

# Transcription factor 21 gene polymorphism in patients with coronary artery disease

Waleed Abdou Hamed<sup>1</sup>  
Ghada E Hammouda<sup>2</sup>  
Sally M El-Hefnawy<sup>2</sup>  
Eman M Abd El-Gayed<sup>2</sup>

<sup>1</sup>Cardiology Department,

<sup>2</sup>Biochemistry Department, Faculty of Medicine, Menoufia University, Shebin El-Kom, Egypt

**Background:** CAD is the most common type of heart disease and the leading cause of death in both men and women. Transcription factor 21 (*TCF21*) is required for normal epicardial development and regulates epicardium-derived cell (EPDC) differentiation into smooth muscle and fibroblast lineages.

**Objectives:** The aim of this study was to investigate the distribution of *TCF21* (12190287G/C) gene polymorphism in patients with CAD and its association with other clinical and laboratory variables in these patients.

**Patients and methods:** This study was carried out in 100 patients with CAD, and 50 healthy subjects served as controls. All studied subjects underwent laboratory investigations, including measurement of total cholesterol (TC), triglycerides (TG), low-density lipoprotein cholesterol (LDL-c), high-density lipoprotein cholesterol (HDL-c), creatinine kinase-MB (CK-MB), troponin I (TnI) and genotyping of *TCF21* (12190287G/C) gene using the TaqMan Allelic Discrimination assay technique.

**Results:** Serum creatinine, TC, TG, LDL-c, CK-MB and TnI were significantly higher in patients with CAD than in the control group. There was a statistically significant difference regarding genotype frequency of *TCF21* (12190287G/C) polymorphism between the two studied groups, with the highest frequency of GG genotype among the control group, while CC genotype had the highest frequency in the patient group. C allele was more frequent in the patient group, while G allele was more frequent in the control group.

**Conclusion:** CC genotype and C allele of *TCF21* (12190287G/C) polymorphism are considered as genetic risk factors for CAD; these findings could have an impact on the future preventive and management protocols in this group of patients.

**Keywords:** coronary artery disease, transcription factor, gene polymorphism

## Introduction

Coronary artery disease (CAD) is a multifactorial disease and considered the main cause of death in many countries.<sup>1</sup> Many risk factors contribute to the pathogenesis of CAD, such as smoking, diabetes, hypertension, hypercholesterolemia, obesity, physical inactivity and stress.<sup>2</sup> Although many patients with CAD share similar risk factors, it is not clear why some individuals are more susceptible to these environmental determinants than others with the same risk factors, pointing to a complex interaction of both environmental and genetic factors.<sup>3</sup>

Recently, genome-wide association studies (GWASs) have identified several common variants that are associated with the risk of CAD and myocardial infarction.<sup>4</sup>

Transcription factor 21 (*TCF21*) is a class II basic helix–loop–helix transcription factor that is produced in the mesenchyme of developing organs such as the lung, kidney and epicardium.<sup>5</sup>

Correspondence: Waleed Abdou Hamed  
Cardiology Department, Menoufia University, 49 Zaki Shabana Street, Shebin El-Kom, Menoufia, 11511, Egypt  
Tel +20 10 0361 8195  
Fax +20 48 223 5679  
Email waleedabdou@yahoo.com

In the heart, *TCF21* is required for normal epicardial growth and regulates epicardium-derived cell (EPDC) differentiation into smooth muscle cells and fibroblast cells. Vascular smooth muscle cells and cardiac fibroblasts are essential for the development of the coronary vasculature and cardiac fibrous matrix, which support myocardial integrity and function in the heart.<sup>6</sup>

*TCF21* attaches certain DNA sequences and regulates cell differentiation by regulating cell cycle arrest and tissue gene expression.<sup>7</sup>

The rs12190287 is present in the exon of the *TCF21* and is considered an expression quantitative trait locus for this gene by regulating *TCF21* gene expression.<sup>8</sup> The gene study indicated that the C allele of rs12190287 enhanced the expression of *TCF21* in vascular smooth muscle cells.<sup>9</sup>

## Aim of the study

The aim of the present study was to investigate the distribution of *TCF21* (12190287G/C) gene polymorphism in patients with CAD and its association with other clinical and laboratory variables in these patients.

## Patients and methods

The present study was carried out in 100 patients with CAD (56 males and 44 females) with a mean age of  $52.3 \pm 4.1$  years; the diagnosis of CAD was based on coronary angiographic demonstration of >50% stenosis in at least one of the major epicardial coronary arteries. The patients were selected from the Cardiology Department, Menoufia University, in the period from April 2015 to October 2015.

This patient group was compared with 50 (32 males and 18 females) age-matched and sex-matched healthy subjects with a mean age of  $51.2 \pm 3.4$  years who had no detectable cardiovascular risk factors and did not receive any medications. Control subjects were volunteers from the hospital staff, medical and nursing students and members of the local community. The studied population was enrolled after obtaining the written informed consent and after approval from the university research ethics committee of Menoufia University Hospitals.

Patients with valvular heart disease, heart failure or hepatic or renal dysfunction were excluded from this study.

All patients and control groups were subjected to complete history taking, complete clinical examination, electrocardiography (ECG) and laboratory investigation, including fasting blood sugar (FBS), lipid profile, urea, creatinine, creatinine kinase-MB (CK-MB), troponin I (TnI) and genotyping of *TCF21* (rs12190287).

## Sample collection and assay

A total of 10  $\mu$ L of venous blood was obtained from all subjects as follows: 2  $\mu$ L into tubes containing EDTA for genotyping of *TCF21* gene and 8  $\mu$ L into tubes without additives for separation of serum for colorimetric measurement of urea,<sup>10</sup> creatinine,<sup>11</sup> total creatine kinase (CK) and CK-MB levels,<sup>12</sup> total cholesterol (TC),<sup>13</sup> triglycerides (TG)<sup>14</sup> and high-density lipoprotein cholesterol (HDL-c).<sup>15</sup> Low-density lipoprotein cholesterol (LDL-c) was calculated according to the Friedewald equation.<sup>16</sup> Enzyme-linked immunosorbent assay (ELISA) was done for the measurement of serum TnI.<sup>17</sup>

## *TCF21* genotyping

DNA was extracted from whole blood using GeneJET Whole Blood Genomic DNA Purification Mini Kit (Thermo Fisher Scientific, Waltham, MA, USA).

*TCF21* gene was genotyped using the TaqMan allelic Discrimination method, which determines variants of single nucleic acid sequence. Using two primer/probe pairs in each reaction allows genotyping of the two possible variants at the single nucleotide polymorphism in a target template sequence.

Using the maxima probe qPCR Master Mix (2X), primers and probes were applied from Thermo Fisher Scientific; the forward primer was 5'-CTCCAAGGGCT-GAGAACTTCGGTGA-3' and the reverse primer was 5'-TTCATCCACCTGTCTATTTGCACAT-3'.

The genotyping reaction was done by adding 10  $\mu$ L of master mix, 1.25  $\mu$ L of the genotyping ready-made assay mix (probes and primers) and 3.75  $\mu$ L of DNAase-free water. For every unknown reaction, 5  $\mu$ L of genomic DNA template and 5  $\mu$ L of DNAase-free water for the negative control reaction were applied.

The cycling parameters were as follows: first, denaturation was done at 95°C for 10 minutes, followed by 40 cycles of denaturation at 94°C for 15 seconds, annealing at 50°C for 60 seconds and then extension at 72°C for 2 minutes and the last extension at 72°C for 1 minute. Data analysis was performed using 7500 Real-Time PCR System, software version 2.0.1.

## Statistical analysis

Results were collected, tabulated and statistically analyzed by IBM personal computer and statistical package SPSS version 20 (IBM Corporation, Armonk, NY, USA). Chi-square test ( $\chi^2$ ) was used to study the association between two qualitative variables. Odds ratio (OR) describes the probability that people who are exposed to a certain factor will have a

disease compared to people who are not exposed to the factor. Student's *t*-test was used for comparison between two groups having quantitative variables. Analysis of variance (ANOVA) (*F*) test was used for comparison between three or more groups having quantitative variables. Multiple regression analysis was performed to calculate the effects of risk factors as independent ORs with the effects of other confounders removed. *P*-value < 0.05 was statistically significant.

## Results

The clinical, echocardiographic and baseline characteristics of the study population are depicted in Table 1. There was a statistically significant difference between the two studied groups regarding heart rate, systolic blood pressure ejection fraction and smoking (*P* < 0.05), while there was no significant difference as regards other clinical data, medical history, age and sex.

Furthermore, there was a statistically significant difference between the two studied groups regarding creatinine, TC, TG, LDL-c, HDL-c, CK-MB, TnI (*P* < 0.05) and FBS (*P* < 0.001), while there was no significant difference regarding blood urea (Table 2).

Interestingly, *TCF21* genotype distribution between the two studied groups showed a significant difference, with increased frequency of the CC and GC genotypes and C allele in the patient group and increased GG genotype frequency in the control group (*P* < 0.05; Table 3 and Figure 1).

The results also showed that the GC genotype of *TCF21* increases the risk of CAD by 2.3-fold and CC genotype increases the risk by 5.9-fold, while the C allele increases the same risk by 2.9-fold, as shown in Table 3.

There was also a statistically significant difference among different genotypes of *TCF21* in patients with CAD

**Table 1** Demographic characteristics and clinical data of the studied groups

Parameter	Control (n = 50)	Patients (n = 100)	P-value
Age (years), mean±SD	51.2 ± 3.4	52.3 ± 4.1	>0.05
Sex, n (%)			
Male	32 (64)	56 (56)	>0.05
Female	18 (36)	44 (44)	
HR (beat/min), mean±SD	73.5 ± 5.3	88.6 ± 4.6	<0.05
Systolic blood pressure (mmHg), mean±SD	125.2 ± 3.1	160.1 ± 6.8	<0.05
Diastolic blood pressure (mmHg), mean±SD	80.7 ± 7.8	81.2 ± 3.6	>0.05
EF (%), mean±SD	60.6 ± 4.3	50.4 ± 3.2	<0.05
Smoking, n (%)			
+ve	16 (32)	62 (62)	<0.05
-ve	34 (68)	38 (38)	

**Abbreviations:** EF, ejection fraction; HR, heart rate; SD, standard deviation.

regarding the lipid profile and TnI; patients with CC genotype predominance showed higher levels of TC, TG, LDL-c and TnI (*P* < 0.05), while HDL-c was significantly higher in patients with a predominance of GG genotype, as shown in Table 4.

Using multivariate analysis for predicting risk factors for CAD, the most significant risk factors were CC genotype (OR 44.7, *P* < 0.001), smoking (OR 22.1, *P* < 0.001) and FBS (OR 15.4, *P* < 0.001), followed by LDL-c (OR 13.1, *P* < 0.05), TG (OR 9.9, *P* < 0.05) and TC (OR 7.1, *P* < 0.05) as shown in Table 5.

## Discussion

CAD is an important cause of death and disability in both developed and developing countries.<sup>18,19</sup> CAD is a complex disease caused by the inheritance of multiple genetic variants together with environmental factors promoting the disease state.<sup>20</sup> Genome-wide associations (GWAs) have explored several genetic variant associations with CAD.<sup>21,22</sup>

**Table 2** Comparison between the studied groups regarding laboratory parameters

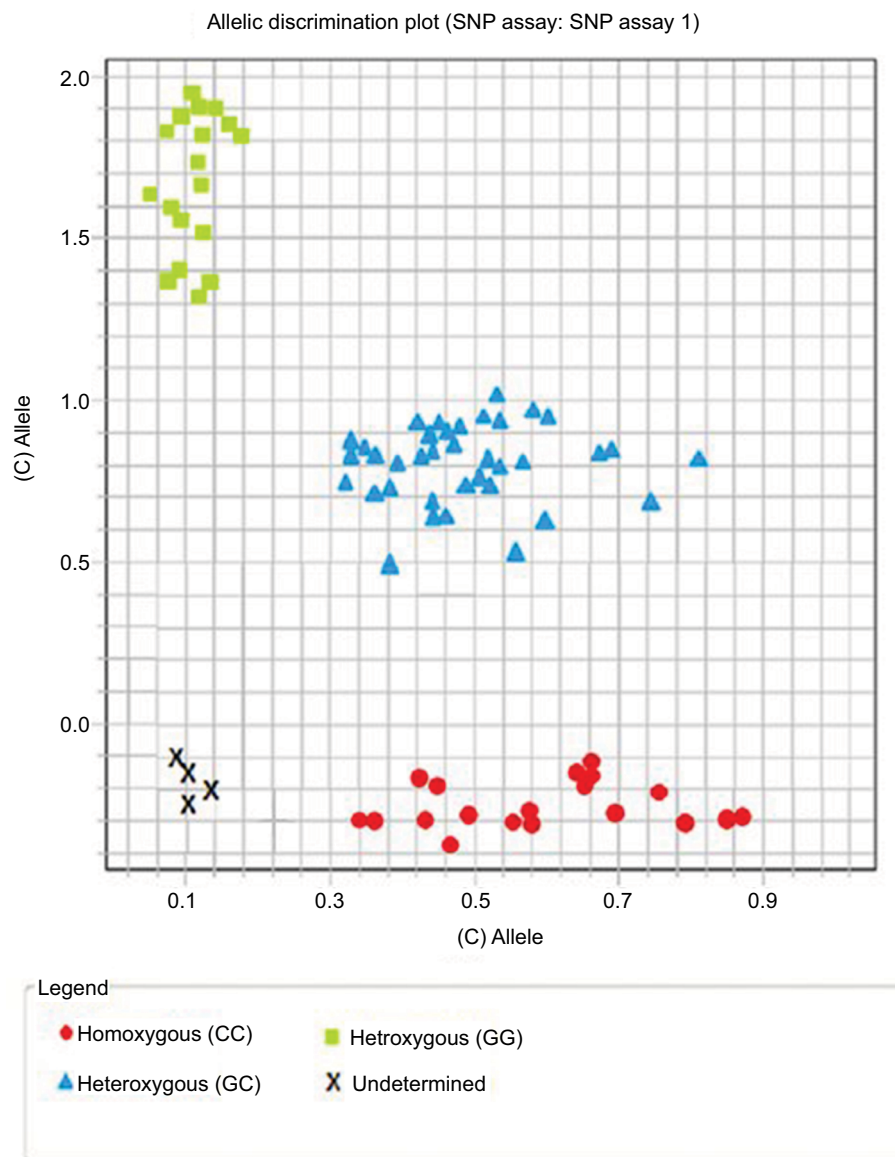
Parameter	Control (n = 50) (mean±SD)	Patients (n = 100) (mean±SD)	P-value
Serum creatinine (mg/dL)	0.98 ± 0.3	1.66 ± 0.4	<0.05
Blood urea (mg/dL)	28.6 ± 4.6	30.4 ± 5.2	>0.05
TC (mg/dL)	152.4 ± 23.6	172.3 ± 22.4	<0.05
TG (mg/dL)	105.1 ± 19.6	130.5 ± 20.2	<0.05
LDL-c (mg/dL)	84.5 ± 13.2	105.2 ± 16.7	<0.05
HDL-c (mg/dL)	47.6 ± 10.1	37.4 ± 9.5	<0.05
CK-MB (IU/μL)	15.2 ± 5.6	35.5 ± 8.9	<0.05
TnI (μg/L)	0.18 ± 0.1	0.62 ± 0.2	<0.05
FBS (mg/dL)	106.5 ± 18.7	201.3 ± 64.2	<0.001

**Abbreviations:** CK-MB, creatine kinase-MB; FBS, fasting blood sugar; HDL-c, high-density lipoprotein cholesterol; LDL-c, low-density lipoprotein cholesterol; SD, standard deviation; TC, total cholesterol; TG, triglycerides; TnI, troponin I.

**Table 3** Comparison of *TCF21* genotypes between the studied groups

Genotypes	Control (n = 50)		Patients (n = 100)		χ <sup>2</sup>	P-value	OR (95% CI)
	n	%	n	%			
<i>TCF21</i> genotype							
GG	22	44	14	14	8.4	<0.05	–
GC	20	40	56	56			2.3 (0.7–8.1)
CC	8	16	30	30			5.9 (1.4–25.2)
Allele							
G	64	64	84	42	9.2	<0.05	–
C	36	36	116	58			2.9 (1.4–5.8)

**Abbreviations:** CI, confidence interval; OR, odds ratio; *TCF21*, transcription factor 21.



**Figure 1** Allelic discrimination plot showing CC, GC and GG genotypes.

**Abbreviation:** SNP, single nucleotide polymorphism.

In the heart, *TCF21* is required for normal epicardial development and controls EPDC differentiation into smooth muscle and fibroblast cell lines.<sup>6</sup>

The present study revealed that smoking is a strong risk factor for CAD, which is consistent with many studies.<sup>23–26</sup> Smoking is associated with excess oxidative stress, free radicals and release of cytokines.<sup>26</sup>

Furthermore, this study showed that systolic blood pressure was significantly higher in patients with CAD as compared to the control group; these results are in agreement with Cannon<sup>27</sup> who found that hypertensive patients have more risk of CAD in relation to patients with normal blood pressure. McCann et al<sup>23</sup> showed that systolic blood pressure was higher in patients with CAD compared to controls. Surprisingly, we

revealed no significant difference in diastolic blood pressure between patients and controls in contrast to that reported by Santos et al<sup>28</sup> who indicated high diastolic blood pressure in patients with CAD when compared to the control group. The explanation of our results may be because most of the hypertensive individuals in the patient group were suffering from only systolic hypertension.

The current study showed that serum creatinine was higher in patients with CAD compared to the control group. These results agree with those of Santos et al.<sup>28</sup>

Inrig et al<sup>29</sup> reported that ~40% of patients who present with CAD have some degree of renal impairment.

Furthermore, our study showed that serum TC and serum TG were increased in patients with CAD. These results run in

**Table 4** Biochemical parameters of the studied patients with CAD in different genotypes of *TCF21*

Parameter	GG (n = 14) (mean±SD)	GC (n = 56) (mean±SD)	CC (n = 30) (mean±SD)	P-value
Urea (mg/dL)	34 ± 12.6	32 ± 18.7	37 ± 10.3	>0.05
Creatinine (mg/dL)	0.88 ± 0.22	0.99 ± 0.20	1 ± 0.30	>0.05
TC (mg/dL)	132.5 ± 22.8	172.4 ± 27.9	202.8 ± 35.1	<0.05
TG (mg/dL)	120.6 ± 14.5	134.7 ± 17.2	139.6 ± 18.1	<0.05
LDL-c (mg/dL)	99.4 ± 24.1	103.6 ± 28.5	139.4 ± 37.2	<0.05
HDL-c (mg/dL)	43 ± 3.3	38.8 ± 4	35.4 ± 5.4	<0.05
CK-MB (IU/μL)	28.9 ± 8.2	35.3 ± 14.9	45.2 ± 17.1	<0.05
TnI (μg/L)	0.29 ± 0.11	0.37 ± 0.30	0.59 ± 0.30	<0.05

**Abbreviations:** CAD, coronary artery disease; CK-MB, creatine kinase-MB; HDL-c, high-density lipoprotein cholesterol; LDL-c, low-density lipoprotein cholesterol; SD, standard deviation; TC, total cholesterol; TG, triglycerides; TnI, troponin I.

**Table 5** Multivariate analysis of predicting risk factors in CAD

Risk factor	OR (CI)	P-value
CC genotype	44.7 (14.9–133.7)	<0.001
Smoking	22.1 (9.9–188.2)	<0.001
FBS (mg/dL)	15.4 (8.6–121.7)	<0.001
LDL-c (mg/dL)	13.1 (1.5–112.9)	<0.05
TG (mg/dL)	9.9 (1.2–99.8)	<0.05
TC (mg/dL)	7.1 (1.18–32.9)	<0.05

**Abbreviations:** CAD, coronary artery disease; CI, confidence interval; FBS, fasting blood sugar; LDL-c, low-density lipoprotein cholesterol; OR, odds ratio; TC, total cholesterol; TG, triglycerides.

parallel with McCann et al<sup>23</sup> who reported that disturbance of lipoprotein metabolism with high-fat diets, obesity and physical inactivity have led to atherosclerotic disease in the developed countries. The common genetic disorders of lipoproteins with adverse environmental factors cause premature atherosclerosis.

Type 2 diabetes is linked with atherogenic dyslipidemia. It was found that the incidence of atherosclerosis is increased in diabetic patients compared to the general population.<sup>30,31</sup> Subjects with diabetes mellitus have other risk factors such as obesity and hypertension; these risk factors appear to act in synchrony to increase plaque formation.<sup>32</sup> Oxidative stress due to hyperglycemia may explain association between diabetes and CAD, as it leads to endothelial dysfunction and atherogenesis.<sup>33</sup>

This study showed that the mean concentration of CK-MB and TnI in the CAD group was significantly higher than that in the control group. Similar results were reported by Pas-aoglu et al<sup>34</sup> though TnI is more sensitive for minor degrees of myocardial injury.<sup>34–36</sup>

This study revealed that single nucleotide polymorphism (SNP) rs12190287 was significantly associated with CAD and the frequencies of CC and GC genotypes are associated with 5.9 and 2.3, respectively, higher risk of CAD than GG genotypes. Furthermore, C allele was associated with 2.9 higher risk of CAD than G allele. In agreement with

our results, Schunkert et al<sup>21</sup> and Wang et al<sup>37</sup> reported a significant association of SNP rs12190287 of *TCF21* with CAD. In contrast to our results, Chang et al<sup>38</sup> did not find any association between rs12190287 and CAD.

This difference in results of previous studies indicates that *TCF21* may have different disease determinants in CAD development, and association between *TCF21* and CAD susceptibility may have an impact on patients' risk levels.

## Study limitations

First, the maintenance medications for the patients were not standardized before randomization; they were on different types and doses of statin therapy, antiplatelet drugs and beta blockers; such a therapy may have a potential effect on the clinical and laboratory variables. Second, there was a lack of assessment of microRNAs (miRNAs), which are known to be involved in post-transcriptional regulation of gene expression and may affect both the stability and translation of mRNAs. Third, the study lacked a large validation population. Further prospective studies are thus needed to confirm our results.

## Conclusion

CC genotype and C allele of *TCF21* (12190287G/C) polymorphism are genetic risk factors for CAD; this association could provide a new method to identify patients at higher cardiovascular risk.

## Disclosure

The authors report no conflicts of interest in this work.

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