Beneficial Effects of Chlorogenic Acids on Essential Hypertension

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Abstract
Hypertension is a worldwide condition considered as the most important risk factor in the development of the cardiovascular disease. The contribution of the oxidative stress in its pathophysiology is a well known fact. Polyphenols are potent antioxidant compounds able to mitigate the damage produced by reactive oxygen species. Chlorogenic acids are polyphenols occurring in many dietary sources but primarily in green coffee extracts. Several studies have shown their antioxidant and anti-inflammatory properties. In the last decades there have been performed several clinical trials showing that the use of these compounds decrease the systolic and diastolic blood pressures probably by increasing the bioavailability of nitric oxide and improving the endothelium function. These properties could be the bases for supporting an important adjuvant therapy for the treatment of essential hypertension.

Keywords: Chlorogenic acids, Essential hypertension, Reactive oxygen species, Nitric oxide, Endothelium, Oxidative stress, Antioxidants.

Introduction
Increasing body of evidence has involved the contribution of oxidative stress in the mechanism of essential hypertension[1]. Accordingly, it has been demonstrated that patients with hypertension have increased production of reactive oxygen species (ROS) [2,3] and lower levels of antioxidant species[4]. In addition, it has been found that ROS production is enhanced, redox-dependent signaling is amplified, and antioxidant bioactivity is reduced in the arteries of hypertensive humans[5], all accounting for a role of vascular oxidative stress in the pathogenesis of essential hypertension[6-8].

The ROS play a physiological role in the homeostasis of the vascular wall by regulating the vascular tone and endothelial function[6,9]. This regulation is related to the activation/down-regulation of metabolic pathways modulated by low-to-moderate concentration of ROS[8,9]. However, when there is an imbalance between ROS production and antioxidant species the endothelium shifts its actions generating a reduction in the vasodilation of the wall and an increasing in the proinflammatory state and prothrombotic setting[10].

In the vascular wall different sources of ROS coexist, which can be classified as enzymatic and nonenzymatic compounds. NADPH oxidase is the primary biochemical source of superoxide in the vascular endothelium. Nevertheless, there are other enzymes that contribute to the oxidative stress including xanthine oxidase, mitochondrial enzymes and uncoupled endothelial nitric oxide synthase (eNOS)[11].

The main generator of ROS in the vasculature is NADPH oxidase (NOX) that produces superoxide anion by reducing molecular oxygen. This enzyme is up-regulated in hypertension primarily by molecular signs[11]. The superoxide anion can interact
with some molecules to form other free radicals or can react with biomolecules generating a disruption in the biological processes of the cell. For example, L-arginine and tetrahydrobiopterin (BH4) are two essential cofactors of eNOS. Their deficiency or oxidation (BH4) is associated with uncoupling of the L-arginine-nitric oxide (NO) pathway resulting in decreased formation of NO and increased eNOS-mediated generation of superoxide instead. Furthermore, superoxide can combine with NO to form peroxynitrite that can also oxidize and destabilize eNOS to produce more superoxide\cite{12,13} leading to a positive feedback reaction. In turn, the xanthine oxidase enzyme catalyzes the last two steps of purine metabolism. During this reaction, oxygen is reduced to superoxide. There is evidence that in spontaneously hypertensive rats (SHR) the xanthine oxidase is up-regulated, thus leading to increased ROS production and increased vascular tone\cite{14}.

The mitochondria are a major source of ROS. It has been studied that a part of the superoxide that is produced in the intermembrane space may be carried to the cytoplasm\cite{15}, being the complex I the main source of superoxide generated in the mammalian mitochondria. Normally, the other complexes do not produce significant levels of superoxide. Aside from this increased ROS source, it has been reported a reduction in antioxidant enzymatic activity in patients with hypertension\cite{16}.

There are some factors able to protect the tissues against oxidative stress, being NO one of the most important endogenous factors. The latter is known to play an important role as a key paracrine regulator of vascular tone.

Physiologically, NO inhibits leukocyte–endothelial cell adhesion, vascular smooth muscle cell (VSMC) proliferation and migration, and platelet aggregation, all contributing to maintain the health of the vascular wall. Therefore, taking into account the diverse beneficial effects of NO, it is conceivable that decreased NO bioavailability in the vasculature reduces vasodilatory capacity, thereby contributing to the development of hypertension.

Except the vasorelaxing and antiproliferative properties per se, NO plays an important role in antagonizing the effects of angiotensin II (AT-II), endothelins and ROS\cite{17}.

NO not solely antagonizes the effects of AT-II on vascular tone, cell growth, and renal sodium excretion, but also down-regulates the synthesis of angiotensin I converting enzyme (ACE) and AT1 receptors expression\cite{1,11}.

It is noteworthy that exogenous antioxidants are also able to abrogate the effect of increased ROS in the vascular wall, thereby leading to beneficial effects. On this line, polyphenols can be used to decrease the oxidative stress. In the last decades there were studies demonstrating that these compounds have antioxidant and anti-inflammatory properties. They can participate in the modulation of different cellular pathways to improve the endothelial function\cite{17}. For example they can enhance the bioavailability of NO by different mechanisms including activation of eNOS by the PK3-kinase/Akt pathway\cite{19}. In addition, they can inhibit ROS generators enzymes such as NADPH and xanthine oxidase and increase the levels of glutathione\cite{17}. On the other hand, there is also evidence that they can prevent the COX-dependent formation of endothelium-derived contracting factors\cite{19}.

Chlorogenic acids (CGAs) are one of the polyphenols that have gained interest during the last years. There are several studies that related CGAs properties with an improvement of glucose and insulin metabolism\cite{20}. Knowing that hypertension is the most important risk factor in the development of cardiovascular diseases (CV)\cite{21}, it could represent a relevant target of reducing the cardiovascular risk; in agreement with several studies showing that the use of green coffee extracts (GCEs) causes decreased values of blood pressure in humans\cite{22}. Those effects have been attributed to the polyphenols present in the GCEs. As CGAs are the major polyphenols present in coffee, this effect was attributed primarily to them. All of these properties could be probably explained by their capacity to improve the bioavailability of nitric oxide and endothelial function. However it is important to considerer that even though there are not many studies about the possible negative effects of CGAs in humans, some experiments in rats have shown more incidence in carcinomic effects when treated with caffeic acid\cite{23}. That fact probably indicates that it should be considerer a balance between the positive and negative effects of these compounds.

The aim of this chapter was to present an update of the studies supporting a role of CGAs to counteract the oxidative stress and herein the development of essential hypertension.

The Role of Chlorogenic Acid in the Essential Hypertension

The CGAs are a family of polyphenols formed through a reaction of esterification between a trans-cinnamic acid and a quinic acid (Figure 1). They are found in many dietary sources such as fruits, vegetables and seed. The most commonly CGA found in coffee, is 5-O-cafeoylquinic acid, often called “chlorogenic acid”, being one of the most important source of CGAs\cite{24}. Moreover, due to the high consumption of coffee worldwide, CGAs are one of the major founts of polyphenols in the human diet.

Recently, there have been several studies that analyze the importance of these compounds in some diseases. One of the most interest areas is their possible protection in cardiovascular diseases.

There is evidence that the use of CGAs may improve the glucose metabolism and insulin sensitivity\cite{25}, thereby reducing the relative risk of type 2 diabetes\cite{25,26}. In addition, it probably improves the reduction of weight in obesity\cite{27,28} and it is related with the reduction of the relative risk of stroke in women patients\cite{29,30}.
Mechanism of CGAs against Essential Hypertension

Like polyphenols CGAs have antioxidant and anti-inflammatory properties but their mechanism is not completely understood\[22\]. The total effect in the blood pressure is produced by a combination of various CGAs metabolites with different power of action\[31\]. With a common dietary intake of GCEs, CGAs are hydrolyzed in the small intestine to form the two principal components of the CGAs which are trans-cinnamic acid (such as caffeic and ferulic acids) and quinic acid. Some studies have analyzed that the major percentage of CGAs present in an infusion of coffee are formed by a caffeoylquinic acids (86 %)\[32-44\]. The hydroxylation takes place in the small intestine but only one third of the CGAs is absorbed there. The majority of the CGAs metabolites are absorbed in the large intestine after interacting with the microflora where caffeic acid can be converted to ferulic acid by a transferase or be reduced to dihydrocaffeic acid\[45\].

Studies in SHR suggested that the most powerful hypotensive effect is produced by ferulic acid with an approximately 9 times better decrease than caffeic acid and 17 times better decrease than quinic acid\[31\]. For that reason we will emphasize our analysis in the ferulic acids (FAs).

The hypotensive effect of FAs could be probably explained from their antioxidant properties. They can act not solely as scavengers\[35\] but also as non-selective NOX antagonists\[36,37\], contributing to a minor production of superoxide and less formation of ROS. This condition will protect the eNOS from disruption and increase the bioavailability of NO\[38\], leading to an improvement in the endothelial function. In addition, there is evidence that they can enhance acetylcholine induced endothelial dependent vasodilation\[38\] and inhibit the vascular proliferation of SMVC induced by angiotensin II\[39\].

Caffeic acid shows similar properties of FAs like their regulation of some function of angiotensin II. In SHR it was analyzed that these effects were due to a blocking of metabolic pathways involved in the process like JAK/STAT cascade or Ras/Raf signaling\[40\]. Additionally, it has been observed that caffeic acid could decrease the levels of Rac 1 GTPase, which can participate in the oxidative stress by contributing to generation of ROS\[36,41\].

Furthermore, CGAs might interact with the renin-angiotensin aldosterone system by inhibiting ACE activity as shown both in vitro and in vivo\[41,43\].

As polyphenols, the anti-inflammatory properties of CGAs could not be explained by their antioxidant properties. They probably will regulate some signaling pathways such as iκb/NF-κb or even interfere with the COX-2 activation. In a study of hepatic ischemia reperfusion injury in rats, CGAs shows protection properties. In addition they inhibited the translocation of the factors NF-κb and IRF-1 and induce the Nrf2\[42\].

However this area has been less explored and not well elucidated yet but it may have importance in the long-term regulation and could be an important therapeutic strategy for atherosclerotic disease\[31,44\]. Figure 2

Studies in Human Patients

Numerous studies investigating the effect of CGAs in blood pressure have been performed however, in a recent review of meta-analysis of randomized clinical trials (RCTs), only 5 of them where found eligible and with duration of more than 4 weeks. They demonstrated a significant reduction in the systolic (MD: -4.31 mmHg) and diastolic (MD: -3.68 mmHg) blood pressures compared with the placebo/control group\[46\]. Patients selected in two RCTs were normotensive whereas in three of them they had mild essential hypertension. There were no adverse effects reported in those studies.

However these studies have some limitations that are important to expose. One of them is that all of them took place in Asia (four in Japan and one in India). This shows that it is unclear if the properties of CGAs can be present in occidental people. In addition, the studies have different methodology, design, duration (ranges 4 to 26 weeks) and have small sample size which can probably biased the results\[47-50\].

Perspectives

It seems that CGAs could have some beneficial effects on essential hypertension, based on their biological properties. There is a biomolecular basis that supports this theory and studies that have positive results. However, it is necessary to diminish the limitations of those studies in order to improve their eligibility. For example, it is imperative to analyze these results in people with descent other than Asian, have a bigger sample size, increase the duration of the studies in order to investigate if it is really no adverse effects, determine the minimum effective dose of CGA which can reduce the blood pressure and unified the methodology and design of the studies in order to reduce the big heterogeneity that they have between each other.
Nonetheless, the use of CGA could be very useful in patients with incipient or low hypertension due to the molecular basis of their action that antagonizes with the remodeling mechanisms in blood vessels. This effect is more debatable in patients with high or long term hypertension.

Although there is a reduction in the blood pressure values, the impact of that reduction in the clinical practice is modest at best. It is likely that the use of CGA could be focused on the prevention rather than the treatment of established hypertension.

Conclusion

Cumulated evidence shows that oxidative stress participates in the pathophysiology of the essential hypertension. In this context, is reasonable to propose that polyphenols such as CGAs could have protective effects in this disease. The use of CGAs has been related to a vasodilatory response probably mediated by an increase of NO bioavailability in the vascular wall. This leads to a modest decrease of the systolic and diastolic blood pressure values. Although the results of RCTs are controversial due to the big heterogeneity in the designs between each other, it seems consensual that CGAs can reduce the blood pressure values. The clinical impact of this reduction is still uncertain but it probably could support a prevention strategy or as a complement to the conventional established therapies for treatment of essential hypertension (adjunct therapy).

Finally, it is important to note that the molecular mechanisms involved in the vasodilatory response are not elucidated yet and there are probably more molecular pathways able to participate in the antioxidant function. For example, the regulation of the metabolic Keap1/Nrf2 pathway which is activated in low-to-moderate ROS concentration, thus giving rise to a very important target of study because it promotes the antioxidant response cell. As CGAs acts as free radical scavengers, they could decrease the concentration of ROS and indirectly activate this pathway.

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