MOLECULAR STUDY ON BETA-THALASSEMIA CHILDREN PATIENTS IN A PORTION OF THE ALGERIAN POPULATION (NORTHEAST ALGERIA)

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Objective:- This study was planned to determine the frequency of ß-thalassemia mutations in Batna region (northeast Algeria).

Materials and Methods:- 19 blood samples of clinically thalassemic children patients were collected from department of pediatrics, university hospital of Batna. We carried out the molecular genetics of beta globin gene by the method of minisequencing using Snapshot™ kit (Applied Biosystems) in search of the four most common HBB genetic variants including three ß-thalassemia mutations: codon 39(C>T) (HBB: c.118C>T), IVSI-110(G>A) (HBB: c.93-21G>A) and IVSI-1-2(T>G) (HBB: c.92+2T>G), as well as the hemoglobin S variant (HBB: c.20A>T) and we used direct DNA sequencing to detect the rare mutations of beta globin gene.

Results:- We have revealed the presence of four different ß-globin gene mutations responsible for ß-Thalassemia in region of Batna. According to our results; the nonsense mutation at codon 39 (C>T), is the most frequent mutation type in our province the same as other geographical regions of Algeria, followed by codon 54(T), this molecular lesion was detected in a second Algerian family; the proband was homozygote, and the first association of Hb Knossos: codon 27 (G>T) allele with codon 39 (C>T) in Algerian population. Here we report also association of codon 39(C>T) with IVS-I-110 (G>A).

Conclusion:- our preliminary results show the heterogeneity of the beta-thalassemia mutations in the region of Batna.

Introduction:-
Beta-thalassemia is a recessive monogenic disorder encountered worldwide with a higher prevalence among Mediterranean, Middle Eastern and Indian populations (Weatherall DJ, 1991). The disease is due to mutation in beta globin locus for which more than 200 alleles have been reported (Weatherall DJ and Clegg JB, 2002). Numerous disorders of the β-globin chain of hemoglobin lead to different disease phenotypes.(Thein SL, 2008; Weatherall DJ and Clegg JB , 2001) Of these, ß-thalassemia is a subset of the hemoglobinopathies characterized by a hereditary...
anemia with a wide phenotypic spectrum that can have significant morbidity and mortality. (Weatherall DJ and Clegg JB, 2001; Rund D and Rachmilewitz E, 2005). The β-thalassemia exhibit a range of severities, each corresponding to an absence or reduction of β-globin protein synthesis. These β-thalassemia phenotypes are related to the myriad mutations that affect the β-globin gene (HBB) on chromosome 11p15.5, and different populations have their own mutation spectrum (Thein SL, 2002). In Algeria, the frequency of β-thalassemia gene is 3% (Belhani M, 2009); these diseases are a real public health problem often compounded by rate inbreeding of the population (30-32%) (Bellis G et al, 2001). Previous investigations have disclosed a high molecular heterogeneity of β-thalassemia (Bennani C et al., 1993; Bennani C et al., 1994). This study aims to describe the mutation spectrum from a sample of β-thalassemia patients and from β-thalassemia carriers. In addition the present study was performed to assess the usefulness of the minisequencing technique as an alternative strategy for genetic diagnosis of HBB gene disorders as a screening technique for the detection of unknown β-globin gene mutations.

Materials and Methods:–
This study was realized in β-thalassemia homozygous children and from β-thalassemia carriers (age brackets 4 to 16 year old of male and female sex). These subjects are from the region of Batna, cared in the pediatric ward of the University Hospital Batna. Venous blood samples of 2.5 ml volume were drawn from the study subjects and were collected in EDTA anticoagulant containers. Blood withdrawals were performed a few minutes before the regular blood transfusion for patients homozygous whereas heterozygous children the sampling is performed during family investigations.

DNA Extraction:–
The molecular analysis of the HBB gene was carried out after taking informed written consent from all the parents of the minors. Genomic DNA was extracted from peripheral blood leukocytes using the FlexiGene-DNA Kit (Cat # 51206; Qiagen Inc., Valencia, CA, USA) and stored at 4 °C.

Minisequencing reaction of HBB gene:–
The minisequencing assay was developed for the detection of the four most common HBB genetic variants including three β-thalassemia mutations: codon 39(C>T) (HBB: c.118C>T), IVS-I-110(G>A) (HBB: c.93-21G>A) and IVS-I-2(T>G) (HBB: c.92+2T>G), as well as the hemoglobin S variant (HBB: c.20A>T). To detect these four mutations, an allele specific PCR was performed, followed by highly multiplexed minisequencing reaction. The specific primer sequences of the HBB gene and PCR conditions are available upon request. Polymerase chain reaction products were purified using QIAquick PCR Purification by Kit (Qiagen Inc.). Purified fragments were used as template in a primer extension reaction containing the mutation-specific primer cocktail (see Table I).

For the extension reaction, we used the SNapshot Multiplex Kit (Applied Biosystems, Foster City, CA, USA), following manufacturer's instructions. After extension, the samples were treated with shrimp alkaline phosphatase according to the manufacturer protocol. Multiplex minisequencing products were resolved by automated capillary electrophoresis ABI PRISM 310 Genetic Analyzer (Applied Biosystems). Briefly, 12 ml of HiDi ™ formamide and 0, 5 ml size GeneScan 120 LIZ-calibrator (Applied Biosystems) were added to 1 ml of multiplex minisequencing product. The mixture was denatured at 95 °C for 3 min. next transferred to ice for 2 min. and loaded on an ABI PRISM ® 310 Genetic Analyzer capillary.

Table I:– Primers for multiplex minisequencing analysis.

<table>
<thead>
<tr>
<th>Investigated mutations</th>
<th>Minisequencing primers (sequences in 5’ &gt;3’ direction)</th>
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<tbody>
<tr>
<td>HbS (HBB: c.20A&gt;T)</td>
<td>T(45) ATG GTG CAC CTG ACT CCT G</td>
</tr>
<tr>
<td>IVS-I-2 (T&gt;G) (HBB: c.92+2T&gt;G)</td>
<td>T(55) GTG AGG CCC TGG GCA GG</td>
</tr>
<tr>
<td>IVS-I-110 (G&gt;A) (HBB: c.93-21G&gt;A)</td>
<td>T(65) ACT GAC TCT CTC TGC CTA TT</td>
</tr>
<tr>
<td>Codon 39 (C&gt;T) (HBB: c.118C&gt;T)</td>
<td>T(75) GTG GTC TAC CCT TGG ACC</td>
</tr>
</tbody>
</table>

* The variants are described using Human Genome Variation Society nomenclature.
Figure 1: GeneScan analysis of the multiplex minisequencing reaction. Electropherograms: (a) shows the analysis results of a HBB wild-type sample; (b) illustrates the pattern of peaks for all mutant positions in the heterozygous state; (c) illustrates the pattern of peaks for all mutant positions in the homozygous state.

Direct DNA sequencing:
If the minisequencing technique presents a snapshot normal, we use the direct DNA sequencing. The β-globin gene was amplified using couples of primers: HBB F: 5' - CTG ACA CAA CTG TGT TCA CT-3' and HBB R: 5' - TTC ACC TTA GGG TTG CCC -3'.

The β-thalassemia mutation was identified by automated sequence analysis performed on an ABI Prism 310 Genetic Analyzer (Applied Biosystems, Foster City, CA, USA) using the fluorescent dideoxy-termination method (Big Dye-Terminator Cycle Sequencing Kit, Applied Biosystems, Foster City, CA, USA).

Results:
Here we apply the minisequencing technique as an alternative strategy for genetic diagnosis of HBB gene disorders in children with β-thalassemia, provided by the pediatrics department, Hospital Batna (Northeast of Algeria). We chose this method as it allows quick search of the 4 most common and frequent mutations of the HBB gene. The GeneScan electropherograms of our subject’s samples after multiplex minisequencing primers are shown in figures 2, 3, 4 and 5 and the results of direct DNA sequencing are shown in figures 6 and 7.
Table II shows the results of molecular diagnosis for the 38 chromosomes from 9 individuals with beta-thalassemia trait and 10 children with beta-thalassemia major analyzed with 4 different beta-thalassemia mutations. This study confirms the observations that the frequency of several mutations varies from one ethnic group to another. The four different beta-thalassemia mutations have been identified in this study, were codon 39 (C>T) the most frequent beta-thalassemia mutations. In addition, two genetic variants without disease association, the polymorphisms codon 39 (C>T) and IVS-I-110 (G>A), a first association of Hb Knossos: HBB: c.82 G>T with HBB: c.118 C>T mutation causes thalassemia homozygous in the Algerian population, was found in one subject. Codon 54(-T) mutation was found in one subject who was homozygous for this molecular lesion.

**Table II:** Spectrum of beta-thalassemia mutations in northeast Algeria (Batna).

<table>
<thead>
<tr>
<th>HBB mutation name or variant</th>
<th>Phenotype</th>
<th>Localization at HBB</th>
<th>Genomic variation (HGVS)</th>
<th>No. of Alleles</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Codon 39 (C&gt;T)</td>
<td>β0</td>
<td>Exon 2</td>
<td>c.118C&gt;T</td>
<td>32</td>
<td>84.21</td>
</tr>
<tr>
<td>IVS-I-110 (G&gt;A)</td>
<td>β+</td>
<td>Intron 1</td>
<td>c.93−21G&gt;A</td>
<td>3</td>
<td>7.89</td>
</tr>
<tr>
<td>Codon 27 (G&gt;T) Hb Knossos</td>
<td>β+</td>
<td>Exon 1</td>
<td>c.82G&gt;T</td>
<td>1</td>
<td>2.63</td>
</tr>
<tr>
<td>Codon 54(-T)</td>
<td>β0</td>
<td>Exon 1</td>
<td>c.165delT</td>
<td>2</td>
<td>5.26</td>
</tr>
</tbody>
</table>

**Figure 2:** Electropherogram shows a peak for codon 39(C>T) mutation in the heterozygous state.

**Figure 3:** Electropherogram shows a peak for codon 39(C>T) mutation in the homozygous state.

**Figure 4:** Electropherogram shows a peak for codon 39(C>T) mutation and IVS-I-110 (G>A) in the heterozygous composite state.

**Figure 5:** Electropherogram shows a peak for IVS-I-110 (G>A) in the heterozygous state.
Figure 6:- Part of electropherogram obtained by sequencing the genomic DNA from the beta-thalassemic patient shows the presence of codon 27 (G> T) mutation (Hb Knossos).

Figure 7:- Part electropherogram obtained by sequencing the genomic DNA from the homozygous beta-thalassemia patient shows the presence of c.165delT mutation.

Discussion:-
This is the first study investigating the molecular level basic of β-thalassemia in the region of Batna (Northeast Algeria). We conducted the identification and characterization of the molecular basis of β-thalassemia among children born in Batna region.

The β-thalassemia in Algeria, are by their frequency and severity of a health problem, mainly in transfusion support (Addour NB, 2008). According (Lemsaddek W et al., 2004) the prevalence of β-thalassemia allele in North Africa would increase from west to east Mediterranean countries (Morocco 0, 94%, 1, 4% of Algeria, Tunisia 3% and Egypt 4, 5%). (Barragan E et al., 2006, Tadmouri A et al., 2001).
The great heterogeneity of molecular defects at the origin of β-thalassemia in Algeria, the number of β-thalassemia mutations in Algeria is 25 mutations (Addour NB, 2008). On the other hand, the Algerian population is characterized by four dominant mutations, which represent over 80% of β-thalassemia alleles. These are the mutation nonsense codon 39 C → T; IVS-I-110 substitution G → A; frashift the codon 6 (-A) and mutation IVS-I-1 G → A (16).

The nonsense mutation codon 39 (C>T) is widespread in Algeria with a frequency 25, 94% (Addour NB, 2008), 27,6% (Bennani C, Bouhass R Perrin- Pécontal P et al., 1994) and is more common in the west and decreases in center to be predominant in the East (Bouhass R et al., 1993).

The IVS-I-110 was discovered in 1981 (Spritz RA, Jagadeeswaran P, Choudary PV et al., 1981; Westaway D and Williamson R., 1981) and is caused by the replacement of a guanine by adenine in the consensus sequence, located in the first intron of the β-globin gene 19 nucleotides away from the site of splicing AG. (Orkin SH et al., 1982). The IVS-I-110 mutation found in Turkey (Perrin et al., 1998), represents 40% beta thalassemia alleles; it is predominant in central Algeria and located in a low frequency in the west (16%) (Bennani C et al., 1994). His consistent distribution with the extent of the Ottoman Empire between the 16th and 19th century. In Tunisia, it is (21%) (22 Fattoum S et al., 2004) and Egypt (26%) (Jiffri EH et al., 2010). It is rare in Morocco (3.2%) (Lemsaddek W et al., 2004). For Codon 54(-T) mutation: this is the second Algerian family which presents this mutation; the proband is homozygote also in a swedish family. The deletion of T from codon 54 result in framshift with a nonsense codon at codon 60 (TGA) and premature termination of translation (Landin BA, 1996). In this study, we detect a rare hemoglobin variant caused by a mutation in beta-globin gene, HBB: c.82G>T (Hb Knossos), which produces the classical phenotype of intermedia beta--thalassemia in association with HBB: c.118C>T) mutation causes thalassemia homozygous in an Algerian children patient (Fessas PH et al., 1982; Baklouti F et al., 1986). Hb Knossos in the heterozygous state has been recently recognized as the underlying abnormality in atypical beta-thalassemia trait. (Fessas Ph et al., 1982; Arous N, et al., 1982). First identified in a family from Crete, Hb Knossos was discovered soon afterwards in two families from northeast Algeria and in a family from the French West Indies. (Rouabhi F et al., 1983; Morl L et al., 1984).

Conclusion:
In this study, we used the minisequencing assay as a rapid screening procedure to identify four most common HBB genetic variants including three beta-thalassemia mutations and direct DNA sequencing to detect the rare mutations of beta-globin gene. Our data show the complexity of the beta-thalassemia mutations in our area due to the historical aspects and geographical location of region of Batna. This study confirms the heterogeneity of the beta-thalassemia mutations in Algerian population. Four different β-thalassemia mutations have been identified in the Batna population. Codon 39 (C>T) is the most frequent mutation type in our province; followed by codon 54(-T) and the first association of Hb Knossos: codon 27 (G>T) with codon 39 (C>T) in Algerian population. Here we report also association of codon 39 (C>T) with IVS-I-110 (G>A). Molecular genetic testing will be increasingly important in the future because it is anticipated that the advances in the understanding of the molecular basis of the disease may lead to specific pharmacologic therapies in the future.

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Conflict of Interest:
The authors of this paper have no conflicts of interest, including specific financial interests, relationships, and/or affiliations relevant to the subject matter or materials included.
References:


