Research paper

Genetic polymorphism of MTHFR C677T and premature coronary artery disease susceptibility: A meta-analysis

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A B S T R A C T

The association between 5,10-methylenetetrahydrofolate reductase (MTHFR) C677T gene polymorphism and premature coronary artery disease (PCAD) is controversial. To explore a more precise estimation of the association, a meta-analysis was conducted in the present study. The relevant studies were identified by searching PubMed, EMBASE, the Web of Science, Cochrane Collaboration Database, Chinese National Knowledge Infrastructure, Wanfang Database and China Biological Medicine up to November, 2014. The meta-analysis was performed by STATA 11. 21 studies with a total of 6912 subjects, including 2972 PCAD patients and 3940 controls. The pooled analysis showed that MTHFR C677T gene polymorphism was probably associated with PCAD (CT vs. CC: OR = 1.13, 95% CI = 1.01–1.27; dominant model: OR = 1.16, 95% CI = 1.04–1.29; recessive model: OR = 1.19, 95% CI = 1.00–1.40; allele analysis: OR = 1.17, 95% CI = 1.01–1.34). Subgroup analysis by plasma homocysteine concentration showed a significant association in the homocysteine >15 μmol/L subgroup (CT vs. CC: OR = 1.44, 95% CI = 1.10–1.88; TT vs. CC: OR = 2.51, 95% CI = 1.12–5.63; dominant model: OR = 1.51, 95% CI = 1.16–1.96; recessive model: OR = 2.33, 95% CI = 1.05–5.20; allele analysis: OR = 1.48, 95% CI = 1.18–1.87). Subgroup analysis by continent displayed a significant association among the Asian population (CT vs. CC: OR = 1.51, 95% CI = 1.23–1.86; TT vs. CC: OR = 2.81, 95% CI = 1.87–4.23; dominant model: OR = 1.65, 95% CI = 1.35–2.01; recessive model: OR = 2.22, 95% CI = 1.53–3.21; allele analysis: OR = 1.61, 95% CI = 1.37–1.89). The statistical stability and reliability was demonstrated by sensitivity analysis and publication bias outcomes. In conclusion, the meta-analysis suggests that MTHFR C677T gene polymorphism may be associated with PCAD.

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1. Introduction

Coronary artery disease (CAD) is still one of the vital causes of death and disability in the world. Premature coronary artery disease (PCAD) is a special form of CAD, which occurs in men <55 years old and women <65 years old (Tonstad and Westheim, 2002). Compared with conventional CAD, PCAD may cause heavier burden on the health system of various countries (Dogra et al., 2012).

Evidence has suggested that high homocysteine (Hcy) level is a risk factor for PCAD (Sadeghian et al., 2006; Puri et al., 2003; Nikfardjam et al., 2001). The most common genetic defect resulting in high Hcy level is the 677C → T mutation in 5,10-methylenetetrahydrofolate reductase (MTHFR). In the past few decades, several studies about MTHFR C677T gene polymorphism were performed. Some studies demonstrated that MTHFR C677T gene polymorphism was associated with PCAD (Tomatuloo et al., 2012; Sarecka-Hujar et al., 2012). However, others drew the opposite conclusion (Saffari et al., 2013; Eftychiou et al., 2012). To date, no meta-analysis has been conducted to obtain an accurate evaluation. Therefore, in the present study, a meta-analysis was conducted to evaluate the association between MTHFR C677T gene polymorphism and PCAD.

2. Methods

2.1. Search strategy

A critical review of literatures from PubMed, EMBASE, the Web of Science, Cochrane Collaboration Database, Chinese National Knowledge Infrastructure, Wanfang Database and China Biological Medicine (up to November, 2014) was performed to identify the relevant studies, using the following terms: (‘MTHFR’ OR ‘methylenetetrahydrofolate reductase’) AND (‘genetic polymorphism’ OR ‘allele’ OR ‘genotype’) AND (‘young’ OR ‘premature’) AND (‘coronary artery disease’ OR ‘myocardial infarction’ OR ‘angina’). Meanwhile, the references used in the eligible articles or textbooks were also reviewed as sources to find potential
2.5. Statistical analysis

Studies were included in our meta-analysis had to be consistent with the following criteria: (1) the association between \textit{MTHFR} C677T gene polymorphism and PCAD was involved; (2) the study should be a case–control study; (3) all patients had been diagnosed with PCAD. In addition, PCAD was defined as CAD occurred in men < 55 years old and women < 65 years old (Tonstad and Westheim, 2002); (4) the study should provide the total number of cases and controls, and also the number of cases and controls for each genotype; (5) the study was of sufficiently high quality.

Studies were excluded when they were: (1) reviews, letters or case reports; (2) duplicate publications of data from the same study.

2.2. Inclusion and exclusion criteria

Studies included in our meta-analysis had to be consistent with the following criteria: (1) the association between \textit{MTHFR} C677T gene polymorphism and PCAD was involved; (2) the study should be a case–control study; (3) all patients had been diagnosed with PCAD. In addition, PCAD was defined as CAD occurred in men < 55 years old and women < 65 years old (Tonstad and Westheim, 2002); (4) the study should provide the total number of cases and controls, and also the number of cases and controls for each genotype; (5) the study was of sufficiently high quality.

Studies were included in this meta-analysis (n=21)

3. Result

3.1. Study and data included in the meta-analysis

A flow diagram summarizing the process of study selection was shown in Fig. 1. Of the 192 potential relevant studies identified, only 21 case-control studies met all inclusion criteria. The characteristics of all included studies were listed in Table 1. The dataset represented 2972 PCAD patients and 3940 controls. No study was excluded in the meta-analysis on grounds of quality.

3.2. Pooled analysis

There were 21 studies concerning \textit{MTHFR} C677T gene polymorphism in the present meta-analysis. Based on the values of heterogeneity (CT vs. CC: P = 0.351, $I^2 = 8.3$%; TT vs. CC: P = 0.001, $I^2 = 55.3$%; dominant model: P = 0.014, $I^2 = 44.8$%; recessive model: P = 0.012, $I^2 = 45.6$%; allele analysis: P < 0.001, $I^2 = 63.1$%), fixed effects models were used under the CT vs. CC, dominant and recessive genetic inheritance models, and a random effects model was used under the TT vs. CC genetic inheritance model and allele analysis. Overall, a significant association between \textit{MTHFR} C677T gene polymorphism and PCAD was found under the CT vs. CC (OR = 1.13, 95% CI = 1.01–1.27), dominant (OR = 1.16, 95% CI = 1.04–1.29) and recessive (OR = 1.19, 95% CI = 1.00–1.40) genetic inheritance models, and allele analysis (OR = 1.17, 95% CI = 1.01–1.34). In addition, borderline significant association was also shown in the TT vs. CC genetic inheritance model (OR = 1.34, 95% CI = 0.99–1.82) (Figs. 2–5).

3.3. Subgroup analysis

In the subgroup analysis, studies were categorized by plasma Hcy concentration, continent, and source of controls. The results of the
subgroup analysis were shown in Table 2. In the subgroup analysis by plasma Hcy concentration, a significant association was shown between MTHFR C677T gene polymorphism and PCAD in the Hcy > 15 μmol/L subgroup (CT vs. CC: OR = 1.49, 95% CI = 1.10–1.98; TT vs. CC: OR = 2.51, 95% CI = 1.12–5.63; dominant model: OR = 1.51, 95% CI = 1.16–1.96; recessive model: OR = 2.33, 95% CI = 1.05–5.20; allele analysis: OR = 1.48, 95% CI = 1.18–1.87). Nevertheless, the other subgroup showed no obvious association. In the subgroup analysis by continent, a significant association between MTHFR C677T gene polymorphism and PCAD was observed in the Asian population (CT vs. CC: OR = 1.51, 95% CI = 1.18–1.87). However, no obvious association was found in the other populations.

3.4. Sensitivity analysis

The sensitivity analysis was performed to evaluate the influence of each individual study on the pooled OR by omitting every single study. The analysis results reflected that our results were statistically stable and reliable.

3.5. Publication bias

No significant publication bias was found in the meta-analysis, reflected by P values from Begg’s correlation (CT vs. CC: P = 0.097, TT vs. CC: P = 0.349, dominant model: P = 0.139, recessive model: P = 0.487, allele analysis: P = 0.349) and Egger’s regression (CT vs. CC: P = 0.073, TT vs. CC: P = 0.322, dominant model: P = 0.084, recessive model: P = 0.448, allele analysis: P = 0.332). The shapes of the funnel plots did not show any strong evidence of asymmetry.

4. Discussion

High Hcy level has been accepted as a risk factor for PCAD (Sadeghian et al., 2006; Puri et al., 2003; Nikfardjam et al., 2001). MTHFR plays an important role in the process from 5,10-methylenetetrahydrofolate to 5-methyltetrahydrofolate. 5-methyltetrahydrofolate is the methyl donor for Hcy remethylation. Therefore, MTHFR can adjust the plasma Hcy level indirectly. A common mutation in MTHFR gene (677C → T) may cause the enzyme thermolabile and less active, resulting in the elevation level of Hcy (Frosst et al., 1995; Engbersen et al., 1995; Jacques et al., 1996). Besides the genetic factor, Hcy level is also influenced by nutritional conditions. The adequate status of xfolate, riboflavin, Vitamin B12 and Vitamin B6 are indispensable to maintain the normal level of Hcy (Krishnaswamy and Lakshmi, 2002; Imamura et al., 2010).

Although a few meta-analyses have been carried out to identify whether MTHFR C677T gene polymorphism is associated with CAD (Xuan et al., 2011; Li, 2012; Zhiwen et al., 2004), the paper has been the first meta-analysis to evaluate the association between MTHFR C677T gene polymorphism and PCAD which only occurs in young populations.
population. In the meta-analysis, a significant association between \textit{MTHFR} C677T gene polymorphism and PCAD was found in the CT vs. CC, dominant and recessive genetic inheritance models, and allele analysis. In addition, borderline significant association was also obtained in the TT vs. CC genetic inheritance model. The results indicated that C → T common mutation in \textit{MTHFR} C677T might be a risk factor for PCAD.

In the subgroup analysis by Hcy concentration, a significant association between \textit{MTHFR} C677T gene polymorphism and PCAD was shown in the Hcy \( \geq 15 \) μmol/L subgroup. However, no obvious association was found in the Hcy \( < 15 \) μmol/L subgroup. The effect of high Hcy level on PCAD susceptibility may be responsible for this result. Individuals with higher level of Hcy should be considered at higher risk for PCAD and should be treated seriously. Subgroup analysis by continent showed a significant association between \textit{MTHFR} C677T gene polymorphism and PCAD susceptibility among the Asian population. Nevertheless, no obvious correlation was found in other populations. This geographical variability may be attributed to less folate intake in the Asian population (Nath et al., 1998; Sato et al., 2013; Shohag et al., 2012). Wald et al. (2001) reported that dietary supplementation with folate could reduce elevated Hcy level in order to slow down the process of CAD. Thus, it is
Table 2

<table>
<thead>
<tr>
<th>Subgroup</th>
<th>No. of study</th>
<th>CT vs. CC</th>
<th>TT vs. CC</th>
<th>Dominant model</th>
<th>Recessive model</th>
<th>Allele analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>OR (95% CI)</td>
<td>P</td>
<td>Het</td>
<td>OR (95% CI)</td>
<td>P</td>
</tr>
<tr>
<td>Homocysteine (mol/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N≤15</td>
<td>8</td>
<td>1.01 (0.85,1.21)</td>
<td>0.98</td>
<td>0.23</td>
<td>2.29 (1.77,2.98)</td>
<td>0.002</td>
</tr>
<tr>
<td>N≥15</td>
<td>8</td>
<td>1.01 (0.85,1.21)</td>
<td>0.98</td>
<td>0.23</td>
<td>2.29 (1.77,2.98)</td>
<td>0.002</td>
</tr>
<tr>
<td>Continents</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Asian</td>
<td>8</td>
<td>1.51 (1.23,1.86)</td>
<td>0.795</td>
<td>0.67</td>
<td>2.81 (1.87,4.23)</td>
<td>0.795</td>
</tr>
<tr>
<td>North American</td>
<td>4</td>
<td>1.07 (0.82,1.39)</td>
<td>0.701</td>
<td>0.01</td>
<td>1.05 (0.73,1.52)</td>
<td>0.643</td>
</tr>
<tr>
<td>African</td>
<td>1</td>
<td>0.72 (0.44,1.17)</td>
<td>0.016</td>
<td>0.01</td>
<td>0.67 (0.41,1.09)</td>
<td>0.016</td>
</tr>
<tr>
<td>HB</td>
<td>8</td>
<td>1.26 (1.01,1.57)</td>
<td>0.041</td>
<td>0.01</td>
<td>1.37 (1.07,1.76)</td>
<td>0.019</td>
</tr>
<tr>
<td>PB</td>
<td>13</td>
<td>1.09 (0.89,1.35)</td>
<td>0.774</td>
<td>0.01</td>
<td>1.13 (0.94,1.37)</td>
<td>0.247</td>
</tr>
</tbody>
</table>

appropriate for the Asian population to take sufficient folate to prevent PCAD.

Some limitations of this meta-analysis should be addressed. Firstly, studies have demonstrated that the effect of \textit{MTHFR} polymorphisms on Hcy concentration is modified by nutrient intake of folate and vitamins (Krishnaswamy and Lakshmi, 2002; Imamura et al., 2010). Unfortunately, the amounts of the nutrient intake of folate and vitamins are almost unavailable in the included studies. Therefore, the neglect of the nutrient intake may bring some bias. Secondly, sample size is still relatively small in the meta-analysis, especially in the subgroup analysis. Thus, studies with larger sample size are needed in the future. Thirdly, only publications were included in the present meta-analysis, so we need more unpublished reports to expand our analysis. Despite the limitations, our meta-analysis significantly increased the statistical power based on substantial data from different studies. The sensitivity analysis and publication bias outcomes both reflected that our results were statistically stable and reliable.

In conclusion, this meta-analysis suggests that \textit{MTHFR} C677T gene polymorphism may be associated with PCAD susceptibility. This finding may provide an important scientific basis for the comprehensive prevention of PCAD. In order to reach a more definitive conclusion, much larger sample size are needed for future studies.

Conflict of interest

None of the authors had any conflicts of interest.

Acknowledgment

We are really appreciative of all the participants in this study.

References


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