Significant association between IRF6 820G→A and non-syndromic cleft lip with or without cleft palate in the Thai population

C Srichomthong, P Siriwan, V Shotelersuk


Background: Previous data have shown an association between DNA sequence variants in the IRF6 gene and an increased risk of non-syndromic cleft lip with or without cleft palate (CL/P) in some populations.

Objective: To investigate Thai CL/P patients and relatives for a 820G→A polymorphism.

Subjects: 192 CL/P Thai patients, 177 of their mothers, 73 of their fathers, and 278 controls.

Results: There were significant differences in the frequency distributions of both genotypes (p = 0.02) and alleles (p = 0.04) among probands as compared with the control group. The odds ratio calculated for the patients having the GG genotype compared with the other two genotypes (GA and AA) was 1.67 (95% confidence interval, 1.13 to 2.47). This pattern is consistent with a recessive effect of the G allele.

No association between any of the parents' genotypes and CL/P was found. The IRF6 820G→A was responsible for 16.7% of the genetic contribution to CL/P.

Conclusions: The findings confirm that IRF6 820G→A is associated with CL/P.

METHODS

The subjects of this study were 161 sporadic cases of non-syndromic CL/P. In addition, 31 cases were collected with a positive family history, bringing the total families in this study to 192. All subjects were self identified as Thai or Thai-Chinese. These groups are further characterised by type, laterality, severity, and sex, as shown in table 1. We also collected 177 of their mothers, 73 of their fathers, and 278 controls. Fifty three were complete trios. All of the patients were studied under the auspices of the Thai Red Cross, a national charity organisation devoted to providing clinical care for the poor. Subjects were recruited between 2000 and 2004 from 10 centres in Thailand (Nakornratchasima, Nan, Uthaithalanee, Maehongsorn, Trang, Sraakew, Kalasin, Nongkhai, Mahasarakam, and Bangkok). As preoperative evaluations, all patients were screened for the presence of associated anomalies or syndromes by a geneticist (VS), and only those determined to have isolated cleft lip with or without cleft palate were included in the study.

Blood samples for DNA analysis were obtained at the time of blood typing and determination of packed cell volume. The control samples were blood donors, self identified as Thai or Thai-Chinese, with no oral clefts, who denied history of oral clefts in other family members, in Nakornratchasima, Kalasin, Nongkhai, and Bangkok, collected over the same period.

The study was approved by the institutional review board of the Faculty of Medicine of Chulalongkorn University, and written informed consent was obtained from each person of blood typing and determination of packed cell volume. The study was approved by the institutional review board of the Faculty of Medicine of Chulalongkorn University, and written informed consent was obtained from each person included in the study.

DNA was extracted by phenol chloroform extraction protocol and was amplified using the polymerase chain reaction (PCR) with primers IRF6E7F: 5'AGTGGCCCTTCCAGATGCTG-3' and IRF6E7R: 5'CCTGACCTCCTCCAGACTAA-3', giving a PCR product of 647 bases pairs (bp). Genotyping for the IRF6 820G→A polymorphism was carried out by restriction digestion of PCR products with DpnII (New England Biolabs, Beverly, MA) and taqI (New England Biolabs, Beverly, MA).

Abbreviations: CL/P, cleft lip/palate

<table>
<thead>
<tr>
<th>Characteristics of the patients with non-syndromic cleft lip with or without cleft palate</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
</tr>
<tr>
<td>No of probands</td>
</tr>
<tr>
<td>Sporadic</td>
</tr>
<tr>
<td>Familial</td>
</tr>
<tr>
<td>Laterality</td>
</tr>
<tr>
<td>Right side cleft</td>
</tr>
<tr>
<td>Left side cleft</td>
</tr>
<tr>
<td>Bilateral cleft</td>
</tr>
<tr>
<td>Severity</td>
</tr>
<tr>
<td>Complete cleft</td>
</tr>
<tr>
<td>Incomplete cleft</td>
</tr>
<tr>
<td>Sex</td>
</tr>
<tr>
<td>Male</td>
</tr>
<tr>
<td>Female</td>
</tr>
</tbody>
</table>

**Table 1**

**Abbreviations:** CL/P, cleft lip/palate
Massachusetts, USA), according to the company’s recommendations. DPnII digests the G variant to five fragments (322, 177, 80, 35, and 33 bp). The A variant adds another restriction site, causing the 322 bp to be digested into two smaller fragments of 235 and 87 bp (fig 1).

Statistical analysis
Standard 2 and p values were calculated by a program available at http://www.unc.edu/~preacher/chisq/chisq.htm. Odds ratio and 95% confidence intervals (95% CI) were calculated from the Epi Info 2000 program. The transmission/disequilibrium test (TDT) analysis was carried out on subjects with heterozygous informative parents. Data from families with one parent missing were excluded. The TDT data were analysed using a k-1/k correction (where k is the number of alleles).7

Attributable risk
We estimated the attributable risk (AR) for the G allele and the GG genotype according to the formula AR = {(P[E/D])(RR−1)}/RR, using the odds ratio as an estimate of the relative risk.5

RESULTS
The observed frequencies of the 820G and 820A alleles, and the GG, GA, and AA genotypes in affected subjects, their mothers, their fathers, and controls are given in table 2. The observed distribution of genotypes among controls was compared with that expected according to Hardy–Weinberg equilibrium: no difference was found (χ² = 0.14, df = 2, p = 0.93). All genotype frequencies of the patients and their parents also followed Hardy–Weinberg equilibrium (data not shown).

The distributions of genotypes and alleles among patients and their parents were compared with those among controls: differences for both genotypic (p = 0.02) and allelic (p = 0.04) distributions between patients and controls were found. There were no differences of the distributions between the patients’ parents and controls.

Analyses were carried out to estimate the risk associated with each genotype. The results of the GG genotype compared with the other two genotypes, calculated as odds ratios and 95% CI are reported in table 2. A significantly higher frequency of the 820GG genotype was observed in the CL/P patients compared with the controls, with an odds ratio of 1.67 (95% CI, 1.13 to 2.47). The odds ratios in the parental groups were not increased. No association was found with the GA and AA genotypes (data not shown).

TDT analysis was carried out on subjects with heterozygous informative parents, but it showed no evidence for association of CL/P, as shown in table 3.
For the comparison of the subjects with CL/P with population based controls, as shown in tables 4 and 5, the estimated attributable risk for the G allele and the GG genotype were 16.69% and 19.47%, respectively.

**DISCUSSION**

CL/P is a multifactorial disorder caused by a combination of genes and environmental interactions. Each factor may contribute differently to CL/P in different populations. In the Thai population, we have previously observed a significantly higher frequency of the 677CT/1298AC genotype in the mothers of CL/P patients compared with controls, with an odds ratio of 4.43 (95% CI, 1.33 to 15.10). The observation was reinforced by other studies in different populations.

In the current study, we have established an IRF6 genotype in 192 CL/P patients, their parents (177 mothers and 73 fathers), and 278 controls.

The frequency of the 820G allele in our controls was 61%, which is comparable to the frequencies of 58–66% in other east Asian (China and Japan) and southeast Asian populations (Cambodia). However, the frequency of this variant is very different among ethnic groups, with the frequency of the G allele in African of 100% and in European of 90–100%.

When we analysed the genotype of our CL/P patients and their parents, we found the odds ratio calculated for the patients having the GG genotype—compared with the other two genotypes (GA and AA)—was 1.67 (95% CI, 1.13 to 2.47). The GA and AA genotypes were not associated with clefting (data not shown). This pattern is consistent with a previous finding of a recessive effect of the G allele.

Our observation of the recessive effect was further supported by the finding of a significant difference in the frequency distributions of both genotypes (p = 0.02) and alleles (p = 0.04) among probands as compared with the control group; the frequencies of the GG genotype and the G allele were increased, and the frequencies of the GA and AA genotypes and the A allele were reduced in CL/P patients as compared with unaffected subjects (table 2). No association between any of the parents’ genotypes and CL/P was found. This is different from variation in the MTHFR gene, in which the maternal genotype is associated with CL/P, rather than the affected subjects themselves.

Of our 192 CL/P patients, 177 of their mothers, and 73 of their fathers, only 53 were complete trios. Transmission disequilibrium calculation was carried out on subjects with heterozygous informative parents. We found a statistically non-significant trend toward a positive association between cleft lip and the G allele (table 3), consistent with a previous study in the Indian and European groups. The statistical non-significance may be explained by the small numbers of the complete trios, reflecting in the small number of heterozygous informative parents in our studies. Therefore, our findings require additional investigation before they can be extrapolated to a clinical setting.

The estimated attributable risk for the G allele of CL/P of 16.69% for the IRF6 820G allele in the Thai population is in accordance with the attributable risk of 11.6% in the Filipino population. Although the IRF6 820A allele is very rare or absent among Europeans (inhibiting the estimation of the attributable risk of the polymorphism), a recent study in an Italian population showed a strong evidence of linkage disequilibrium between polymorphisms at the IRF6 locus and non-syndromic CL/P.

These findings confirm the contribution of the gene in the aetiology of non-syndromic CL/P in several populations.

The IRF6 820G→A is a substitution of an isoleucine for an evolutionarily conserved valine residue at codon 274 in a protein binding domain, termed SMIR (Smad-interferon regulatory factor binding domain). Although the exact functions of IRF6 are still unknown, previous data suggest it plays a role in the transforming growth factor β signalling pathway.

In summary, our findings confirm that IRF6 G→A is associated with CL/P. This improves our understanding of the birth defect, has implications for the genetic counselling of families with children who have cleft lip, and provides a lead for identifying other genes or environmental factors linked to CL/P.

**ACKNOWLEDGEMENT**

We wish to thank the patients and their families for their participation in the study, and the medical staff of the Thai Red Cross, the Craniofacial Centre of King Chulalongkorn Memorial Hospital, and the Provincial Hospitals of Nakornratchasima, Nan, Uthai than, Maehongsorn, Trang, Sra Kaew, Kalasin, Nong Khai, and Mahasarakam for their excellent care of their patients. We are also indebted to professors Jeffrey C Murray and Mary L Marazita for their helpful comments and suggestions. This study was supported by the Research Unit Grant from Chulalongkorn University, the National Centre for Genetic Engineering and Biotechnology, and the Thailand Research Fund.

**Authors’ affiliations**

C Srichomthong, V Shotelersuk, Division of Medical Genetics and Metabolism, Department of Paediatrics, Faculty of Medicine, Chulalongkorn University, Bangkok, Thailand

P Siriwan, Division of Plastic Surgery, Department of Surgery, Faculty of Medicine, Chulalongkorn University

Competing interests: none declared

Correspondence to: Dr Vorasuk Shotelersuk, Head of Division of Medical Genetics and Metabolism, Department of Paediatrics, Sor Kor Building 11th floor, King Chulalongkorn Memorial Hospital, Bangkok 10330, Thailand; vorasuk.s@chula.ac.th

Received 18 February 2005

Revised version received 16 March 2005

Accepted for publication 16 March 2005

**REFERENCES**


Significant association between \textit{IRF6} 820G$\rightarrow$A and non-syndromic cleft lip with or without cleft palate in the Thai population

C Srichomthong, P Siriwan and V Shotelersuk

\textit{J Med Genet} 2005 42: e46
doi: 10.1136/jmg.2005.032235