Changes of Platelet Indices in Juvenile Idiopathic Arthritis in Acute Phase and After Two Months Treatment

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Abstract

Background: Various indices have been raised as predictors of activity and severity of juvenile idiopathic arthritis.

Objectives: This study was conducted to investigate the changes of platelet indices in acute phase and two months after treatment in these patients.

Patients and Methods: In a cohort study, platelet count, mean platelet volume (MPV), platelet distribution width (PDW), plateletcrit (PCT) were evaluated in children referred to children’s medical center, Tehran due to juvenile idiopathic arthritis from March 2013 to March 2014 during the acute phase and two months after standard treatment. The statistical data were analyzed by SPSS 19 software, and the significance level was set as P < 0.05.

Results: In this study, 55 children (24 boys and 31 girls) with mean ± SD age of 7.50 ± 3.35 years were studied. The mean ± SD value of platelet count was 441827.7 ± 551836.9 in the acute phase and reached 395418.2 ± 196601.6 two months after treatment (P = 0.01). The mean ± SD PCT in the acute phase of various subtypes of the disease was 0.32 ± 0.11, which reached 0.29 ± 0.10 after treatment (P = 0.09). However, the PDW range in different subtypes of the disease reached 11.4 ± 8.0 from 13.9 ± 2.9 and MPV reached 8.7 ± 0.9 from 8.8 ± 1.1 after treatment, but they were not significantly different from the results in the acute phase (P = 0.5).

Conclusions: Platelet count is one of the most remarkable indices in JIA. Evaluation of PCT can also help determine the severity of the inflammatory process in the follow-up and treatment process.

Keywords: Juvenile Idiopathic Arthritis, Acute Phase Reactants, Platelet Count, Platelet Indices, Mean Platelet Volume

1. Background

As the most common type of rheumatologic disease during childhood, juvenile idiopathic arthritis (JIA) is in fact a set of arthritis conditions before the age of 16 with unknown etiology that lasts longer than 6 weeks (1, 2). Although the etiology and pathogenesis of JIA is unknown; it seems that various types of HLA class I and II, polymorphism in genes related to Tumor necrosis factors-α (TNF-α), interlukines IL-6, and IL-1 secreted from monocyte cells and macrophages, some environmental factors and trauma to joints contribute to emergence of the disease (3). However, autoimmunity has a main role in the progression of this disease. In fact, T cells release proinflammatory cytokines such as L-6, TNF-α, and IL-1, and thus cause hypertrophy and hyperplasia of villous with hyperemia and synovial tissue edema (4).

One of the effects of proinflammatory cytokines produced in response to systemic inflammation in the liver is megakaryocyte stimulation and thus increases in the number of platelets in response to inflammation (3, 5). This results in the production of larger and more reactive platelets (6). There is evidence that platelets penetrate to synovial fluid in rheumatoid arthritis (RA) (7, 8). These platelets can bind to endothelial cells and leukocytes in inflamed synovial blood vessels causing thrombosis and abnormalities in synovial perfusion and joint destruction (9). Therefore, platelets may also contribute to the pathogenesis of JIA.

Few studies have been conducted on changes in platelet indices in adult rheumatoid arthritis. Mean platelet volume (MPV) is one of the markers that can be suggestive of platelet response. In physiological terms, increased MPV may suggest the entry of greater number of young platelets due to their higher production in blood. Some studies suggest MPV changes in the active phase of rheumatoid arthritis. In a few studies, MPV was used for monitoring response to treatment in rheumatoid arthritis (10, 11). In a study, treatment with anti-TNFα was associated with a significant increase in MPV (10). Another study was conducted on the relationship between MPV and clinical activity indices of RA and ankylosing spondylitis and the results showed that MPV was lower in the active phase of RA and AS compared to the control group and was significantly increased in the patient group after treatment (12). However, a study on JIA patients showed increased MPV in

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active phase of JIA (11).

2. Objectives

Given that no study has been conducted on the changes in the inflammatory phase reactants in patients with a variety of JIA in the course of the disease, we investigated the change pattern of platelet indices in the acute phase and after disease control.

3. Patients and Methods

This cohort study evaluated children whose JIA was diagnosed according to international league of associations for rheumatology (ILAR) from March 2013 to March 2014 (13). All patients were selected from pediatric rheumatology division in a tertiary children’s medical center in Tehran, Iran. JIA patients, systemic type and pauci- and polyarticular form (with RF negative) enrolled in this study. Other types include RF positive polyarticular form, psoriasis arthritis; enthesitis related arthritis and undifferentiated arthritis were not enrolled in this study due to low sample size in each group. Patients were excluded from the study if they failed to take part in follow-up examinations, the diagnosis of JIA changed during the first month or they simultaneously had the symptoms of infectious diseases. In addition to treatment with methotrexate, all patients were treated with prednisolone or nonsteroidal anti-inflammatory drugs (NSAIDs), based on the severity of diseases. In addition to treatment with methotrexate, all patients were examined of whom 24 (43.6%) were boys and 31 (54.4%) girls. Patients’ mean ± SD age was 7.50 ± 3.35 years (from 1 to 14 years). In terms of disease subtype, 27 (49.1%) patients had oligoarthritis, 19 patients (34.5%) polyarthritis and 9 (16.4%) had systemic arthritis.

Table 1 shows mean ± SD values of platelet count and platelet indices in patients. The difference between platelet count and PCT was significant in all subtypes of the disease in the acute phase (P = 0.01 and P = 0.02, respectively). In analysis among the subtypes of JIA, mean difference of MPV and PCT had significant differences between oligoarticular and polyarticular subtype in the acute phase (both P = 0.03). On the other hand significant difference in the acute phase was related to differences between oligoarticular and polyarticular subtype.

After 2 months treatment, there was still a statistically significant difference between the mean platelet count in all three subtypes of the disease (P = 0.03). There was no significant difference between the mean platelet indices in different subtypes of the disease although this difference was significant in PCT as borderline (P = 0.09) (Table 2).

Table 3 presents the comparison of the number of platelets and platelet indices before and two months after treatment. As shown, the total number of platelets and also the number of platelets in the systemic disease declined after treatment (P = 0.01, P < 0.01, respectively). Also systemic PCT significantly reduced after treatment (P = 0.02) and the total value of PCT indicated a significant borderline reduction after treatment (P = 0.09). The index showed no significant change in other subtypes of the disease. Also, other platelet indices i.e. PDW and MPV had no significant changes overall and in subtypes of the disease (P = 0.5 for each one).

4. Results

In this study, a total of 55 patients diagnosed with JIA were examined of whom 24 (43.6%) were boys and 31 (54.4%) girls. Patients’ mean ± SD age was 7.50 ± 3.35 years (from 1 to 14 years). In terms of disease subtype, 27 (49.1%) patients had oligoarthritis, 19 patients (34.5%) polyarthritis and 9 (16.4%) had systemic arthritis.

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5. Discussion

Clinical presentations of JIA are highly variable such that it can be seen in a range of very mild self-limiting arthritis to highly destructive arthritis. However, no measure has been developed so far to evaluate the response rate to treatment. Also, no specific and sensitive marker has been identified to assess disease activity rate (15). Evaluation of acute phase reactants such as platelet indices is
Table 1. Comparison of Indices in Patients With Different Subtypes of JIA in Acute Phase of Disease

| Parameter | Mean ± SD | 95% Confidence Interval |
|-----------|-----------|------------------------|
|           | Lower Bound | Upper Bound |
| PLT      |            |                |
| Oligo-articular, n = 27 | 380629.6 ± 137244.9 | 326337.35 | 434921.91 |
| Poly-articular, n = 19  | 481842.1 ± 122358.0  | 422867.45 | 540816.76  |
| Systemic, n = 9          | 541222.2 ± 182091.2  | 401254.56 | 681189.89  |
| Total, n = 55            | 441872.7 ± 151836.9  | 400825.46 | 482919.99  |
| PDW      |            |                |
| Oligo-articular, n = 27 | 14.2 ± 2.9     | 13.0       | 15.3       |
| Poly-articular, n = 19  | 13.8 ± 3.0     | 12.4       | 15.2       |
| Systemic, n = 9          | 13.5 ± 3.1     | 11.1       | 16.0       |
| Total, n = 55            | 13.9 ± 2.9     | 13.2       | 14.7       |
| PCT      |            |                |
| Oligo-articular, n = 27 | 0.28 ± 0.10    | 0.24       | 0.32       |
| Poly-articular, n = 19  | 0.35 ± 0.10    | 0.31       | 0.40       |
| Systemic, n = 9          | 0.36 ± 0.11    | 0.27       | 0.44       |
| Total, n = 55            | 0.32 ± 0.11    | 0.29       | 0.35       |
| MPV      |            |                |
| Oligo-articular, n = 27 | 9.2 ± 1.1      | 8.8        | 9.7        |
| Poly-articular, n = 19  | 8.4 ± 0.9      | 8.0        | 8.8        |
| Systemic, n = 9          | 8.5 ± 0.9      | 7.8        | 9.2        |
| Total, n = 55            | 8.8 ± 1.1      | 8.5        | 9.1        |

Abbreviations: SD, standard deviation; JIA, juvenile idiopathic arthritis; PLT, platelet; PDW, platelet distribution width; MPV, Mean platelet volume; PCT, plateletcrit.

aSignificant (P < 0.05).

a way to evaluate these patients (3). Although few studies have been conducted on platelet indices in rheumatologic diseases, studies on these factors in JIA are much fewer. In this study, increased mean platelet count in the acute phase of the disease was seen in the systemic and polyarticular subtypes; however, mean platelet count was normal in oligoarticular subtype. Interleukin 6 (IL-6) is the cause of thrombocytosis in autoimmune diseases such as RA (16). As reactive thrombocytosis is a symptom of inflammation in JIA (17, 18), this represents a more severe inflammatory process in the polyarticular and systemic subtypes. In other studies, inflammatory process has been reported in the systemic subtype of JIA (19, 20). Spiegel (17) also found that thrombocytosis in an active systemic disease can be considered as an important factor in the poor prognosis of the patients.

In this study, after 2 months of treatment, the mean platelet count was evaluated again in all three subtypes of the disease, which indicated a decrease in all of them. The reduction in mean platelet count after treatment compared to before treatment was statistically significant. In polyarticular subtype, the mean platelet count was still above the normal range despite two months of treatment, although this variable had reached normal range in the systemic subtype like oligoarticular involvement. This can be a sign of slower remission of inflammatory symptoms in polyarticular subtype with mild treatments compared to systemic disease. Ozturk et al. (14) showed that the mean number of platelets is reduced in Crohn’s and ulcerative colitis remission phase compared to the active phase. Berntson et al. (2) reported that the mean number of platelets in patients with JIA with at least one joint involvement is significantly higher than the mean platelet count in patients who are in complete remission. However, the number of platelets higher than the normal range was not necessarily associated with a symptomatic disease.

In another study conducted by Sandborg et al. (21), predicting measures of joint destruction were evaluated in
Table 2. Comparison of Indices in Patients With Different Subtypes of JIA After 2 Months Treatment

| Parameter | Mean ± SD | 95% Confidence Interval |
|-----------|-----------|-------------------------|
|           | Lower Bound | Upper Bound             |
| PLT       |            |                         |
| Oligo-articular, n = 27 | 358444.4 ± 91415.1 | 322281.8 | 394607.1 |
| Poly-articular, n = 19 | 453105.3 ± 134702.1 | 388180.9 | 518029.6 |
| Systemic, n = 9 | 384555.6 ± 126177.6 | 287566.9 | 481544.2 |
| Total, n = 55 | 395418.2 ± 119601.6 | 363085.3 | 427751.0 |
| PDW       |            |                         |
| Oligo-articular, n = 27 | 13.1 ± 5.7 | 10.8 | 15.310 |
| Poly-articular, n = 19 | 12.7 ± 8.8 | 8.5 | 17.0 |
| Systemic, n = 9 | 16.2 ± 11.9 | 7.0 | 25.3 |
| Total, n = 55 | 13.4 ± 8.0 | 11.2 | 15.6 |
| PCT       |            |                         |
| Oligo-articular, n = 27 | 0.27 ± 0.09 | 0.24 | 0.31 |
| Poly-articular, n = 19 | 0.33 ± 0.12 | 0.28 | 0.39 |
| Systemic, n = 9 | 0.26 ± 0.10 | 0.18 | 0.33 |
| Total, n = 55 | 0.29 ± 0.10 | 0.26 | 0.32 |
| MPV       |            |                         |
| Oligo-articular, n = 27 | 8.9 ± 1.0 | 8.5 | 9.3 |
| Poly-articular, n = 19 | 8.6 ± 0.67 | 8.3 | 8.9 |
| Systemic, n = 9 | 8.6 ± 0.8 | 7.9 | 9.2 |
| Total, n = 55 | 8.7 ± 0.9 | 8.5 | 9.0 |

Abbreviations: SD, standard deviation; JIA, juvenile idiopathic arthritis; PLT, platelet; PDW, platelet distribution width; MPV, Mean platelet volume; PCT, plateletcrit.

*Significant (P < 0.05).

JIA. This study showed that children who had more than 788,000 platelets at least once in the first 3 months, and also children with more than 13 joints involved, were more likely to develop joint destruction within 2 years, despite platelet count less than 788,000. Spiegel et al. (17) also showed that with each 100,000 increase in platelet count in the tests performed 6 months after diagnosis, the risk of severe disease with adverse outcomes in patients with systemic JIA would be 1.45 times greater. Milovanovic et al. (5) found that the mean number of platelets in the active phase of rheumatoid arthritis is significantly higher than in remission phase of the disease, and the increase in the number of platelets is associated with IL-6, but not with thrombopoietin.

MPV is a platelet index that is measured automatically, is associated with platelets function and activity (22, 23), and is affected by inflammatory reactions. Inflammatory mediators stimulate bone marrow precursors in the presence of autoimmune diseases to produce more platelets through shortening their maturation time, so smaller platelets enter the blood stream, while active platelets are destroyed at the site of inflammation (24). However, there is a nonlinear inverse relationship between MPV and platelet count (25).

In this study, MPV was within normal range in the acute phase in all cases. The highest MPV was observed in oligoarticular subtype, and then in the systemic and polyarticular subtype, however, MPV difference was not significant between systemic and polyarticular form in the acute phase. It seems that MPV has an inverse relationship with inflammatory disease severity such that studies by Isik et al. (26) and Gasparyan et al. (27) suggest a decline in MPV in severe inflammatory diseases such as rheumatoid arthritis active phase. In most cases; however, it should be considered that MPV is in the low normal range (26). In contrast, in a study on JIA, MPV increased in acute phase and in correlation to severe disease (11). In our study, after 2 months of treatment, MPV was assessed again in all three disease subtypes,
Table 3. Comparison of Indices in Acute Phase of Disease and 2 Months after treatment in Different Subtypes of JIA

| Parameter | Mean ± SD | 95% Confidence Interval |
|-----------|-----------|------------------------|
|           | Lower Bound | Upper Bound |
| PLT       |            |            |
| Oligo-articular, n = 27 | 22185.2 ± 20674.0 | -20310.8 | 64681.1 |
| Poly-articular, n = 19  | 28736.8 ± 28179.7  | -30466.6 | 87940.3 |
| Systemic, n = 9         | 15666.7 ± 36913.2* | 7544.7   | 24788.6 |
| Total, n = 55           | 46454.5*        | 13531.161 | 79377.930 |
| PDW       |            |            |
| Oligo-articular, n = 27 | 1.1 ± 1.2     | -1.25 | 3.5   |
| Poly-articular, n = 19  | 1.1 ± 1.9     | -3.1 | 5.2   |
| Systemic, n = 9         | -2.6 ± 4.3    | -12.7 | 7.5   |
| Total, n = 55           | 0.49          | -1.77 | 2.76  |
| PCT       |            |            |
| Oligo-articular, n = 27 | 0.004 ± 0.021 | -0.039 | 0.047 |
| Poly-articular, n = 19  | 0.023 ± 0.026 | -0.032 | 0.079 |
| Systemic, n = 9         | 0.100 ± 0.033 | 0.023   | 0.177 |
| Total, n = 55           | 0.03          | 0.00   | 0.06  |
| MPV       |            |            |
| Oligo-articular, n = 27 | 0.33 ± 0.21   | -0.10 | 0.76  |
| Poly-articular, n = 19  | -0.39 ± 0.16  | -0.53 | 0.15  |
| Systemic, n = 9         | 0.00 ± 0.43   | -0.99 | 0.99  |
| Total, n = 55           | 0.10          | -0.18 | 0.37  |

Abbreviations: SD, standard deviation; JIA, juvenile idiopathic arthritis; PLT, platelet; PDW, platelet distribution width; MPV, Mean platelet volume; PCT, plateletcrit.
*Significant (P < 0.05).

and despite a modest rise in the polyarticular subtype, it had a decrease in other two subtypes. However, differences were not statistically significant after treatment compared to MPV before treatment and the rate was within normal limits in all cases. Studies by Isik (26) and Gasparyan (27) also showed that MPV that decreased in the active phase of rheumatoid arthritis as a severe inflammatory disease would increase after remission to a normal range [7.4 - 10.4 fL]. Gasparyan et al. (27) depicted that MPV is decreased in severe inflammatory diseases, but is increased in mild inflammatory diseases. So it seems that increased MPV in subsystemic type after two months of treatment can represent the appropriate effect of treatment in the control of inflammatory response in these patients. Studies of Kapsoritakis et al. (28) and Yuksel et al. (29) also showed that there is an inverse relationship between the severity of inflammatory bowel disease in patients with IBD and MPV. However, Ozturk et al. (14) found that MPV was decreased after the remission of acute phase of ulcerative colitis and increased in remission of Crohn’s disease. So it seems that the evaluation of MPV changes alone is not helpful in the follow-up and evaluation of inflammatory diseases because MPV showed a modest rise in polyarticular subtype and a decrease in other subtypes of the disease two months after treatment in our study. However, in most cases, MPV was in the low normal range in the acute phase and two months after treatment. Milovanovic et al. (5) also believe that because of the overlap of MPV in inflammatory diseases with normal values, it is necessary that changes of this index be individually evaluated and interpreted in the course of the assessment of each patient.

PDW is another indicator of platelet volume directly measured by flow cytometry, which reports platelet distribution by evaluating the top 20% of the distribution curve (30). This study showed that PDW was reduced in all subtypes in the acute phase of the disease, and the highest drop in PDW was observed in systemic subtype, then in polyarticular and oligoarticular subtype. According to the
results of Isik et al. (26), it seems that PDW like MPV has an inverse relationship with inflammatory disease severity and is reduced in severe inflammatory diseases such as the active phase in rheumatoid arthritis. Ozturk et al. (14) also found that PDW has an inverse relationship with ulcerative colitis activity. In our study, after 2 months of treatment, the mean PDW was evaluated again in all three subtypes of the disease, and despite an increase in the systemic subtype, it showed a decrease in other two subtypes, but PDW changes were not statistically significant before and after treatment. The index only reached a low normal range in systemic subtype and was still lower than the normal range in other two subtypes (15.6 - 18.2 fl). Isik et al. (26) also showed that PDW is decreased in the active phase of rheumatoid arthritis as a severe inflammatory disease; however, unlike our study, the mean PDW in the study of Isik was in low normal range. Ozturk et al. (14) reported that PDW which is decreased in the active phase of inflammatory bowel diseases is increased after the remission of acute phase of Crohn’s disease and ulcerative colitis and can be used as a good indicator to track inflammatory bowel diseases from acute phase to remission. In contrast to previous studies on JIA patients, Gunes et al. (11) reported, similar to MPV, higher range for PDW in acute phase and in patients with severe disease. However, in the present study, the evaluation of PDW changes was not helpful in follow-up and disease assessment.

Unlike other platelet-related indices that are measured directly by a machine, PCT is a measure that depends on platelet count and MPV, and shows platelets in a unit of blood volume (31). This study showed that an increase in the mean PCT in the acute phase of the disease is observed in systemic and polyarticular subtypes; however, mean PCT in oligoarticular subtype was within the normal range. Studies of Isik et al. (26) and Santimone et al. (32) suggest an increase in PCT in severe inflammatory diseases such as rheumatoid arthritis active phase; however, it should be considered that despite the increase mentioned, PCT is in high normal range in most cases (26). The study of Ozturk et al. (14) also found that PCT is increased with ulcerative colitis activity. In our study, the mean PCT was evaluated again after 2 months of treatment in all three subtypes of the disease, and despite the lack of significant changes in the oligoarticular subtype, it was decreased in two other subtypes. These changes after treatment were only statistically significant in systemic subtype, compared to the mean PCT before treatment. This index reached a normal range only in the systemic subtype after treatment and was still higher than the normal range in the polyarticular subtype; this can be a sign of slower remission of inflammation symptoms in treatment of polyarticular disease as compared to the systemic subtype. Ozturk et al. (14) showed that PCT which is increased in the active phase of inflammatory bowel diseases can be used as a good indicator to track IBD from the acute phase to remission. The study of Isik et al. (26) also indicated that PCT is higher in the active phase of rheumatoid arthritis than remission period of the disease. According to the results of our study it seems that the evaluation of PCT changes is only helpful in the follow up and assessment of the disease in the systemic and polyarticular subtypes as in our study this measure was within normal limits even before the treatment in oligoarticular subtype.

Confounding factors and severity of the JIA has not been considered in this study. In a recent study, platelets indices were compared with severity of the rheumatoid arthritis and JIA (11, 33). Other studies are recommended for covering these limitations and considering severity in subtype of JIA patients.

5.1. Conclusion

This study showed that JIA leads to some changes in platelet counts and platelets indices, especially in the acute phase of the disease in different subtypes with developing a chronic inflammatory process. On the other hand, although in most cases only platelet count is considered as an indicator of acute inflammation, PCT evaluation can also be helpful in determining the severity of the inflammatory process in the course of follow up and treatment. However, MPV and PDW had no significant changes after treatment. Since platelet indices are non-specific for autoimmune diseases, these findings can be considered and interpreted besides other clinical and laboratory findings.

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Footnote

Authors’ Contribution: Marjan Vakili participated in the acquisition of data and manuscript preparation; Vahid Ziaee participated in the concept and Design carried of the study, data collection, analysis and interpretation and critical revision of the manuscript; Mohammad Hassan Moradinejad participated in the concept and design carried of the study and critical revision of the manuscript; Seyed Reza Raeekarami and Farzad Kompani participated in the design of the study and interpretation of findings; Tayebeh Rahamooz participated in the acquisition of data. All authors read and approved the final manuscript.
