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Herbal beverages: Bioactive compounds and their role in disease risk reduction - A review

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ABSTRACT

There is a renewed interest in non-nutritive bioactive compounds of foods and beverages as 'lifespan nutrients' in the risk reduction of non-communicable diseases. Herbal beverages, consumed as part of a balanced diet, may improve the antioxidant status and enhance the overall health status. Herbal teas/beverages are rich sources of natural bioactive compounds such as carotenoids, phenolic acids, flavonoids, coumarins, alkaloids, polyacetylenes, saponins and terpenoids, among others. A wealth of available scientific evidence demonstrates that natural bioactive compounds render a number of diversified biological effects, such as antioxidant, antibacterial, antiviral, antiinflammatory, antiallergic, antithrombotic and vasodilatory actions, as well as antimutagenicity, anticarcinogenicity and antiaging effects. A number of herbal beverages are consumed globally and some beverages have gained more popularity than others depending on their geographical origin. However, in the era of globalization, ethnic barriers have gradually been removed and such commodities although from different areas, are now universally available as international health-pro products.

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1. Introduction

Excessive generation of reactive oxygen species (ROS) in the body causes oxidative stress, an injurious process leading to the oxidation of biomolecules such as proteins, lipids, carbohydrates and DNA. Oxidative stress is well known for its pivotal role in the etiology of several non-communicable diseases (NCDs) such as cardiovascular diseases, arthritis, type 2 diabetes, different types of cancer, autoimmune diseases and neurodegenerative disorders, among others.¹ The human body has endogenous antioxidant defense mechanisms those act simultaneously against ROS. These include enzymes (catalase, superoxide dismutase, glutathione reductase, glutathione peroxidase), low-molecular-weight antioxidants (uric acid, glutathione, albumin, protein-SH groups, bilirubin) and certain vitamins (ascorbic acid, α -tocopherol) as well as carotenoids.² However, external sources of antioxidants are needed

to prevent oxidative damage in the human body once internal antioxidant defense systems are challenged by over exposure to free radicals and other ROS.

There is a renewed interest in natural non-nutrient antioxidative compounds in reducing the incidence and severity of NCDs. Antioxidant compounds are widely distributed in plant materials, animal tissues and microorganisms. Fruits, vegetables, cereals, legumes, oilseeds, teas and certain spices are important sources of plant-derived antioxidants.³

Herbal beverages, commonly known as teas, have gained popularity among health conscious consumers. They have penetrated into an emerging niche market along with other popular beverages such as tea, coffee and cocoa which are also prepared using plant materials. In addition, a rapidly growing segment of the population uses herbal beverages for slimming, weight loss and a number of other cosmetic purposes.

In general, herbal beverages are prepared from natural ingredients of different morphological plant parts, namely leaves, stems, roots, fruits, buds and flowers. Herbal teas/beverages are rich sources of natural bioactive compounds such as carotenoids, phenolic acids, flavonoids, coumarins, alkaloids, polyacetylenes, saponins and terpenoids, among others. Scientific evidence shows

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that these bioactive compounds render a myriad of biological effects, such as antioxidant, antibacterial, antiviral, antiinflammatory, antiallergic, antithrombotic and vasodilatory action, as well as antimutagenicity, anticarcinogenicity and antiaging effects, among others.^{4–6} This contribution provides an overview of constituent antioxidants, and bioactivities of herbal beverages.

1.1. Antioxidants

Antioxidants are known for their ability to inhibit or delay the oxidation of other molecules in food and biological systems. They are protective against oxidative stress via different mechanisms and modes of action that are often independent of their antioxidant effect and may render their effects cooperatively via several mechanisms. These modes of action include free radical scavenging, singlet oxygen quenching, inactivation of peroxides and other ROS, metal ion chelation, quenching of secondary oxidation products, and inhibition of pro-oxidative enzymes, among others.⁷

Antioxidants are naturally present in many foods. Further, they can be synthesized, similar to their natural counterparts, such as synthetic vitamins C and E. Synthetic antioxidants generally contain of a phenolic ring and one or more hydroxyl substituents. Synthetic antioxidants that are still used by the food industry include butylated hydroxyanisole (BHA), butylated hydroxytoluene (BHT), propyl gallate (PG), and *tert*-butylhydroquinone (TBHQ). However, there are concerns associated with potential carcinogenic effect of some of these synthetic antioxidants at high concentrations in animal models which limits their use in food applications and a desire by the consumers to have green label products devoid of synthetic additives.⁷

1.2. Herbal beverages

The commonly used tea is a beverage made from leaves and buds or twigs of the plant *Camellia sinensis*, and is only second to water, the most consumed beverage in the World. There are basically four major types of true teas, namely black tea, oolong tea, green tea, and white tea. Nevertheless, the term herbal tea/beverage usually refers to infusions with fruit or other herbs that do not contain *Camellia sinensis*. Herbal beverages are also called tisane, herbal infusion, or botanical infusion to avoid confusion with true teas. Herbal beverages can be made with fresh or dried flowers, immature fruits, leaves, seeds, and/or roots by steeping (infusion) or boiling (decoction) of the source materials including herbs.

Herbal beverages when consumed within a balanced diet, may improve the antioxidant status, and reduce oxidative stress in humans.⁸ In addition, many commonly consumed herbal beverages do not contain any detectable caffeine levels as in coffee and tea. Health Canada categorizes herbal beverages under natural health products (NHPs). However, according to Health Canada moderate consumption (2–3 cups/day) of selected herbal teas such as citrus peel, lemon balm, ginger, orange peel and rosehip is recommended during pregnancy and breastfeeding.⁹

Herbal beverages have been used as natural part of the food culture in countries where traditional medicines are widely used. For instance, herbal teas prepared from *Aegle marmelos* (Bael), *Cassia auriculata* (*Ranawara*), *Aerva lanata* (*Polpala*), and *Hemidesmus indicus* (*Iramusu*) are common social beverages of food cultures in India and Sri Lanka. China is another country where combined herbal teas are often drunk on a daily basis to promote health and reduce the risk of certain health-related issues of different severity as simple as cold to diseases of liver and other organs.

Herbal teas consist of one or more herbal substances intended

for oral consumption and prepared by means of decoction, infusion or maceration. Generally the tea is prepared immediately before use. However, ready-to-serve bottled herbal beverages are becoming popular. Herbal teas are usually supplied in bulk form or in sachets. The herbal substance (s) used in tea formulations may be processed in advance by means of drying, comminuting and crushing. Therefore, commercially available products may be in different forms such as whole dried plant parts, dried powder, dried particles within tea bags, as well as granulates, and solutions which can be consumed directly. Table 1 presents selected examples of herbal beverages commonly used by populations around the world to boost optimum health as well as for reducing the risk of a number of disease conditions such as hyperglycemia, dyslipidaemia, cancer, and hypercholesterolemia.

2. Antioxidant compounds in herbal beverages

2.1. Phenolic compounds

Phenolic compounds are ubiquitous in plant organs. They are secondary metabolites consisting of an aromatic ring with different degrees of hydroxylation.³ Phenolics are derived from biosynthetic precursors such as pyruvate, acetate, aromatic amino acids such as phenylalanine and tyrosine, acetyl CoA and malonyl CoA following the pentose phosphate, shikimate, and phenylpropanoid metabolism pathways.¹⁰ Phenolic compounds occurring in herbal beverages include phenolic acids, coumarins, flavonoids, tannins, lignans and lignins.

2.2. Phenolic acids

Two classes of phenolic acids, hydroxybenzoic acids and hydroxycinnamic acids are found in plants.³ Hydroxybenzoic acids (C₆-C₁) include gallic, *p*-hydroxybenzoic, vanillic, syringic, and protocatechuic acids, among others. The hydroxycinnamic acids, better known as phenylpropanoids (C₆-C₃), include *p*-coumaric, caffeic, ferulic, and sinapic acids.³ Herbal beverages have been reported to include a number of phenolic acids.

2.3. Flavonoids

Flavonoids are synthesized by condensation of a phenylpropanoid compound with three molecules of malonyl coenzyme A. This reaction is catalyzed by the enzyme chalcone synthase that leads to the formation of chalcones. The chalcones are subsequently cyclized under acidic conditions to form flavonoids.³ There are different subclasses of flavonoids, namely flavones, flavonols, flavonones, flavononols, isoflavones, anthocyanidins and flavanols. Flavones and flavonols are present as aglycones in foods.³ They have similar C ring structures with a double bond at the 2–3 positions. Flavones lack a hydroxyl group at the third position.³ Flavonols (quercetin, kaempferol, and myricetin), flavones (luteolin, apigenin and chrysin), flavanols (catechin, epicatechin, epigallocatechin, epicatechin gallate, and epigallocatechin gallate), flavanones (narigenin, hesperitin, and eriodictyol), anthocyanidins (cyanidin, malvidin, peonidin, petunidin, pelargonidin and delphinidin), and isoflavones (genistein, daidzein, and glycitein) are the commonly found flavonoids in the human diet. Flavonoids, namely catechin, quercetin, kaempferol, rutin, apigenin, and isorhamnetin are constituents of the aqueous infusions of flower, leaf and seed of *Sideritis condensata*¹¹ (Table 2). In addition, flower infusion prepared within 10 min time contained 15 mg of isorhamnetin per gram of dry matter.

Table 1
Selected Herbal beverages commonly consumed in different parts of the world.

Local name	Scientific name	Family	Plant part	Health benefits
Beli, bael, bengal quince	<i>Aegle marmelos</i>	Rutaceae	Dried leaves, buds, flowers, immature fruits, bark	Anticancer, antidyslipidaemia, antihyperglycemic, antidiabetic, antiinflammatory, Antihyperglycemic
Tanner's Cassia, <i>Avartaki</i> , <i>Ranawara</i>	<i>Cassia auriculata</i>	Leguminosae	Dried leaves, flowers	Antihyperglycemic, antiinflammatory
<i>Polpala</i>	<i>Aerva lanata</i>	Amaranthaceae	Dried whole plant	Antioxidant, antithrombotic, antiplatelet aggregation
Indian Sarasaparilla, <i>Iramusu</i>	<i>Hemidesmus indicus</i>	Asclepiadaceae	Dried whole plant	Anticancer, increase the activity of antioxidant enzymes
Pegaga, Indian pennywort, <i>Gotukola</i>	<i>Centella asiatica</i>	Apiaceae	Dried whole plant	Antioxidant, hypocholesterolemic, anticancer, antiinflammatory
Chamomile	<i>Matricaria chamomilla</i> <i>Chamaemelum nobile</i>	Compositae	Dried flowers	Antidiabetic, antiinflammatory
Heart leaved mooseed	<i>Tinospora cordifolia</i>	Menispermaceae	Stem, roots	Antioxidant, antibacterial, hypocholesterolemic, anticancer, antiinflammatory
Coriandum	<i>Coriandrum sativum</i>	Apiaceae	Dried fruits	Antioxidant, antibacterial
Dag cayi	<i>Sideritis condensate</i>	Lamiaceae	Dried aerial parts	Antioxidant, antitumor
Peppermint tea	<i>Mentha piperita</i>	Lamiaceae	Dried leaves	hypocholesterolemic, hepatoprotective, cardiovascular system protective
Yerba mate	<i>Ilex paraguariensis</i>	Aquifoliaceae		Increase liver antioxidant status
Sage, adacayi, minchi	<i>Salvia officinalis</i>	Lamiaceae		Antiinflammatory
Rosehips	<i>Rosa canina</i>		Fruits	Antioxidant, anticancer
Rooibos	<i>Aspalathus linearis</i>			Antihepatocellular carcinoma, antioxidant
Borututu	<i>Cochlospermum angolensis</i>		Roots	
Ginger	<i>Zingiber officinale</i>		Rhizome	Antiinflammatory, hypoglycemic

2.4. Lignans

Lignans are compounds that comprise of two coupled phenylpropanoid units linked by the central carbons of their side chains.³ The common plant lignans found in the human diet include secoisolariciresinol, matairesinol, lariciresinol, pinoresinol and syringaresinol.¹² Secoisolariciresinol, and matairesinol are readily converted to mammalian lignans, enterodiol and enterolactone, respectively, by intestinal microflora in the human gut and are known to exert strong antioxidant and estrogenic activities.¹²

2.5. Lignins

Lignins are formed via polymerization of a mixture of the three monolignols, namely *p*-coumaryl, sinapyl and coniferyl alcohols.¹³ Additional compounds are incorporated into lignin in small quantities. They include coniferaldehyde, sinapaldehyde, dihydroconiferyl alcohol, 5-hydroxyconiferyl alcohol, tyramine ferulate and *p*-hydroxy-3-methoxybenzaldehyde, among others.¹⁴

Table 2
Phenolic acids and flavonoid contents of *Sideritis condensate* steeped at 100 °C for 10 or 30 min.

Phenolic compound	Plant part	µg/g of dry weight
Phenolic acids		
Protocatechuic	flower	200
<i>p</i> -Hydroxybenzoic	flower	1178
Vanillic	flower	1569
<i>p</i> - Coumaric	seed	249
Caffeic	leaf	601
Ferulic	flower	59
Flavonoids		
Catechin	leaf	209
rutin	leaf	879
Quercetin	flower	1902
kaempferol	leaf	1057 ^a
Isorhamnetin	flower	15284 ^a

Source: Data adapted from Kara et al.¹¹.

^a Steeping for 10 min.

2.6. Tannins

Tannins are composed of a group of compounds with a wide diversity in structure and have ability to bind and precipitate proteins.³ Tannins are classified into three groups, namely condensed tannins, hydrolysable tannins and complex tannins.¹⁵

2.7. Coumarins

Coumarins are lactones of *cis*-*O*-hydroxycinnamic acid derivatives and exist in the free form or as glycosides. In foods, simple coumarins, furanocoumarins (psoralens) and pyranocoumarins are found.³

2.8. Terpenes

Terpenes and terpenoid derivatives are secondary metabolites which originate from isoprene (2-methylbutadiene) units.¹⁶ The C₅H₈ isoprene units polymerise and subsequently produce different classes of terpenoids that include hemiterpenes consisting of a single C₅ isoprene unit, monoterpenes (C₁₀), sesquiterpenes (C₁₅), diterpenes (C₂₀), sesterterpenes (C₂₅), triterpenes (C₃₀), carotenoids (C₄₀) and polyterpenes consisting of long chains of many isoprene units. Triterpene group of compounds include sterols and triterpenes, which are accumulated as glycosides (saponins) in plants. Saponins are glycosylated (aglycone named sapogenin) secondary metabolites with surface-active properties. Some of these saponins are valuable starting materials for the synthesis of steroidal drugs.¹⁷

2.9. Carotenoids

Carotenoids are widespread pigments with yellow, orange, and red colours. They have pro-vitamin A and antioxidant activities. Carotenoids belong to hydrocarbons (carotenes) with a 40-carbon atom skeleton of 8 isoprene units. Their structures may be cyclized at one or both ends, and have different number of hydrogen atoms, or possess oxygen-containing functional groups,

the latter named as xanthophylls. Common carotenes in the human diet include β -carotene, α -carotene and lycopene, whereas xanthophylls include lutein, zeaxanthine, cryptoxanthin, canthaxanthin, astaxanthin and fucoxanthin. Major provitamin A active carotenoids are β -carotene, α -carotene and cryptoxanthin.

2.10. Polyacetylenes

Polyacetylenes are a group of bioactive compounds consisting of carbon-carbon triple bond or alkynyl functional group.¹⁸ Aliphatic C₁₇-polyacetylenes of the falcarinol type such as falcarinol and falcarindiol are widely distributed in the Apiaceae and Araliaceae families. Polyacetylenes of the falcarinol-type are formed from oleic acid by dehydrogenation that leads to the formation of C₁₈-acetylenes crepenynic acid and dehydrocrepenynic acid, which are then transformed to C₁₇-acetylenes upon β -oxidation.¹⁸ Health promoting properties and the content of polyacetylenes of a number of traditional medicinal herbs have been investigated during past few decades. Herbs such as American ginseng root (*Panax quinquefolium*), *Peucedanum praeruptorum*, *Echinacea pallida*, *Bupleurum spinosum* and *Actractylodes lancea* have been reported to contain polyacetylenes^{19–22}.

3. Antioxidant activities and bioactivities of herbal beverages

Herbal teas have been consumed as social drinks for centuries. In addition they are also used as alternative herbal medicines to treat a number of ailments. In the modern societies they are used for reducing the risk conditions of non-communicable diseases such as type 2 diabetes, hypertension, dyslipidemia and cancer.^{23–36} Table 3 summarizes selected information on medicinal claims for herbal concentrates of *A. marmelos*.

3.1. Asian herbal teas

Centella asiatica is an herbal tea commonly used by Asian populations. The dried whole herb as a single ingredient or mixed with other products such as garlic, coriander or ginger is used. This herb is known to increase the activity of antioxidant enzymes, namely superoxide dismutase, catalase, and glutathione peroxidase.³⁷ *C. asiatica* consists of a number of bioactive compounds such as alkaloids, terpenoids and saponins.³⁸ The principle triterpenoids identified in this herb include asiatic acid, madecassic acid, asiaticoside and madecassoside, among others.³⁹ In addition, *C. asiatica* is a rich source of bioactive polyacetylenes.¹⁸ Govindan et al.⁴⁰ identified a polyacetylene compound, cadiyenol, from the areal part of the plant. They further demonstrated that cadiyenol was capable of inducing cell apoptosis by 63% at 28 μ M concentration within 24 h in mouse lymphoma cells (P388D1). In addition, the

compound also reduced nitric oxide production by 70% in lipopolysaccharide (LPS) activated mouse macrophages at 24 μ M level.

In Japan, a number of herbal teas such as Arabian jasmine, Balsam pear, barley grass, guava, hardy rubber tree, Japanese persimmon, Jobs tears, and Wolof berry tea are consumed for health promotion.⁴¹ Total phenolic content of Arabian jasmine, balsam pear, barley grass, chameleon plant, guava, hardy rubber tree, Japanese persimmon, jobs tears, and wolf berry tea were 101, 15, 11, 83, 43, 41, 4 and 17 mg tannic acid equivalents (eq)/g of herb (dry weight), respectively. The antioxidant activity of these herbal teas ranged from 7 to 173 mmol/L as copper reducing power.

A. marmelos, commonly known as bael, is a plant native to Southeast Asia. It is a slow growing, subtropical tree and is grown in India, Sri Lanka, Pakistan, Bangladesh, Burma, Thailand, and other Southeast Asian countries. Bael fruit is a rich source of coumarins, vitamin C, and riboflavin. The leaves, stems, and bark as well as fruits are known in traditional medicine for dysentery and various other intestinal complaints (Plate 1). The leaves are also widely used to treat diarrhoea. In addition, bael is a potent radioprotective, analgesic, antihyperglycemic, antidyslipidemic, anticancer, and antidiabetic agent.^{42–47} The total phenolic content (TPC) and total flavonoid content (TFC) of bael fruit were reported to be 87 mg gallic acid eq/g of dry weight (dw) and 15 mg catechin eq/g dw, respectively.⁴⁸ In addition, the content of carotenoids and ascorbic acid of bael fruit were reported to be 3.3 and 26 mg/100 g dw, respectively. Furthermore, monoterpenes (limonene, pulegone) and sesquiterpenes (cubebene) were reported as its dominant volatile compounds.⁴⁸ Limonene is the major constituent producing the characteristic bael fruit flavor, among others. In addition, coumarins such as marmelosin, marmesin, and imperatorin, as well as alkaloids, namely aeglin, and aegelinine were also isolated from different parts of bael.⁴⁹ The extract of leaf, root, stem and the fruit demonstrated high antioxidant activities as determined by the 2,2-diphenyl-1-picrylhydrazyl (DPPH) scavenging activity, and ferric reducing antioxidant activity power (FRAP) assays. Furthermore, several bioactivities such as antidiabetic, antihyperlipidemic, antiviral, antifungal, antibacterial, anticancer, and anti-inflammatory were reported in association with different parts of bael tree.^{26,50} The antidiabetic effect of the fruit extract is probably due to the presence of coumarins, which potentiate insulin secretion from existing beta cells of the isles of langerhans.⁵¹ In Sri Lanka, dried buds and flowers and immature fruits are prepared as herbal tea infusions to replace the popular tea prepared from *Camelia sinensis*.

Generally bael is considered to be safe but only few studies have been conducted on its toxicity. Das et al.⁵² examined the toxic effects of aqueous extract of the leaves of bael and reported that 50 mg/100 g of body weight of male albino Wistar strain rats did not exhibit any toxicity in the liver and kidneys. Further, it was demonstrated that neither gross abnormalities nor

Table 3
Evidences on medicinal claims for *Aegle marmelos*.

Plant part	Outcome	Reference
Leaf extract	antidiabetic action in hyperglycaemic rats	Sachdewa et al. ²³ ; Ponnachan et al. ²⁴ ; Upadhyaya et al. ²⁵
	Reduced cholesterol level in diabetic patients	Gohil et al. ²⁶
	Radioprotective effects in mice	Jagetia et al. ²⁷
	Antihyperlipidaemic effect in rats with isoproterenol -induced myocardial infarction	Rajadurai and Prince ²⁸
Root extract	Anti-inflammatory activity in animals	Benni et al. ²⁹
	Diuretic activity in rats	Singh et al. ³⁰
Fruit extract	Antidyslipidemic effects in rats	Krushna et al. ³¹
	Reduces intraocular pressure, a cause for glaucoma in New Zealand white rabbits	Agarwal et al. ³²
	Hypoglycaemic activity in diabetic rats	Kamalakkannan and Prince ³³
	antidiabetic action in rats	Kamalakkannan and Prince ³⁴
	Chemopreventive effect in Swiss albino mice	Agarwal et al. ³⁵
Bark extract	Antifertility activity in male Wistar rats	Agarwal et al. ³⁶

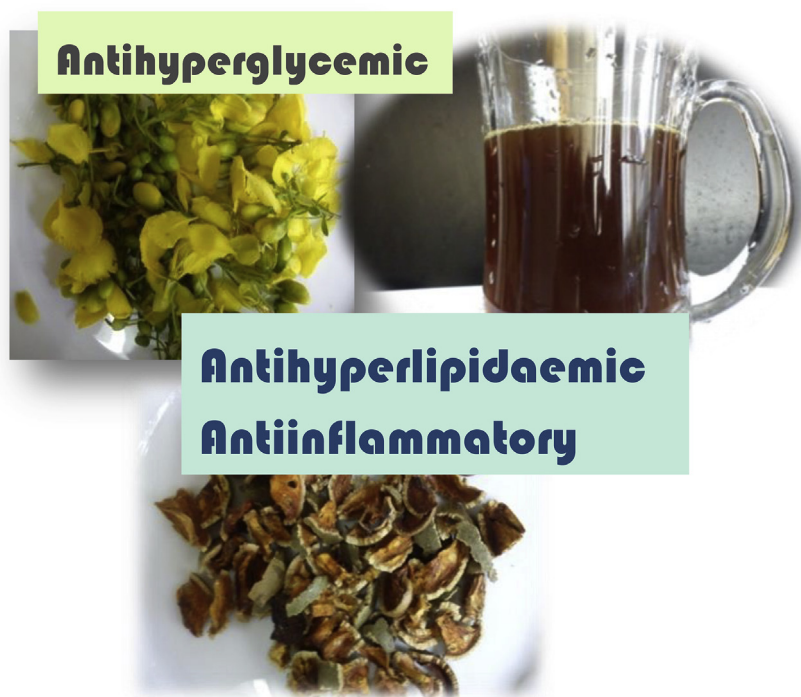


Plate 1. Fresh flower and buds of Tanner's Cassia and cuttings of dehydrated immature fruit and preparation of herbal tea of bael.

histopathological changes were observed in the heart, liver, kidneys, testis, spleen and brain after continuous administration of 50 mg/kg body weight of the extracts of *A. marmelos* intraperitoneally for 14 d.⁵³ In addition, the aqueous extract of bael fruit was not reported to be mutagenic to *Salmonella typhimurium* strain TA 100 in the Ames assay.⁵⁴

Dried flower buds, flowers and leaves of Tanner's Cassia (*Cassia auriculata*) have gained wide popularity as herbal teas in countries such as India and Sri Lanka. Furthermore, plant parts of Tanner's Cassia have been used as an alternative therapeutic agent particularly in controlling hyperglycemic conditions in traditional medical system. Methanolic extracts of Cassia flowers inhibited α -glucosidase activity *in vivo* and *in vitro*.⁵⁵ Further, the flower and leaf extracts of Cassia exhibited antihyperglycemic effects in streptozotocin-induced experimental diabetes.^{56,57} In addition, the aqueous extract of Cassia inhibited lipid peroxidation in the brain of diabetic rats.⁵⁸ Several phenolic compounds, namely (-) catechin, (-) epicatechin and procyanidin B₁ were identified in aqueous alcoholic extract of cassia seeds.⁵⁹ Puranik et al.⁵⁹ reported on cardiovascular safety and good tolerance of Cassia seed extracts without any adverse effects in male and female rats. However, Cassia seed extracts obtained using supercritical fluid extraction interfered with the absorption of metformin when co-administered with extracts in a rat model.⁵⁹

3.2. Herbal teas popular in Africa

African rooibos (*Aspalathus linearis*), borututu (*Cochlospermum angolensis*) and honey bush tisanes are popular South African herbal teas. Traditional medicinal uses of rooibos in South Africa include alleviation of infantile colic, allergies, asthma and dermatological problems. Furthermore, decoction of honey bush is used as a restorative and as an expectorant in chronic catarrh and pulmonary tuberculosis.⁶⁰ McKay and Bulmberegar⁶¹ reported that rooibos is rich in polyphenols and a rare source of the dietary

dihydrochalcones, namely aspalathin and nothofagin. In addition, major polyphenols in honey bush are xanthone mangiferin and the flavonones, hesperitin and isokuranetin.⁶² Both rooibos and honey bush teas have demonstrated potent antioxidant and antimutagenic activities *in vitro*. Rooibos tea also renders beneficial effects for heart health. It was reported that chrysoeriol, which is present at low levels in Rooibos is a potential agent in preventing and treating vascular diseases in humans.⁶³ Chrysoeriol is able to inhibit the migration of smooth muscle cells inside the aorta, a key cause of atherosclerosis. Furthermore, Rooibos tea has demonstrated ACE (angiotensin converting enzyme) inhibitory activities and reducing several of the pertinent biomarkers associated with cardiovascular disease.^{64,65}

3.3. Herbal teas popular in South America

Yerba mate (*Ilex paraguariensis*) is widely consumed by native people as social and medicinal beverage in South America for centuries and unlike other herbal beverages it contains caffeine. Several health promoting properties such as hepatoprotective, diuretic as well as central nervous system stimulating effects have been reported for yerba mate.⁶⁶ Furthermore, other studies have demonstrated its antioxidant, anti-inflammatory, antimutagenic and lipid lowering activities.^{67–70} The main bioactive compounds in yerba mate responsible for its activities were identified as caffeoyl derivatives such as caffeic acid, mono- and dicaffeoylquinic acids, methylxanthines (caffeine and theobromine) and flavonoids, namely rutin, quercetin and kaempferol.⁷¹ In a separate study, it was demonstrated that yerba mate and its bioactive compounds regulate the expression of genes related to adipogenesis.⁷² Yerba mate extract down-regulated the expression of genes responsible for adipogenesis, such as Creb-1 and C/EBP α . Furthermore, the extract up-regulated the expression of genes related to the inhibition of adipogenesis, including Dlk1, Gata2, Gata3, Klf2, Lrp5, Ppar γ 2, Sfrp1, Tcf7l2, Wnt10b, and Wnt3a.⁷²

Table 4
Oxygen radical absorbance capacity (ORAC) and cellular antioxidant activity (CAA) of herbal beverages.

Herbal beverage	ORAC ($\mu\text{mol trolox eq/g dw}$)	CAA ($\mu\text{mol quercetin eq/g dw}$)
Peppermint tea	1438	27.9
Sage	1351	35.3
Yerba mate	1195	46.5
Rosehip	330	2.9

Source: Data adapted from Bender et al.⁸³.

Kombucha tea is a health beverage prepared by fermentation of black tea brew and sugar with a symbiotic culture of acetic acid bacteria and yeasts reported to have potential health effects.⁷³ Aloulau et al.⁷³ demonstrated that compared to black tea, kombucha tea was an effective inhibitor of α -amylase and lipase activities in the plasma and pancreas of diabetic rats. Further, kombucha tea demonstrated higher suppressor activity of increased blood glucose levels than that of black tea. In addition, kombucha induced a marked delay in the absorption of LDL-cholesterol and triglycerides and a significant increase in HDL-cholesterol. According to histological analysis it exerted an ameliorative action on the pancreases and efficiently protected the liver-kidney functions of diabetic rats. This was further evidenced by significant decreases in aspartate transaminase, alanine transaminase, and gamma-glytamyl transpeptidase activities in the plasma, as well as in the creatinine and urea contents of diabetic rats.⁷³

3.4. Herbal teas popular in Europe

Chamomile is a member of Asteraceae or Compositae family and is represented by several varieties, namely *Chamomilla recutita*, *Matricaria chamomilla* and *Chamaemelum nobile*. Chamomile tea is widely consumed in Europe. Chamomile tea, is brewed from dried flower heads. The main constituents include phenolic compounds, primarily the flavonoids apigenin, quercetin, patuletin, and luteolin.⁷⁴ The principal compounds of the essential oil are the terpenoids alpha-bisabolol and azulenes, including chamazulene. Chamomile has moderate antioxidant and antimicrobial activities, and significant antiplatelet activity *in vitro*.⁷⁴ In addition, animal studies have shown potent anti-inflammatory, antimutagenic and cholesterol-lowering activities for chamomile. Furthermore, some antispasmodic and anxiolytic effects of chamomile tea have also been demonstrated.⁷⁴

Several studies have shown the antioxidant, hypocholesterolemic, anti-parasitic, anti-aging, and anticancer properties of chamomile.^{75–77} Chamomile has long been known as a treatment for inflammatory diseases. It was demonstrated the chamomile treatment inhibited the release of LPS-induced prostaglandin E₂ in RAW 264.7 macrophages *in vitro*.⁷⁷ This effect was found to be due to inhibition of COX-2 enzyme activity by chamomile extracts. Furthermore, chamomile caused reduction in LPS-induced COX-2 mRNA and protein expression, without affecting COX-1 expression.⁷⁷ A small percentage of people are sensitive to chamomile and develop allergic reactions.⁷⁸ Evidence of herb-drug interactions of chamomile is not well documented, and further studies are needed to reach any final conclusions.⁷⁸

'Dagcayi' is a popular herbal beverage in Turkey prepared from infusion of *S. condensate*. In a recent study, phenolic composition of aqueous infusions of *Dagcayi* was reported.¹¹ Dried aerial part of the plant is generally used to infuse the tea. Hydroxybenzoic as well as hydroxycinnamic acids were identified in flower, leaf and seed extracts of *S. condensate*. Phenolic composition of *S. condensate* changed based on the infusion temperature, time duration and part of the plant used for infusion. The phenolic acid identified in all plant parts examined was *p*-coumaric acid and its content ranged

from 10 to 398 $\mu\text{g/g}$ of dry matter of different plant parts, namely flower, leaf and seed. Chlorogenic acid was detected only at the steeping temperature of 60 °C.¹¹

Peppermint tea, brewed from *Mentha piperita* leaves, is a commonly consumed tisane that gives calming effect in the body and is popular in Europe and North Africa. Health benefits reported for peppermint tea include *in vitro* antibacterial activity against a range of pathogenic bacteria and antioxidant activity.^{79,80} The phenolic compounds reported in the leaves of *M. piperita* include rosmarinic acid and flavonoids.

Several preparations of rosehip fruits and seeds (*Rosa canina*) demonstrated antioxidant and antiinflammatory activities.⁸¹ *R. canina* L. fruits have a high content of ascorbic acid, and phenolics, including flavonoids, which render antioxidant activity as well as several other beneficial bioactivities.⁸²

According to Bender et al.,⁸³ *in vitro* oxygen radical absorbance capacity (ORAC) and *ex vivo* cellular antioxidant capacity (CAA) of yerba mate, peppermint tea, sage and rosehip fruit infusions in HaCat human keratinocytes cell line were found to vary considerably (Table 4). Sage (*Salvia officinalis*) is commonly used to prepare beverages as well as a flavoring agent in foods. Several health benefits such as antitumoral, antibacterial and antiinflammatory properties of sage have been reported.^{84,85}

Herbal beverages are habitually used as part of the normal diet in some populations in the world and trend of using them among others is progressively increasing. However, very limited published information is available on safety of herbs and herbal beverages, and herb-herb as well as herb-therapeutic drug interactions. It should be noteworthy that herbal beverages are prepared to maintain palatable characteristics of a social beverage thus they are less strong in their flavour and their bioactive compounds are present in smaller amounts due to the dilution as compared to those of herbal preparations intended to use as medicine. For instance, in Sri Lankan traditional and Ayurvedic medical system decoctions to be used as medicine are prepared by boiling dried or fresh herbs in 8 cups of water until volume is reduced to one cup.

According to a review by Singh et al.⁸⁶ the herb *C. asiatica* might cause liver problems, stomach upset, nausea, and drowsiness. Choi et al.⁸⁷ reviewed the herb drug interactions and specially focused on the effect of herbs on metabolic enzymes and transporters. Active compounds of some herbs may inhibit phase 1 and 2 metabolic enzymes thus affect the drug metabolism. In addition, concentrated form of some herbs, such as green tea can cause detrimental effects, namely liver damage, interaction with medications, interaction with metabolic enzymes and other natural ingredients.^{88,89}

4. Summary

Herbal beverages are potential rich sources of phytochemicals that may help in reducing disease risk conditions and therefore in the management of NCDs. A number of herbal beverages are consumed globally. Though bioactivities of some herbal beverages are known through preclinical studies, further analytical and clinical research is warranted in order to investigate the bioactive compounds that render such effects and their mode(s) of actions in

disease risk reduction and health promotion needs to be verified.

Conflict of interest

There are no conflict of interest to report.

References

- Iannitti T, Palmier B. Antioxidant therapy effectiveness: an up to Date. *Eur Rev Med Pharmacol Sci*. 2009;13:245–278.
- Halliwell B, Gutteridge JMC. The antioxidants of human extracellular fluids. *Arch Biochem Biophys*. 1990;280:1–8.
- Shahidi F, Nacz M. *Phenolics in Food and Nutraceuticals*. Boca Raton, FL: CRC press; 2004.
- Craig WJ. Health-promoting properties of common herbs. *Am J Clin Nutr*. 1999;70:491–499.
- Mckay DL, Blumberg JB. The role of tea in human health: an update. *J Am Coll Nutr*. 2002;21:1–13.
- Wargovich MJ, Woods C, Hollis DM, Zander ME. Herbs, Cancer prevention and health. *J Nutr*. 2001;131:3034–3036.
- Shahidi F, Zhong Y. Lipid oxidation and the improving the oxidative stability. *Chem Soc Rev*. 2010;39:4067–4079.
- Cabrera C, Artacho R, Giménez R. Beneficial effects of green tea-A review. *J Am Coll Nutr*. 2006;25:79–99.
- Public Health Agency of Canada. 2016, www.phac-aspc.gc.ca/hp-gs/known-savoir/cafeine-eng.php. (Accessed 1 May 2016).
- Randhir R, Lin Y, Shetty K. Phenolics, their antioxidant and antimicrobial activity in dark germinated fenugreek sprouts in response to peptide and phytochemical elicitors. *Asia Pac J Clin Nutr*. 2004;13:295–307.
- Kara M, Shain H, Turumtay H, Dinc S, Gumuscu A. The phenolic composition and antioxidant activity of tea with different parts of *Sideritis condensate* at different steeping conditions. *J Food Nut Res*. 2014;2:258–262.
- Liu RH. Whole grain phytochemicals and health. *J Cereal Sci*. 2007;46:207–219.
- Lewis N, Yamamoto E. Lignins: occurrence biosynthesis and biodegradation. *Annu Rev Plant Physiol*. 1990;41:455–496.
- Ralph J, Lapierre C, Marita JM, et al. Elucidation of new structures in lignins of CAD- and COMT- deficient plants by NMR. *Phytochemistry*. 2001;57:993–1003.
- Khanbabaee K, Van Ree T. Tannins: classification and definition. *Nat Prod*. 2001;18:641–649.
- Gershenzon J, Kreis W. Biosynthesis of monoterpenes, sesquiterpenes, diterpenes, sterols, cardiac glycosides and steroid saponins. In: Wink M, ed. *Biochemistry of Plant Secondary Metabolites*. Sheffield, UK: Sheffield Academic Press; 1999:222–299. Annual Plant Reviews.
- Liu J, Henkel T. Traditional Chinese medicine (TCM): are polyphenols and saponins the key ingredients triggering biological activities? *Curr Med Chem*. 2002;9:1483–1485.
- Minto RE, Blacklock BJ. Biosynthesis and function of polyacetylenes and allied natural products. *Prog Lipid Res*. 2008;47:233–306.
- Pellati F, Calo S, Benvenuti S, Adinolfi B, Nieri P, Melegari M. Isolation and structure elucidation of cytotoxic polyacetylenes and polyene from *Echinacea pallida*. *Phytochemistry*. 2006;67:1359–1364.
- Baranska M, Schulz H, Christensen LP. Structural changes of polyacetylenes in American ginseng root can be observed in situ by using Raman Spectroscopy. *J Agric Food Chem*. 2006;54:3629–3635.
- Christensen LP, Jensen M, Kidmose U. Simultaneous determination of Ginsenosides and polyacetylenes in American ginseng root (*Panax quinquefolium* L.) by high performance liquid chromatography. *J Agric Food Chem*. 2006;54:8995–9003.
- Chicca A, Adinolfi B, Martinotti E, et al. Cytotoxic effects of Echinacea root hexanic extracts on human cancer cell lines. *J Ethnopharmacol*. 2007;110:148–153.
- Sachdewa A, Raina D, Srivatsava A, Khemani LD. Effect of Aegle marmelos and Hibiscus rosa sinensis leaf extract on glucose tolerance in glucose induced hyperglycemic rats (Charles foster). *J Environ Biol*. 2001;22:53–57.
- Ponnachan PTC, Paulose CS, Panikkar KR. Hypoglycaemic effect of alkaloid preparation from leaves of *Aegle marmelos*. *Amala Res Bull*. 1993;13:37–41.
- Upadhyaya S, Shanbhag KK, Sunetha G, Balachandra Naidu M, Upadhyaya S. A study of hypoglycemic and antioxidant activity of *Aegle marmelos* in alloxan induced diabetic rats. *Indian J Physiol Pharmacol*. 2004;48:476–480.
- Gohil T, Pathak N, Jivani N, Devmurari V, Patel J. Treatment with extracts of *Eugenia jambolana* seed and *Aegle marmelos* leaf extracts prevents hyperglycemia and hyperlipidemia in alloxan induced diabetic rats. *Afr J Pharm Pharmacol*. 2010;4:270–275.
- Jagetia G, Venkatesh P, Baliga M. Evaluation of the radioprotective effect of bael leaf (*Aegle marmelos*) extract in mice. *Int J Radiat Biol*. 2004;80:281–290.
- Rajadurai M, Prince PS. Comparative effects of *Aegle marmelos* extract and alpha tocopherol on serum lipids, lipid peroxides and cardiac enzyme levels in rats with isoproterenol-induced myocardial infarction. *Singap Med J*. 2005;46:78–81.
- Benni JM, Jayanthi M, Suresha R. Evaluation of the antiinflammatory activity of *Aegle marmelos* (Bilwa) root. *Indian J Pharmacol*. 2011;43:393–397.
- Singh S, Singh SK, Srivastava S, et al. Experimental evaluation of diuretic activity of *Aegle marmelos* in rats. *Int J Pharm Biol Sci*. 2013;3:98–102.
- Krushna GSS, Kareem MA, Reddy VD, Padmawathi P, Hussain SA, Kodidhela LD. *Aegle marmelos* fruit extract attenuates isoproterenol- induced myocardial oxidative stress in rats. *J Clin Biochem Nutr*. 2009;6:199–204.
- Agarwal R, Gupta SK, Srivastava S, Saxena R, Agrawal SS. Intraocular pressure-lowering activity of topical application of *Aegle marmelos* fruit extract in experimental animal models. *Ophthalmic Res*. 2009;42:112–116.
- Kamalakkannan N, Prince PS. Hypoglycaemic effect of water extracts of *Aegle marmelos* fruits in streptozotocin diabetic rats. *J Ethnopharmacol*. 2003;87:207–210.
- Kamalakkannan N, Prince PS. The effect of *Aegle marmelos* fruit extract in streptozotocin diabetes: a histopathological study. *J Herb Pharmacother*. 2005;5:87–96.
- Agrawal A, Verma P, Goyal P. Chemomodulatory effects of *Aegle marmelos* against DMBA-induced skin tumorigenesis in Swiss albino mice. *Asian Pac J Cancer Prev*. 2010;11:1311–1314.
- Agrawal SS, Kumar A, Gullaiya S, et al. Antifertility activity of methanolic bark extract of *Aegle marmelos* (L.) in male wistar rats. *Daru*. 2012;20:94.
- Chin HW, Lin CC, Tang KS. The hepatoprotective effects of Taiwan folk medicine ham-hong-chho in rats. *Am J Chin Med*. 1996;24:231–240.
- Singh D, Singh P, Gupta A, Solanki S, Sharma E, Nema R. Qualitative estimation of the presence of bioactive compound in *Centella Asiatica*: an important medicinal plant. *Inter J Life Sci Med Sci*. 2012;2:5–7.
- Bonfill M, Mangas S, Cusido RM, Osuna L, Pinol MT, Palazon J. Identification of triterpenoid compounds of *Centella asiatica* by thin-layer chromatography and mass spectrometry. *Biomed Chromatogr*. 2005;20:151–153.
- Govindan G, Sambandan TG, Govindan M, Rha CK. A bioactive polyacetylene compound isolated from *Centella asiatica*. *Planta Med*. 2007;73:597–599.
- Toda S. Polyphenol content and antioxidant effects in herb teas. *Chin Med*. 2011;2:29–31.
- Jagetia G, Venkatesh P. Inhibition of radiation-induced clastogenicity by *Aegle marmelos* (L.) correa in mice bone marrow exposed to different doses of gamma-radiation. *Hum Exp Toxicol*. 2007;26:111–124.
- Shankarananth V, Balakrishnan N, Suresh D, Sureshpandian G, Edwin E, Sheeja E. Analgesic activity of methanol extract of *Aegle marmelos* leaves. *Fitoterapia*. 2007;78:258–259.
- Narendar T, Shweta S, Tiwari P, et al. Antihyperglycemic and antidyslipidemic agent from *Aegle marmelos*. *Bioorg Med Chem Lett*. 2007;17:1808–1811.
- Costa-Lotufo LV, Khan MT, Ather A, et al. Studies of the anticancer potential of plants used in Bangladeshi folk medicine. *J Ethnopharmacol*. 2005;99:21–30.
- Subramaniam D, Giridharan P, Murmu N, et al. Activation of apoptosis by 1-hydroxy-5, 7-dimethoxy-2-naphthalene-carboxaldehyde, a novel compound from *Aegle marmelos*. *Cancer Res*. 2008;68:8573–8585.
- Sabu MC, Kuttan R. Antidiabetic activity of *Aegle marmelos* and its relationship with its antioxidant properties. *Indian J Physiol Pharmacol*. 2004;2004(48):81–88.
- Narendhirakannan RT, Subramanian S, Kandaswamy M. Biochemical evaluation of anti-diabetogenic properties of some commonly used Indian plants on streptozotocin-induced diabetes in experimental rats. *Clin Exp Pharmacol Physiol*. 2006;33:1150–1157.
- Suvmol C, Praneer A. Bioactive compounds and volatile compounds of Thai bael fruit (*Aegle marmelos* (L.) Correa) as a valuable source for functional food ingredients. *Int Food Res*. 2008;15:45–63.
- Maity P, Hansda D, Bandyopadhyay U, Mishra DK. Biological activities of crude extracts and chemical constituents of bael, *Aegle marmelos* (L.) Corr. *Indian J Exp Biol*. 2009;47:849–861.
- Dhankhar S, Ruhil S, Balhara M, Dhankhar S, Chhillar AK. *Aegle marmelos* (Linn.) correa: a potential source of Phytomedicine. *J Med Plants Res*. 2011;5:1497–1507.
- Das UK, Maiti R, Jana D, Ghosh D. Effect of aqueous extract of leaf of *Aegle marmelos* on testicular activities in rats. *IJPT*. 2006;5:21–25.
- Veerappan A, Miyazaki S, Kadarkaraisamy M, Ranganathan D. Acute and sub-acute toxicity studies of *Aegle marmelos* Corr., an Indian medicinal plant. *Phytomedicine*. 2007;14:209–215.
- Kruawan K, Kangsadalampai K. Antioxidant activity, phenolic compound contents and antimutagenic activity of some water extract of herbs. *Thai J Pharm Sci*. 2006;30:28–35.
- Abesundara KJM, Matsui T, Matsumoto K. α -Glucosidase inhibitory activity of some Sri Lanka plant extracts, one of which, *Cassia auriculata*, exerts a strong anti-hyperglycemic effect in rats comparable to the therapeutic drug Acarbose. *J Agric Food Chem*. 2004;52:2541–2545.
- Latha M, Pari L. Antihyperglycemic effect of *Cassia auriculata* in experimental diabetes and its effects on key metabolic enzymes involved in carbohydrate metabolism. *Clin Exp Pharmacol Physiol*. 2003;30:38–43.
- Gupta S, Sharma SB, Bansal SK, Prabhu KM. Antihyperglycemic and hypolipidemic activity of aqueous extract of *Cassia auriculata* L. leaves in experimental diabetes. *J Ethnopharmacol*. 2009;123:499–503.
- Latha M, Pari L. Preventive effects of *Cassia auriculata* L. flowers on brain lipid peroxidation in rats treated with streptozocin. *Mol Cell Biochem*. 2003;243:23–28.
- Puranik AS, Halade G, Kumar S, et al. *Cassia auriculata*: Aspects of safety pharmacology and drug interaction. *J Evid Based Complement Altern Med*. 2011. <https://doi.org/10.1093/ecam/nep237>.
- Joubert E, Gelderblom WCA, Louw A, de Beer D. South African herbal teas: *Aspalathus linearis*, *yclopia* spp. and *Athrixia phylicoides*-A review.

- J Ethnopharmacol.* 2008;119:376–412.
61. McKay DL, Blumberg JB. The role of tea in human health: an update. *J Am Coll Nutr.* 2002;21:1–13.
 62. McKay DL, Blumberg JB. A review of the bioactivity of South African herbal teas: rooibos (*Aspalathus linearis*) and honeybush (*Cyclopia intermedia*). *Phytother Res.* 2007;21:1–16.
 63. Cha B, Shi WL, Yonezawa T, Teruya T, Nagai K, Woo J. An inhibitory effect of chrysoeriol on platelet-derived growth factor (PDGF)-induced proliferation and PDGF receptor signaling in human aortic smooth muscle cells. *J Pharmacol Sci.* 2009;110:105–110.
 64. Marnewick JL. Rooibos and honeybush: recent advances in chemistry, biological activity and pharmacognosy. In: Juliani HR, Simon JE, Ho CT, eds. *African Natural Plant Products: New Discoveries and Challenges in Chemistry and Quality*. Washington DC, USA: American Chemical Society; 2010:277–294. ACS Symposium Series Volume 1021.
 65. Persson IA, Persson K, Hägg S, Andersson RGG. Effects of green tea, black tea and Rooibos tea on angiotensin-converting enzyme and nitric oxide in healthy volunteers. *Public Health Nutr.* 2010;3:730–737.
 66. Heck CI, de Mejia EG. Yerba Mate Tea (*Ilex paraguariensis*): a comprehensive review on chemistry, health implications, and technological considerations. *J Food Sci.* 2007;72:R138–R151.
 67. Lobato R, Matsumoto T, Mendonça S, Moura de Oliveira D, Souza MF, Markowicz Bastos DH. Effects of maté tea intake on ex vivo LDL peroxidation induced by three different pathways. *Nutrients.* 2009;1:18–29.
 68. Vanderjagt TJ, Ghattas R, Vanderjagt DJ, Crossey M, Glew RH. Comparison of the total antioxidant content of 30 widely used medicinal plants of New Mexico. *Life Sci.* 2002;70:1035–1040.
 69. Bracesco N, Sanchez AG, Contreras V, Menini T, Gugliucci A. Recent advances on *Ilex paraguariensis* research: minireview. *J Ethnopharmacol.* 2011;136:378–384.
 70. Gao H, Long Y, Jiang X, et al. Beneficial effects of Yerba Mate tea (*Ilex paraguariensis*) on hyperlipidemia in high-fat-fed hamsters. *Exp Gerontol.* 2013;48:572–578.
 71. Isobella S, Cogoi L, Lopez P, Anesini C, Ferraro G, Filip R. Study of the bioactive compounds variation during yerba mate (*Ilex paraguariensis*) processing. *Food Chem.* 2010;122:695–699.
 72. Arcari DP, Santos JC, Gambero A, Ribeiro ML. The in vitro and in vivo effects of yerba mate (*Ilex paraguariensis*) extract on adipogenesis. *Food Chem.* 2013;141:809–815.
 73. Aloulou A, Hamden K, Elloumi D, et al. Hypoglycemic and antilipidemic properties of kombucha tea in alloxan-induced diabetic rats. *BMC Complement Altern Med.* 2012;12:63.
 74. McKay DL, Blumberg JB. A review of the bioactivity and potential health benefits of chamomile tea (*Matricaria recutita* L.). *Phytother Res.* 2006;20:519–530.
 75. Babenko NA, Shakhova EG. Effects of *Chamomilla recutita* flavonoids on age-related liver sphingolipid turnover in rats. *Exp Gerontol.* 2006;41:32–39.
 76. Lee KG, Shibamoto T. Determination of antioxidant potential of volatile extracts isolated from various herbs and spices. *J Agric Food Chem.* 2002;50:4947–4952.
 77. Srivastava JK, Gupta S. Health promoting benefits of chamomile in the elderly population. In: Watson R, ed. *Complementary and Alternative Therapies in the Aging Population*. Elsevier Inc., Academic Press; 2009:135–158.
 78. Budzinski JW, Foster BC, Vandenhoeck S, Arnason JT. An in vitro evaluation of human cytochrome P450 3A4 inhibition by selected commercial herbal extracts and tinctures. *Phytomedicine.* 2000;7:273–282.
 79. Bupesh G, Amutha C, Nandagopal S, Ganeshkumar A, Sureshkumar P, Murali KS. Antibacterial activity of *Mentha piperita* L. (peppermint) from leaf extracts – a medicinal plant. *Acta Agric Slov.* 2007;89:73–79.
 80. Sharafi SM, Rasooli I, Owlia P, Taghizadeh M, Darvish S, Astaneh A. Protective effects of bioactive phytochemicals from *Mentha piperita* with multiple health potentials. *Pharmacogn Mag.* 2010;6:147–153.
 81. Chrubasik C, Roufogalis BD, Müller-Ladner U, Chrubasik S. A systematic review on the *Rosa canina* effect and efficacy profiles. *Phytother Res.* 2008;22:725–733.
 82. Roman I, Stanila A, Stanila S. Bioactive compounds and antioxidant activity of *Rosa canina* L. biotypes from spontaneous flora of Transylvania. *Chem Cent J.* 2013;2013(7):73.
 83. Bender C, Graziano S, Zimmerman BF, Weidlich HH. Antioxidant potential of aqueous plant extracts assessed by the cellular antioxidant activity assay. *Am J Biol Life Sci.* 2014;2:72–79.
 84. González A, Abad T, Jiménez I, Ravelo A. A first study of antibacterial activity of diterpenes isolated from some *Salvia* species. *Biochem Syst Ecol.* 1987;17:293–296.
 85. Vladimir-Knežević S, Blažeković B, Kindl M, Vladić J, Lower-Nedza AD, Brantner AH. Acetylcholinesterase inhibitory, antioxidant and phytochemical properties of selected medicinal plants of the *Lamiaceae* family. *Molecules.* 2014;19:767–782.
 86. Singh D, Gupta R, Saraf SA. Herbs-are they safe enough? An overview. *Crit Rev Food Sci Nutr.* 2012;52:876–898.
 87. Choi YH, Chin YW, Kim YG. Herb-drug interactions: focus on metabolic enzymes and transporters. *Arch Pharm Res.* 2011;34:1843–1863.
 88. Schönthal AH. Adverse effects of concentrated green tea extracts. *Mol Nutr Food Res.* 2011;55:874–885.
 89. Colalto C. Herbal interactions on absorption of drugs: mechanisms of action and clinical risk assessment. *Pharmacol Res.* 2010;62:207–227.