Adverse Health Consequences of Dietary Advanced Glycation End Products (ages) and Inhibitory Effects of Natural Ingredients on Ages

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Abstract

Dietary advanced glycation end products (AGEs) may result in various adverse health consequences, such as inflammation, cancer, liver diseases, obesity, cardiovascular diseases, diabetes, proteostasis impairment and memory decline. Natural ingredients may be a promising strategy to inhibit AGEs generation and ameliorate health risks of AGEs. The mechanisms involved in inhibition of AGEs with use of natural ingredients include antioxidant activity, reactive carbonyl trapping, and suppressing pathways of RAGE (receptor of advanced glycation end product), MAPK (mitogen-activated protein kinases) and NF-κB (nuclear factor-kappa B). This mini review summarizes the present understanding of the adverse impact of AGEs on health and the inhibitory effect of natural ingredients on AGEs.

Keywords: Maillard reaction; Dietary advanced glycation end products; Adverse health consequences; Inhibitor; Natural ingredients

Abbreviations: AGEs: Advanced Glycation End Products; NAFLD: Non-Alcoholic Fatty Liver Disease; CML: Carboxy Methyl Lysine; RAGE: Receptor of Advanced Glycation End Product; MAPK: Mitogen Activated Protein kinases; LSOPC: Lotus Seedpod Oligomeric Pro Cyanidins

Introduction

Maillard reaction, a form of non-enzymatic browning, is a chemical reaction between proteins and reducing sugars in vitro and in vivo [1,2]. The Maillard reaction consists of three stages: the early, advanced and final stages. Briefly speaking, the initial glycosylation reaction occurs during the early stage of the Maillard reaction, and amadori products are formed through rearrangement of Schiff bases [3]. For the advanced stage, the Amadori products subsequently form cross-linked structures with a high degree of complexity named advanced glycation end products (AGEs) through fission, dehydration and Strecker degradation [1]. In terms of final stage, nitrogenous polymers and co-polymers termed melanoidins are formed [1]. Although Maillard reaction may aid in development of attractive color, flavor and aroma in food products, it is worth pointing out that AGEs in Maillard reaction products may lead to adverse health consequences [4,5]. In recent years extensive research reveals that AGEs are associated with various diseases such as cancer and diabetes [4,6]. Foods are major sources of AGEs, and commonly consumed foods such as grilled chicken, broiled steak and French fries contain relatively high content of AGEs [7]. Meanwhile, a recent study indicates that a high level of dietary AGEs is associated with a high level of AGEs in plasma and urine [8]. Because contents of AGEs derived from diet cannot be ignored and AGEs in the diet are associated with induction and progression of many chronic diseases, it is essential to comprehensively understand adverse health consequences of dietary AGEs.

In view of numerous health risks of dietary AGEs, multiple measures are required to suppress AGEs. Medicine such as metformin and irbesartan can be applied to relieve adverse health effects induced by AGEs [9]. However, the synthetic pharmaceutical ingredients may bring about short-term health risks. Metformin may lead to digestive disorders such as vomiting and diarrhoea and irbesartan can sometimes result in symptoms such as dizziness and headache [10,11]. Therefore, considering the side effects of synthetic ingredients, natural ingredients should be employed as AGEs inhibitors in place of synthetic ingredients. During recent decade’s inhibition of AGEs with natural ingredients have drawn substantial scholarly attentions. More and more natural ingredients prove to possess inhibitory influence on AGEs, and significant achievements about the underlying molecular mechanisms have been made [3,12,13]. Thus, the present mini review aims to address the current understandings towards adverse health consequences of dietary AGEs and inhibitory influence of natural ingredients on AGEs.
Adverse Health Consequences of Dietary Advanced Glycation End Products

Extensive biomedical studies about health effects of dietary AGEs on animals and human beings have been conducted over the past few years [14,15]. As shown in Figure 1, relevant research indicates that dietary AGEs are closely linked to induction and progression of diverse diseases such as inflammation, cancer, liver diseases, obesity, cardiovascular diseases, diabetes, proteostasis impairment and memory decline. It is worthwhile to note that the adverse health consequences of dietary AGEs depend on actuation duration. At times short-term intake of dietary AGEs may not have profound health risks. A recent study revealed that short-term ingestion of advanced AGEs could possibly exert little health influence on rats [16].

![Figure 1: Adverse health consequences of dietary AGEs.](image)

Inflammation

Several studies support the association between dietary AGEs and inflammation. Based on accurate analysis of AGEs content in the diet, a cross-sectional study divided participants into two groups: high and low intake of dietary AGEs. During the 3-month period a correlation between intake of dietary AGEs and inflammation was observed, indicating that a chronic AGEs-rich diet could result in inflammation [5]. In addition, it was reported that long-term consumption of AGEs in thermally processed red meat could lead to chronic inflammation [4].

Cancer

Recent studies reveal that dietary AGEs can increase risk of certain cancers. After analysis of more than 2000 human pancreatic cancer cases during an average of 10.5-year follow-up, it was found that higher intake of dietary AGEs contributed to higher risk of pancreatic cancer in men, and consumption of dietary AGEs was not linked to risk of pancreatic cancer in women [4].

Liver diseases

AGEs in food may induce various liver diseases including liver injury, liver fibrosis, liver inflammation and so on. In mice model of NAFLD (non-alcoholic fatty liver disease), long-term exposure to high levels of dietary AGEs worsened liver pathology through modulating liver injury and liver fibrosis, and reducing intake of dietary AGEs might weaken the harmful effects [14]. Nε-(carboxymethyl) lysine (CML) is one of the major dietary AGEs, and oral acute and sub acute toxicity of CML was evaluated in mice models. It was found that CML presented negligible acute hazards, but repeated ingestion of CML could induce liver damage via oxidative stress [15]. Influence of dietary AGEs on liver was studied using a mice model, and the results suggested that diets with high levels of AGEs could induce liver inflammation in the absence of steatosis [17].

Obesity and cardiovascular diseases

Obesity is closely associated with cardiovascular diseases, and they are both major health problems [18]. Recent studies have demonstrated that dietary AGEs can bring about obesity and cardiovascular diseases. Different groups of mice were fed with diets containing high or low levels of AGEs, and weight gain of these mice was analyzed after 6 weeks. It was observed that a high-AGEs diet could contribute to weight gain [19]. Although evidence is still lacking about links between dietary AGEs in children and adolescents, sufficient data demonstrate that dietary AGEs can induce vascular dysfunction in adults, contributing to occurrence of cardiovascular diseases [20].

Diabetes and proteostatis impairment

A large amount of evidence suggests that dietary AGEs may interfere with metabolism of sugars and proteins, leading to diabetes and proteostatis impairment. Mice were treated with oral AGEs, and a series of indexes such as glucose tolerance, glucose uptake and gene modulation were analyzed. It was found that chronic ingestion of AGEs promoted insulin resistance and diabetes, and the relevant mechanism was possibly depleting the antioxidant defenses AGE receptor-1 and sirtuin-1 [6]. In vivo studies revealed that chronic intake of dietary AGEs may cause disruption of proteostasis through inhibiting proteasome peptidase activity [21].

Memory decline

Correlations exist between dietary AGEs and cognitive decline. High oral intake of AGEs was reported to be associated with faster rate of memory decline in young elderly. To avoid disruption of daily life by memory loss, the intervention method of taking a low-AGEs diet may reduce risk of cognitive compromise [22].

Inhibiting formation of Advanced Glycationend Products with Natural Ingredients

Given that AGEs can bring about potential health risks, various measures should be taken to suppress formation of AGEs. During recent years natural ingredients prove to be a reliable means to reduce contents of AGEs. Due to high biosafety and biocompatibility of natural ingredients, inhibiting AGEs with natural ingredients has attracted scientific and industrial interests in many fields. In general, natural ingredients can suppress AGEs formation via mechanisms such as antioxidant activity, reactive carbonyl trapping, and disruption of RAGE-MAPK signaling and NF-κB activation.

Antioxidant Activity

It is reported that many natural ingredients have powerful antioxidant activity, which helps to scavenge free radicals. Multiple studies demonstrate that inhibitory influence of some natural ingredients is ascribed to their antioxidant activity. Influence of
ferulic acid on in vitro glycation of soy proteins was investigated utilizing Nε-(carboxymethyl)lysine (CML) analysis and fluorescence measurement. It was found that use of ferulic acid could reduce formation of CML and fluorescent AGEs by around 90% [3]. Recent work investigated inhibitory effects of ethanolic extracts from 17 medicinal plants on glycation of bovine serum albumin, and total phenolic content and antioxidant capacity of the medical plant extracts were also studied. The results demonstrated that influence of medical plant extracts on AGEs inhibition was positively correlated with free radical scavenging activity, indicating that the antioxidant activity of medical plant extracts contributed largely to their antiglycation effects [23].

A 4-week study revealed that red grape skin extract at different concentrations (0.031–0.500 mg/mL) could suppress formation of AGEs during relatively long-term exposure to AGEs [24]. In comparison with red grape skin extract, Mesona Chinensis Benth extract required higher effective concentrations. Mesona Chinensis Benth extract at different concentrations (0.25–1.00 mg/mL) significantly inhibited AGEs formation during 4 weeks of study [25]. In addition, olive leaf extract could inhibit formation of AGEs in a biscuit model, and the main way of action was possibly inhibiting oxidation of reaction intermediates towards the AGEs formation [26]. Apart from these in vitro studies, in vivo studies also indicate that antioxidant activity of certain natural ingredients is associated with their inhibitory effects on AGEs. Influence of Cuminum cyminum monon AGEs inhibition in diabetic rats was systematically investigated, and the results showed that Cuminum cyminum could control oxidative stress and suppress formation of AGEs. The in vivo inhibitory effects of Cuminum cyminum on AGEs proved to be correlated with free radicals scavenging of Cuminum cyminum [27].

Reactive Carbonyl Trapping

Among current published literature reactive carbonyl trapping is the major mechanism involved in inhibition of natural ingredients on AGESs. Methylglyoxal and glyoxal are major carbonyl compounds during Maillard reaction and identified as well-known precursors of AGESs, and trapping these reactive carbonyl compounds may reduce formation of AGESs. A recent study demonstrated that anthocyanins in blackcurrant could react with 50 percent of methylglyoxal when blackcurrant anthocyanins were incubated with methylglyoxal, suggesting that blackcurrant anthocyanins could effectively prevent AGES formation by trapping methylglyoxal [29].

Polyphenols in Houttuynia cordata consisted of components quercetin, chlorogenic acid and rutin. It was found that polyphenols in Houttuynia cordata could decrease AGES production by 91-94.6% [28]. Genistein is a natural isoflavone derived from soy products, and genistein could inhibit AGES production via trapping reactive dicarboxyl species efficiently under in vitro physiological conditions (pH 7.4, 37°C) [13]. Rutin metabolites, which were formed through metabolism of rutin by gut microflora, could serve as effective natural inhibitor of AGESs by trapping both methylglyoxal and glyoxal [30]. Since rutin is a commonly consumed flavonoid that is rich in many fruits and vegetables, the use of rutin metabolites may be a feasible way to attenuate AGESs formation. Pomegranate phenolics and their in vivo metabolites had inhibitory effects on all three stages of Maillard reaction (early, middle and late stages), and pomegranate phenolics could significantly suppress AGESs production via scavenging reactive carbonyl species such as methylglyoxal [31].

Since pomegranate is a fruit with good flavors and odors, oral intake of pomegranate appears an attractive dietary strategy to prevent and offset adverse health consequences of AGESs. Anti glycation effects of phenolic phyto chemicals extracted from blueberries, blackberries, strawberries, raspberries, cranberries and grapes were investigated, and the results showed that these sugar-free phyto chemicals could suppress AGESs generation by scavenging reactive carbonyls [32]. In addition, natural ingredients such as chlorogenic acid, ellagic acid, thiamin, quercetin, and oligomeric procyanidins of lotus seedpod, herbal teas and thymoquinone could effectively inhibit AGESs generation through reactive carbonyl trapping [12,33-38].

Disruption of RAGE-MAPK Signaling and NF-κ B Activation

Receptor of advanced glycation end product (RAGE), mitogen-activated protein kinases (MAPK) and nuclear factor-kappa B (NF-κB) are detrimental pathways in animals and people, and a number of evidence reveals that inhibitory effects of natural ingredients on AGESs are associated with disruption of RAGE-MAPK signaling and NF-κB activation. High-fat diet rats were selected as model to study inhibition of AGESs by lotus seedpod oligomeric procyanidins (LSOPC), and expression of RAGE, MAPK and NF-κB was investigated. It was found that LSOPC could inhibit formation of AGESs in rats through suppression of RAGE-MAPK-NF-κB signaling [39]. Diabetic cardiomyopathy rats were applied to investigate influence of mangiferin on formation of AGESs, and the results demonstrated that mangiferin could effectively inhibit AGESs formation via NF-κB deactivation and decreasing expression of RAGE [40].

Concluding Remarks and Future Perspectives

In summary, dietary AGESs may lead to the induction and progression of various health problems such as inflammation, cancer, liver diseases, obesity, cardiovascular diseases, diabetes, proteostasis impairment and memory decline. Natural ingredients may serve as an approach to reducing AGESs generation, and the involved inhibition mechanisms are antioxidant activity, reactive carbonyl trapping, and disruption of RAGE-MAPK signaling and NF-κB activation. Generally speaking, the natural ingredients are promising intervention tools to control content of AGESs. Meanwhile, we need to realize that there are many unexplored areas and ongoing challenges. Molecular mechanisms involved in health risks of dietary AGESs should be fully understood in details, and more in vivo animal and human studies should be conducted to confirm the inhibitory effects of natural ingredients on AGESs generation.

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