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Is adiponectin associated with acute myocardial infarction in Iranian non obese patients?

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Abstract

Backgrounds: Adiponectin is an adipose tissue-derived mediator with significant anti-atherogenic properties. A few studies were done in acute phase of myocardial infarction especially in non obese patients. We design a study to investigate the association between adiponectin concentration and acute phase of myocardial infarction in non obese patients.

Methods: This case-control study was done in Paymaneah Hospital (Jahrom, Iran) from Feb 2007 to May 2008. Plasma adiponectin levels were measured in 43 patients with AMI (mean age: 62.7 ± 13.3 years, male: 67.4%) at the first 24 hours of admission and 43 normal controls (mean age: 62.1 ± 12.3 years, male: 55.8%) matched for age, sex and other CAD risk factors.

Results: Adiponectin levels in patients with AMI (3.36 µg/mL) were significantly lower than that of the control group (5.03 µg/mL) ($p < 0.0001$). Lower adiponectin were independently associated with higher risk of AMI (odds ratio = 8.97; 95% CIs: 2.3–34.5; $p = 0.001$). Adiponectin levels negatively correlated with triglyceride ($r = -0.46$, $p = 0.002$) and total cholesterol ($r = -0.32$, $p = 0.03$) in the case group and with body mass index (BMI) in control subjects.

Conclusion: The present study showed that adiponectin was associated with AMI in non obese patients but it is not related to sex, age and other CAD risk factors.

Introduction

Adiponectin is a new member of adipocyte derived proteins belonging to the soluble defense collagens that produces specifically by adipose tissue and is present in the circulation [1]. Lower levels of plasma adiponectin have been detected in obesity [1], Type 2 diabetes mellitus [2] and coronary artery disease (CAD) [3]. Adiponectin appears to play an important role in glucose and lipid metabolism, vascular biology, and energy homeostasis [4,5] and low adiponectin level has been shown as a risk factor for cardiovascular events [6,7].

Although several studies have been done on adiponectin and CAD, but most of them have been conducted on patients with chronic ischemia and the studies in acute phase of myocardial infarction (MI) especially in developing countries are limited and rare [8-14]. Yet there is controversy about the association of adiponectin and sex [8,11], CAD risk factors [8,11,14] and race [13]. Although adiponectin decreased in both obesity and AMI, some study [9] considered this for explanation of obesity effect on AMI, but with attention to direct effect of adiponectin

on atherosclerosis [15], it seems that hypo adiponectinemia in AMI is independent of obesity.

The purpose of present study was to investigate the association between plasma adiponectin levels and AMI in Iranian non obese patients.

Methods

Subjects

This case-control study conducted on 43 consecutive patients with acute myocardial infarction (AMI) (as case group) including 29 men and 16 women with the mean age of 62.72 ± 13.4 y/o out of 850 patients who were referred with chest pain to emergency room at Paymaneah Hospital, a referral center affiliated to Jahrom Medical University (Iran) from Feb 2007 to May 2008. We selected all patients after approved diagnosis of AMI and rule out of other patients with exclusion criteria. We also selected 43 individuals with atypical chest pain and normal coronary angiography as control group and matched them with case group for age, sex and other CAD risk factors such as hypertension (HTN), diabetes mellitus (DM), hyperlipidemia (HLP), body mass index (BMI), smoking and renal function.

The study protocol was approved by research ethics committee of Jahrom Medical University and informed consent was obtained from all participants before enrollment.

A questionnaire including information about the past medical and drug history (HTN, HLP, DM, smoking, chronic diseases such as rheumatologic disorder, renal failure and asthma), family history of CAD and demographic information was completed for each subjects.

The exclusion criteria were the history of infectious diseases, collagen vascular disease such as systemic lupus erythematosus, scleroderma, rheumatoid arthritis, cirrhosis, hepatitis, renal disease, malignancy, septicemia, angioplasty and stable or unstable angina.

Definitions

AMI was defined as resting chest pain lasting more than 30 minutes accompanied by ischemic electrocardiographic changes and was confirmed by the presence of total creatine kinase or creatine kinase isoenzyme (CK-MB) fraction levels of more than twice the upper normal limit. We considered ST elevation MI (STEMI) according to ST segment elevation > 1 mm in two neighbor electrocardiographic leads and non ST elevation MI (NSTEMI) without this finding [16]. The absence of any narrowing in coronary artery diameter was considered as normal coronary angiography. Unstable angina was defined as angina pectoris (or equivalent type of ischemic discomfort) with at least one of three features: occurring at rest

(or minimal exertion) and usually lasting > 20 minutes (if not interrupted by nitroglycerin administration); being severe and described as frank pain, and of new onset (i.e., within 1 month; and occurring with a crescendo pattern (i.e., more severe, prolonged, or frequent than previously) [17]. Height and weight were measured with light clothes and without shoes using a stadiometer and electronic scale in the cardiac care unit after confirming patient stable condition. Body mass index was calculated [weight (kg)/height (m)²] and used as the criteria of overall adiposity. We defined BMI less than 25 as normal, 26–30 as overweight and above 30 as obese [18]. Blood pressure was measured two times in sitting position after 5 min of rest using a mercury sphygmomanometer (Riester, Germany). Hypertension was defined as blood pressure more than 130/85 mmHg or use of any antihypertensive medication [19]. Diabetes mellitus was defined by a physician's diagnosis, a fasting plasma glucose level of ≥ 126 mg/dl or use of diabetes medications [20]. All subjects completed a research questionnaire on smoking habits.

Laboratory analysis

Fasting levels of plasma total cholesterol, high density lipoprotein cholesterol (HDL), low density lipoprotein cholesterol (LDL), and triglycerides were determined in Research Laboratory of Jahrom Medical University. Total cholesterol and triglyceride levels were measured by enzymatic techniques using an Selectra E biochromatic analyzer. HDL and LDL level was measured after precipitation of the other lipoproteins with heparin and manganese chloride (Human, Germany). Plasma glucose levels were measured by the glucose oxidase method and serum creatinine by the Jaffe reaction method.

Blood samples (5 cc) for adiponectin were obtained by venipuncture from the patients immediately after admission before starting any IV medication by trained staff and for lipid profile and fasting blood sugar at the first 24 hours of AMI after 12 hours of fasting. In control subjects all blood sample were obtained after 12 hours of fasting then plasma was separated and frozen at -70°C for later processing. Then we determined the plasma adiponectin concentration by enzyme linked immunosorbent assay (ELISA) (Biovendor Laboratory Medicine, Inc., Brno, Czech Republic) (REF: RD195023100) in patients and control group. This assay measures the total adiponectin that is all molecular forms. The sensitivity and the intra- and interassay coefficients of variation were 0.8 mg/liter, 6% and 7%, respectively.

Statistical analysis

Statistical analysis was performed by SPSS (version 11.5; SPSS, Inc., Chicago, IL). Data were expressed as mean \pm 1 SD. Continuous variables were summarized as mean \pm SD and compared using Student's *t*-test. Discrete variables

were presented as frequencies and group percentages. Nominal variables were tested with Pearson's χ^2 test and Binary variables were tested with the Mann-whitney test. Pearson correlation coefficients were calculated to evaluate unadjusted (univariate) associations between adiponectin and other variables. All tests were two-tailed with a 0.05 type I error rate. Logistic regression models were used to estimate odds ratios (ORs) between three different adiponectin levels and AMI. We considered adiponectin level greater than 5 $\mu\text{g/ml}$ as reference level.

Results

The demographic and clinical characteristics of the study groups, as well as laboratory variables are shown in Table 1. In patient group 6 cases (14%) had non-ST elevation MI (NSTEMI) and 37 (86%) had ST elevation MI (STEMI).

Plasma adiponectin levels in the patients with AMI on admission were significantly lower than those in the control group ($3.3 \pm 1.7 \mu\text{g/ml}$ vs. $5 \pm 1.5 \mu\text{g/ml}$, $p < 0.001$).

We examined the association between plasma adiponectin levels and selected cardiovascular risk factors among cases and controls. There was a significant negative correlation between plasma adiponectin levels and triglyceride ($r = -0.46$, $p = 0.002$) and total cholesterol ($r = -0.33$, $p = 0.03$) in the case group as well as BMI in control group ($r = -0.44$, $p = 0.003$). No significant correlation was found between HTN, Type 1 DM, Type 2 DM, age, sex, smoking, serum creatinine, LDL and HDL and adiponectin levels (Table 2). There was no significant difference in plasma adiponectin between patients with STEMI and NSTEMI ($3.4 \pm 1.7 \mu\text{g/ml}$ vs. $3 \pm 1.5 \mu\text{g/ml}$, $p = 0.49$).

Table 1: Demographic and clinical characteristics of the study groups

Variable	Case group n = 43	Control group n = 43	P-value
Age	62.7 \pm 13.3	62.1 \pm 12.3	0.84
Male, n (%)	29 (67.4%)	24 (55.8%)	0.27
BMI (kg/m ²)	22.3 \pm 3	23.6 \pm 3.15	0.44
Current smoker, n(%)	10 (23.3%)	4 (9.3%)	0.14
HTN, n(%)	9 (20.9%)	14 (32.6%)	0.33
Adiponectin ($\mu\text{g/ml}$)	3.3 \pm 1.7	5 \pm 1.5	< 0.001*
Type 1 DM, n (%)	2 (4.7%)	4(9.3%)	0.68
Type 2 DM, n (%)	7(16.3%)	4(9.3%)	0.52
Total Cholesterol (mg/dL)	189.3 \pm 45.2	171.8 \pm 33	0.05
LDL-C (mg/dL)	113.7 \pm 36.6	103.9 \pm 28.7	0.13
HDL-C (mg/dL)	46.7 \pm 11.4	41.9 \pm 9.5	0.05
LDL/HDL ratio	4.23	4.24	0.96
Triglyceride (mg/dL)	146 \pm 99.6	144 \pm 99.8	0.93
Serum creatinine (mg/dl)	1 \pm 0.1	1.1 \pm 0.1	0.68

BMI: body mass index, LDL-C: low density lipoprotein- cholesterol, HDL-C: high density lipoprotein- cholesterol, BUN: blood urea nitrogen

Values are presented as mean \pm SD or %.

Table 2: Pearson Correlation Coefficients between Plasma Adiponectin Level and selected variables in the Case and Control groups

Variable	Case group		Control group	
	Correlation	P-value	Correlation	P-value
Age	0.24	0.04*	0.04	0.79
BMI	-0.03	0.86	-0.44	0.003**
Serum creatinine	0.13	0.41	0.27	0.08
Total Cholesterol	-0.33	0.03*	-0.11	0.48
LDL-C	-0.03	0.84	-0.02	0.87
HDL-C	-0.06	0.72	-0.09	0.55
Triglyceride	-0.46	0.002**	-.09	0.57
TC/HDL	-0.22	0.14	-0.40	0.80

*P < 0.05, **P < 0.001

BMI: body mass index, IDDM: insulin dependent diabetes mellitus, NIDDM: non insulin dependent diabetes mellitus, LDL-C: low density lipoprotein- cholesterol, HDL-C: high density lipoprotein- cholesterol, BUN: blood urea nitrogen

Patients were divided into three groups based on adiponectin split level. The results are shown in Table 3. The group with the adiponectin levels greater than 80th percentile (5 $\mu\text{g/mL}$) was used as the reference. Odds ratio for AMI increased with decreasing adiponectin levels. Lower adiponectin were independently associated with higher risk of AMI (odds ratio = 8.97; 95% CIs: 2.3–34.5; $p = 0.001$). The greatest increase in AMI risk was seen at adiponectin levels $\leq 3 \mu\text{g/mL}$.

Discussion

In the present case-control study, in line with previous studies [7-10,21], we found that lower plasma adiponectin levels was associated with AMI among Iranian non obese patients independent of traditional cardiovascular risk factors. Of note, the relationship was also independent of hypertension, diabetes, age, sex, BMI and smoking; the factors that were closely related to adiponectin levels in previous studies [10].

Cavusoglu et al. [22] showed that high baseline plasma adiponectin levels are independently associated with an increased risk of both death and MI at 2-year follow-up in a cohort of men with stable angina, unstable angina, and non ST elevation MI referred for coronary angiography. Our study also showed significant association between adiponectin levels in patients with ST elevation MI and non-ST elevation MI in spite of their different pathophysiology [23,24]. These findings indicated that adiponectin may be directly involved in the pathophysiology of atherosclerosis and thrombosis at the vascular wall level.

The exact mechanism of adiponectin decrease immediately after the onset of AMI remains unknown. Although some authors [25] believed that the changed adiponectin levels are associated with presence of risk factors of meta-

Table 3: Odds ratio association of different adiponectin levels with acute myocardial infarction (AMI)

Adiponectin levels	Case group (AMI) (n, %)	Control group (n,%)	P value	OR (95% CI)
> 5 µg/mL	9 (20.9%)	19 (44.2%)		Reference group (1.00)
3.01–5 µg/mL	17 (39.5.6%)	20 (46.5%)	0.26	1.97 (0.65–4.99)
≤ 3 µg/mL	17 (39.5%)	4(9.3%)	0.001*	8.97 (2.3–34.5)

CI: Confidence Interval, OR: odds ratio

*p < 0.001 significant with single logistic regression analysis

bolic syndrome but our findings in line of other studies emphasize that its deficiency may lead to thrombus formation and platelet aggregation [15]. Plasma adiponectin may decrease as a result of rupture of coronary plaques. Adiponectin has been detected in the injured vessels but not in the intact ones in humans and rodents [26,27]. Although it seems that low adiponectin in AMI is secondary to plaque rupture and inflammation but if according to previous cohort studies [14,22,28] adiponectin consider as risk factor for coronary artery disease, measurement of adiponectin levels may be helpful in the stratification of patients at risk of AMI. In our study the greatest increase in MI risk occurred at adiponectin levels ≤ 3 µg/mL in contrast to the Wolk et al. study [9] that this rise occurred at adiponectin levels ≤ 5.5 µg/mL. This difference may be due to exclusion of the patients with major CAD risk factors in Wolk et al. study that were not excluded from us. Our present data, consistent with those of other authors [8,11,14,22,25] showed that adiponectin levels was negatively correlated with triglycerides in patients and with BMI only in the control subjects, but in contrast to the others we found negative correlation with total cholesterol in patients and any correlation with other CAD risk factors such as smoking[14], age, DM, HTN, HDL and LDL.

About non significant correlation between plasma adiponectin levels and BMI in case group, it should be noted that in the most of previous studies [8,11,25,29] there were significant but weak correlation ($r = -0.18$ to -0.33) between them, So, in confirming Pliz et al opinion [11], it should be noted that the adiponectin levels may be more associated with blood lipid profile and insulin resistance instead of obesity. Also this difference may be due to reduction of adiponectin secondary to acute MI.

Most studies [8,10-12,14] showed higher adiponectin levels in women in comparison to men but our results showed no statistically significant difference in adiponectin levels between men and women. Nakamura et al. [8] found similar results but they believed that solid phase ELISA was the reason for this difference, because this assay cannot distinguish between the lower weight trimer-dimer forms of adiponectin and the high molecular weight complexes; one of the factors associated with sex

difference. We measured all forms of adiponectin and this difference in our study is not related to this reason. It may be due to less obesity in our subjects and/or mismatch between BMI in men and women in other studies.

Another remarkable clinical finding in our study was that plasma adiponectin levels in our study were much lower than those of other studies [7-9,14] such that this level in our normal subjects was approximately equal to that of patients with AMI. This may be due to lower BMI in our subjects, high carbohydrate diet [30] or race differences.

Kanaya et al. [13] showed that Whites had higher median adiponectin levels than Blacks and high circulating levels of adiponectin were associated with a higher risk of CAD in older Blacks. Significant differences in our results indicate that before application of adiponectin in clinical setting, further prospective and cohort studies in other countries and races are required to confirm our results.

Although some studies [30,31] showed increased adiponectin level in patients with renal dysfunction but our study showed as the same of Shibata et al. [32] that there is no association between adiponectin and renal function in atherosclerotic patients with normal renal function.

Study limitations

Serum adiponectin levels were measured in a single sample, which might limited the observed associations. Weak or any correlation may be due to small number of participants in this study. We didn't study the association of adiponectin concentration with amount of smoking, diet and previous medication because of limited data. Some authors such as Inoue et al. [33] suggest that measurement of high molecular weight (HMW) adiponectin alone may be a clinically useful marker of CAD but we measured only total form of adiponectin. Although abdominal obesity can give better insight than BMI, but because some studies use only BMI, for comparison we only used this parameter too.

Conclusion

Lower plasma adiponectin levels are associated with AMI in Iranian non obese patients, independent of other traditional cardiovascular risk factors. Our findings showed

although obesity is a reducing factor of adiponectin but this decreased in AMI is not related only to obesity.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

MS had substantial contributions to conception and design and interpretation of data and writing the manuscript. AS carried out the biochemical analysis. GS had contributions to data analysis. All authors read and approved the final manuscript.

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References

- Arita Y, Kihara S, Ouchi N, Takahashi M, Maeda K, Miyagawa J, Hotta K, Shimomura I, Nakamura T, Miyaoka K, Kuriyama H, Nishida M, Yamashita S, Okubo K, Matsubara K, Muraguchi M, Ohmoto Y, Funahashi T, Matsuzawa Y: **Paradoxical decrease of an adipose-specific protein, adiponectin, in obesity.** *Biochem Biophys Res Commun* 1999, **257**:79-83.
- Hotta K, Funahashi T, Arita Y, Takahashi M, Matsuda M, Okamoto Y, Iwahashi H, Kuriyama H, Ouchi N, Maeda K, Nishida M, Kihara S, Sakai N, Nakajima T, Hasegawa K, Muraguchi M, Ohmoto Y, Nakamura T, Yamashita S, Hanafusa T, Matsuzawa Y: **Plasma concentrations of a novel, adipose-specific protein, adiponectin, in type 2 diabetic patients.** *Arterioscler Thromb Vasc Biol* 2000, **20**:1595-9.
- Ouchi N, Kihara S, Arita Y, Maeda K, Kuriyama H, Okamoto Y, Hotta K, Nishida M, Takahashi M, Nakamura T, Yamashita S, Funahashi T, Matsuzawa Y: **Novel modulator for endothelial adhesion molecules: adipocyte-derived plasma protein adiponectin.** *Circulation* 1999, **100**:2473-6.
- Havel PJ: **Update on adipocyte hormones: regulation of energy balance and carbohydrate/lipid metabolism.** *Diabetes* 2004, **53**(suppl 1):S143-51.
- Chandran M, Phillips SA, Ciaraldi T, Henry RR: **Adiponectin: more than just another fat cell hormone?** *Diabetes Care* 2003, **26**:2442-50.
- Efstathiou SP, Tsioulos DI, Tsiakou AG, Gratsias YE, Pefanis AV, Mountokalakis TD: **Plasma adiponectin levels and five-year survival after first-ever ischemic stroke.** *Stroke* 2005, **36**:1915-1919.
- Pischon T, Girman CJ, Hotamisligil GS, Rifai N, Hu FB, Rimm EB: **Plasma adiponectin levels and risk of myocardial infarction in men.** *JAMA* 2004, **291**:1730-1737.
- Nakamura Y, Shimada K, Fukuda D, Shimada Y, Ehara S, Hirose M, Kataoka T, Kamimori K, Shimodono S, Kobayashi Y, Yoshiyama M, Takeuchi K, Yoshikawa J: **Implications of plasma concentrations of adiponectin in patients with coronary artery disease.** *Heart* 2004, **90**(5):528-33.
- Wolk R, Berger P, Lennon RJ, Brilakis ES, Davison DE, Somers VK: **Association between plasma adiponectin levels and unstable coronary syndromes.** *Eur Heart J* 2007, **28**(3):292-298.
- Kojima S, Funahashi T, Sakamoto T, Miyamoto S, Soejima H, Hokamaki J, Kajiwara I, Sugiyama S, Yoshimura M, Fujimoto K, Miyao Y, Suefuji H, Kitagawa A, Ouchi N, Kihara S, Matsuzawa Y, Ogawa H: **The variation of plasma concentrations of a novel, adipocyte derived protein, adiponectin, in patients with acute myocardial infarction.** *Heart* 2003, **89**:667.
- Pilz S, Horejsi R, Möller R, Almer G, Scharnagl H, Stojakovic T, Dimitrova R, Weihrauch G, Borkenstein M, Maerz W, Schauenstein K, Mangge H: **Early Atherosclerosis in Obese Juveniles Is Associated with Low Serum Levels of Adiponectin.** *J Clin Endocrinol Metab* 2005, **90**(8):4792-4796.
- Kojima S, Funahashi T, Otsuka F, Maruyoshi H, Yamashita T: **Future adverse cardiac events can be predicted by persistently low plasma adiponectin concentrations in men and marked reductions of adiponectin in women after acute myocardial infarction.** *Atherosclerosis* 2006, **194**(1):204-213.
- Kanaya AM, Wassel FC, Vittinghoff E, Havel PJ, Cesari M: **Serum adiponectin and coronary heart disease risk in older Black and White Americans.** *J Clin Endocrinol Metab* 2006, **91**(12):5044-50.
- Laughlin GA, Barrett-Connor E, May S, Langenberg C: **Association of adiponectin with coronary heart disease and mortality: the Rancho Bernardo study.** *Am J Epidemiol.* 2007, **165**(2):164-174.
- Kato H, Kashiwagi H, Shiraga M, Tadokoro S, Kamae T, Ujiiie H, Honda S, Miyata S, Ijiri Y, Yamamoto J, Maeda N, Funahashi T, Kurata Y, Shimomura I, Tomiyama Y, Kanakura Y: **Adiponectin acts as an endogenous antithrombotic factor.** *Arterioscler Thromb Vasc Biol* 2006, **26**:224-230.
- Antman EM, Braunwald E: **ST-elevation Myocardial Infarction: pathology, pathophysiology and clinical feature.** In *Braunwalds Heart Disease* 8th edition. Edited by: Libby P, Bonow RO, Mann DL, Zipes DP. Elsevier Saunders company; 2008:1207.
- Cannon CP, Braunwald E: **UnStable Angina and Non ST- elevation Myocardial Infarction.** In *Braunwalds Heart Disease* 8th edition. Edited by: Libby P, Bonow RO, Mann DL, Zipes DP. Elsevier Saunders company; 2008:1243.
- Flier JS, Maratos-Flier E: **Biology of obesity.** In *Harrison's principle of internal medicine* 17th edition. Edited by: Fauci AS, Braunwald E, Kasper DL, Haase SL, Longo DL, et al. Mc Grow Hill company. Philadelphia: USA; 2008:262-3.
- Victor RG, Kaplan NM: **Systemic Hypertension: Mechanisms and Diagnosis.** In *Braunwalds Heart Disease* 8th edition. Edited by: Libby P, Bonow RO, Mann DL, Zipes DP. Elsevier Saunders company; 2008:1027.
- Powers AC: **Diabetes Mellitus.** In *Harrison's principle of internal medicine* 17th edition. Edited by: Fauci AS, Braunwald E, Kasper DL, Haase SL, Longo DL, et al. Mc Grow Hill company. Philadelphia: USA; 2008:2275.
- Iwashima Y, Horio T, Kumada M, Suzuki Y, Kihara S, Rakugi H, Kawano Y, Funahashi T, Ogihara T: **Adiponectin and renal function, and implication as a risk of cardiovascular disease.** *Am J Cardiol* 2006, **15**, **98**(12):1603-8.
- Cavusoglu E, Ruwende C, Chopra V, Yanamadala S, Eng C, Clark LT, Pinsky DJ, Marmur JD: **Adiponectin is an independent predictor of all-cause mortality, cardiac mortality, and myocardial infarction in patients presenting with chest pain.** *Eur Heart J* 2006, **27**(19):2300-9.
- Braunwald E: **Un stable angina: An etiologic approach to management.** *Circulation* 1998, **98**:2219.
- Marrow DA, Braunwald E: **Future of Biomarkers in acute coronary syndrome: Moving toward a multi marker strategy.** *Circulation* 2003, **108**:250.
- Stejskal D, Bartek J: **Adiponectine in patients with various stages of coronary heart disease – comparison of its concentration in coronary arteries and peripheral venous circulation.** *Biomed Papers* 2003, **147**(2):161-166.
- Ouchi N, Kihara S, Arita Y, Nishida M, Matsuyama A, Okamoto Y, Ishigami M, Kuriyama H, Kishida K, Nishizawa H, Hotta K, Muraguchi M, Ohmoto Y, Yamashita S, Funahashi T, Matsuzawa Y: **Adipocyte-derived plasma protein, adiponectin, suppresses lipid accumulation and class A scavenger receptor expression in human monocyte-derived macrophages.** *Circulation* 2001, **103**:1057-1063.
- Okamoto Y, Arita Y, Nishida M, Muraguchi M, Ouchi N, Takahashi M, Igura T, Inui Y, Kihara S, Nakamura T, Yamashita S, Miyagawa J, Funahashi T, Matsuzawa Y: **An adipocyte-derived plasma protein, adiponectin, adheres to injured vascular walls.** *Horm Metab Res* 2000, **32**:47-50.
- Frystyk J, Berne C, Breglund L, Jenesvik K, Flyvbjerg A, Zethelius B: **Serum Adiponectin Is a Predictor of Coronary Heart Disease: A Population-Based 10-Year Follow-Up in Elderly Men.** *J Clin Endocrinol Metab* 2007, **92**(2):571-576.
- Iwashima Y, Horio T, Kumada M, Suzuki Y, Kihara S, Rakugi H, Kawano Y, Funahashi T, Ogihara T: **Adiponectin and Renal Function, and Implication as a Risk of Cardiovascular Disease.** *Am J Cardiol* 2006, **98**:1603-1608.
- Pischon T, Girman CJ, Rifai N, Hotamisligil GS, Rimm EB: **Association between dietary factors and plasma adiponectin concentrations in men.** *Am J Clin Nutr.* 2005, **81**(4):780-786.

31. Saito O, Saito T, Okuda K, Okuda K, Kotoda A, Akimoto T, Ando Y, Muto S, Ishikawa SE, Kusano E: **Serum adiponectin and markers of endothelial injury in hemodialysis patients with arteriosclerosis obliterans.** *Clin Exp Nephrol* 2008, **12**:58-64.
32. Shibata R, Numaguchi Y, Matsushita K, Sone T, Kubota R, Ohashi T, Ishii M, Kihara S, Walsh K, Ouchi N, Murohara T: **Usefulness of Adiponectin to Predict Myocardial Salvage Following Successful Reperfusion in Patients With Acute Myocardial Infarction.** *Am J Cardiol* 2008, **101**:1712-5.
33. Inoue T, Kotooka N, Morooka T, Komoda H, Uchida T, Aso Y, Inukai T, Okuno T, Node K: **High Molecular Weight Adiponectin as a Predictor of Long-Term Clinical Outcome in Patients with Coronary Artery Disease.** *Am J Cardiol* 2007, **100**:569-574.

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