**ABSTRACT**

**Introduction:** Tuberculosis (TB) is a common and a deadly infectious disease which is caused by Mycobacterium tuberculosis. Inflammatory cytokines play an important role during the course of the disease and they may be responsible for tissue damage which is caused by lipid peroxidation.

**Method:** The present study was conducted in the Department of Biochemistry and patients were selected from Department of TB and Chest Sri Aurobindo Institute of Medical Sciences and also from Manorama Raje Tuberculosis Hospital, Indore. 32 healthy controls and 35 pulmonary TB patients were compared initially for their serum Malondialdehyde (MDA) and tumour necrosis factor alpha (TNF α) levels. Serum TNF α and MDA levels were correlated.

**Result:** TNF α and MDA levels in serum were significantly increased (p<0.001) in pulmonary tuberculosis patients as compared to those of controls. Increased Serum TNF α was positively correlated to MDA levels and it was found to be statistically not significant (correlation coefficient r =0.282, p>0.05).

**Conclusion:** The present study supports the view that there may be a link between lipid peroxidation and cytokine response and relative roles of cytokines and lipid peroxidation in the pathogenesis of tuberculosis.

**Key words:** Cytokine, Lipid peroxidation, Tuberculosis

**INTRODUCTION**

Tuberculosis is a common and a deadly infectious disease which is caused by mycobacteria, mainly by Mycobacterium tuberculosis. Globally, there were an estimated 9.27 million incidences of TB [1]. One-third of the world’s current population has been infected by TB, and new infections occur at a rate of one per second. The precise clinical manifestations of tuberculosis (TB) are likely to result from a complex interaction between the host and the pathogen. Cytokines are primarily involved in host response to disease or infection [2]. TNF α is a monocyte-activating cytokine which stimulates antimycobacterial activity and helps in maintaining the integrity of the tuberculous granulomas in which Mycobacterium tuberculosis is contained [3]. TNF α is believed to play multiple roles in the immune and pathological responses in tuberculosis [4]. Serum TNF α measurement might play an important role in the evaluation of the inflammatory phenomena in TB.

During pulmonary inflammation, increased amounts of reactive oxygen species (ROS) and reactive nitrogen intermediates (RNI) are produced as a consequence of a phagocytic respiratory burst [5]. These ROS and RNI induce lipid peroxidation (LP), a general mechanism of tissue damage which is caused by free radicals, that is known to be responsible for cell damage and it may induce many pathological events [6]. Inflammatory cytokines play an important role during the course of the disease and they may be responsible for tissue damage which is caused by lipid peroxidation [7]. Exposure to Mycobacterium leads to cytokine production by macrophages. These cells increase free radical release as a part of bactericidal action, which may, in turn, enhance cytokine secretion, thus creating a positive feedback loop. The present study was carried out to study levels of lipid peroxidation and serum cytokines in tuberculosis patients and to correlate lipid peroxidation and serum cytokine levels.

**MATERIAL AND METHODS**

**Subject:** All patients were recruited from the TB and Chest Department of Shri Aurobindo Institute of Medical Sciences and also from Manorama Raje Tuberculosis Hospital, Indore. Thirty-five patients (20 males and 15 females) with pulmonary TB (PTB) and 32 healthy controls (17 males and 15 females) participated in the study. Their mean age was 31.62 ± 6.37 (range 18-55 years). On their entry into the study, all patients of pulmonary tuberculosis who were found to be new sputum positive, and belonged to category I (two sputum specimens were positive for acid-fast bacilli by direct microscopy or one sputum specimen was positive for culture) were included. No extrapulmonary involvement was found in any of the patients. Those with diabetes mellitus, pregnancy or immunological or autoimmune diseases other than tuberculosis and subjects with a history of smoking were excluded from the study. None of the subjects had any serological evidence of HIV infection. All patients were administered anti-TB therapy according to standard antitubercular regimen. A control group of 32 healthy volunteer subjects (17 women and 15 men) with a mean age of 27.74 ± 7.30 (range18-55 years) was also studied. Consent to participate in the study was obtained from each individual and the study protocol was approved by the institutional and human ethical committees of Sri Aurobindo Institute of Medical Sciences, Indore.

**Blood collection:** Fasting blood samples (5 ml) were collected by antecubital venipuncture in plain tubes and they were left to clot. They were then centrifuged at 1000 rpm for 10 min. The sera were then aliquoted and stored until they were used for assay of parameters. Blood samples were collected from all controls and from the PTB patients.

**Estimation of serum TNF alpha:** Serum TNF alpha was estimated by using a commercial immunossay kit which was available with DIACLONE United Kingdom. The kit was used according to the manufacturer’s instructions. The concentrations of TNF alpha in the samples were determined by using a standard curve and the results were expressed as pg/ml. [Normal range = < 8pg / ml]

**Estimation of serum MDA:** Estimation of MDA in the serum was done by thiobarbituric acid method of Wilbur KM et al., [8]. Thiobarbituric acid reacts with serum malondialdehyde which is produced by hydrolysis of lipid hydroperoxides to form a pink

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**Cytokine in Tuberculosis Patients**

**A Study of Serum Malondialdehyde and Cytokine in Tuberculosis Patients**

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red colour complex with high absorbance that can be measured spectrophotometrically at 532 nm. (The results were expressed as nmoles/ml.) This complex is usually quantified against MDA standards which are generated from 1, 1, 3, 3 tetraethoxypropane under the same reaction conditions [Normal range = 1-3.42 nmo/ ml].

### STATISTICAL ANALYSIS

SPSS, version 10 was used for statistical assessments, to evaluate mean levels of variables between study groups and healthy controls by using Unpaired t-test. Correlations were calculated by Pearson’s correlation coefficients (two-tailed). A p value of ≤ 0.05 was used as a threshold of significance.

### RESULTS

There was no significant difference between mean ages of controls and pulmonary tuberculosis patients (p>0.05). The serum levels of TNF alpha and MDA were highly significant in pulmonary tuberculosis patients as compared to those of controls (p<0.001). Among 35 patients, 28 were found to be having increased levels of MDA (ranges 2.90-7.01 nmo/ml) and TNF levels were found to be increased in all patients (range 12-73pg/ml) [Table/Fig-1].

Increased Serum TNF α was positively correlated with MDA levels and this was found to be statistically not significant (r =0.282, p>0.05) [Table/Fig-2].

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Tuberculosis patients</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>31.62 ± 6.37</td>
<td>29.37 ± 8.2</td>
<td>p&gt;0.05</td>
</tr>
<tr>
<td>Serum MDA nmo/ml</td>
<td>2.1 ± 0.82</td>
<td>5.04 ± 1.15</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Serum TNF α Pg/ml</td>
<td>14.59 ± 8.88</td>
<td>53.14 ± 19.06</td>
<td>p&lt;0.001</td>
</tr>
</tbody>
</table>

[Table/Fig-1]: Comparison: Mean Age & Serum Levels of TNF α & MDA in & Pulmonary Tuberculosis Patients Control Group.

![Graph showing correlation between TNF alpha & MDA](image)

In the present study, a positive correlation was observed between serum TNF–α and MDA levels, which proposed that increased TNF–α production by activated macrophages could lead to an enhanced production of oxygen radicals and lipid peroxidation. However, the correlation between TNF–α and MDA was not significant, which suggested that inflammation and lipid peroxidation were not parallel occurrences in this study.

The present study supports the fact that there is a link between lipid peroxidation and cytokine response and the relative role of cytokines and lipid peroxidation in the pathogenesis of tuberculosis. Also, serum TNF α and MDA measurements may play an important role in the evaluation of the inflammatory phenomena in TB.

### REFERENCES

1. World Health Organization (WHO). Global Tuberculosis Control; Epidemiology, strategy, financing. 2009.
7. Cern EH, Yusuf TR, Mustafa Cekmen. Serum levels of TNF-a, sIL-2R, IL-6, and IL-8 are increased and associated with elevated lipoxidation in patients with Behcet’s disease. Mediators of Inflammation. 2002; 11 : 87–93.
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[18] Yuan-Chun Liao. IL-19 Induces Production of IL-6 and TNF and Results in Cell Apoptosis through TNF. The Journal of Immunology 2002; 169: 4288-4297.


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