Relationship between calculated total antioxidant status and atherosclerotic coronary artery disease

Maryam Sotoudeh Anvari, Maryam Mortazavian Babaki, Mohammad Ali Boroumand, Bahareh Eslami*, Arash Jalali*, Hamidreza Goodarzynejad*

Department of Surgical and Clinical Pathology, *Cardiac Research, Tehran Heart Center, Tehran University of Medical Sciences; Tehran-Iran

ABSTRACT

Objective: Antioxidants play a major role in the cellular protection cascade against oxidative damage. Oxidative stress has been linked to the pathogenesis of coronary atherosclerosis. Our aim was to evaluate the association between calculated serum total antioxidant status (cTAS) and the presence and severity of coronary artery disease (CAD).

Methods: One hundred and seventy-four patients with angiographically documented significant (≥50%) luminal stenosis (n=123) or with minimal (<50%) luminal stenosis (n=51) in at least one coronary artery or major branch segment in the epicardial coronary tree were categorized as CAD+ group; 88 patients with no luminal stenosis were considered as the control group. The level of cTAS (mmol/L) was evaluated using the following equation: (0.63×albumin concentration)+(1.02×uric acid concentration)+(1.53×bilirubin concentration).

Results: In univariate analyses, mean levels of cTAS, uric acid, and creatinine were significantly higher in CAD+ group than in controls. However, adjusted cTAS level was not found to be a CAD predictor in the total population [odds ratio (OR)=1.20; 95% confidence interval (CI): 0.81–1.76; p=0.364] or in men (OR=1.25; 95% CI: 0.73–2.12; p=0.420) and women (OR=1.20; 95% CI: 0.66–2.19; p=0.553). A weak but statistically significant correlation was found between cTAS and Gensini score (Spearman’s ρ=0.16, p=0.019).

Conclusion: In patients with suspicious CAD, the level of cTAS was not found to be an independent predictor for the presence of CAD. Further studies with larger sample size are required to confirm the results. (Anatol J Cardiol 2016; 16: 689-95)

Keywords: calculated total antioxidant status, coronary angiography, coronary artery disease, Gensini score

Introduction

Coronary artery disease (CAD) is among the leading causes of death worldwide. Traditional and established risk factors of atherosclerosis, such as hypertension, hyperlipidemia, diabetes, age, obesity, and cigarette smoking, play a major role in a person’s chances of developing CAD. However, approximately half of the patients with CAD do not have any of the established risk factors, indicating that other potential CAD risk factors are yet to be identified (1). Over the last decade, several studies have linked the excessive generation of reactive oxygen species (ROS) and resulting oxidative stress with the pathogenesis of coronary atherosclerosis (2–4).

Antioxidants, including various agents such as enzymes (glutathione peroxidase, superoxide dismutase, and catalase), large molecules (albumin and ferritin), and small molecules (uric acid, glutathione, bilirubin, vitamin C, and vitamin E), play an important role in the cellular protection cascade against oxidative damage (5). The total antioxidant status (TAS) mirrors the activity potential of the antioxidant system. Several methods have been introduced to measure the total antioxidant capacity (TAC) in different biological specimens (6). The measurement of TAC reflects the antioxidative status of plasma because antioxidative effects of the plasma antioxidant components are additive (7). The assessment of plasma TAC can be more useful than the measurement of individual antioxidant levels in cells and plasma because it could determine the synergistic interaction among different individual antioxidants (8, 9).

We hypothesized that there is a relationship between coronary occlusion and TAS. The aim of the present study was to investigate the levels of calculated serum total antioxidant status (cTAS) in patients with angiographically documented CAD as compared with those in the control group.
Methods

Study design and population

This cross-sectional study consisted of 262 consecutive patients (187 men, 71.4%), who underwent elective coronary angiography at our institution from December 2010 to July 2012 because of symptoms related to CAD. Patients with acute illness or with a history of renal failure; heart failure; liver and hematologic disease; inflammatory and rheumatic disease, particularly gout; alcohol use; and malignancy as well as those on antioxidant drugs including aspirin, beta blockers, and statins or any drug affecting uric acid, bilirubin, or albumin serum concentrations were excluded. The study protocol was approved by the Ethics Committee of Tehran Heart Center (Approval number: 2011/08/0490) affiliated to Tehran University of Medical Sciences, and written informed consent was achieved from all patients who approved the collection of blood samples for scientific research. Patients with angiographically documented significant (≥50%) luminal stenosis (n=123) or with minimal (<50%) luminal stenosis (n=51) in at least one coronary artery or major branch segment in the epicardial coronary tree were categorized as CAD+ group. Patients with no luminal stenosis (n=88) were considered as the control group.

Anthropometric indices, physical examination, and definitions of CAD risk factors

Height, waist circumference, and weight were measured to the nearest 0.1 cm and 0.1 kg, respectively, by qualified and trained staff. Body mass index was calculated as weight (kg) divided by height squared (m²). The participants remained at rest for at least 10 min and then the staff measured blood pressure using standard calibrated sphygmomanometers. Two measurements were performed for each patient, with at least 1 min interval, and the mean of the two measurements was reported as the patient’s blood pressure. Definitions for analyzed risk factors of CAD have been previously reported (10, 11). In brief, hyperlipidemia was defined as plasma total cholesterol level ≥200 mg/dL and/or low density lipoprotein (LDL)-cholesterol level ≥130 mg/dL or being on lipid-lowering drugs at the time of the study. Patients were considered to have hypertension if they had arterial blood pressure >140/90 mm Hg, or they were being treated with antihypertensive drugs. Patients were considered to have diabetes if they were taking plasma glucose-lowering drugs, including insulin and oral tablets, or had a previous history of diabetes. Patients unaware of their previous histories of diabetes were defined as those who meet the new World Health Organization criteria for diagnosing diabetes mellitus.

Coronary angiography

Coronary angiography was performed using standard angiographic techniques from the percutaneous femoral approach. The angiograms were categorized as either showing no coronary lesions (Absent), no coronary lesions with >50% luminal stenosis (Minimal) or as having one (Mild), two (Moderate), or three (Severe) major epicardial coronary arteries with >50% luminal obstructions. Left main stem (LMS) stenosis was regarded as one vessel. If LMS and the left anterior descending and/or left circumflex arteries were affected, this was counted as two points. The degree of stenosis was visually determined by comparing the greatest percentage reduction of luminal diameter in any view with the nearest normal segment. Gensini score was used for the measurement of CAD severity. This severity score has been previously described (12). In brief, the coronary arterial tree was divided into segments with multiplying factors according to geographic functional importance of any given segment (5 for the left main stem to 0.5 for the most distal segments) as well as the percentage reduction in lumen diameter. The roentgenographic appearance of concentric lesions and eccentric plaques was assigned a score (0, 1, 2, 4, 8, 16, or 32 according to the degree of luminal stenosis). The sum of the segmental scores gives the Gensini score which puts emphasis on the severity of the disease.

Biochemical tests

After an overnight fasting of 10 h, venous blood samples from participants were collected from an antecubital vein in plain tubes and immediately used for biochemical analyses. Biochemical measurements, such as total cholesterol, LDL-cholesterol, high density lipoprotein-cholesterol, triglycerides, and fasting blood sugar levels, were performed with an auto analyzer (Beckman Synchron CX4, Beckman Coulter Inc., Fullerton, CA, USA) using standard methods and commercial kits. All samples were continuously processed. Assay performance was monitored after every 50 tests using the lipid control serum commercial kit (Pars Azmon Inc., Tehran, Iran). Quality control data were plotted on Levey–Jennings chart, and Westgard rules were applied to determine whether the results from the samples can be released, or if they need to be rerun.

Albumin and uric acid were colorimetrically measured using the bromocresol green and uricase, respectively; serum bilirubin level was measured by the diazo method. The plasma concentrations of albumin, bilirubin, and uric acid were measured in mg/dL then converted to mmol/L using related molecular weights. To determine the serum TAS, cTAS (mmol/L) was evaluated using the following equation: (0.63×albumin concentration)+(1.02×uric acid concentration)+(1.53×bilirubin concentration) (13), wherein the concentrations of albumin, uric acid, and bilirubin must be expressed in mmol/L.

As recommended by the American Heart Association (14), we used eGFR (estimated glomerular filtration rate) with the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) formula rather than serum creatinine to assess renal function. CKD-EPI equation is expressed as a single equation (15):

\[
edGFR = 141 \times \min(\text{Scr}/\kappa, 1)^{1.154} \times \max(\text{Scr}/\kappa, 1)^{-1.209} \times (0.993)^{\text{Age}} \times 1.018 \times \text{(female)} \times 1.159 \text{(African descent)}
\]

Where:

Scr is serum creatinine in mg/dL,
\(\kappa\) is 0.7 for females and 0.9 for males,
\(\alpha\) is –0.329 for females and –0.411 for males,
min indicates the minimum of Scr/\(\kappa\) or 1, and max indicates the maximum of Scr/\(\kappa\) or 1.
Statistical methods

Continuous data were presented as mean with standard deviation or median (25th to 75th percentiles) except for Gensini score which is presented as median (25th to 75th percentiles). Categorical variables were expressed as frequency (n) or percentage (%). The mean age of the study patients was 54.4±10.6 years and 187 (71.4%) were men. Overall, the cardiovascular risk factors including hypertension, diabetes mellitus, cigarette smoking, hyperlipidemia, and family history of CAD were common in our study population. Cigarette smoking was much more common in men than in women (32.6% vs. 0, p<0.001). The serum levels of creatinine, uric acid, and cTAS were significantly higher in men than in women (0.9±0.2 vs. 0.7±0.1, 7.6±2.1 vs. 6.8±2.3, and 0.4±0.0 vs. 0.3±0.1, respectively; p<0.001).

The clinical and laboratory characteristics of the participants with and without CAD in total and individually in men and women are presented in Table 1. Mean levels of cTAS, uric acid, and creatinine in men were significantly higher than in women (0.9±0.2 vs. 0.7±0.1, 7.6±2.1 vs. 6.8±2.3, and 0.4±0.0 vs. 0.3±0.1, respectively; p<0.001). The clinical and laboratory characteristics of the participants with and without CAD in total and individually in men and women are presented in Table 1. Mean levels of cTAS, uric acid, and creatinine in men were significantly higher than in women (0.9±0.2 vs. 0.7±0.1, 7.6±2.1 vs. 6.8±2.3, and 0.4±0.0 vs. 0.3±0.1, respectively; p<0.001).

Association of cTAS levels with other covariates was measured using Pearson’s or Spearman’s correlation coefficient. Variables that were simultaneously associated with cTAS and CAD with p<0.1 were considered as potential confounders; the effect of cTAS on CAD, adjusted for detected possible confounders, was assessed using binary logistic regression analysis. p<0.05 was considered statistically significant. All statistical analyses were performed using the SPSS software version 18.0 for windows (SPSS Inc., Chicago, IL, USA).

Results

The mean age of the study patients was 54.4±10.6 years and 187 (71.4%) were men. Overall, the cardiovascular risk factors including hypertension, diabetes mellitus, cigarette smoking, hyperlipidemia, and family history of CAD were common in our study population. Cigarette smoking was much more common in men than in women (32.6% vs. 0, p<0.001). The serum levels of creatinine, uric acid, and cTAS were significantly higher in men than in women (0.9±0.2 vs. 0.7±0.1, 7.6±2.1 vs. 6.8±2.3, and 0.4±0.0 vs. 0.3±0.1, respectively; p<0.001).

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Creatinine were significantly higher in CAD patients than in those without CAD. However, after stratification of the study groups by gender, there were no significant differences in men and women subgroups in this regard (p>0.1).

Spearman’s correlation coefficient demonstrated a weak but statistically significant correlation between the serum cTAS level and the severity of CAD by using Gensini score (Spearman’s $\rho=0.16$, p=0.015). Moreover, mean cTAS levels in different Gensini score quartiles were tested and revealed no significant difference (Fig. 1).

In a multiple logistic regression model, adjusted effects of cTAS levels on CAD presence were assessed. As shown in Table 2, after adjusting for other covariates, cTAS level was not found to be an independent predictor for CAD occurrence in the total population (OR=1.20; 95% CI: 0.81–1.76; p=0.364) or in men [OR=1.25; 95% confidence interval (CI): 0.73–2.12; p=0.420] and women (OR=1.20; 95% CI: 0.66–2.19; p=0.553). We did not find any significant difference in cTAS levels between CAD+ and non-CAD groups and were unable to find a cut-off for cTAS measure to predict the presence or severity of CAD.

**Discussion**

The main finding of the present study is that the mean cTAS level in the CAD+ group was significantly higher than that in the control group in univariate analyses, but not after adjusting for other covariates, such as age, sex, hyperlipidemia, and diabetes mellitus. ROS are part of unspecified defense system of various organisms. However, excessive ROS levels may cause cellular damage, resulting in oxidative stress from an imbalance in the ratio of ROS production to degradation. The role of oxidative stress in the development of CAD is well known (16). Several previous investigators have reported that oxidative stress is associated with vulnerable plaque and occurrence of acute coronary syndrome (17–20). Results from clinical trials using antioxidant supplementation for improving cardiovascular outcomes in humans have, however, not been as promising as expected (21, 22), and whether antioxidant interventions actually succeeded in decreasing ROS levels and oxidative stress was never ascertained (23).

The evaluation of a single parameter may cause misinterpretations, and oxidative stress should be evaluated as a whole including both total oxidative stress and total antioxidant capacity.

![Figure 1. The association between different Gensini score quartiles and serum cTAS level; cTAS, calculated total antioxidant score; GS, Gensini score](image)

Table 2. Binary logistic regression model for identifying the independent effect of cTAS level on CAD in all participants as well as in male and female participants

| Predictors                   | All participants |            | Male |            | Female |            |
|------------------------------|------------------|-----------|------|-----------|--------|-----------|
|                              | OR (95% CI)      | P         | OR (95% CI) | P      | OR (95% CI) | P      |
| cTAS, per 0.1 mmol/L increase| 1.2 (0.8–1.8)    | 0.375     | 1.3 (0.7–2.1) | 0.395 | 1.2 (0.7–2.2) | 0.548 |
| Age, years                   | 1.1 (1.0–1.1)    | <0.001    | 1.1 (1.0–1.1) | 0.001 | 1.1 (0.9–1.1) | 0.248 |
| Male sex                     | 2.5 (1.1–6.5)    | 0.030     | –    | –         | –      | –         |
| Hypertension                 | 1.2 (0.6–2.4)    | 0.609     | 1.3 (0.5–3.0) | 0.599 | 1.6 (0.5–5.0) | 0.438 |
| Hyperlipidemia               | 2.1 (1.1–3.9)    | 0.025     | 3.1 (1.4–6.9) | 0.004 | 1.0 (0.3–3.2) | 0.935 |
| Cigarette use                | 0.699            | 0.727     | 0.738 |
| Non-smoker                   | 1*               | –         | 1*   | –         | 1*     | –         |
| Ex-smoker                    | 1.1 (0.4–3.2)    | 0.887     | 1.1 (0.3–3.5) | 0.907 | 1.7 (0.9–30.8) | 0.738 |
| Current smoker               | 1.4 (0.6–3.3)    | 0.402     | 1.4 (0.6–3.5) | 0.434 | –        | –         |
| FH of CAD                    | 2.1 (0.7–5.7)    | 0.172     | 3.6 (0.7–18.6) | 0.120 | 1.3 (0.3–6.2) | 0.752 |
| Diabetes                     | 5.2 (1.6–16.8)   | 0.006     | 2.1 (0.5–8.4) | 0.280 | 20.4 (2.1–198.3) | 0.009 |
| eGFR                         | 1.0 (0.9–1.0)    | 0.397     | 1.0 (0.9–1.0) | 0.833 | 1.0 (0.9–1.0) | 0.292 |

*Reference group  
CAD - coronary artery disease; CI - confidence interval; cTAS - calculated total antioxidant status; eGFR - estimated glomerular filtration rate; FBS - fasting blood glucose; FH - family history; OR - odds ratio
Although the combined antioxidant activity of albumin, bilirubin, and uric acid does not necessarily reflect the whole scenario of antioxidant capacity, it majorly contributes to the total antioxidant activity in plasma. Although patients with acute disease were not included in the study, the interval between the onset of symptoms and angiographic intervention could have affected oxidative status, and it would have been much better to investigate new patients who had just started to show symptoms.

Each measure of antioxidant status has its own limitations. Direct measurement of ROS has been described, but these species are transient in nature; the procedure is complex, and the results have not always shown to be reliable (24–26). The methods for calculating measured TAS (mTAS), measurement of TAS levels in the plasma using a spectrophotometer, are relatively inexpensive and usually straightforward. However, colorimetry as one of the most widely used methods for measuring total oxidant status involves either fluorescence or chemiluminescence, which requires sophisticated techniques; these technologies are unavailable in many routine clinical biochemistry laboratories, or even if available, their routine use is limited (27). In order to detect the antioxidative status of plasma, instead of mTAS, we calculated TAS using Bonnefont-Rousselot et al. (13) method that is an adaptation of a method initially described by Miller et al. (28).

We chose this method because it is an easier and cheaper method (or formula), and as Lassnigg et al. (29) stated, albumin, uric acid, and total bilirubin are robust markers for assessing TAS.

Our finding that TAS levels were significantly higher in univariate analyses in CAD patients than in the control group may be explained by higher levels of uric acid and other antioxidants in these patients. Indeed, cTAS is calculated based on uric acid, bilirubin, and albumin concentrations in the serum. In a previous study, high uric acid level was found to be independently associated with the development of CAD (11); however, it is still unknown whether high serum uric acid is causally an independent risk factor, a consequence, or merely a marker for CAD (30). Another finding of our study is that the level of cTAS was positively but negligibly correlated with Gensini score. Several other studies suggested that the severity of coronary atherosclerosis calculated by Gensini score was positively correlated with oxidative stress markers (31–33).

Because cigarette smoking, renal function, and diabetes mellitus are important conditions of increased oxidative stress that should be addressed (34, 35), multiple logistic regression models were established to compare outcome variables across groups, with the presence of other covariates, to find the independent effect of cTAS on the presence and severity of CAD. The level of cTAS was not found to be an independent predictor for CAD occurrence. Consistent with our result, the TAS level showed no significant independent contribution to CAD among middle-aged men in a study by Nojiri et al. (36). Several other studies also found no significant difference in TAS levels between CAD patients and control group (37–39). However, the relationship between CAD and TAS is controversial. In a recent study, Aydin et al. (31) showed that in patients with CAD, the plasma TAC levels were increased; they ascribed increased TAC levels to the use of drugs with anti-oxidative impacts or to the intake of dietary antioxidant nutrients in patients with severe CAD. Nieto et al. (30) also studied 150 subclinical cases of carotid atherosclerosis identified by carotid ultrasound and reported that the levels of serum TAC were significantly higher than in controls; they almost entirely explained this difference by the rise in serum uric acid level in the case group. In contrast, Fazendas et al. (40) demonstrated that plasma TAS was decreased in patients with myocardial infarction; numerous other studies have also shown a significant reduction in the level of TAC among CAD patients (41–46). Such disparity in study results may be because of variation in study design, small sample sizes, age, gender, and differences in other genetic and environmental risk factors among various populations.

The estimated renal function in our patients with CAD was worse than in the control group in univariate analysis; however, multiple regression models showed that the association between eGFR and CAD was strongly affected by confounding variables. The levels of different markers of oxidative stress have been shown to increase in patients with different degrees of renal function, including patients with end-stage renal failure (47, 48). It has been suggested that the level of renal function significantly correlates with TAS; however, it appears to be dependent on several confounding variables, including increased uric acid levels (47). On the other hand, Karamouzis et al. (48) suggested that antioxidant capacity remains rather stable with the loss of renal function and changes only in patients with end-stage renal failure.

**Study limitations**

The results of the present study should be interpreted with caution because in our study, TAS level was merely calculated (i.e., cTAS) and this equation is neither very sensitive nor specific. In addition, the serum TAS measured or calculated in circulating blood cannot necessarily reflect antioxidant concentrations in atherosclerotic plaques as the target tissue. In addition, a history of coronary revascularization and presence of collaterals could influence oxidant/anti-oxidant balance in CAD patients. But related data were not available. Finally, the small sample size that restricts the power to detect associations with statistical analyses may be considered a potential limit of this study. Because a large percentage of patients with CAD were on antioxidant medications and excluded from our study, this should be considered a relatively large sample size; however, the results need to be confirmed in larger studies.

**Conclusion**

Our findings demonstrated that serum cTAS level was significantly associated with the presence of CAD in univariate analyses.
However, after controlling for other covariates, the level of cTAS was not found to be an independent predictor for CAD. Further studies with larger sample size are required to confirm the results.

**Conflict of interest:** None declared.

**Peer-review:** Externally peer-reviewed.

**Authorship contributions:** Concept- M.S.A.; Design- M.M.B.; Supervision- M.A.B.; Data collection &/or processing – B.E., A.J.; Analysis and/or interpretation- A.J., M.M.B.; Literature search- B.E.;Writing – H.G., M.S.A.; Critical review- M.S.A., M.A.B.

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