Characterization of beta-thalassemia mutations in patients from the state of Rio Grande do Norte, Brazil

Zama Messala Luna da Silveira1, Maria das Vitórias Barbosa1, Thales Allyrio Araújo de Medeiros Fernandes2, Elza Miyuki Kimura3, Fernando Ferreira Costa4, Maria de Fátima Sonati3, Ivanise Marina Moretti Rebecchi1 and Tereza Maria Dantas de Medeiros1

1Departamento de Análises Clínicas e Toxicológicas, Universidade Federal do Rio Grande do Norte, Natal, RN, Brazil.
2Departamento de Ciências Biomédicas, Universidade do Estado do Rio Grande do Norte, Mossoró, RN, Brazil.
3Departamento de Patologia Clínica, Faculdade de Ciências Médicas, Universidade Estadual de Campinas, Campinas, SP, Brazil.
4Hemocentro, Universidade Estadual de Campinas, Campinas, SP, Brazil.

Abstract

35 unrelated individuals were studied for characterization as either heterozygous or homozygous for beta-thalassemia. Molecular analysis was done by PCR/RFLP to detect the mutations most commonly associated with beta-thalassemia (β+IVS-I-1, β+IVS-I-6, and β39). In the patients who showed none of these mutations, the beta-globin genes were sequenced. Of the 31 heterozygous patients, 13 (41.9%) had the β+IVS-I-6 mutation, 15 (48.4%) the β0IVS-I-1 mutation, 2 (6.5%) the β+IVS-I-110 mutation and 1 (3.2%) the β+IVS-I-5 mutation. IVS-I-6 was detected in the four homozygotes. The mutation in codon 39, often found in previous studies in Brazil, was not detected in the present case. This is the first study aiming at identifying mutations that determine beta-thalassemia in the state of Rio Grande do Norte.

Key words: hereditary hemoglobinopathies, beta-thalassemia, mutations, PCR-RFLP, Brazilian population.

Received: November 30, 2010; Accepted: May 2, 2011.

Beta-thalassemia is a group of hemoglobin diseases caused by a reduction (β- thalassemia) or absence (β0 thalassemia) in the synthesis of beta-globin chains. More than 200 different types of mutations have been described as being responsible for this disease. Affected individuals can be heterozygous, compound heterozygous, or homozygous for beta-thalassemia, or even have interactions with other hemoglobinopathies. Their phenotypes include microcytic and hypochromic anemia, raised HbA2 levels, and various syndromes caused by the combination of β0 and β+ alleles (Thein, 1998; Weatherall, 2001).

The frequency of heterozygotes in Brazil is around 1% (Freitas and Rocha, 1983; Ramalho et al., 1999). Studies in the South and Southeast of Brazil have shown that the most frequent mutations are β039 (C→T) and β+ IVS-I-110 (G→A) (Martins et al., 1993; Reichert et al., 2008), whereas in the Northeast, an entirely different pattern was observed, the most frequently encountered allele being β+IVS-I-6 (T→C), followed by β+IVS-I-1 (G→A) (Araújo et al., 2003).

Due to the lack of information regarding the types of beta-thalassemia mutations encountered in the state of Rio Grande do Norte, an effort was made to characterize the disease in patients who were homozygous and heterozygous for beta-thalassemia, by way of hematological and molecular tests.

The sample consisted of 35 unrelated individuals (13 males and 22 females) with HbA2 levels above 3.5%. Age was between 1 and 70 years and all were born in the state of Rio Grande do Norte. The subjects were recruited from individuals referred to the Integrated Laboratory Clinical Analysis, at Rio Grande do Norte Federal University, between April, 2008 and October, 2009, by hematologists from the public and private sectors for the diagnosis of possible anemia. The study was approved by the Research Ethics Committee of Rio Grande do Norte Federal University (CEP-UFRN, protocol number 015/08) in accordance with
the standards laid down in the National Health Council resolution 196/96. All the participants or their legal guardians were informed of the aim of the study and signed a voluntary informed-consent form.

Samples of venous blood from each patient were collected in two sterile tubes, one with EDTA and the other without an anti-coagulant. The aliquot of blood containing EDTA was used for measurement of red blood cell indices (ABX Diagnostics, Montpellier, France), alkaline hemoglobin electrophoresis (Dacie and Lewis, 1995), measurement of HbA2 by elution (Bezerra, 1984), quantification of HbF by alkaline denaturation (Betke et al., 1959), and DNA extraction using the illustra blood genomicPrep Mini Spin commercial kit (GE Healthcare, Little Chalfont, Buckinghamshire, UK). The aliquot without anti-coagulant was used to measure serum ferritin by chemiluminescence (Immulite, Diagnostics Products Co., Los Angeles, CA, USA).

The mutations most commonly associated with the disease, viz., β0IVS-I-1 (G→A), β+IVS-I-6 (T→C) and β+39 (C→T), were investigated by PCR-RFLP. All the 39 thalassemia alleles investigated, 53.8% bore the β0IVS-I-6(T→C) mutation and 15.4% the β+IVS-I-1 (G→A) mutation, whereas in 3 (9.7%) no mutation could be identified by PCR-RFLP. DNA from the latter three patients was submitted to beta-globin gene sequencing, thereby revealing two to be heterozygous for the β+IVS-I-110 (G→A) mutation and one for the β+IVS-I-5 (G→C) mutation. All the four homozygous patients bore the β+IVS-I-6 (T→C) mutation.

Of the 39 thalassemia alleles investigated, 53.8% bore the IVS-I-6 mutation, 38.5% the IVS-I-1, 5.1% the IVS-I-110, and 2.6% the IVS-I-5 mutation.

Comparison of hematological analysis between thalassemia patients without iron deficiency and heterozygous for β0 IVS-I-6 (T→C) and β+ IVS-I-1 (G→A) mutations, revealed a significant difference as regards MCV (p = 0.023), MCH (p = 0.007), and Hb A2 (p = 0.001) quantification. Nevertheless, the comparison of laboratory analyses between patients heterozygous and homozygous for the IVS-I-6 mutation, revealed a statistically significant difference (p < 0.05) for all the parameters analyzed (Table 1).

Table 1 - Comparison of β+IVS-I-1 (G→A) and β0 IVS-I-6 (T→C) mutations found in this study with blood indices

<table>
<thead>
<tr>
<th>Blood indices</th>
<th>Type of mutation</th>
<th>p value (1,a)</th>
<th>p value (1,b)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Heterozygotes</td>
<td>Homozygotes</td>
<td>Heterozygotes</td>
</tr>
<tr>
<td></td>
<td>β+IVS-I-1 (N = 12)*</td>
<td>β+IVS-I-6 (N = 4)</td>
<td>β+IVS-I-6 (N = 9)*</td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>10.0 ± 1.0</td>
<td>7.8 ± 0.8</td>
<td>11.8 ± 1.1</td>
</tr>
<tr>
<td>Red blood cells (x10^12/L)</td>
<td>5.21 ± 0.71</td>
<td>4.39 ± 0.51</td>
<td>5.47 ± 0.31</td>
</tr>
<tr>
<td>MCV (fL)</td>
<td>63.3 ± 3.8</td>
<td>60.0 ± 5.8</td>
<td>68.0 ± 4.5</td>
</tr>
<tr>
<td>MCH (pg)</td>
<td>19.3 ± 1.7</td>
<td>17.9 ± 1.5</td>
<td>21.7 ± 2.0</td>
</tr>
<tr>
<td>Hemoglobin A (%)</td>
<td>93.3 ± 1.8</td>
<td>83.4 ± 5.0</td>
<td>95.1 ± 0.5</td>
</tr>
<tr>
<td>Hemoglobin A2 (%)</td>
<td>5.3 ± 0.6</td>
<td>5.7 ± 0.6</td>
<td>4.1 ± 0.2</td>
</tr>
<tr>
<td>Hemoglobin F (%)</td>
<td>1.4 ± 1.8</td>
<td>10.9 ± 5.2</td>
<td>0.8 ± 0.5</td>
</tr>
</tbody>
</table>

1) p value in the Student t-test for independent samples.
2) heterozygous for β+IVS-I-1 and β0 IVS-I-6 mutations.
3) heterozygous and homozygous for the β+IVS-I-6 mutation.
4) Seven patients (3 heterozygotes with the IVS-I-1 mutation and 4 with the IVS-I-6 mutation) with iron deficiency or whose serum ferritin levels had not been measured, were excluded.
Orkin et al. (1982) identified the pattern of beta-thalassemia mutations in individuals of Mediterranean origin, wherein $\beta^+\text{IVS-I-1}10 (G \rightarrow A)$, $\beta^3\text{IVS-I-1} (C \rightarrow T)$, $\beta^+\text{IVS-I-1} (G \rightarrow A)$, and $\beta^+\text{IVS-I-6} (T \rightarrow C)$ mutations were the most common. In Portugal, studies have shown that the most frequently found mutations are $\beta^3\text{IVS-I-1} (C \rightarrow T)$, $\text{IVS-I-1} (G \rightarrow A)$ and $\text{IVS-I-6} (T \rightarrow C)$, with frequency varying in accordance with the region (Tamagnini et al., 1983; Cabeda et al., 1999; Faustino et al., 1999).

In Brazil, the pattern of beta-globin mutations varies according to the region. In the South and Southeast, the $\beta^3\text{IVS-I-1} (C \rightarrow T)$ and $\beta^+\text{IVS-I-1}10 (G \rightarrow A)$ mutations are very frequent (Martins et al., 1993; Bertuzzo et al., 1997; Fonseca et al., 1998; Reichert et al., 2008), whereas in the Northeast, the most frequent mutations are $\beta^+\text{IVS-I-6} (T \rightarrow C)$ and $\beta^\text{+IVS-I-1} (G \rightarrow A)$ (Araújo et al., 2003).

This is the first study to determine the profile of beta-thalassemia mutations in the population of Rio Grande do Norte State. Through PCR/RFLP analysis, IVS-I-6 and IVS-1-1 mutations were detected in 36 (92.3%) of the 39 thalassemia alleles analyzed. The remaining three (7.7%) were characterized by beta-globin gene sequencing.

The Brazilian population is one of the most heterogeneous in the world, due to five centuries of interethnic crossing of peoples from three continents, namely, European colonizers, mainly represented by the Portuguese, African slaves and autochthonous Amerindians (Reichert et al., 2008). As regards European immigration, it is estimated that about 500,000 Portuguese arrived in the country between 1500 and 1808. Significantly, in the approximately 100-year-period from 1872 to 1975, Brazil received ever increasing numbers of immigrants from various parts of the world, viz., Italians (34%), Portuguese (29%), Spanish (14%), Japanese (5%), Germans (4%), Lebanese and Syrians (2%), and others (12%) (Pena et al., 2009). The population of Rio Grande do Norte is the result of miscegenation between Amerindians, sub-Saharan Africans, and European colonizers, the African influence having been insignificant. Of the Europeans, the Portuguese exerted the greatest influence, followed by the French. Although the Dutch were also present in the state, their contribution to the genetic makeup of the population was not significant (Cascudo, 1980).

The high frequency of IVS-I-6 (53.8%) and IVS-I-1 (38.5%) mutations appears to have resulted from the Portuguese contribution to the genetic makeup of the population of Rio Grande do Norte. Both these mutations were also frequent in Pernambuco, thereby demonstrating the thus far heterogeneity of beta-thalassemia in Brazil (Araújo et al., 2003).

In the present study, the IVS-I-110 (G \rightarrow A) mutation was identified in two out of the three patients whose $\beta$ genes were sequenced. In Brazil, this mutation was the most commonly found in the southern and southeastern region (Martins et al., 1993; Bertuzzo et al., 1997; Reichert et al., 2008). The $\beta^+\text{IVS-I-1}5 (G \rightarrow C)$ mutation, found in the third patient, and which could not be characterized by PCR-RFLP, was also encountered, with a frequency of 9.3%, in a population studied by Araújo et al. (2003). This mutation is very common in Southeast Asia, especially Malaysia and Indonesia, as well as in various regions of India (Thein, 1998).

PCR-RFLP, through its usefulness in identifying the most common beta-thalassemic alleles in the population studied, represents a practical alternative in situations where sequencing is unavailable. As there are no records to date of the types of beta-thalassemia mutations found in Rio Grande do Norte, and as studies have shown that the pattern of such mutations varies according to ethnic influence in the different regions of Brazil, the authors believe this study will play an important role in acquiring a greater understanding of the molecular profile of beta-thalassemia in Brazil.

Acknowledgments

Research was supported by the Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq, Grant n. 409766/2006-2) and the Fundação de Amparo à Pesquisa do Estado de São Paulo (FAPESP, grant n. 2008/57441-0).

References


