Correlation of Serum Urate Levels and Lichen Planus in Male and Female Patients

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Authors’ contributions
This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

ABSTRACT

Aim: To correlate serum urate level in male and female patients with lichen planus (LP) presenting at Dermatology department, Liaquat University Hospital Jamshoro.
Methodology: This Cross-sectional study was conducted from 2019 to 2020. Total 126 patients, divided into two equal groups male (n=63) and female (n=63), between the age of 20-50 years and duration of disease from one month after eruption of lesion to 18 months were incorporated in this study. The patients having gout, obesity, chronic kidney disease, Pregnant ladies, smoker and those taking serum urate lowering drugs or steroids were excluded from this study. After taking history with clinical examination, patients were subject to height, weight, BMI, and duration of disease in months was stated.
Results: Total 126 patients with age range between 20-50 years and the mean age was 32.5 ± 4.47 years in male and 35.3±6.74 years in females were analyzed. The mean BMI was 23.6 ± 2.68 kg/m² in male and 22.5 ± 3.47 in kg/m² females were recorded. The mean duration of disease was 35.4 ± 16.92 and 37.12 ± 18.92 weeks were recorded in male and female patients.
respectively. The mean serum urate level in Idiopathic Lichen Planus patients was 3.82 ± 0.79 mg/dL in male and 2.3+0.43 mg/dL in female were observed. **Conclusion:** Our study results indicate that there is a positive correlation between LP and serum urate levels in male and female patients.

Keywords: Serum uric acid; lichen planus; BMI.

1. INTRODUCTION

Lichen planus (LP) is seen in approximately 1% of all new patients visiting health care clinics. There is no considerable racial, geological and predisposition noted for LP. In most of the patients age ranges from 30-60 years; however, it can appear at any age [1]. The exact etiology is unknown but is likely an interaction between autoimmunity and environmental triggers. Environmental factors comprises medication, viruses (particularly hepatitis C), allergic reactions to a foreign substance including metallic dental materials and stress also contributing in the development of LP [2].

The typical lesions of lichen planus are described by 6 ps: Polygonal, Purple, pruritic, papules, planar and/or plaque [3]. It mostly involves wrist and ankles but can be anywhere and even generalized. Mucosal LP usually effects buccal mucosa or vulva. Cutaneous lesions of LP are not associated with risk of skin cancer, but erosive LP involving oral mucosa particularly in men and vulvar lesions in women, do carry a small risk of malignant transformation [4].

In humans, uric acid makes about half of the antioxidant capacity of blood plasma [5]. Uric acid possesses strong antioxidant and scavenging activity for reactive oxygen species (ROS) and peroxynitrite. High concentration of uric acid are found in the cytoplasm of human and mammalian cells, especially in the hepatocytes [6], vascular endothelial cells, and in nasal secretions, where it acts as an antioxidant [5]. In fact, uric acid may exert a title role in healing of tissues by initiating the inflammatory response that is required for tissue repair, hunting oxygen free radicals, and activating progenitor endothelial cells [7].

Low serum urate levels, which ultimately lead to a decrease in antioxidants, were found in patients having multiple sclerosis. Peroxynitrites and ROS are supposed to be liable for myelin sheath degeneration in multiple sclerosis (MS) which can be blocked by achieving high levels of serum urate. In support of this view, gout patients almost never developed multiple sclerosis (MS) [8]. There are a number of reports documenting association of low serum uric acid levels with MS [8,9] and other degenerative neurological disorders [10], i.e. Parkinson [11], and Alzheimer [12] disease. Similarly, Pemphigus vulgaris, (an autoimmune blistering disorder involving skin and mucous membranes), and lichen planus (an autoimmune inflammatory disease involving skin, mucosa, even hair and nails) [13], were also found to be associated with low urate levels in serum and saliva [14].

2. MATERIALS AND METHODS

This Cross-sectional study was conducted from 2019 to 2020. Total 126 patients, divided into two equal male (n=63) and female (n=63) groups, between the age of 20 to 50 years and duration of disease from one week after eruption of lesion to 18 months were included in this study. The patients on steroid or immune-suppression drugs or NSAIDs for last one month, gouty, obesity (BMI> 27), chronic kidney disease, Pregnant ladies, smoker for >1 year and smoking more than 1 pack per day, and those taking serum urate lowering drugs were excluded from this study. After taking detailed history with full clinical examination, patients were subject to relevant investigations i.e. the patient’s height, weight, BMI, and duration of disease in months was stated. The data was analyzed by using Statistical Package for Social Sciences (SPSS) Version.22.

3. RESULTS

Total 126 patients with age range between 20 to 50 years and the mean age was 32.5 ± 4.47 years in male and 35.3±6.74 years in females were analyzed. The mean BMI was 23.6± 2.68 kg/m² in male and 22.5 ± 3.47 in kg/m² females were recorded. The mean duration of disease was 35.4 ± 16.92 and 37.12± 18.92 weeks were recorded in male and female respectively. The mean serum urate level in Idiopathic Lichen Planus patients was 3.82 ± 0.79 mg/dL in male and 2.3±0.43 mg/dL in female were observed. When the outcome variable was stratified with
In respect to age and duration of disease, significant difference was observed, shown in Table 1. Similarly, when the outcome variable was stratified with respect to BMI and gender, no significant difference was observed, shown in Table 2.

4. DISCUSSION

Lichen planus is a chronic inflammatory disease. The skin has a number of defense mechanisms to prevent their deleterious effect, this interacts with ROS [13]. Serum urate is a natural end product for purine metabolism and is made in mammalian systems. There are evidence that higher serum urate levels are a risk factor for cardiovascular disease where oxidative stress plays a major role in pathophysiology. This pollutes free radicals by inhibiting endothelial function under oxidative stress conditions inside a cell that discharges glutathione. There is growing evidence that the serum urate has an in vitro impact as antioxidant and antioxidant plasma potential increases with serum urate administration [15].

Our analysis showed a significant reduction in serum serum urate levels in patients, i.e. 4.32±0.79 mg/dL. This study was related to Chakraborti et al, medium serum urate levels, 3.6 mg/dL in patients and 3.94 mg/dL controls [16]. The mean difference is 0.34 mg/dl. There was also a significant decrease in serum urate levels in the sample of Italian LP patients [17]. On the contrary, Israel's report found that hyperuricemia was more common than the general population, although LP was not found to be a source of overproduction of uric acid. Saawarnet al stated that oxidative stress can play a role in oral LP, meanwhile, the strong antioxidant lycopene was found to be effective in another study in the oral LP's management. This therapeutic effect shows indirectly oxidative stress's function in LP pathogenesis [18].

### Table 1. Mean serum uric acid level stratification in age-related patients with LP in male and female

| Serum uric acid level (Mean ± SD) | Age | P-value |
|----------------------------------|-----|---------|
|                                 | 20-35 (n=36) | 35-50 (n=27) |
| Male                            | 3.59 ±0.1   | 3.01±0.07   | < 0.001 |
| Female                          | 1.9±0.43    | 2.4±0.03    | < 0.001 |

### Table 2. Mean serum uric acid level stratification in gender-specific patients with LP

| Serum uric acid level (Mean ± SD) | Gender | P-value |
|----------------------------------|--------|---------|
|                                  | Male   | Female (n=27) |
|                                  | 3.82 ± 0.79 | 2.3±0.43 | < 0.001 |

### Table 3. Stratification of the level of mean serum uric acid in weeks of disease duration in patients with idiopathic lichen planus

| Serum uric acid level | Duration of disease (weeks) | P-value |
|-----------------------|-----------------------------|---------|
|                       | 1-40 (weeks)                | 40-72 (weeks) |
| Male                  | 3.94±0.1                    | 3.68 ± 0.2     | < 0.001 |
| Female                | 2.23±0.1                    | 2.06±0.3       | < 0.001 |

### Table 4. Stratification of the level of mean serum uric acid in idiopathic lichen planus patients with respect to BMI

| Serum uric acid level | BMI | P-value |
|-----------------------|-----|---------|
|                       | 18-25 | >25 to ≤27 |
| Male                  | 3.46±0.2  | 3.87±0.4     | < 0.001 |
| Female                | 2.28±0.3  | 2.46±0.5     | < 0.001 |
This study demonstrates a significant correlation with serum urate levels between age of the patient and duration of the disease. However, no significant gender and BMI relationship with serum urate was identified. Compared to other studies, there is no significant correlation between age and gender with serum urate in one study. Although a significant association between serum urate and disease period was found similar to this research [16].

Indirect evidence of increased oxidative stress in LP is confirmed by the fact that saliva serum urate is decreased in oral LP patients [19]. In addition, vitamin E and C levels in LP are decreased and supplementation of these may play a role in the management of LP [20].

Serum urate is a potent free radical scavenger and, using two methodologically distinct assays, it has been demonstrated that systemic administration of serum urate improves ex vivo serum-free radical scavenging to a significantly greater extent than vitamin C, another effective aqueous physiological antioxidant. Administering antioxidants also raises serum urate levels [21,22,19].

5. CONCLUSION

Our study results show that LP can be correlated with serum urate depletion in male and female patients. Serum urate can be considered as a useful antioxidant biomarker in LP for the production and monitoring of treatment strategies.

CONSENT

All authors declare that written informed consent was obtained from the patient.

ETHICAL APPROVAL

Ethical approval was taken prior to conducting study from ethical review committee of CPSP.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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