



Association of circulating omentin-1 level with lung cancer in smokers

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Abstract

Background: Omentin has recently been considered as an adipokine secreted from visceral fat and is expressed in the lungs, heart, ovary etc. Various studies have shown that omentin may have an anti-inflammatory role in the inflammatory process and the amount of omentin alters in some cancers, such as colorectal, prostate and renal cells cancers, changes. The serum level of omentin, however, remains unknown in non-patient smokers and the smokers afflicted with lung cancer. Therefore, this study examines the serum levels of omentin in smokers suffering from lung cancer.

Methods: The amount of serum omentin was measured in 45 patients with lung cancer and 61 age - and sex - matched controls (30 smokers and 31 non-smokers) using enzyme-linked immunosorbent assay (ELISA) kit. Data were analyzed using SPSS-16, and one-way analysis of variance and Scheffe post hoc test were used to determine and compare the serum levels of omentin in different types of lung malignancies. Significance level was set at $p \leq 0.05$.

Results: The amount of circulating omentin for healthy non-smokers and non-patient smokers was 3.55 ± 0.57 ng/l and 5.43 ± 1.95 ng/l, respectively ($p < 0.001$). The serum level of omentin was 3.63 ± 0.70 ng/l for smokers afflicted with cancer ($p < 0.001$ compared with non-patient smokers: 5.43 ± 1.95).

Conclusion: The meaningful decrease in omentin levels in smokers with lung cancer can be considered as a risk factor in smokers and can use as a significant factor in the prognosis of lung cancer in these people.

Keywords: Smoker, Omentin, Lung cancer

Conflicts of Interest: None declared

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Introduction

Lung cancer is one of the pulmonary diseases and is the uncontrolled growth of the epithelial cells of the lung and its prevalence in smokers is higher as compared to non-smokers. Around the world, lung cancer is the most common cancer in terms of outbreaks and death. In 2012, 1.8 million new items (12.9% of all people with all types of cancers diagnosed in this year) and 1.6 million deaths from lung cancer were reported (1). Being among the five major cancers in Iran, the lung cancer afflicts a large number of

people about 80% of which included by smokers (2). Adipose tissue is known as an endocrine and immune organ (3). Until recently, adipose tissue was thought to affect only lipid metabolism and glucose homeostasis. This tissue is now identified as the secretion source of more than 20 types of different hormones and molecules called adipocytokine or adipokine which play significant biological roles in the vascular system, glucose homeostasis, reproduction, bone metabolism, immune system and cancer (4-5).

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↑What is “already known” in this topic:

There are studies about the role of omentin hormone in a variety of diseases, such as cancer; however, its serum levels in patients with lung cancer are unknown.

→What this article adds:

Cigarettes can increase the level of hormones in the serum and the level of serum omentin hormone changes in smokers who have lung cancer.

As one of the adipokines, omentin (or intelectin) was first found in the intestinal paneth cells, which reacts with galactofuranose in the carbohydrate part of the cell wall of the bacteria and is involved in intestinal defense mechanisms against pathogenic bacteria (6). Omentin-1 is the type of omentin which is mostly addressed in research papers. There is another homolog of omentin which is called omentin-2. About 83% of its amino acids are identical to that of omentin-1. Omentin 1 is also expressed in the lungs, heart, ovary, and placenta (7-10) and is secreted as a new adipokine in visceral adipose tissue. The Laboratory experiments show that omentin increases the level of insulin sensitivity (i.e. increases the glucose uptake by insulin) (10) and activates Akt signaling pathway. Akt is a serine / threonine protein kinase that plays an important role in multiple cellular actions such as glucose metabolism, cell proliferation and apoptosis as a secondary messenger (7). Also, the increase in omentin leads to a decrease in the level of CRP in serum and decreases the effects of TNF- α (tumor necrosis factor alpha that increases in inflammatory conditions). So there can be a connection between omentin and inflammation (7, 11-12). Chronic inflammation increases the risk of cancer (12), and in some cancers, inflammation occurs before the onset of cancer and malignancy (1, 13). Cigarette smoking causes chronic inflammation in the lungs, and inflammatory mechanisms also cause a lung tumor under the influence of smoking (12).

The amount of gene expression of this adipokine is reduced in the epithelium of smokers' airways (14), and though it is known that omentin is secreted from the epithelial cells of the lung, it is not clear whether there is an increase or decrease in the serum level of this hormone in smokers with lung cancer.

Methods

Study population

The patient group of this study were 45 smoker male patients referred to the pulmonary clinic of the Imam Khomeini hospital in Urmia, Iran. The control group consisted of 61 males with no history of illness and with matched of age and BMI (Body mass index) (30 smokers and 31 non-smokers). The protocol of the study followed the principles of the Declaration of Helsinki. All the subjects signed the informed consent form. Initially, careful lung examinations were conducted and then the tissue biopsy specimens were sent to the histopathology laboratory for more accurate diagnosis. The blood and urine parameters for all the individuals in both groups were within normal ranges.

Laboratory analysis

Before starting any type of therapeutic treatment in patients, peripheral blood was drawn from subjects after

measuring their height and weight to calculate BMI. After separation of the serum of the samples in a cooling centrifuge and spinning at 3000 rpm for 10 min, the biochemical parameters such as fasting blood glucose (FBG), triglyceride (TG) and low density lipoprotein – cholesterol (LDL-c) were detected via the standard protocols using a clinical chemistry autoanalyzer (BT 3000, Italy) in the laboratory of Imam Khomeini Hospital in Urmia in the sampling day. Then serum was frozen at -80 °C until the analysis of omentin. Serum omentin was measured using commercially available enzyme-linked immunoassay (ELISA) kits (Bioassay technology laboratory, China) according to manufacturer's instructions. The sensitivity of the assay was 1.03 ng/l (assay range: 2-600 ng/l). The intra- and inter-assay coefficients of variance were <8% and <10%, respectively.

Statistical analysis

All data were analyzed using the SPSS software package version 16. The results are presented as means \pm S.D. The analyses of normally distributed variables (age and BMI and biochemical analysis) were conducted using independent-sample t-test. To determine and compare the serum levels of omentin in different types of lung malignancies, one-way analysis of variance was used and finally, Scheffe post hoc test was used as well. Kolmogorov-Smirnov test were used to determine whether or not they were normally distributed and the analyses of abnormally distributed variables were conducted with the Mann-Whitney U test. Comparisons with $p < 0.05$ were considered to be statistically significant.

Results

The baseline characteristics of the subjects are summarized in Table 1. The mean age in the control group who did not smoke was 58.87 ± 7.19 whereas it was 58.65 ± 7.82 for the smokers in the control group. The mean BMI of these the two groups were 24.58 ± 2.93 and 24.10 ± 2.93 , respectively. The mean age of the patient group was 65.13 ± 9.32 , and the mean BMI of the patient group was 23.82 ± 4.28 . The values of other biochemicals are given in Table 1.

Figure 1 shows the changes in the level of omentin hormone in the healthy and cancerous groups as follows: 3.55 ± 0.57 ng/l for nonsmokers and 5.43 ± 1.95 ng/l for smokers in the control group and 3.63 ± 0.70 ng/l for the patient group.

The amount of serum omentin was different in a variety of pulmonary malignancies. It was 3.04 ± 0.38 ng/l in SCC (squamous cell carcinoma), 4.08 ± 0.31 ng/l in adenocarcinoma, 3.88 ± 0.50 ng/l in small cell carcinoma and 4.27 ± 0.65 ng/l in other lung diseases (for example poorly differentiated carcinoma) according to Figure 2.

Table 1. The baseline characteristics of subjects

Variables	Non- smoker, non- patient (n=31)	Smoker, non- patient (n= 30)	Smoker, patient (n= 45)	p
Age (years)	58.87 ± 7.1	58.65 ± 7.82	65.13 ± 9.32	0.061
BMI (kg /m ²)	24.58 ± 2.93	24.10 ± 2.93	23.82 ± 4.28	0.917
FBS (mg/dl)	81.35 ± 9.58	82.16 ± 9.45	82.32 ± 10.44	0.946
TG (mg/dl)	95.83 ± 7.91	94.30 ± 8.94	95.86 ± 8.97	0.438
LDL-c (mg/dl)	79.19 ± 7.73	79.26 ± 9.31	80.13 ± 11.27	0.717

Data are presented as mean \pm SD. BMI (body mass index), FBS (fasting blood sugar), TG (triglyceride), LDL-c (low-density lipoprotein-cholesterol).

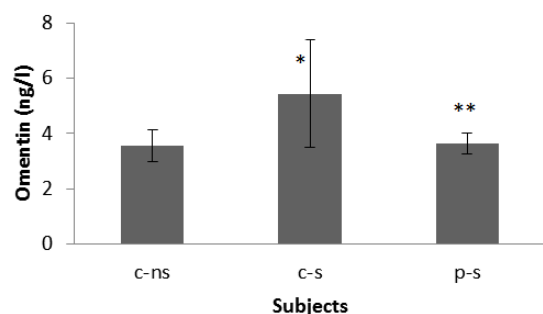


Fig. 1. The levels of omentin in the serum of subjects

Data are presented as mean \pm SD. Control-nonsmoker (c-ns), control-smoker (c-s) and patient-smoker (p-s). *Indicates a statistically significant difference vs. c-ns ($p < 0.05$). ** Indicates a statistically significant difference vs. c-s ($p < 0.05$).

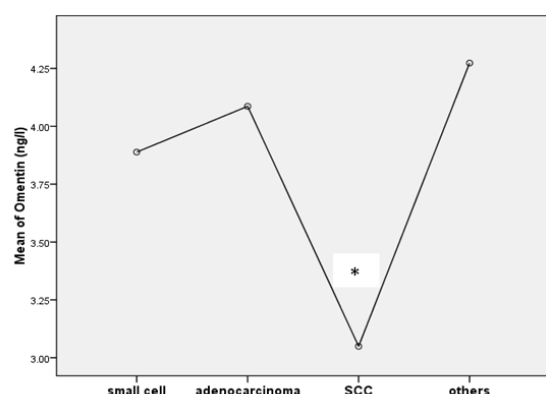


Fig. 2. Serum level of omentin in the variety of lung malignancies

Decreasing plasma omentin in patients with SCC shows a significant decrease compared with other patients. Data are expressed as mean. *indicates a statistically significant difference relative to the other patients ($p < 0.05$) (SCC vs adenocarcinoma, SCC vs small cell and SCC vs others).

Discussion

The present study demonstrates that the serum level of omentin increases in smokers compared to non-smokers (neither with a history of illness), whereas Brendan and colleagues in 2008 proved that the expression of the omentin gene is reduced in the epithelial cells of the airways of smokers (14). Smoking can predispose a person to inflammatory diseases (15). The level of serum omentin decreases in inflammatory diseases such as Crohn's disease (chronic inflammatory bowel disorder) and rheumatoid arthritis. In these inflammatory diseases, omentin is reduced as an anti-inflammatory factor (7) and on the other hand in patients with chronic kidney disease that do not undergo dialysis as well as in non-alcoholic fatty liver patients, serum levels of omentin increase. The cause of this increase in the amount of omentin in these patients is its defective degradation and excretion in renal patients or lack of synthesis of some inflammatory factors such as CRP in patients with liver disease (16-17). In 2013, a study by Wang and his colleagues showed that omentin is decreased in the serum of OSAS (obstructive sleep apnea syndrome) patients, while a study by Kar Kurt in 2014 showed the opposite result. In the Wang et al. study, cases had no history of any other illness (such as diabetes, tumors, or alcohol and drug abuse) other than OSAS, but in the other study performed by Kar Kurt et al. only OSAS patients without specific comorbidities in-

cluding diabetes, CVD (cardiovascular diseases) and metabolic syndrome were included (18-19). The use of omentin reduces the inflammatory effects and also improves muscle cell proliferation in the culture of human smooth muscle cells of the aorta (20) and reducing the amount of omentin increases the formation of atherosclerotic ulcers by inducing inflammatory responses of macrophages through (5'-AMP-activated protein kinase) AMPK-Akt signaling pathway (21) and omentin also reduces inflammatory responses in endothelial and vascular smooth muscle cells that are stimulated with TNF- α (22-23). Thus, omentin has anti-inflammatory properties (17). Based on the present study it can be said that a rise in the serum level of omentin in non-patient smokers is due to its secretion from other sources. The result may be supported by the study conducted by Watanabe in 2016 that an increase in the amount of omentin in serum of patients with CAD (coronary artery disease) in spite of the decrease in the expression of omentin in the epicardial adipose tissue in these patients is due to the secretion of omentin from the macrophages in the atherosclerotic plaques (20).

Since the discovery of omentin, its role in tumorigenesis has been shown in a large number of studies. In women with EEC (Endometrioid Endometrial Cancer) and CAH (Complex Atypical endometrial Hyperplasia), the levels of serum omentin decrease in relation to BMI-matched healthy people. EEC is an inflammatory disease that is associated with obesity and diabetes (24). Also, in women with ovarian cancer the levels of serum omentin decrease. Besides, the power of proliferation, motility, and cellular invasion in these cancer cells is reduced by the effect of omentin (25). So that patients with high serum omentin have a longer life span than those with lower serum omentin. In this case, it is hypothesized that there may be a negative effect on cell proliferation (24). It has been proved in various studies that the level of serum omentin increases in prostate, colorectal and pancreatic adenocarcinoma cancers, while it decreases in RCC (Renal Cell Carcinoma) and Bladder cancer. It seems that increasing the amount of omentin in some cancers stimulates cell growth by triggering genomic instability and PI3K/Akt (phosphatidylinositol-3 kinase downstream effector) signaling pathways thereby contributing to

the pathogenesis of these cancers (26-29). On the other hand, omentin plays a major role in inflammatory responses and cellular differentiation and has a significant contribution to inducing apoptosis in cancer cells (30-31). Omentin stimulates apoptosis in cells through the activation of JAK signaling pathway and P53 upregulation mechanisms (30). It also inhibits inflammatory diseases via suppression of JNK (Jun N-terminal kinase) activation through the AMPK/eNOS (endothelial nitric oxide (NO) synthase) signaling pathway (22). Also omentin reduces the expression of VCAM-1 (vascular cell adhesion molecule-1) and ICAM-1 (intercellular cell adhesion molecule-1) (the molecules involved in the pathogenesis of cancer cells) adhesion molecules through inhibition of P38, JNK pathway and blocking of ERK/NF- κ B (extracellular signal-regulated kinase/nuclear transcription factor kappa B) pathway (23, 31). Another mechanism of omentin in triggering apoptosis

in cells is increasing the bax/bcl-2 protein ratio and activation of the caspase-3 signaling pathway (30). Therefore, the reduction of omentin in RCC and ECC increases cellular proliferation and decreases apoptosis in cancer cells. In this study, we demonstrated that the circulation of omentin is decreasing in the lung cancer patients and the decrease in its amount in the smoker patients is significant compared to non-patient smokers.

About 85% of the total lung cancer cases were non-small cell lung cancer (NSCLC) which mainly includes squamous cell carcinoma (32) and adenocarcinoma, and these contain approximately 400,000 deaths each year in the world (32-33). Smoking is the main cause (85-90%) of lung cancer (34), and the prevalence of SCC is more than of adenocarcinoma in smokers (33). Metastasis and recurrence are very common in SCC (32) and in the invasive types of the tumor with rapid initial growth (35). As shown in this study, the amount of serum omentin in SCC patients decreases more significantly than in patients afflicted with other lung malignancies.

Conclusion

Omentin-1 levels were found to be significantly lower in smoker lung cancer patients, irrespective of their general clinical conditions. These findings suggest that this adipokine might play some crucial role in the tumorigenesis of lung cancer via mechanisms that are active in lung cancer. Thus, the measurement of omentin in smokers may be considered as a significant factor in the prognosis of lung cancer in smokers and should be further investigated.

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Conflict of Interests

The authors declare that they have no competing interests.

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