Major Components of Metabolic Parameters and Nutritional Intakes in Different Genotypes of Adiponectin +276 G>T Gene Polymorphism in Non-Diabetes and Non-Alcoholic Iranian Fatty Liver Patients

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Abstract

Background: Genetic and environmental factors are both involved in the etiology of Non-Alcoholic Fatty Liver Disease (NAFLD). Among the genetic factors, certain polymorphisms of adiponectin gene are associated with NAFLD. In the current study, we investigated the association between metabolic parameters with different genotypes of adiponectin +276 G>T polymorphism among the Iranian NAFLD patients, and the effect of nutritional intake with development of NAFLD.

Methods: In this study, 75 patients with NAFLD and 76 healthy individuals were enrolled. Dietary intakes were assessed using a semi-quantitative Food-Frequency Questionnaire (FFQ). Body Mass Index (BMI) and Waist to Hip Ratio (WHR) were calculated. Biochemical assays including FSG (Fasting Serum Glucose), liver enzymes, lipid profiles, Malondialdehyde, insulin resistance and Total Antioxidant Capacity (TAC) were measured after 12 hr fasting. Gene polymorphism study was done by using of sequencing method.

Results: Although, T allele frequency was more prevalent in patients with NAFLD than control, adiponectin +276 G>T polymorphism was not associated with risk of NAFLD. Among the metabolic parameters, TAC in TT genotype was significantly lower 1.44 (0.69 to 2.81) p>0.05, AST in GT, GG genotypes, and ALT in all three genotypes were higher in NAFLD patients in compared to healthy subjects (p<0.05). Patients with GT genotype have significantly lower fat consumption and vitamin E intake as compared to control group with the same genotype (p<0.05).

Conclusion: In this study, we showed the association of different genotypes of +276 G>T polymorphism in adiponectin gene with some metabolic parameters.

Keywords: Adiponectin, Nonalcoholic fatty liver disease (NAFLD), Polymorphism

Introduction

Non-Alcoholic Fatty Liver Disease (NAFLD) is one of the most common hepatic disorders with unusual lipid deposition in hepatocytes 1. NAFLD is an epidemic metabolic liver disease observed in many countries and its prevalence is increasing worldwide 2. According to the US National Health and Nutrition Examination Survey (NHANES), the prevalence of NAFLD among chronic liver diseases grew from 47 to 75% between 1988 and 2008 3. The prevalence of NAFLD in the adult general population was reported 21.5% in a large population-based study in southern regions of Iran in 2011 4.

NAFLD is the hepatic manifestation of the metabolic syndrome because obesity and insulin resistance are in close inter-relationship with NAFLD 1,6. Numerous studies have suggested that insulin resistance is characteristic of NAFLD 5. Insulin resistant patients with NAFLD show decreased insulin sensitivity either at the level of muscle or at the level of liver and adipose tissue 6,7. Insulin resistance is related to increase of Free Fatty Acids (FFAs) flux that increases TG production and secretion of Very Low-Density Lipoprotein (VLDL) is stimulated in hepatocytes. Fat accumulation in liver is linked with lipid peroxidation and oxidative stress 5. Oxidative stress phenomenon induces imbalance in the pro-oxidant/antioxidant equilibrium; a condition that may influence a number of pathophysiological events in the liver 8.

Genetic and environmental factors are both involved in etiology of NAFLD as a multifactorial disease; genetic polymorphisms and dietary intake have been identified as influencing factors in NAFLD development 9. The long-term excessive intake of dietary composition in food groups is associated with NAFLD pro-
Adiponectin +276 G>T Gene Polymorphism, Metabolic Parameters, Dietary Intake and NAFLD

Abstract

Background: Adiponectin is an adiposity-derived hormone that plays a protective role against the development of fatty liver disease. This study aimed to investigate the association of +276 G>T polymorphism of adiponectin gene with metabolic parameters, dietary intake and prevalence of non-alcoholic fatty liver disease (NAFLD) in a sample of Iranian adults.

Materials and Methods

The present case-control study was performed among 151 volunteers aged 20-50 with Body Mass Index (BMI) between 25 and 39.9 kg/m² including 75 patients with NAFLD and 76 healthy individuals; these two groups were matched by age and gender with group matching. Diagnosis was confirmed by the physician based on the findings of ultrasonography. The patients had simple steatosis (grade 1-2). Control group was composed of the volunteers from university staff and relatives of patients. They were matched by age and gender with case group. Also ultrasonography was carried out in standing position at the level of the umbilicus and Hip Circumference (HC) was measured at the maximum circumference between the hip and the buttock with a non-elastic tape.

Biochemical assessments

After an overnight fast, 7 ml venous blood samples were obtained from subjects. Approximately 2 ml of the blood was transferred into tubes containing Ethylene Diamine Tetra Acetic acid (EDTA) for genetic assays. Serum samples were extracted for biochemical assays from remaining blood. Measuring of Fasting Serum Glucose (FSG), Alanine aminotransferase (ALT), Aspartate aminotransferase (AST), Total Cholesterol (TC), Triglyceride (TG) and High Density Lipoprotein cholesterol (HDLC) was assessed by Abbott ALCYON™ 300 auto analyzer using commercial ELISA kits (Pars-Azmoon, Tehran, Iran). Serum LDL-C was calculated by Friedewald formula. Insulin resistance was estimated by the homeostasis model (HOMA-IR) by following equation: fasting serum insulin (µU/ml) × fasting glucose (mg/dl)/405. Measuring of serum Total Antioxidant Capacity (TAC) was performed by colorimetric method using Randox Kit (Randox laboratories ltd., UK). Malondialdehyde (MDA) levels were measured by Thiobarbituric Acid Reactive Substances (TBARS) method. Samples were heated with 0.6% thiobarbituric acid under acidic condition; the colored product was extracted into n-butanol after cooling. The color absorbance was measured at 530 nm. MDA standards were made with 1, 1, 3, 3 - tetraethoxypropane. All of the biochemical assays were conducted by a trained lab assistant who was blinded to group assignments.

Dietary intake

Dietary intakes were assessed using a semi-quantitative Food-Frequency Questionnaire (FFQ) adapted to the Iranian society. The FFQ included 168 food items with specified serving sizes commonly consumed by Iranians. Participants reported their average frequency of consumed foods and portion sizes for each food item during the previous year in terms of the number of specified serving sizes consumed per day/week/month/year, or never. The reported frequency of consumed foods and portion sizes for each food item were converted to a daily intake.

DNA extraction and genotyping

Genomic DNA was isolated from the blood cells by salting out method. DNA fragment analogous to the polymorphisms of +45T>G (rs2241766) and +276G>T (rs1501299) were amplified by primers, 5'- ATCAAG GTGGGCTGCAATA -3' as reverse primer and 5'-TGGGAATAGGGATGAGGGT -3' as forward primer, respectively. For doing Polymerase Chain Reaction (PCR), 1 µl of genomic DNA, 0.2 µl of Taq DNA polymerase and 1 µl from each primers were added to 22
μl of 1×PCR master-mix. PCR procedure included a primary denaturation at 95°C for 2 min followed by 30 cycles of denaturation at 95°C for 30 s, annealing at 61.3°C for 30 s, extension at 72°C for 20 s and a final extension at 72°C for 5 min. Sequencing of PCR products (PCR products were directly used for sequencing) was carried out according to Sanger method using ABI 3730XL Capillary Sequencer. Sequencing results were compared with the sequence of normal adiponectin gene obtained from NCBI website: http://www.ncbi.nlm.nih.gov; also, sequence traces were assembled using Chromas software (version 2.4).

**Statistical analysis**

All data analysis was performed using the SPSS software (Statistical Package for the Social Sciences, SPSS Inc., Chicago, IL, USA). The normality of variables was assessed by Kolmogorov-Smirnov test. Variables are expressed as means±Standard Deviation (SD) or numbers and percentages. The comparison of continuous variables between two groups was performed by independent sample t-test. Comparison of continuous variables between different genotypes was performed by Analysis of Covariance (ANCOVA) with adjustment for the confounder effect of age and sex. Categorical variables were also compared using the χ² test. Logistic regression analysis was used to assess the relationship between anthropometric and biochemical variables with NAFLD genotypes adjusted for the confounding role of sex and age. p-value of less than 0.05 was considered to be significant.

**Results**

The demographic characteristics are presented in table 1. WC and WHR in NAFLD patients were significantly higher as compared to healthy group (p<0.05). Serum HDL-C and LDL-C concentrations were lower and serum AST, ALT and TG concentrations were significantly higher as compared to healthy group (p<0.05). WC and WHR in NAFLD patients were significantly higher than in control, but this high frequency was not achieved between groups. In accordance to our finding, T allele frequency was more prevalent in patients with NAFLD than in control, but this high frequency was not significant (p>0.05), (Table 3).

As shown in table 4, ALT level in patients with NAFLD was significantly higher than control group in all three genotypes (p=0.022); meanwhile AST level was higher in GT and GG genotypes in NAFLD group as compared to control (p<0.05). TAC level in TT genotype was significantly lower in patients as compared to healthy subjects with the same genotype (p<0.05), (Table 4).

Mean dietary intake including energy, protein, fat, vitamins C and E have not shown any significant differences between NAFLD and control group (p>0.05), (Table 5).

The comparison of dietary intakes according to +276 G>T adiponectin gene polymorphism between case and control groups is presented in table 6. Patients with GT genotype have significantly lower fat consumption and vitamin E intake as compared to control group with the same genotype (p<0.05). No significant difference was observed for other nutrients according to the genotypes of -276 G>T adiponectin gene polymorphism (p>0.05).

**Discussion**

In the present study, we evaluated the possible association between +276G>T adiponectin gene polymorphism, metabolic parameters and nutritional intake among the Iranian NAFLD population. To our review of literature, this is the first report, which evaluates the association between different genotypes of +276 G/T polymorphism in adiponectin gene with metabolic parameters among the Iranian NAFLD patients. Our results showed no significant association between (+276 G/T) polymorphism and risk of NAFLD in the studied groups. Several studies have evaluated the effect of gene polymorphism of adiponectin gene on NAFLD in different populations.

Musso et al 23 found an association between +45 T/G and +276 G/T polymorphisms of adiponectin gene

**Table 1. Demographic characteristics of study subjects**

<table>
<thead>
<tr>
<th>Variable</th>
<th>NAFLD (n=75) n(%)</th>
<th>Control (n=76) n(%)</th>
<th>Mean Difference (95%CI)</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male n (%)</td>
<td>36 (48%)</td>
<td>29 (38.2%)</td>
<td>-</td>
<td>0.252</td>
</tr>
<tr>
<td>Female n (%)</td>
<td>39 (52%)</td>
<td>47 (61.8%)</td>
<td>-</td>
<td>0.252</td>
</tr>
<tr>
<td>Age (years)</td>
<td>40.65(8.41)</td>
<td>38.87(8.2)</td>
<td>1.78(-0.89 to 4.46)</td>
<td>0.189</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>31.78 (4.17)</td>
<td>31.38(4.04)</td>
<td>0.40(-0.92 to 1.72)</td>
<td>0.549</td>
</tr>
<tr>
<td>WC (cm)</td>
<td>103.12(9.46)</td>
<td>100.14(8.72)</td>
<td>2.98(0.55 to 5.90)</td>
<td>0.046</td>
</tr>
<tr>
<td>WHR</td>
<td>0.92(0.06)</td>
<td>0.89(0.06)</td>
<td>0.02(0.002 to 0.04)</td>
<td>0.031</td>
</tr>
</tbody>
</table>

BMI, body mass index; WC, waist circumference; WHR, waist to hip ratio.

*p-value for gender based on Chi-Square Tests and p-value for other variables based on 2- tailed independent T-test using equal variable.
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and risk of NAFLD in their study; however, their features included non-obese and normo lipidemic subjects. In another case-control study by Zhou et al 24, the G/T variant at +276 may decrease the susceptibility to NAFLD. In the line of our study, Wong et al 4 found no association between polymorphism of adiponectin at position +276 and NAFLD in Chinese patients. Also, in Japanese population different genotypes of adiponectin +276 G/T polymorphism did not show significant difference between NAFLD patients and control group 20.

Insulin resistance and central obesity are well-known characteristic associated with intra-abdominal fat accumulation which has been positively correlated with liver fat 25. In our study, insulin resistance and HOMA-IR did not show significant association with different genotypes of +276 G/T polymorphism in NAFLD patients; these findings were also supported by previous studies 26,27. However, these results were not consistent with Melistas et al 18 study and the mentioned parameters were significant according to genotype of TT in +276 G/T gene polymorphism. It can be explained that the subjects in cross-sectional study by Melistas et al 18 included non-diabetic women without any other disease.

In this study we aimed to evaluate the environmental impact, along with genetic factors in the outbreak of NAFLD. As we showed in table 6, normal samples with TT genotype had low calorie, fat and high vitamins E and C consumption (although this difference was not significant due to small sample size). So, we can say that individuals with mutant and pathogen genotype which have proper diet, can overcome the disease. But to evaluate any additional impact we should study this issue in larger sample sizes. In addition, we observed that NAFLD patients with GT genotype consumed small amounts of vitamin E in their usual diet. In this regard, Musso et al 27 reported a low dietary ascorbic acid and tocopherol intake in patients with non-alcoholic steatohepatitis. Also, Erhardt et al 28 reported reduced tocopherol intake as dietary antioxidant compounds in patients with NASH compared to healthy group. In fact, oxidative stress is one of the most common factors involved in the pathogenesis of NAFLD; moreover vitamins C and E are well-known antioxidants capable in blocking distribution of radical

<table>
<thead>
<tr>
<th>Variable</th>
<th>NAFLD (n=5) mean (SD)</th>
<th>Control (n=76) mean (SD)</th>
<th>Mean Difference (95%CI)</th>
<th>p †</th>
</tr>
</thead>
<tbody>
<tr>
<td>TC (mg/dl)</td>
<td>183.44(36.91)</td>
<td>187.96(28.89)</td>
<td>-4.52(-15.17 to 6.13)</td>
<td>0.403</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>43.24(11.4)</td>
<td>48.29(11.6)</td>
<td>-5.05(-8.75 to -1.35)</td>
<td>0.008</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>104.11(34.62)</td>
<td>111.52(26.43)</td>
<td>-6.81(-16.71 to 3.09)</td>
<td>0.030</td>
</tr>
<tr>
<td>FSG (mg/dl)</td>
<td>90.59(11.24)</td>
<td>89.59(9.93)</td>
<td>0.63(-2.78 to 4.03)</td>
<td>0.717</td>
</tr>
<tr>
<td>ALT (IU/l)</td>
<td>49.96(25.958)</td>
<td>26.84(9.814)</td>
<td>23.12(16.82 to 29.41)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>AST (IU/l)</td>
<td>32.99(14.86)</td>
<td>23.08(6.12)</td>
<td>9.91(6.26 to 13.55)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>152.00(114.00-225.00)</td>
<td>118.50(79.50-198.00)</td>
<td>-</td>
<td>0.004</td>
</tr>
<tr>
<td>MDA (nmol/ml)</td>
<td>2.82±1.01</td>
<td>2.75±1.03</td>
<td>-0.056(-0.38-0.27)</td>
<td>0.734</td>
</tr>
<tr>
<td>TAC (µm/L)</td>
<td>1.45±0.26</td>
<td>1.58±0.28</td>
<td>0.12(0.03-0.21)</td>
<td>0.005</td>
</tr>
<tr>
<td>Insulin (µU/ml)</td>
<td>19.10(13.80-27.30)</td>
<td>17.70(13.00-22.10)</td>
<td>-</td>
<td>0.107</td>
</tr>
<tr>
<td>HOMA-IR*</td>
<td>4.01(3.21-6.21)</td>
<td>3.72(2.71-5.33)</td>
<td>-</td>
<td>0.139</td>
</tr>
</tbody>
</table>

TC, total cholesterol; TG, triglyceride; HDL, high density cholesterol; LDL, low density cholesterol; FSG, fasting serum glucose; ALT, alanine amino transferase; AST, aspartate amino transferase; MDA, malondialdehyde; TAC, total antioxidants; HOMA-IR, homeostasis model assessment insulin resistance.

† P-value for TG, Insulin and HOMA-IR based on Mann-Whitney; otherwise based on independent T-test using equal variable.

*TG, Insulin and HOMA-IR are presented based on median (P25-P75) and other variables data are presented based on mean (SD).

<table>
<thead>
<tr>
<th>Polymorphism</th>
<th>NAFLD (n=75) a (%)</th>
<th>Control (n=76) a (%)</th>
<th>OR †† (95% CI)</th>
<th>p †</th>
</tr>
</thead>
<tbody>
<tr>
<td>Genotype</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>GG</td>
<td>33(44.0)</td>
<td>39(51.3)</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>GT</td>
<td>32(42.7)</td>
<td>28 (36.8)</td>
<td>1.40(-8.75 to 4.99)</td>
<td>0.489</td>
</tr>
<tr>
<td>TT</td>
<td>10(13.3)</td>
<td>9(11.8)</td>
<td>1.44(0.69 to 2.81)</td>
<td>0.344</td>
</tr>
<tr>
<td>+276 G&gt;T</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>allele</td>
<td>G</td>
<td>98(65.3)</td>
<td>1.00</td>
<td>0.321</td>
</tr>
<tr>
<td></td>
<td>T</td>
<td>106(69.7)</td>
<td>1.28(0.78 to 2.10)</td>
<td></td>
</tr>
</tbody>
</table>

† P-value based on Chi-Square Tests.

†† Odds Ratio (OR) based on logistic regression analysis adjusted for age and gender (p>0.05).
These antioxidants play important roles in histological improvement of inflammation in NAFLD. On the other hand, hepatic de novo lipogenesis can be affected by dietary macronutrients; high fat intake reduced hepatic de novo lipogenesis in obese hyperinsulinemic subjects compared to obese normoinsulinemic subjects. Therefore, dietary recommendation should be based on individual status and even genetic background. It is clear that dietary modification is...
the easiest and even the most efficient way to reduce risk factors of chronic disease.\(^3\)

### Conclusion

Overall, among the metabolic parameters, TAC in TT genotype was significantly lower, AST in GT, GG genotypes, and ALT in all three genotypes were higher in NAFLD patients as compared to healthy subjects. Also non-statistical significance of other results might be attributed to the difference in the stage of disease; on the other hand, this is the first study to compare nutritional intakes according to adiponectin +276G>T gene polymorphism in patients with NAFLD. One limitation of this study was its relatively small sample size. Therefore, further studies with larger sample size and interventional designs are needed to confirm the effect of dietary compounds in different +276 G/T genotypes in nonalcoholic fatty liver disease.

### Acknowledgement

We appreciate all the participants in the current study. We also thank Nutrition Research Center of Tabriz University of Medical Sciences and Cellular and Molecular Research Center of Qazvin University of Medical Science for providing laboratory facilities.

### Conflict of Interests

The authors declare that there is no conflict of interest.

### Funding

The study has been supported by a grant from Qazvin University of Medical Sciences.

### Ethical approval

Study protocol was approved by the Ethics committee of Qazvin University of Medical Sciences (Identifier code: 11013).

### References


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**Table 5. Comparison of energy, macro and micronutrient intakes between study groups**

<table>
<thead>
<tr>
<th>Variable</th>
<th>NAFLD (n=75) mean (SD)</th>
<th>Control (n=76) mean (SD)</th>
<th>Mean Difference (95%CI)</th>
<th>p *</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calories (kcal)</td>
<td>2815.06 (536.35)</td>
<td>2794.93 (448.64)</td>
<td>-20.13(-179.29 to 139.03)</td>
<td>0.803</td>
</tr>
<tr>
<td>Protein (g/day)</td>
<td>93.88 (21.62)</td>
<td>88.86 (16.60)</td>
<td>-4.99(-11.20 to 1.20)</td>
<td>0.113</td>
</tr>
<tr>
<td>Carbohydrate (g/day)</td>
<td>437.38 (89.72)</td>
<td>438.49 (79.58)</td>
<td>1.10(-26.18 to 28.39)</td>
<td>0.936</td>
</tr>
<tr>
<td>Total fat (g/day)</td>
<td>89.70 (27.03)</td>
<td>91.52 (24.35)</td>
<td>1.82(-6.45 to 10.10)</td>
<td>0.664</td>
</tr>
<tr>
<td>Vitamin E (mg/day)</td>
<td>14.08 (5.60)</td>
<td>14.63 (4.11)</td>
<td>0.80(-1.03 to 2.12)</td>
<td>0.496</td>
</tr>
<tr>
<td>Vitamin C (mg/day)</td>
<td>127.14 (66.09)</td>
<td>140.59 (64.18)</td>
<td>13.44(-7.50 to 34.39)</td>
<td>0.207</td>
</tr>
</tbody>
</table>

* P-value of variables based on independent T-test using equal variable.


