Original Research Article

Analytical study of reactive thrombocytosis prevalence in microcytic hypochromic anemia

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ABSTRACT

Background: In routine hematology blood samples, every now and then when reporting peripheral smears, we come across cases of microcytic hypochromic anemia with increased platelet count more often in iron deficiency anemia. We have to differentiate is it the cases are just reactive thrombocytosis or else we are neglecting any platelet disorders. Iron deficiency is the commonest cause of secondary thrombocytosis.

Aims and Objectives: To know the prevalence of reactive thrombocytosis in microcytic hypochromic anemia cases. To analyze the range of hemoglobin levels and MCV values in which reactive thrombocytosis is more consistent.

Materials and Methods: The study includes 500 blood samples of patients of all the age groups and both the gender with microcytic hypochromic anemia. The samples were scrutinized for the corresponding thrombocytosis by using a hematology analyzer and confirmed by peripheral smear examination. Serum iron studies were done in confirmed samples of microcytic hypochromic anemia with valid thrombocytosis.

Results: Out of 500 cases of microcytic hypochromic anemia analysed 115 cases had associated thrombocytosis. Out of which 17(14.78%) cases were males and 98 (85.22%) cases of females showed thrombocytosis. The majority of cases showed mild thrombocytosis which is frequently seen in cases with MCV 60-70 fl and moderate degree of anemia with hemoglobin level 7-9.9g/dl.

Conclusion: The utmost prevalent cause of reactive thrombocytosis in microcytic hypochromic anemia is iron deficiency. It is rational to distinguish between reactive and clonal thrombocytosis. However, the distinction cannot be always made with certainty, and the diagnosis often depends on watching the platelet count over a while. By identifying the etiology of increased platelet count and type of anemia treatment will be ease.

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1. Introduction

The platelet count > 4,50,000/µl in the peripheral blood sample is referred to as thrombocytosis, by the rampant usage of electronic cell counters diagnosis of thrombocytosis is made easily and more often observed as an unexpected finding.¹,²

In the peripheral blood, numerous disease conditions can lead to increased platelet count. The causes may be primary in elderly patients due to myeloproliferative disorders or secondary to infection or inflammation.³

Reactive thrombocytosis has to be differentiated from primary or clonal platelet disorders which are neglected many times which is of practical clinical importance and a diagnostic challenge. However, the distinction is not always made with certainty, and the diagnosis often depends on observing the platelet count over a period of time following the treatment.⁴

The commonest non-infectious cause of secondary thrombocytosis is iron deficiency anaemia. Thrombocytosis is either due to a reactive process (Secondary) or because of clonal disorders (primary thrombocytosis). It is critical
concern to distinguish, because thrombo-haemorrhagic complications are frequent in clonal rather than reactive thrombocytosis.5,6

2. Aims and objectives of the study

1. It is a prospective study of 500 cases of microcytic hypochromic anemia.
2. How often microcytic hypochromic anemia cases present with thrombocytosis and in what range of hemoglobin level and MCV values it is affecting more consistently.
3. Which age group and sex are affected more commonly.

3. Methods

3.1. Selection of samples

All blood samples of patients of all the age groups were included in the study from January 2019 to December 2019 in a tertiary care center.

All EDTA blood samples were run in electronic cell counter for recording the Haemoglobin levels, MCV, and platelet counts along with platelet indices (PDW, Plateletecrit, and MPV) Patients who have low hemoglobin levels and MCV less than 80fl were screened for microcytic hypochromic anemia on peripheral smears stained by Leishman stain to confirm RBC morphology.

Random 500 blood samples with microcytic hypochromic anemia were selected and analyzed for the corresponding thrombocytosis. Platelet counts were confirmed by peripheral smear examination.

Confirmed cases of microcytic hypochromic anemia associated with thrombocytosis were sent for iron profile.

3.2. Inclusion criteria

Blood samples of all age groups received in the hematology laboratory who are diagnosed with microcytic hypochromic anemia were included in the study

3.3. Exclusion criteria

Blood samples of all age groups presenting with anemia and thrombocytosis for various reasons other than the microcytic hypochromic anemia were excluded.

In our study, the normal values of hemoglobin are considered according to the age and sex of the individual cases concerning WHO criteria for anemia.

World Health Organization classification graded as hemoglobin level 11-12.9 g/dL as mild anemia in males and 11-11.9 g/dL in females.7

Hemoglobin level 8-10.9 g/dL as moderate anemia and <8 g/dL as Severe anemia both in males and females

Dame and Sutor8 have categorized thrombocytosis into mild (5 to 7 lakhs/cumm) moderate (>7 to 9 lakhs/cumm) and severe (9lakhs/cumm) and extreme (> 10 lakhs/cumm) and the same criteria is used in this study

4. Results

In the present study of 500 patients of Microcytic hypochromic anemia, 23% (n =115) of the total cases showed thrombocytosis (Graph 1)

Table 1 In the present study out of 500 cases of microcytic hypochromic anemia, 22.6 % (n = 113) were male patients and 77.4 % (n = 387) were female patients. In males microcytic hypochromic anemia is more prevalent in age group between 41 to 60 and then cases are more in the age group more than 70 years.

In females microcytic hypochromic anemia is more prevalent in the age group between 21 to 50 years, which is common in the reproductive age group.

Table 2 Out of 113 total male patients, 17 (15.04%) of them showed thrombocytosis associated with microcytic hypochromic anemia.

Out of 387 female patients, 98 (25.3%) of them showed thrombocytosis which is almost 10% more when compared to males (15.04%) with thrombocytosis associated with microcytic hypochromic anemia.

Table 3 In male patients microcytic hypochromic anemia with thrombocytosis is more common in the age group above 70 years (23.52%) and then in less than 10 years. (17.64%)

In female patients Microcytic hypochromic anemia with thrombocytosis is more common in the age group above 70 years (29.59%) and then in 31-40 years. (28.57%)

Table 4 In our study of male patients showed mild thrombocytosis (70.58%) in majority of the cases and commonly associated with moderate anemia (47.05 %) and only 5.88% showed severe thrombocytosis with one patient with platelet count more than 10 Lakhs/cumm.

In this study majority of the female patients showed mild thrombocytosis (86.74%) and commonly associated with moderate anemia (74.48%) and only (2.04%) showed severe
**Table 1:** Showing age wise distribution of male and female patients with microcytic hypochromic anaemia.

<table>
<thead>
<tr>
<th>Age (in years)</th>
<th>1-10</th>
<th>11-20</th>
<th>21-30</th>
<th>31-40</th>
<th>41-50</th>
<th>51-60</th>
<th>61-70</th>
<th>&gt; 70</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of cases</td>
<td>10</td>
<td>32</td>
<td>125</td>
<td>112</td>
<td>119</td>
<td>43</td>
<td>30</td>
<td>29</td>
<td>500 (100%)</td>
</tr>
<tr>
<td>Total number of Males</td>
<td>08</td>
<td>07</td>
<td>13</td>
<td>11</td>
<td>23</td>
<td>18</td>
<td>11</td>
<td>22</td>
<td>113 (22.6%)</td>
</tr>
<tr>
<td>Total number of Females</td>
<td>02</td>
<td>25</td>
<td>112</td>
<td>101</td>
<td>96</td>
<td>25</td>
<td>19</td>
<td>07</td>
<td>387 (77.4%)</td>
</tr>
</tbody>
</table>

**Table 2:** Distribution of male and female patients with microcytic hypochromic anaemia associated with thrombocytosis

<table>
<thead>
<tr>
<th>Age (in years)</th>
<th>1-10</th>
<th>11-20</th>
<th>21-30</th>
<th>31-40</th>
<th>41-50</th>
<th>51-60</th>
<th>61-70</th>
<th>&gt; 70</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients presented with thrombocytosis</td>
<td>03</td>
<td>10</td>
<td>23</td>
<td>29</td>
<td>31</td>
<td>08</td>
<td>06</td>
<td>05</td>
<td>115 (100%)</td>
</tr>
<tr>
<td>No of male patients presented with thrombocytosis</td>
<td>03</td>
<td>02</td>
<td>01</td>
<td>01</td>
<td>02</td>
<td>02</td>
<td>02</td>
<td>04</td>
<td>17 (14.78%)</td>
</tr>
<tr>
<td>No of female patients presented with thrombocytosis</td>
<td>02</td>
<td>08</td>
<td>20</td>
<td>28</td>
<td>29</td>
<td>06</td>
<td>04</td>
<td>01</td>
<td>98 (85.22%)</td>
</tr>
</tbody>
</table>

**Table 3:** Showing age wise distribution of male and female patients with microcytic hypochromic anaemia associated with thrombocytosis.

<table>
<thead>
<tr>
<th>Age (in years)</th>
<th>1-10</th>
<th>11-20</th>
<th>21-30</th>
<th>31-40</th>
<th>41-50</th>
<th>51-60</th>
<th>61-70</th>
<th>&gt; 70</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total number of patients presented with thrombocytosis</td>
<td>03</td>
<td>10</td>
<td>23</td>
<td>29</td>
<td>31</td>
<td>08</td>
<td>06</td>
<td>05</td>
<td>115 (100%)</td>
</tr>
<tr>
<td>No of male patients presented with thrombocytosis</td>
<td>03</td>
<td>02</td>
<td>01</td>
<td>01</td>
<td>02</td>
<td>02</td>
<td>02</td>
<td>04</td>
<td>17 (14.78%)</td>
</tr>
<tr>
<td>No of female patients presented with thrombocytosis</td>
<td>02</td>
<td>08</td>
<td>20</td>
<td>28</td>
<td>29</td>
<td>06</td>
<td>04</td>
<td>01</td>
<td>98 (85.22%)</td>
</tr>
</tbody>
</table>

**Table 4:** Distribution of severity of thrombocytosis in males and females in relation to severity of anaemia.

<table>
<thead>
<tr>
<th>Males</th>
<th>Severity of anaemia (gm/dl)</th>
<th>No of male patients presented with mild thrombocytosis</th>
<th>No of male patients presented with moderate thrombocytosis</th>
<th>No of male patients presented with severe thrombocytosis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mild (Hb 11-12.9)</td>
<td>2</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>Moderate (Hb 8-10.9)</td>
<td>6</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Severe (Hb &lt; 8)</td>
<td>4</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Total (n = 17)</td>
<td>12 (70.58 %)</td>
<td>4 (23.52 %)</td>
<td>1 (5.88 %)</td>
</tr>
<tr>
<td>Females</td>
<td>Severity of anaemia (gm/dl)</td>
<td>No of Female patients presented with mild thrombocytosis</td>
<td>No of Female patients presented with moderate thrombocytosis</td>
<td>No of Female patients presented with severe thrombocytosis</td>
</tr>
<tr>
<td></td>
<td>Mild (Hb 11-11.9)</td>
<td>7</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Moderate (Hb 8-10.9)</td>
<td>64</td>
<td>7</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td>Severe (Hb &lt; 8)</td>
<td>14</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>Total (n = 98)</td>
<td>85 (86.7 %)</td>
<td>11 (11.22 %)</td>
<td>2 (2.04 %)</td>
</tr>
</tbody>
</table>

Table 5 In male patients most cases of microcytic hypochromic anaemia with thrombocytosis presented within the range of 60.1 to 70fL MCV (52.94%) and majority of the cases showed mild thrombocytosis (58.8%). In female patients most cases of microcytic hypochromic anaemia with thrombocytosis presented within the range of 60.1 to 70fL MCV (68.36%) and majority of the cases showed mild thrombocytosis (84.69%).

Platelet indices analysed like MPV (Mean platelet volume), PDW (Platelet Distribution width), and PCT (Plateletcrit) were well within the normal range and no significant variations noted in all the cases of thrombocytosis associated with microcytic hypochromic anemias in the present study.

Most of the cases of microcytic hypochromic anaemia with thrombocytosis (115) were sent for Iron studies and the majority of them were confirmed as iron deficiency anemia, which again proved that iron deficiency anemia is the most prevalent cause of reactive thrombocytosis.

5. **Discussion**

There are various disease states that can bring about the elevated levels of platelets. The causes may be primary...
Table 5: Showing severity of thrombocytosis in males and females in relation to MCV.

<table>
<thead>
<tr>
<th>MCV (fl)</th>
<th>Male</th>
<th>Female</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No of male patients</td>
<td>No of Female patients</td>
</tr>
<tr>
<td></td>
<td>presented with mild</td>
<td>presented with moderate</td>
</tr>
<tr>
<td></td>
<td>thrombocytosis</td>
<td>thrombocytosis</td>
</tr>
<tr>
<td>70.1-79.9</td>
<td>2</td>
<td>12</td>
</tr>
<tr>
<td>60.1-70</td>
<td>6</td>
<td>57</td>
</tr>
<tr>
<td>&lt;60</td>
<td>4</td>
<td>14</td>
</tr>
<tr>
<td>Total (n=17)</td>
<td>10 (58.8 %)</td>
<td>83 (84.69 %)</td>
</tr>
</tbody>
</table>

in elderly patients due to myeloproliferative disorders or secondary to infectious or inflammatory disorders.

Reactive thrombocytosis needs to be differentiated from primary or clonal platelet disorders, which are neglected many a times. Reactive thrombocytosis needs to be differentiated from primary or clonal platelet disorders, which are neglected many a times.8

It is extremely challenging to understand the reactive and clonal thrombocytosis process based upon clinical findings and laboratory test results even though there are underlying pathophysiological differences and clinical implications. The various physiological and pathological processes which contributes to the elevated platelet count has become clinically significant in routine practice. However, the distinction cannot be made with certainty, and the interpretation often depends on observing the platelet count variations periodically following the treatment.9

Iron deficiency anemia is the most common cause among the non-infectious causes of secondary thrombocytosis. This approach nevertheless is not without dangers, because thrombohemorrhagic complications are more common in clonal rather than reactive thrombocytosis.10

Study done by Ramu. R et al11 about iron deficiency anemia with thrombocytosis and erythropoietin levels also showed that majority of the cases in their study had moderate degree of anemia and presented frequently with mild thrombocytosis which is similar to our study. They also depicted that novel data advocate the symbiotic effect of both erythropoietin and thrombopoietin on the bipotential progenitor cells of the erythroid/megakaryocyte precursors. Despite, this decline to clarify that not all the patients with elevated levels of erythropoietin, and iron-deficiency anemia cases present with thrombocytosis. Hence, there may be boundless added mechanisms that aid in the evolution of thrombocytosis in few cases of iron deficiency anemia.

Tania O et al12 described that bone marrow examination cannot prove the causes of chronic myeloproliferative disorders, especially essential thrombocytemia. In iron deficiency anemia with associated thrombocytosis after an increase in hemoglobin level closer to normal values with packed red cell transfusion, platelet counts become normal with lowering erythropoietin levels. The probable mechanism in iron-deficiency anemia associated thrombocytosis is still a matter of deliberation of the concept of thrombopoietin and erythropoietin amino acid structural homology of might justify the cause of thrombocytosis in children associated with iron-deficiency anemia.

Mehri T et al13 reported that in normal individuals, MPV is inversely related to Platelet count. The reference values of MPV may differ with the platelet count. In majority of the cases of myeloproliferative disorders, MPV is usually increased. Platelet counts are unusually increased in primary myeloproliferative disorders than secondary thrombocytosis cases. PDW in both primary and secondary thrombocytosis also varies accordingly. In our present study all the cases of thrombocytosis, the platelet indices were within normal limits.

Sandoval.C.14 and Nathiya S et al15 defined that amidst of all the anemia types, anemia with iron deficiency most often presents with thrombocytosis and is more persistent in children below 2 years of age, because of the greater prevalence of iron deficiency in this age group. In this study more the severity of anemia was directly proportional to severity of thrombocytosis. The mean MPV and PDW were found to be higher in primary thrombocytosis when compared to secondary thrombocytosis. In the present study most cases were in the reproductive age group in females.

Jonathan S et al16 and Akan et.al determined in his study that cytokines IL-6, IL-11, and TPO were not increased in patients associated with iron deficiency anemia associated with thrombocytosis, in correlation with iron deficiency anemia with normal platelet counts. This shows that the cytokine levels do not have any significant portrayal in iron deficiency associated thrombocytosis.17
Secondary thrombocytosis is usually favorable and platelet becomes normal with prompt treatment of the underlying cause without any thrombotic complications.18

6. Conclusions
In our routine reporting of peripheral smear examination, we come across many cases of increased platelet count in microcytic hypochromic anemia more often in iron deficiency anemia. It is of practical clinical importance to be readily able to distinguish between reactive and clonal thrombocytosis.

However, the distinction cannot be always made with certainty, and the diagnosis often depends on watching the platelet count over a while. By knowing the cause of increased platelet count and type of anemia treatment will be ease

To conclude, as there are only fewer articles published regarding the prevalence of thrombocytosis in microcytic hypochromic anemias involving all the age groups

This study was embarked upon, hemoglobin levels, MCV, and platelet count in all the age groups and both the genders. From the current study, we conclude that mild thrombocytosis is frequently seen in microcytic hypochromic anemia cases in females in the age group 21-50 years in which the majority of them proved to be iron deficiency anemia.

Ruling out the secondary causes of thrombocytosis should be the priority, then primary clonal disorders, whenever there is a suspicion. We have to differentiate that the cases are just reactive thrombocytosis or else we are neglecting any platelet disorders.

Among the non-infectious causes of secondary thrombocytosis, iron deficiency is a common one, since it is the single most common nutritional deficiency worldwide which can be treated efficiently next to Vitamin B12 and folic acid deficiency.

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No financial support was received for the work within this manuscript.

8. Conflict of Interest
The authors declare they have no conflict of interest.

References

Author biography
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Harish S.G, Professor