Hematological profile of sickle cell disease in Chhattisgarh

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ABSTRACT

Background: Sickle cell disease hemoglobinopathy gets inherited in autosomal recessive pattern. In sickle cell disease substitution of amino acid valine for glutamic acid at the sixth position on beta globin chain takes place resulting in sickled hemoglobin which is a hemoglobin tetramer.

Methods: A prospective observational study was conducted in the Sickle Cell Institute, Raipur, India, and Department of Pharmacology in collaboration with Department of Biochemistry, Pt. J.N.M. Medical College, Raipur, Chhattisgarh, India, from February 2018 to June 2018. Patients included were in the steady state for a long period of time without any symptoms related to sickle cell disease or any other diseases which could affect hematological parameters. Subjects transfused in the last three months were excluded. Student t test and Pearson Correlation Coefficient test was done on stat pages and socscistatistics calculators. P-value<0.05 was considered as statistically significant.

Results: A total of 50 subjects of sickle cell disease homozygous (SS) were studied for hematological parameters. The mean age±SD of 50 subjects were 13.3±9.24 years. Out of 50 subjects, 35 were males and 15 were females. Total RBC count, mean corpuscular volume (MCV), mean corpuscular haemoglobin (MCH) and mean corpuscular hemoglobin concentration (MCHC) was low in present study. Significant inverse correlation was found in females between HbA2 and HbF, p=0.01, while it was insignificant and negatively correlated in males being 0.23.

Conclusions: Sickle cell disease homozygous is a common and challenging health problem of Chhattisgarh population.

Keywords: Chhattisgarh, Hematological profile, Sickle cell disease

INTRODUCTION

Sickle cell disease hemoglobinopathy gets inherited in autosomal recessive pattern.¹ In sickle cell disease substitution of amino acid valine for glutamic acid at the sixth position on beta globin chain takes place resulting in sickled hemoglobin which is a hemoglobin tetramer. Sickled haemoglobin (HbS) which is produced is low soluble and polymerized when deoxygenated.² Sickle cell disease in India is characterized by higher levels of foetal hemoglobin (HbF), more frequent alpha thalassemia, higher total hemoglobin and lower retic counts and persistence of splenomegaly compared to Jamaican homozygous sickle cell disease.³ In contrast to this benign picture, studies from central India report severe disease (defined as >3 bone pain crises, >3 transfusions/year) in 30% children.⁴ In a study done for finding sickle cell anemia prevalence in Kanker, Dantewada and Raigarh districts of Chhattisgarh, the prevalence of sickle cell trait (HbAS) was 10.6% and of sickle cell disease (HbSS) and inconclusive band was 0.66%.³ There are scarcity of data in the current topic. With the above background, the present study was conducted to assess the haematological profile of sickle cell disease patients.
METHODS

A prospective observational study was conducted in the Sickle Cell Institute, Raipur, Chhattisgarh and Department of Pharmacology in collaboration with Department of Biochemistry, Pt. J.N.M. Medical College, Raipur, Chhattisgarh, India from February 2018 to June 2018. Study permission was obtained from Institutional Ethical Committee. Patients included were in the steady state for a long period of time without any symptoms related to sickle cell disease or any other diseases which could affect hematological parameters. Subjects transfused in the last three months were excluded. All the sickle cell disease patients are under hydroxyurea therapy at present. Blood samples were collected in the Sickle Cell Institute OPD, Raipur. Five ml of blood was collected in EDTA from patients. Complete blood count (CBC) was measured using hematology autoanalyzer. Quantitative assessment of HbS and HbF was done by high performance liquid chromatography (HPLC) technique, using variant II. Data was recorded on a predesigned proforma and was analyzed on Excel Sheet 2007. Student t test and Pearson Correlation Coefficient test was done on Statpages and Socsci statistics calculators. P value <0.05 was considered as statistically significant.

RESULTS

A total of 50 subjects of sickle cell disease homozygous (SS) were studied for hematological parameters. The mean age±SD of 50 subjects were 13.3±9.24 years. Out of 50 subjects, 35 were males and 15 were females. Male outnumbered females and the sex ratio was 2.3:1. The age of the subjects ranged from 2 years to 43 years. Mean age±SD of males were 12.25±9.99 years while mean age±SD of females were 15.73±6.90.

Hematological profile of study subjects are shown in the Table 1. There were more males as compared to females in our study. Total hemoglobin is low in sickle cell disease patients in present study, more so in females in comparison to males, although the difference is not statistically significant, p=0.61. Total RBC count, mean corpuscular volume (MCV), mean corpuscular haemoglobin (MCH) and mean corpuscular hemoglobin concentration (MCHC) were low in present study. In this study RBC count in males was 2.92±0.87, while in females it was 2.91±0.61, a significant positive correlation was found between total hemoglobin and total RBC count, p-value for male was <0.00001 i.e. significant and was 0.0023 in females Table 2. Mean MCV in males was 87.16±8.42 fl, while it was 83±11.86 fl in females, difference in male and females was insignificant, p value being 0.16. Correlation between MCV and total hemoglobin in both sexes were not statistically significant, p value for males was 0.08 and for females it was 0.96. Mean MCHC in males was 32.87±4.07 and in females it was 32.23±3.77 g/dl, with p value between the two sexes being 0.60, i.e. insignificant. Mean HbA2 values in males and females were 2.26±0.37 and 2.60±0.97% respectively.

Hematological parameters | Male (n=35) Mean±SD | Female (n=15) Mean±SD | Total (n=50) Mean±SD | P value (significant/insignificant)
--- | --- | --- | --- | ---
Hb (g/dl) | 8.18±1.94 | 7.88±1.95 | 8.09±1.93 | 0.61, Insignificant
Hct (%) | 24.84±6.29 | 24.38±5.03 | 24.70±5.89 | 0.80, Insignificant
RBC (million/mm^3) | 2.92±0.87 | 2.91±0.61 | 2.92±0.79 | 0.96, Insignificant
MCV (fl) | 87.16±8.42 | 83±11.86 | 85.92±9.65 | 0.16, Insignificant
MCH (pg) | 28.47±4.07 | 26.65±4.38 | 27.92±4.21 | 0.16, Insignificant
MCHC (g/dl) | 32.87±4.07 | 32.23±3.77 | 32.64±3.95 | 0.60, Insignificant
HbF (%) | 18.79±5.21 | 16.86±6.76 | 18.21±5.72 | 0.27, Insignificant
HbA2 (%) | 2.26±0.37 | 2.60±0.97 | 2.36±0.62 | 0.07, Insignificant

Table 2: Correlation between various hematological parameters in males and females.

<table>
<thead>
<tr>
<th>Hematological parameters</th>
<th>r</th>
<th>P value</th>
<th>Significant/insignificant</th>
</tr>
</thead>
<tbody>
<tr>
<td>Males</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hb vs RBC</td>
<td>0.7231</td>
<td>0.0023</td>
<td>Significant</td>
</tr>
<tr>
<td>Hb vs MCV</td>
<td>-0.0144</td>
<td>0.96</td>
<td>Insignificant</td>
</tr>
<tr>
<td>MCV vs HbA2</td>
<td>-0.6267</td>
<td>0.01</td>
<td>Significant</td>
</tr>
<tr>
<td>HbA2 vs HbF</td>
<td>-0.6265</td>
<td>0.01</td>
<td>Significant</td>
</tr>
<tr>
<td>MCV vs HbF</td>
<td>0.2197</td>
<td>0.43</td>
<td>Insignificant</td>
</tr>
<tr>
<td>Females</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hb vs RBC</td>
<td>0.8693</td>
<td>&lt;0.00001</td>
<td>Significant</td>
</tr>
<tr>
<td>Hb vs MCV</td>
<td>-0.2965</td>
<td>0.0842</td>
<td>Insignificant</td>
</tr>
<tr>
<td>MCV vs HbA2</td>
<td>-0.2081</td>
<td>0.2305</td>
<td>Insignificant</td>
</tr>
<tr>
<td>HbA2 vs HbF</td>
<td>-0.3321</td>
<td>0.05</td>
<td>Insignificant</td>
</tr>
<tr>
<td>MCV vs HbF</td>
<td>-0.0864</td>
<td>0.62</td>
<td>Insignificant</td>
</tr>
</tbody>
</table>

It was observed in present study that between MCV and HbA2 a significant negative inverse correlation exists in the females, p-value being 0.01 and an insignificant negative correlation was seen in the males, with p value was 0.23. Mean HbF value in males was 18.79±5.21 and it was 16.86±6.76 in females. No significant correlation between MCV and HbF in females, p value being 0.43 and in males, p value 0.62 was found in present study.
Significant inverse correlation was found in females between HbA2 and HbF, p value was 0.01, while it was insignificant and negatively correlated in males, p=0.23.

**DISCUSSION**

In the present study hematological profile of 50 cases of sickle cell disease patients are being reported from Chhattisgarh, India. Males outnumber females in 35:15 ratio. Male predominance could be due to the gender bias in our society, in which females pay less attention towards their illness and are discriminated from males. Age of females varied from 4 to 27 years and their mean±SD being 15.73±6.90 years. Age of males varied from 2 to 43 years and the mean±SD being 12.25±9.99 years.

Total mean±SD of haemoglobin (g/dl) in both the sexes was 8.09±1.93. When compared between males and females, it was 8.18±1.94 in males and 7.88±1.95 in females, p value being 0.61, insignificant (Table 1). The difference between hemoglobin levels of males and females were due to the androgenic effect in males and menstruation in females. Low total hemoglobin could be due to increased hemolysis and frequent recurrent infections in the sickle cell disease patients. Similar haemoglobin values were seen in studies done by Jadhav et al, Nagose et al, Rao et al and Boasiako et al, while in study by Kohchale et al it was low.6-10

Total Hct(g/dl) was 24.70±5.89, it was 24.84±6.29 in males and 24.38±5.03 in females, p value being 0.80, i.e. insignificant. Similar values of Hct was found in the studies of Boasiako et al, while higher hematocrit was found in the studies of Jadhav et al, Jawalkar et al, Rao et al and Nagose et al, while it was lower in study by Kohchale et al.6-11

Total RBC counts (million/mm³) was low in our study, it was 2.92±0.79. It was 2.92±0.87 in males and was 2.91±0.61 in females, p value being 0.96, insignificant. Similar values of RBC counts were seen in the study by Boasiako et al.9 Higher RBC counts were reported in studies by Jawalkar et al, Nagose et al, Jadhav et al and Rao et al, whereas lower RBC counts were seen in the study by Kohchale et al.6,8,10,11 Significant positive correlation was found in present study in both sexes between total RBC count and total hemoglobin, p value being 0.0023, in males and <0.00001 in females. These values were similar to the study by Shrikhande et al.12

Mean±SD of mean corpuscular volume (MCV), was 85.92±9.65 and it was 87.16±8.42 in males and 83±11.86 in females, p value between the two sexes being 0.16, insignificant. Similar values of MCV were seen in studies by Nagose et al and Boasiako et al.9 Lower MCV values were reported in the studies by Rao et al, Jawarkar et al and Kohchale et al whereas higher MCV values were reported by Jadhav et al.6,8,10,11 An insignificant and negative correlation was seen in present study in both sexes between haemoglobin and mean corpuscular volume, with a p value of 0.96 in males and 0.08 in females, similar were findings in study by Shrikhande et al.12 In sickle cell disease Vit B12 and folic acid are at a critical borderline levels and increased demands on erythropoiesis due to the chronic haemolytic process leads to macrocytosis and higher MCV. Lower MCV values in some of the studies could be due to coexisting Iron deficiency anemia and alpha thalassemia.6

Mean±SD of mean corpuscular hemoglobin (pg) (MCH) was 27.92±4.21, it was 28.47±4.07 in males and 26.65±4.38 in females. The p value between males and females came out to be 0.16, insignificant. Similar values of MCH was seen in the studies by Nagose et al, Boasiako et al, Rao et al and Jadhav et al.5,8 Lower values in MCH were reported by Kohchale et al and Jawarkar et al.9,10

Mean±SD of mean corpuscular hemoglobin concentration (MCHC) (g/dl) was 32.64±3.95, it was 32.87±4.07 in males and 32.23±3.77 in females, p value between females and males came out to be 0.60, again insignificant. Similar were values of MCHC was reported in studies by Nagose et al, Boasiako et al, Rao et al and Jawarkar et al.6,9,11 Higher MCHC were reported in the studies by Kohchale et al and lower MCHC was reported in study by Jadhav et al.8,10

Mean±SD of HbF (%) was 18.21±5.72, it was 18.79±5.21 in males and 16.86±6.76 in females. The p value of HbF between males and females came out to be 0.27 i.e. insignificant. Almost similar values were seen in the studies by Jadhav et al.6 Lower HbF (%) value was reported in study by Rao et al.8 An insignificant but positive correlation was seen in present study between MCV and HbF in males with p value being 0.43, whereas an insignificant but negative correlation was found in this study between MCV and HbF in females with p value being 0.62. Insignificant but positive correlation was reported from study by Shrikhande et al in both the sexes between MCV and HbF.12

Mean±SD of HbA2 was 2.36±0.62, it was 2.26±0.37 in males while it was 2.60±0.97 in females, p value being 0.07, insignificant. Significant and negative correlation was seen in present study between MCV and HbA2 in males with p value being 0.01. It was insignificant and negatively correlated in females in our study with p value being 0.23. Significant negative correlation was found in study by Shrikhande et al, in both sexes, thus in respect of males present study is similar to the study by Shrikhande et al.12 Insignificant and negative correlation was seen in our study between MCV and HbA2 in females with p value being 0.23, thus in this regards our study differs from study by Shrikhande et al.12 Significant and negative correlation was seen in males in present study between HbA2 and HbF with p value being 0.01, similar were findings in study by Shrikhande et al in males.12
Insignificant and negative correlation was seen in our study in females, between HbA2 and HbF, with p value of 0.05, while in study by Shrikhande et al they reported significant and negative correlation between HbA2 and HbF in females, thus our study differs from study by Shrikhande et al in this regards.12

CONCLUSION

Present study revealed low levels of hemoglobin, hematocrit and RBC count in sickle cell disease homozygous population of Chhattisgarh. A higher level of foetal hemoglobin was found in both sexes in present study. Difference in hematologic parameters was seen between male and female sickle cell disease homozygous population in our study, but the difference was not statistically significant. Large scale studies are required further in the area for guiding the clinicians for the better management of sickle cell disease and for defining a baseline hematologic profile of sickle cell disease in the region.

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Ethical approval: The study was approved by the Institutional Ethics Committee

REFERENCES
