Evaluation of Neutrophil-to-Lymphocyte Ratio, Platelet-to-Lymphocyte Ratio, and C-Reactive Protein in Tension-Type Headache Patients

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Abstract

Objectives A tension headache is the most common type of headache, and its causes are multifactorial. A relationship has been shown between migraine headaches and neutrophil-to-lymphocyte ratio (NLR), platelet-to-lymphocyte ratio (PLR), and C-reactive protein (CRP). In this study, we investigated the NLR, PLR, and serum CRP levels in frequent episodic tension-type headache (FETTH) and chronic tension-type headache (CTTH) patients.

Materials and Methods This retrospective study included 64 patients with FETTH, 80 patients with CTTH, and 60 healthy controls who were followed up in the neurology clinic. Hematological parameters were compared between the patient and control groups.

Results In CTTH patients, platelets, NLR, PLR, and CRP values were statistically higher than in FETTH patients and patients in the control group. In FETTH patients, the PLR value was higher than in patients in the control group, but there was no statistically significant difference in NLR and CRP values between FETTH patients and patients in the control group. Also, there was no correlation between these values and age and gender.

Conclusion Increase platelet count might have an effect on tension-type headache pathophysiology. Systemic inflammation parameters were shown to be significantly higher in CTTH patients. More comprehensive studies are needed to evaluate the effect of systemic inflammation on the chronicity of tension headaches.

Keywords ► neutrophil-to-lymphocyte ratio ► platelet-to-lymphocyte ratio ► tension-type headache

Introduction

Tension-type headache (TTH) is a common primary headache type whose causes are multifactorial. Genetic and environmental factors play a critical role.1 There are episodic and chronic forms of TTH, and there are multiple distinctions between forms. Episodic-type headaches often evolve into chronic-type headaches over time.2 Previous studies have reported that peripheral and central sensitization mechanisms may play a role in the chronification of headaches.3 The effects of local and systemic inflammation on tension headaches are not clearly known.

Recent studies have demonstrated that neutrophil-to-lymphocyte ratio (NLR) and platelet-to-lymphocyte ratio...
In the preceding 72 hours were excluded. Headache at the time of examination or had used analgesics, local or systemic infection. In addition, patients who had a diabetes mellitus, malignant hypertension, trauma, malignancies, such as migraine, pregnancy, autoimmune diseases, exclusion criteria included a history of other neurologic diseases. When the patient presented to the neurology polyclinic. The blood samples were collected. Patients with intracranial pathology in the MRI of the brain was performed. The diagnosis of TTH was based on the International Classification of Headache Disorders-III diagnostic criteria. The patients were divided into FETTH and CTTH groups according to their clinical characteristics. Duration of disease was determined for the patient groups.

All of the headache patients underwent physical, neurological, and fundus examinations. Kidney and liver function test levels and electrolyte levels were measured, and magnetic resonance imaging (MRI) of the brain was performed. Patients with intracranial pathology in the MRI of the brain were not included in the study. Blood samples were collected when the patient presented to the neurology polyclinic. The exclusion criteria included a history of other neurologic diseases, such as migraine, pregnancy, autoimmune diseases, diabetes mellitus, malignant hypertension, trauma, malignancies, thyroid diseases, liver and kidney dysfunction, and local or systemic infection. In addition, patients who had a headache at the time of examination or had used analgesics in the preceding 72 hours were excluded.

C-reactive protein (CRP), which is an acute phase reactant associated with acute and chronic inflammation, plays a role in pathological processes, and studies have found that increased CRP levels are involved in neurological diseases. In particular, CRP has been associated with migraine headaches. It has been noted that platelets play a role in inflammatory processes and immune responses. A study of migraine patients and the biological functions of platelets showed that platelets can play an active role in migraine headache etiology. However, the effect of platelets on the pathophysiology of tension headache disease is not fully known. Also, NLR, PLR, and CRP have not been previously evaluated in TTH. This study aimed to investigate the role of platelets, NLR, PLR, and CRP in frequent episodic tension-type headache (FETTH) and chronic tension-type headache (CTTH) patients.

Materials and Methods

In this study, 144 patients who had a diagnosis of TTH between January 2017 and August 2020 in the Department of Neurology, “A” Hospital were retrospectively evaluated. A control group of 60 healthy volunteers who were admitted to the checkup in the polyclinic was included. Demographic features, including age and sex, clinical features, frequency of pain, drugs being used, and medical history were assessed. The diagnosis of TTH was based on the International Classification of Headache Disorders-III diagnostic criteria. The patients were divided into FETTH and CTTH groups according to their clinical characteristics. Duration of disease was determined for the patient groups.

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Hematological and Biochemical Analyses

Serum CRP level was measured with a Beckman Coulter CX9 (Beckman Coulter, Inc., Brea, California, United States) chemistry analyzer. Hematologic indices were measured using an automated hematology analyzer system (Abbott Cell-Dyn 3700; Abbott Laboratory, Abbott Park, Illinois, United States). All the subsequent analyses were based on absolute cell counts. The baseline NLR was calculated by dividing the neutrophil count by the lymphocyte count, and the PLR was calculated by dividing the platelet count by the lymphocyte count.

Statistical Analyses

Statistical analyses were performed using the Statistical Package for the Social Sciences 21.0 for Windows software program (SPSS Inc., Chicago, United States). Continuous data are presented as mean ± standard deviation. The Kruskal–Wallis’ test was used to compare more than two groups. In comparison to the Kruskal–Wallis’ test, Bonferroni correction was performed, and the Mann–Whitney’s U test was used to compare the two groups. The cutoff values and corresponding sensitivity and specificity values for the prediction between the CTTH group and the FETTH group based on serum PLR, NLR, and CRP were estimated by receiver operating characteristic (ROC) curve analysis. A result was accepted as statistically significant with a p-value < 0.05.

Results

Sixty-four patients with FETTH, 80 patients with CTTH, and 60 healthy controls were enrolled in this study. The mean age of the FETTH patients was 34.62 ± 6.74 years, and the mean disease duration of the FETTH patients was 9.16 ± 6.69 months. The mean age of the CTTH patients was 34.67 ± 6.41 years, and the mean disease duration of the CTTH patients was 12.20 ± 4.03 months. The mean age of the control group was 35.60 ± 10.14 years. There was no statistically significant difference between the groups in terms of sex and age.

While the mean white blood cell (WBC) count in the FETTH patients was found to be 7.97 ± 0.75, it was 7.95 ± 2.21 in the CTTH patients and 8.38 ± 0.69 in the control group. When the WBC count was compared between the two headache groups, it was found to be statistically significantly higher in the CTTH patients (p < 0.001). The platelet count in the CTTH patient group was higher than in the other groups. Furthermore, the platelet counts in both the FETTH and CTTH patients were higher than in the control group. The NLR, PLR, and CRP values in the CTTH patients were statistically higher than in the FETTH patients and control groups. There was no correlation between age, disease duration, NLR, PLR, and CRP values between the two patient groups. The hematological parameters of the FETTH, CTTH, and control groups are presented in Table 1.

A cutoff NLR of 2.58 predicted the difference between the CTTH group and the FETTH group, with 19.4% sensitivity and 93.3% specificity (ROC area under the curve [AUC] of 0.61 and 95% confidence interval [CI] of 0.500–0.734). A cutoff PLR of 236.6 predicted the difference between the CTTH group and the FETTH group, with 9.7% sensitivity and 96.7% specificity (ROC AUC of 0.66 and 95% CI of 0.551–0.782).
A cutoff CRP of 0.42 predicted the difference between the CTTH group and the FETTH group, with 33.3% sensitivity and 86.7% specificity (ROC AUC of 0.64 and 95% CI of 0.528–0.768, p > 0.05; ►Fig. 1).}

Discussion

In this study, we first determined that platelet counts and NLR, PLR, and CRP values were statistically higher in the CTTH patients than in the other two groups. There was no difference in NLR and CRP values between the TTH patients and the control group. There was no correlation between age, frequency and duration of pain, and inflammatory markers in the patient groups. This study shows that increased platelet counts may play a role in the pathophysiology of TTH, and systemic inflammation may be a factor that affects the chronicity of tension headaches.

Peripheral mechanisms are predominant in FETTH, but central mechanisms are involved in CTTH. Recent studies have shown that pericranial myofascial pain sensitivity is increased in TTH and may play an important role in the pathophysiology of these headaches. On the contrary, sensitization of second-order neurons at the level of the spinal dorsal horn, trigeminal nucleus, or supraspinal neurons, as well as decreased descending inhibition from supraspinal structures may be involved in the pathophysiology of CTTH. Also, drug overuse and substance abuse suggest that a genetic factor is involved in the process of headache chronification. There are limited studies evaluating tension headache and peripheral inflammation. Domingues et al showed that the level of interleukin (IL)-8 and monocyte chemoattractant protein-1 increased in TTH types, and they suggested that proinflammatory mechanisms may participate in TTH pathophysiology. NLR is one of the important indicators of inflammation that is associated with many diseases. Studies have been conducted, especially in patients with migraine headaches during attacks, and found that NLR and PLR were higher in the patients compared with the controls. A study by Eryigit et al identified higher NLR levels in patients with subarachnoid hemorrhage compared with patients with migraine or other headache syndromes.
our study, no statistically significant change was observed in the neutrophil and lymphocyte counts in the patient groups. We only found NLR levels to be significantly higher in CTH patients. No difference was observed between the FETTH patients and the control group.

The importance of platelets in headaches has been studied for many years. In migraine cases, essential thrombocytopenia supports the fact that increased platelets can lead to headaches. There are limited studies evaluating the relationship between platelets and TTH. Leira et al showed that platelet-rich plasma serotonin levels were higher in patients with TTH. Blood serotonin levels reflect intraplatelet serotonin because most blood serotonin is stored inside platelets. Also, increased platelet levels are an important indicator of an inflammatory response. Ulusoy showed that mean platelet volume (MPV) and MPV-to-platelet ratio increased in a statistically significant manner in migraine patients, but MPV and MPV-to-platelet ratio had similar levels between FETTH patients and the healthy population. In this study, the platelet counts in patients with TTH were significantly higher. Benedick et al found that platelet counts in children with TTH were higher than in migraine patients. In our study, platelet counts were higher in both FETTH and CTH patients. We also found that the PLR value was higher in patients than in the control group. The higher platelet counts may have been affected by the serotonin levels, or it may be an indicator of the inflammatory response in the chronicity of the disease. Detection of increase in the platelets in the regular follow-up of TTH patients may indicate that the disease may have become chronic. Large-scale studies are needed to support these findings.

The relationship between CRP and migraine has been evaluated in many studies. CRP is an acute phase protein that is produced predominantly by the action of cytokines, such as IL-6 and tumor necrosis factor-α. CRP levels were found to be higher in patients admitted with the complaint of migraine attacks compared with controls. However, the relationship between CRP and CTTH is not clearly known. Bougea et al found no significant differences in CRP, IL-1β, and IL-6 levels between migraine patients and TTH patients. In our literature review, we first found high CRP levels in CTTH patients. Also, the sensitivity of our NLR, PLR, and CRP tests was found to be very low due to the area below the ROC curve, but the likelihood ratios of NLR, PLR, and CRP were good.

The principal limitation of this study is that platelet and serotonin levels were not evaluated together. These findings may uncover the importance of platelets in TTH pathophysiology.

**Conclusion**

In conclusion, increased platelet counts may have an effect on TTH pathophysiology. In addition, systemic inflammation parameters, such as NLR, PLR, and CRP, increased in the chronic phase of the disease. Therefore, we think that more comprehensive studies are needed to evaluate the effect of platelets on TTH and systemic inflammation in the chronicity of TTH.

**Conflict of Interest**

None declared.

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