Congenital methemoglobinemia misdiagnosed as polycythemia vera: Case report and review of literature

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Abstract

Methemoglobinemia is a rare overlooked differential diagnosis in patients presented with cyanosis and dyspnea unrelated to cardiopulmonary causes. Our patient is 29 year old Indian non-smoker male, his story started 6 months prior to presentation to our center when he had generalized fatigue and discoloration of hands. He presented with persistent polycythemia with elevated hemoglobin level. The patient was misdiagnosed in another center as polycythemia and treated with Imatinib. The diagnosis of PV was revisited and ruled out in view of negative JAK2, normal erythropoietin level and absence of features of pancytopenia. Clinical cyanosis and low oxygen saturation in the presence of normal arterial oxygen tension was highly suggestive of methemoglobinemia. Arterial blood gas revealed a methemoglobin level of 38% (normal: 0-1.5%). Cytochrome B5 reductase (Methemoglobin reductase B) was deficient at level of <2.6 U/g Hb) (normal: 6.6-13.3), consistent with methemoglobin (metHb) levels are as high as 40% of the total hemoglobin (Hb) value.1

Hereditary congenital methemoglobinemia due to deficiency of nicotinamide adenine dinucleotide (NADH) cytochrome b5 reductase enzyme is even extremely rare recessively inherited disorder that is not well documented in literature. Here, we report a rare case of methemoglobinemia presented with persistent polycythemia in absence of cardiopulmonary causes led to false diagnosis of polycythemia vera for which the patient needlessly undergone bone marrow examination and received Imatinib.

Introduction

Methemoglobinemia is a rare overlooked differential diagnosis in patients presented with cyanosis and dyspnea unrelated to cardiopulmonary causes. Methemoglobinemia is usually asymptomatic, even when methemoglobin (metHb) levels are as high as 40% of the total hemoglobin (Hb) value.1

Hereditary congenital methemoglobinemia due to deficiency of nicotinamide adenine dinucleotide (NADH) cytochrome b5 reductase enzyme is even extremely rare recessively inherited disorder that is not well documented in literature. Here, we report a rare case of methemoglobinemia presented with persistent polycythemia in absence of cardiopulmonary causes led to false diagnosis of polycythemia vera for which the patient needlessly undergone bone marrow examination and received Imatinib.

Case Report

Our patient is 29 year old Indian non-smoker male, his story started 6 months prior to presentation to our center when he had generalized fatigue and discoloration of hands (Figure 1). His CBC revealed high red cell count 6.7 (4.5-5.5 × 1012/L), elevated hemoglobin at 20 (13-17 g/dL) with normal leukocyte and platelets counts and unremarkable peripheral smear.

Based on his persistent polycythemia, the patient was evaluated in other center and diagnosed as polycythemia vera (PV), bone marrow examination revealed erythropoietin level and absence of features of pancytopenia. Although molecular studies revealed negative JAK-2 mutation; the patient was treated with Imatinib (Indian version) based on the assumed diagnosis of PV.

In our institute, the patient was re-evaluated; where was found to have dusky discoloration (cyanosis) of the face and both upper limbs of long standing history back to nearly age of thirteen but became more evident in last one year after treatment with Imatinib. He had no history of chest pain, syncope or palpitations with no history of medications intake or exposure to oxidant chemicals. There was no history of growth or developmental retardation or neurological symptoms and no hepatosplenomegaly.

Oxygen saturation by pulse oximetry showed saturation of 92%, while the patient showed no evidence of respiratory distress and his clinical examination was unremarkable.

The diagnosis of PV was revisited and ruled out in view of negative JAK2, normal erythropoietin level and absence of features of pancytopenia in the bone marrow biopsy, which was repeated in our center four months after stopping treatment with Imatinib.

The patients’ clinical picture including cyanosis in absence of any evidence of cardiopulmonary disease; raised the possibility...
of an alternative diagnosis including rare hemoglobinopathies such as hemoglobin M disease. Hemolysis work up including hemoglobinopathies screening by high performance liquid chromatography (HPLC) and manual hemoglobin electrophoresis revealed normal hemoglobin pattern.

Arterial blood gas showed the following results: pH 7.380 (7.35-7.45), PC O₂: 38 mmHg (35-45 mmHg), Normal P O₂ at: 96-101 mmHg (83-108), normal arterial O₂ saturation at 100% (95-99.0%), low HCO₃ 21.8 (23-29 mmol/L) and low oxygen saturation (O₂ Hb): at 62.4% (94-98%).

Clinical cyanosis and low oxygen saturation in the presence of normal arterial oxygen tension was highly suggestive of methemoglobinemia. Arterial blood gas (ABG) to check for methemoglobinemia performed and revealed a methemoglobin level of 38% (normal: 0-1.5%). Cytochrome B₅ reductase (Methemoglobin reductase B) was measured spectrophotometrically at Mayo Clinic Medical Laboratories; and came deficient at level of <2.6 U/g Hb (normal: 6.6-13.3), consistent with methemoglobin reductase (cytochrome b₅) deficiency and hence the diagnosis of congenital methemoglobinemia was established. Karyotyping was done to screen for chromosomal abnormalities and came normal. Other family members were asymptomatic with no family history of cyanosis or hemoglobin disorders. Family screening for methemoglobinemia cannot be conducted. Due to lack of oral tablets of methylene blue, the patient started on Vitamin C 500 mg once daily. After one month of treatment, the cyanosis became clinically less evident with significant reduction of methemoglobin concentration measured by ABG (methemoglobin levels 19.2%).

Discussion

Methemoglobin results from oxidation of ferrous iron to ferric iron within the heme moiety of hemoglobin.1 In healthy individuals, less than 1% of hemoglobin is present in the oxidized form (methemoglobin), which has limited ability to carry oxygen, however with increased oxygen affinity at the remaining binding sites.3 This results in decrease of oxygen delivery to tissue leading to hypoxemia and lactic acidosis.

Acquired methemoglobinemia caused by oxidizing agents and usually presents in older age groups is not uncommon; however, congenital deficiency of the methemoglobin reductase enzyme is extremely rare that only a few cases are reported in the English medical literature around the world. That is why the incidence and clinical characteristics of congenital Methemoglobinemia is basically unknown, due to underreporting and of these cases.

One possible hint to the diagnosis of methemoglobinemia is the presence of a saturation gap. This occurs when there is a difference between the SO₂ that has been measured by means of pulse oximetry (the lower value) and the SO₂ that has been calculated by means of arterial blood-gas analysis. Typically, this saturation gap is greater than 5% in cases of methemoglobinemia.3 The failure of very high oxygen saturation to correct cyanosis is very indicative of methemoglobinemia.

Although methemoglobin at level of 20% indicates enzyme deficiency, patients with congenital disease develop physiological compensatory mechanisms and can tolerate elevated levels of methemoglobin (up to 40%) without symptoms. Levels of >70% are usually incompatible with life.1 These adaptation mechanisms include changes in the concentration of 2,3-Diphosphoglycerate and pH, synthesis of globin chains, and secondary polycythemia.4,5

Compensatory elevation of hemoglobin concentration is observed in patients with recessive hereditary methemoglobinemia, because of the methemoglobinemia-induced left shift in the oxyhemoglobin dissociation curve.6 Congenital methemoglobinemia is further classified into two main types with one due to methemoglobin reductase enzyme deficiency and the other due to an abnormal oxygen affinity hemoglobin termed hemoglobin M.7 Methemoglobin reductase enzyme deficiency is either type I or type II; type I: Cytochrome b₅ reductase deficiency, demonstrable only in the erythrocytes, presents as uncomplicated, benign methemoglobinemia, and associated with a normal life expectancy with only fatigue and dyspnea being the most commonly reported symptoms.

Hereditary type II methemoglobinemia, is global (affects both red and white blood cells) and is associated with severe neurologic dysfunction and reduced life expectancy.8 The enzyme’s activity is less than 20% of normal.

Type II congenital methemoglobinemia constitutes approximately 10% of all cases of congenital methemoglobinemia and usually causes death within the first few years of life.9

There is a significant delay in the onset of symptoms in a patient with type I hereditary methemoglobinemia, as most studies indicate that patients are asymptomatic in infancy and childhood. There is a single reported case wherein the onset of symptoms in the patient started at the age of 8 years.10 With such late presentations, treating physicians would usually think of acquired methemoglobinemia, investigating exposure to an oxidative stress as the most likely cause of the findings.

Clinical cyanosis is diagnostically challenging as causes are multiple; especially in absence of cardiopulmonary causes. In the case reported here, the cyanosis was originally overlooked and hematology work-up was misdirected towards investigating polycythemia. In heterozygous cytochrome b₅ reductase deficiency, methemoglobin reductase activity is low, and the patient will have a lower threshold for acquired methemoglobinemia in response to exogenous oxidative stress.

The role of Imatinib in provoking methemoglobinemia is questionable and association between Imatinib and methemoglobinemia never described before. In our case, there were no other offending drugs in aggravating the patients’ symptoms and cyanosis.

When congenital methemoglobinemia is suspected, enzyme activity in all immediate family members should be evaluated. As
Table 1. Clinical characteristics of reported cases of Congenital Methemoglobinemia due to (NADH)-cyto b5 Reductase deficiency reported in English literature.

| Author [ref] | Age/Sex Nationality | Hb (gm/dL) | Presentation | Type | MetHb level | Family study | (NADH)-cyto b5 reductase | Management (MB); ascorbic acid (vit C); not mentioned (N/M); recurrent early pregnancy loss (REPL). |
|--------------|----------------------|------------|--------------|------|-------------|---------------|--------------------------|--------------------------------------------------------------------------------------------------|
| Londhe [10]  | 18/M Indian          | 19         | Cyanosis Headache Bilateral ptosis | I    | 37          | Pos (sister)  | 14 (30-35)               | Oral Vit C thrice/day Aspirin Phlebotomy                                                              |
| Shonola [11] | 24 days/M            | -          | Mild central cyanosis               | I    | 26          | Pos           | 4.2 (10.1-19.4)          | Oral MB (1.5) Vit C (5)                                                                  |
| Takishe [13] | 38/F Japanese        | -          | Cyanosis Convulsion                 | I    | 33          | Neg           | N/M                     | N/M                                                                                              |
| Yasmin [14]  | 21/Indian            | -          | Mild cyanosis exertional dyspnea    | I    | 29          | Neg           | Deficient N/M           | N/M                                                                                              |
| Canturk [15] | 20/M                 | -          | Cyanosis Bronchial asthma symptoms  | I    | 40          | Pos           | N/M                     | MB                                                                                               |
| Badawi [16]  | 16/ M Arabic         | 17         | Cyanosis fatigue dyspnea on exertion | I    | 40          | Pos           | <2.6 (6.6-13.3)         | Oral Vit C N/M                                                                                   |
| Ramanamurty [17] | 55/M Indian     | 15         | Weakness Syncope Cyanosis           | I    | 30          | Neg           | 16 (35.9±5.00)          | N/A                                                                                              |
| Dhiraj [18]  | 33/M Indian          | 18         | Incidental cyanosis                 | I    | 20-30       | Neg           | N/A                     | N/A                                                                                              |
| Gerli [19]   | 64/M Italian         | -          | Marked cyanosis                     | I    | 36          | Pos           | 9% normal value         | Oral Vit C                                                                                       |
| Prabhakar [20] | 1 month/M Indian  | 14         | Central cyanosis                    | I    | 29          | Pos           | 9.36 (30-40)           | MB                                                                                               |
| Prabhakar [20] | 32 y/F Mother Indian | 14       | REPL                                   | I    | 7.5 (n<1.5%) | Pos           | 20                      | Not needed                                                                                     |
| Prabhakar [20] | 34/ M Father Indian | 15         | No symptoms                           | I    | 2.8         | Pos           | 24                      | Not needed                                                                                     |
| Melanie [21] | F/English Mother     | -          | -                                      | I    | -           | Pos           | <0.11                   | -                                                                                               |
| Melanie [21] | English/ M Father    | -          | -                                      | I    | -           | Pos           | 6.37                    | -                                                                                               |
| Melanie [21] | English/ M           | -          | -                                      | I    | -           | Pos           | 7.75 (11.51-20.9)       | -                                                                                               |
| Vives [22]   | 2 year/F Spanish     | -          | Neurological symptoms                | II   | -           | Pos           | -                      | -                                                                                               |
| Yawata [23]  | M Japanese           | -          | Mental retardation skeletal anomalies | II   | -           | -             | RBC: 0.5-4 PLT (10-27), lymphocytes (18-31), fibroblast (50) | -                                                                                               |
| Prabhakar [24] | 5/F                   | 11         | Cyanosis cerebral palsy, Microcephaly generalized hypertonia | II   | 51          | Pos           | 6.0                      | MB (0.5) followed by vit C 500 twice a day                                                                 |
| Prabhakar [24] | 15/M Indian          | 9          | -                                      | II   | 13          | Pos           | 6.21                    | -                                                                                               |
| Prabhakar [24] | 8/M Indian           | 14         | -                                      | II   | 42          | Pos           | 9.60                    | -                                                                                               |
| Nellicka [25] | 1/F Indian           | -          | -                                      | II   | 24          | -             | Vit C high dose                   | Symptoms improved                                                                                     |

Methylene blue (MB); ascorbic acid (vit C); not mentioned (N/M); recurrent early pregnancy loss (REPL).
a result of autosomal recessive transmission. In heterozygous deficiency, methemoglobin reductase activity is low and hence heterozygotes will have a lower threshold for acquired methemoglobinemia in response to exogenous oxidative stress. However, the level of enzyme activity is not low enough to produce clinical disease under normal circumstances. 

This category of rare hemoglobin disease is underreported and often clinically missed. In 1943, the first description of familial idiopathic methemoglobinemia in the United Kingdom was reported in 2 members of one family.12

Upon extensive review of English literature for cases diagnosed as congenital Methemoglobinemia due to deficiency of cytochrome b5 reductase, we found 23 cases diagnosed as type I (including the case reported here). The details of the cases are listed in Table 1.10-25

Seventeen cases (~74%) of type I and 6 cases (27%) of type II. There is male predominance, 73% versus 26% in females. Almost half of reported cases 12 cases (52%) are Italian, 2 Japanese, 3 English, 2 Arabic, one case Spanish and one case Italian.

For type I, the median calculated age is 31 years with cyanosis and shortness of breath being the most common sign and symptoms.

For type II: Only six cases were reported in English literature, all in pediatric age group with median calculated age at presentation is 6 years with neurologic manifestations and mental retardation are type II associated symptoms.

The modalities of treatment vary between: Methylene blue (MB) alone, MB with Ascorbic acid or Ascorbic acid alone. Ascorbic acid either alone or in combination with MB was used in seven cases within this category with no response in one case,4 while the other six cases (including our case) responded well.

Conclusions

Due to lack of systematic epidemiological studies, congenital methemoglobinemia is under diagnosed as it is under investigated and usually overlooked especially when presenting in adulthood and in absence of obvious acquired agents.

In the present report, we highlight this clinically hidden case of methemoglobinemia which is misdiagnosed as polycythemia vera and led to needlessly patient’s exposure to invasive bone marrow procedures and even therapy with Inmatinib.

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