disease pathogenesis but also in sub-categorization of Hansen’s disease patients.

Keywords: Leprosy; Hansen’s disease; Cytokines; Interferon-γ; Interleukin-1β

Introduction

Hansen’s disease is a chronic infectious granulomatous disease caused by Mycobacterium leprae, which is an obligate intracellular pathogen. [1] Prevalence of leprosy in India by the end of 2016 was 0.66/10,000 population. Based on degree and type of immune response, leprosy is classified into tuberculoid (TT), borderline tuberculoid (BT), borderline lepromatous (BB), lepromatous leprosy (LL). [2] There can be downgrading of immunity towards lepromatous pole, and upgrading towards the tuberculoid pole may occur within central groups (BT, BB, BL) but the two polar groups (TT, LL) are stable. [3]

Cytokines are glycoproteins with low molecular weight produced by immune as well as non-immune cells which acts as molecular signals for communication between cells in immune system. [4] In leprosy the final immune response depends on predominant effector pathway, M. leprae bacilli are presented by antigen presenting cells (APC- Dendritic cells and Macrophages) to CD4+ T cells with subsequent release of Interleukin -12 (IL-12) leading to TH1 response. [5]

Release of interferon gamma (IF-γ) promotes further TH1 response development. IF-γ along with tumor necrosis factor (TNF) and IL-2 leads to recruitment of monocytes and their transformation into epithelioid cells and granuloma formation. [6] Onset of lepromatous leprosy (LL) correlates with cytokines produced by TH2 cells on the background of patient’s poor cell mediated immunity (CMI) i.e., increased IL-4, IL-5, IL-10 and deficient IL-2 and IF-γ. On the other hand, tuberculoid leprosy (TT) is associated with a TH1 response i.e., increased IL-2, IL-12, IL-18, IFN-γ, granulocyte monocyte colony stimulating factor (GM-CSF) and TNFα. It has also been shown that in leprosy patients IFN-γ and TNFα are immunoprotective whereas IL-10, IL-4, IL-5 and IL-1β are immunosuppressive. [7]

The strong immune responses result in recruitment of Langerhans cells to the site skin lesions in tuberculoid leprosy. On the other hand, the TH2 helper cells counteract by inhibiting activity of TH1 helper cells and down-regulating the activation of Langerhans cells in Lepromatous leprosy. [8] Very few studies have correlated levels of serum cytokines with clinical and histopathological findings. [9]

Methods

A total of 50 newly diagnosed cases of leprosy attending the dermatology outpatient department (OPD) were included in this study. Patients already undergoing treatment or having comorbid conditions were excluded. Samples were collected before the start of drug treatment. Consent was taken from the patient, followed by detailed clinical history and physical examination. Diagnosis of leprosy was...
done clinicopathologically and for patients towards the lepromatous pole Wade-Fite staining was done. Cases were classified into two broad categories based on bacillary index as multibacillary and paucibacillary groups. The cases comprised of 20 multibacillary [BB, BL and LL, Figure 1] and 30 paucibacillary [BT and TT, Figure 2].

Skin biopsy was taken in the minor O.T and then tissue is processed and stained with H&E as well as Wade Fite’s stain. Based on histopathology, they were classified into BT, TT, BB, BL and LL. Serum was collected from the patient for the measurement of cytokine level (IL-1β and IFN-γ) through ELISA (Boster, USA). Serum sample collection was done in the skin OPD at the time of skin biopsy. Sample was collected in plain vacutainer and left undisturbed in water bath at 37°C for 1/2 hr. Serum was separated by centrifugation of vacutainer for 5 min at 4000 rpm. Serum was stored at − 40°C in deep freezer and thawed before analysis.

Data of age was classified. Mean and standard deviation was calculated for numerical data. Significance of mean difference of age in between the two groups was assessed using Student - t test. Categorical data between two different categories was compared using Pearson chi-square test. Serum cytokine levels of IL-1β and IFN-γ were measured in pg/ml in individual samples and used for analysis. Mann-Whitney U test was used to compare numerical results from ELISA between two different categories. P-value below 0.05 was regarded as significant.

**Results**

Majority of the cases (38%) were in the age group of 21-30 years [Figure 3]. Among paucibacillary patients mean age was 29.23 years with a standard deviation of 13.48 and for multibacillary patients mean age was 36.55 years with standard deviation of 11.26 with difference between mean age of two groups being statistically insignificant (p-value=0.051, Student t- test) [Table 1]. 60.00% cases were males with M-F ratio of 1.5: 1 [Table 2] and difference in M-F percentage between multibacillary and paucibacillary group was not statistically significant (p-value=0.239, Pearson chi-square test).

Significantly high serum levels of IFN-γ were seen in paucibacillary patients (mean=59.39 ± 204.74 pg/ml) as compared to multibacillary patients (mean=53.00 ± 194.54 pg/ml) [Table 3 and Figure 4] and the mean difference was statistically significant (p-value=0.044, Mann-Whitney U test). Although serum IL-1β levels were higher in multibacillary patients (mean=81.77 ± 103.43 pg/ml) as compared to paucibacillary patients (mean=73.77± 98.69 pg/ml) [Table 3 and Figure 4] and the mean difference was statistically significant (p-value=0.044, Mann-Whitney U test). Although serum IL-1β levels were higher in multibacillary patients (mean=81.77 ± 103.43 pg/ml) as compared to paucibacillary patients (mean=73.77± 98.69 pg/ml) [Table 3 and Figure 4].
Transmission showed that during the periods of greatest transmission, in Mexico and in Korea during successive periods of decline in ratio of 1.8:1. A retrospective study conducted by Veeresh et al. (242, 65.05%) and females were 130 (34.95%) with a male to female ratio of 1.3:1. Study conducted by Radhan et al. [13] out of 60 cases studied, majority of them 48 (42.9%) were females with a male to female ratio of 1.3:1. Study at ESIC Medical college, Kalburgi during the period of 2013-2015, of workers. [12-15] In our study the mean age of multibacillary patients is higher than the paucibacillary group. Similar findings were noted by Nobre et al. [14] in Brazil where both the general new case detection rate and the detection of multibacillary cases were higher among 60–79-year-old. The percentage of new cases which were classified as multibacillary progressively increased and reached 67.7% in 60 or more years of age group (180/266). These changes can be attributed to the longer incubation period of multibacillary leprosy that usually recurs in people who were previously infected. [15-19] Recent studies in Mexico and in Korea during successive periods of decline in transmission showed that during the periods of greatest transmission, peak detection already occurred among the elderly. [20,21]

Out of 50 cases included in the present study, majority of the cases (60%) were males with a male to female ratio of 1.5:1. Study done by Moorothy et al. [14] out of 372 patients, majority of patients were males (242, 65.05%) and females were 130 (34.95%) with a male to female ratio of 1:8.1. A retrospective study conducted by Veeresh et al. [22] at ESIC Medical college, Kalburgi during the period of 2013-2015, of the 112 patients studied, majority of them (64, 57.1%) were males and 48 (42.9%) were females with a male to female ratio of 1:3.1. Study conducted by Radhan et al. [13] out of 60 cases studied, majority of them (43, 71.7%) were males and 17 (28.3%) were females with a male to female ratio of 2.52:1. In majority of the studies conducted, the disease burden is more in males than females may be because most of the leprosy workers are males; the examination of women is less complete and less satisfactory particularly in certain cultural situations resulting in under detection of leprosy among females.

Immunity has a major role in the pathogenesis of leprosy. At the tuberculoid pole, TH1 cells are activated leading to a strong CMI response to M. leprae (IL-2, IFN-γ, TNF-α and IL-12 are the predominantly expressed cytokines). [23-28] At the lepromatous pole, TH2 cells are activated which inhibits TH1 response leading to a defective CMI response to M. leprae. IL-10, IL-1β, IL-4, IL-5 are the key cytokines of TH2 response. [16-19]

In the present study, the mean value of IFN-γ is 59.39 pg/ml in 30 patients of paucibacillary group and with standard deviation of 204.74, and in 20 patients of multibacillary group it was 53.00 pg/ml with a standard deviation of 194.54. The difference in IFN-γ was statistically significant (P-value=0.044). Present study showed that paucibacillary leprosy patients coincides with a rise of TH1 cytokines, i.e., IFN-γ.

In the present study, the mean value of IL-1β in 20 patients of multibacillary group was 81.77 pg/ml with a standard deviation of 103.43 and in 30 patients of paucibacillary was 73.77 pg/ml with a standard deviation of 98.69. This difference IL-1β was not statistically significant (P-value=0.259) and thus in the present study the relationship between role of IL-1β in multibacillary leprosy could not be established.

Iyer et al. [31] studied levels of a panel of cytokines i.e., IL-4, IL-6, IL-10 via ELISA technique. IFN-γ, TNF-α, IL-6 peak levels were detected in borderline tuberculoid lesions whereas significantly lower levels were seen in borderline lepromatous and lepromatous leprosy patients.

Parida et al. [32] analyzed serum TNF-α and IL-1β levels in 64 patients spread over the spectrum of leprosy. Elevated levels of TNF (15 to 4500 pg/ml) were detected in lepromatous leprosy cases and low levels (15 to 160 pg/ml) were detected in tuberculoid forms of leprosy. IL-1β levels were found to be significantly higher in Lepromatous Leprosy cases in comparison to other groups or normal controls.

Moubasher et al. [33] who monitored the serum levels of IFN-γ, IL-2, IL-2R, IL-10, TNF-α and IL-1β by ELISA in 55 untreated leprosy patients and 35 reactional leprosy patients in addition to 20 age and sex matched healthy controls. In their study all leprosy patients showed significantly higher serum levels of IFN-γ, IL-2R, IL-10, TNF-α and IL-1β as compared to controls except for IL-2 (all types of Leprosy). Lepromatous leprosy patients showed significantly higher levels of IL-2R, IL-10 and IL-1β. Tuberculoid leprosy patients showed significantly higher serum IFN-γ and TNF-α.

Another study done by Madan et al. [34] studied serum cytokines profile in leprosy found that levels of TNFα, IFNγ were significantly raised in paucibacillary cases and whereas all the other studies cytokines like IL-1β and IL-10 were raised in multibacillary cases, with rise being statistically significant. 34 The above mentioned studies were conducted between healthy controls and multibacillary and paucibacillary patients.

The results of this study may be limited by the small sample size, however, a larger prospective study can be done to identify the role of IL-1β and IFN-γ in pathogenesis of leprosy and categorization of leprosy patients.

In conclusion, serum interleukin levels (IL-1β and IFN-γ) are different among the paucibacillary and multibacillary cases. The fact is useful not only to understand the disease pathogenesis but also in sub categorization of Hansen’s disease patients.

Conflict of Interest
The authors disclose that they have no conflicts of interest.

References


