



E-ISSN: 2320-7078

P-ISSN: 2349-6800

JEZS 2020; 8(1): 1424-1430

© 2020 JEZS

Received: 20-11-2019

Accepted: 24-12-2019

Asik Ikbal

Department of Fish Processing
Technology, Faculty of Fishery
Sciences, West Bengal University
of Animal and Fishery Sciences
Chakgaria, Kolkata,
West Bengal, India

Satarupa Roy

New Zealand Fresh & Natural, 6,
Waterloo Street, Kolkata,
West Bengal, India

Keya Pati

Department of Fish Processing
Technology, Faculty of Fishery
Sciences, West Bengal University
of Animal and Fishery Sciences
Chakgaria, Kolkata, West
Bengal, India

Corresponding Author:**Satarupa Roy**

New Zealand Fresh & Natural, 6,
Waterloo Street, Kolkata, West
Bengal, India

Health benefits of green tea: A mini review

Asik Ikbal, Satarupa Roy and Keya Pati

Abstract

For gaining popularity and greater demand for medicinal plants, various medicinal plants are taken into considerations for therapeutic research. Green tea is the processed natural product consumed as a healthy drink for refreshment worldwide. Its leaf is consumed which has medicinal values such as antioxidant activity, antimicrobial activity and it has several phytochemical constituents. It contains 15-20% protein and 1-4% amino acids, 5-7% carbohydrates and 10-25% major antioxidant catching tannins. Thousands of bioactive components are present in green tea as polyphenols which is the key factor for preventing many diseases. The objective of this manuscript is to review of therapeutic properties of green tea.

Keywords: Green tea, phytochemical constituents, antioxidant properties, antimicrobial properties

Introduction

Green Tea is non-toxic, non-fermented and widespread therapeutic beverages consumed around the world produced from the leaf of the plant called "*Camellia sinensis*" which is found mainly in China and in Southeast Asia (Molan, 2013) ^[1]. It tends to be devoured as a drink, which has many beneficial wellbeing effects or concentrate from its leaves can be made for medicinal use (Namita *et al.*, 2012; Narotzki *et al.*, 2012) ^[2, 3]. Green tea has been consumed over the centuries in India, China, Japan, and Thailand as tea which is a good source of various phytochemicals and the most common drink in the world after water (Fujiki *et al.*, 2002) ^[4]. Recently, green tea has been acquired great attention in pharmacological and food industries due to some beneficial effects including antimicrobial, antioxidant and prebiotic activities and promotes human health (Zou *et al.*, 2014; Su *et al.*, 2008) ^[5, 6].

Tea flavanols have recently received much attention owing to their various biological exercises (Chen *et al.*, 2001) ^[7]. Their beneficial properties are thought to be included antioxidants (Navas *et al.*, 2005) ^[8], antimutagenic (Halder *et al.*, 2005) ^[9], anticarcinogenic (Zhu *et al.*, 2005) ^[10] and antibacterial (An *et al.*, 2004) ^[11] effects (Erol *et al.*, 2009) ^[12]. Antioxidant has a significant job in lessening free radical-intervened debasement of cells and tissues in the organism (Jin *et al.*, 2004; Wongkham *et al.*, 2001; Almajano *et al.*, 2008) ^[13-15].

Green tea, which has mitigating, hostile to tumor, antioxidative and antimicrobial properties, is considered as a healthy product with its utilization connected to bring down rates of all these different obsessive conditions (Bansal *et al.*, 2013) ^[16]. Along with the thousands of bioactive compounds, the most significant bioactive constituents of green tea are polyphenols or flavonoids which play a key role in the prevention and treatment of many diseases. Green tea is framed when the freshly picked leaves are being steamed, rolled, dried and burned, therefore the chemical composition of it, including polyphenols, differs very little from the fresh tea leaves (Wheeler and Wheeler 2004) ^[17].

Tea polyphenols, especially the catechins are potent antimicrobial and antioxidant agents with positive effects on human health as well as fish health too (Boran *et al.*, 2015; Barbosa *et al.*, 2007) ^[18, 19]. Green tea contains between 30 and 40 percent of water-extractable polyphenols, while black tea has 3 to 10 percent which varies according to climate, season or variety (Wanasundara and Shahidi, 1998; Archana *et al.*, 2011) ^[20, 21]. The phytochemical screening of tea revealed the presence of alkaloids, saponins, tannins, catechin, and polyphenols (Mbatia *et al.*, 2008) ^[22]. Studies have shown that the antioxidant property of green tea extract is present because of high quantity of catechins that are chemically defined as flavan-3-ols (Balentine *et al.*, 1997) ^[23] and four polyphenol compounds, Epigallocatechin gallate (EGCG), Epicatechin gallate (ECG), Epigallocatechin (EGC) and Epicatechin (EC) (Frei and Higdon, 2003; Zandi and Gondon, 1999) ^[24, 25]. Among them (-)-Epigallocatechin-3-gallate (EGCG) is the main and biologically most potent and the most luxuriant component. (Yang and Koo, 1997) ^[26].

The tea catechins are free radical scavengers, metal chelators, inhibitors of transcription factors, and enzymes. Therefore, green tea extracts have been used as natural antioxidants, antibacterial and antiviral agents (Manzocco *et al.*, 1998; Tang and Meydani, 2001) [27, 28]. Also, it has been reported that green tea extract has anticarcinogenic and antimutagenic activity (Yang *et al.*, 2000) [29]. Green tea extracts also contain caffeine, vitamins, amino acids, theanine, volatiles, and minerals (da Silva Pinto, 2013; Davidson *et al.*, 2005) [30, 31]. So, Green tea is beneficial to serve as a natural food antioxidant.

During the last decade, the effects of tea and tea polyphenols were extensively investigated and studies showed that tea is capable of lowering the risk of cardiovascular diseases and cancers (Huo *et al.*, 2008; Mukamal *et al.*, 2007) [32, 33], reducing body fat, systolic blood pressure (SBP), and low-density lipoprotein (LDL) cholesterol (Nagao *et al.*, 2007) [34]. Among age-associated pathologies and neurodegenerative diseases, green tea was shown to confer significant protection against Parkinson's disease and Alzheimer's disease (Chang and Jiang, 2007; Rezai-Zadeh *et al.*, 2005) [35, 36]. On the other way, continuous consumption of tea by mothers during pregnancy might be associated with an increased risk of preeclampsia, especially severe preeclampsia (Wei *et al.*, 2009; Dutta *et al.*, 2013) [37, 38].

Phytochemical constituents

There are mainly three types of tea variants available and consumed worldwide. Among them 78% is black, 20% is green, and 2% is oolong tea originates from the same Phyto-species (Chan *et al.*, 2011; Koech *et al.*, 2017) [39, 40]. Green tea has been widely studied, analyzed including all the varieties as it is the non-fermented retaining green color and almost all phytochemical contents (Manning and Roberts, 2003) [41].

The chemical composition of green tea is made up of 15-20% protein and 1-4% amino acids on a dry weight basis. Amino acids like glutamic acid, tryptophan, glycine, serine, aspartic acid, tyrosine, valine, leucine, threonine, arginine, and lysine and carbohydrates (5-7% in dry weight basis) such as cellulose, pectins, glucose, fructose, sucrose are partly included in green tea (Senanayake, 2013) [42]. Other green tea-compounds with interest in human health such as fluorine, caffeine, minerals, trace elements such as chromium and manganese. (Kristanti and Punbusayakul, 2008) [43]. It contains trace elements in lipid form (linoleic acid, alpha-linolenic acid), sterols (stigmasterol), vitamins (B, C, E), xanthine bases (caffeine, theophylline), pigments (chlorophyll, carotenoids), volatile compounds (aldehyde, alcohol, esters, lactones, hydrocarbons).

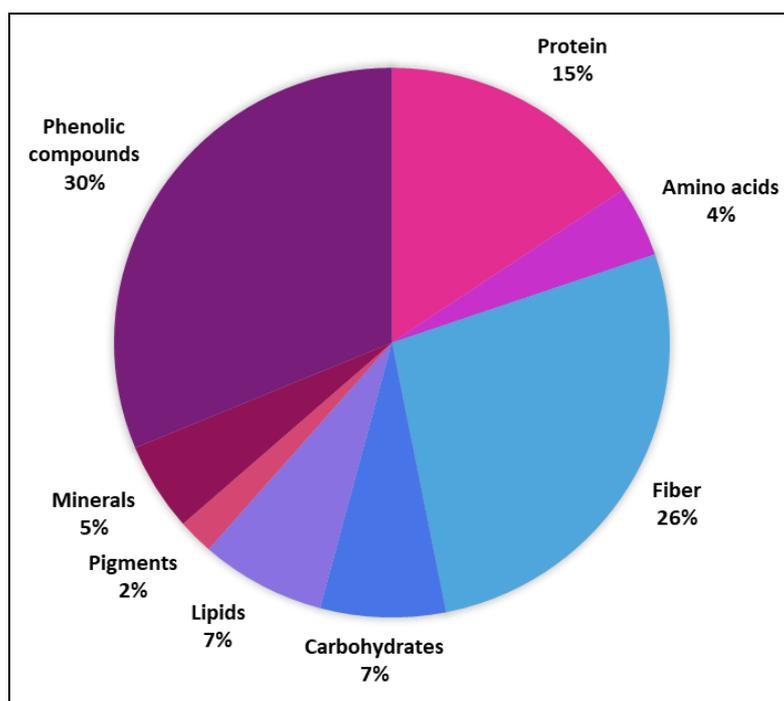


Fig 1: Composition (%) of green tea (Chacko *et al.*, 2010) [44]

Green tea contains polyphenols which include flavonols, flavonoid, flavonoids, and phenolic acids (Jo *et al.*, 2012) [45]. The principal catechins present in green tea such as (-)-epicatechin (EC), (-)-epigallocatechin (EGC), (-)-epicatechin-3-gallate (ECG), (-)-epigallocatechin-3-gallate (EGCG) and (-)-gallocatechin gallate (GCG) are indicated as major part of biologically active substances (Ho *et al.*, 1994; Sharma *et al.*, 2014) [46, 47]. A large number of catechin tannins (10-25%) are available in green tea. EGCG is the most common polyphenol found in green tea and represents up to 10% of its dry weight and includes 60-70% of its total catechins. Many of the green tea has health promotion abilities that are attributed to EGCG (Patil *et al.*, 2016) [48]. EGCG and other catechins show strong antioxidant activities due to their reduction potential of single

electrons. Free radicals are harmful and reactive molecules that become unstable from this unpaired electron. They are involved in diseases from blood clots to cancer (Bancirova, 2010) [49].

Table 1: Phenolic Components of Green Tea (% Dry Weight) (Rice-Evans, 1999) [50]

Catechins	30-42% (i-v)	i. Epigallocatechin gallate (11.2%)
		ii. Epigallocatechin (10.3%)
		iii. Epicatechin gallate (2.3%)
		iv. Epicatechin (2.5%)
		v. Catechin (0.5%)
Flavonols		2%
Simple polyphenols		2%

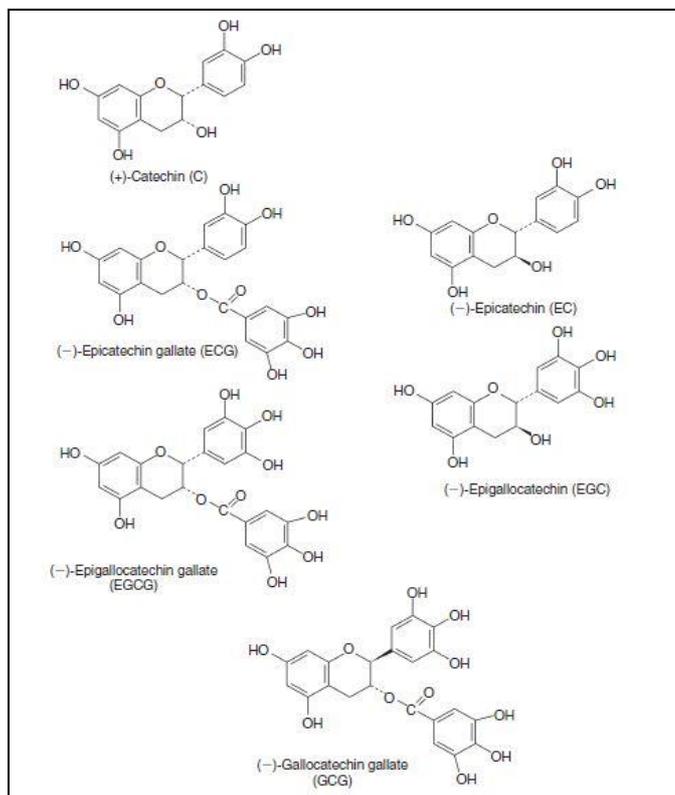


Fig 2: Structures of catechin constituents of green tea (*Camellia sinensis*). (Manning and Roberts, 2003) [41].

Antioxidant properties

An antioxidant is a substance that inhibits lipid oxidation by quenching the formation of free radicals or by producing more free radicals that can spread the oxidation reaction. (Senanayake, 2013) [42]. The antioxidant activity of green tea polyphenols is predominantly due to the combination of aromatic rings and hydroxyl groups, which synthesize their chemical structure and consequently bind and neutralize lipid-free radicals by these hydroxyl groups. Numerous researches have been exhibited in different aspects on antioxidant activity of green tea polyphenols which are exceptional electron donors and are effective scavengers of physiologically reactive oxygen species (ROS) *in vitro*, including superoxide anions (Guo *et al.*, 1999; Yokozawa *et al.*, 2002; Nanjo *et al.*, 1996; Pon and Liu, 2008) [51-54],

peroxyl radicals, and singlet oxygen. Consequently, green tea is the most effective against beta-carotene oxidation as far as the antioxidant property is concerned and can serve as a natural source of free radical scavengers and cancer prevention agent (Senanayake, 2013; Jo *et al.*, 2012) [42, 45]. The most potent antioxidant polyphenol of green tea is EGC (Zuo *et al.*, 2006; Koech *et al.*, 2017) [10, 40]. Significant levels of bioactive catechins are available in green tea that has the ability to stabilize the free radicals by providing hydrogen ions. The high antioxidative effect of polyphenols in green tea is because of the closeness of phenolic hydroxyl groups in their structures that make them potent free radical scavengers (Tariq and Reyaz, 2013) [55]. This hydroxylation built a higher volume of stability on the catechin phenoxyl radicals by participating in electron delocalization that is a significant feature of the anti-radical potential. That is why radical scavenging is high in the gallocatechin including EGCG and EGC (Koech *et al.*, 2017) [40]. Green tea catechins are performed as water-soluble antioxidants as Trolox and ascorbic acid, have been shown to act as active antioxidants in bulk oils and as prooxidants in oil-in-water emulsions (Senanayake, 2013; Frankel *et al.*, 1997) [42, 56].

Green tea extract is a promising wellspring of natural antioxidants that has been effectively utilized not exclusively to enhance flavour in addition to broaden the shelf-life of various food products (Senanayake, 2013) [42]. Green tea otherwise acts as a preventative agent of some common human health disorders (Hossain and Mahmood, 2014) [57]. Green tea contains considerable amounts of catechin tannin which is a powerful, water-soluble polyphenol and antioxidant that is easily oxidized. From a single cup of green tea, one can be benefitted from 10-40 mg of polyphenols (antioxidants) (<http://www.greenteanutritionalfacts.com>, 25 September, 2019) [58]. According to Cao *et al.* (1996) [59], green and black tea had much higher antioxidant activity against free radicals than all the commercially available vegetables (Senanayake, 2013) [42].

Many studies have been reported on green tea that has much higher antioxidant activities against free radicals which are not found in vegetables. It is measured mainly by the Oxygen Radical Absorbance Capacity (ORAC) assay, DPPH (2, 2-diphenyl-1-picrylhydrazyl) radical scavenging assays and Trolox equivalent antioxidant capacity (TEAC) assay [Rice-Evans, 1999; Cao *et al.*, 1996; Tsai *et al.*, 2008) [50-60].

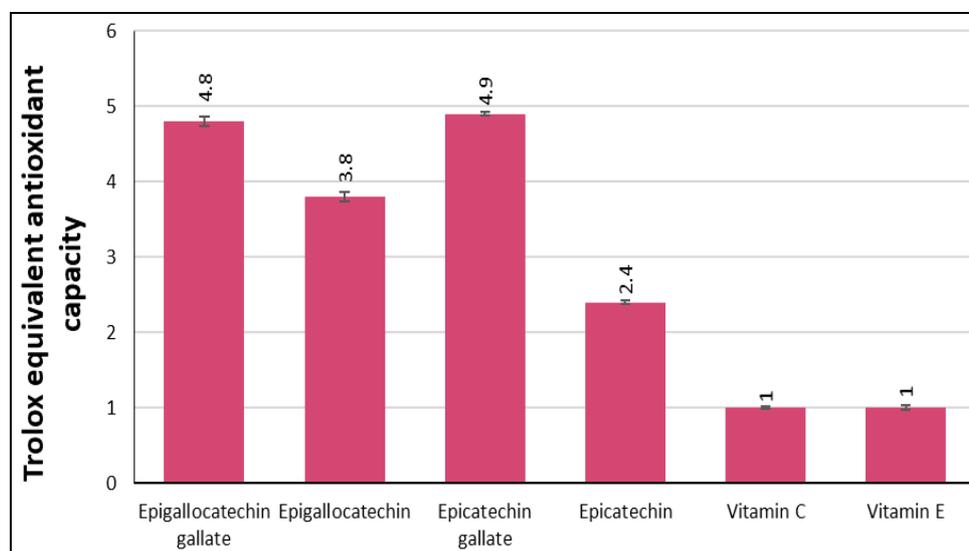


Fig 3: Reactive antioxidant properties of green tea (Rice-Evans, 1999) [50].

*TEAC is the millimolar concentration of Trolox (reference standard) having the equivalent antioxidant activity to a 1 mM concentration of the antioxidant compound or defined concentration or volume of food extract/beverage under investigation.

Antimicrobial properties

Resistance to antimicrobial agents has been becoming harmful to the environment and accelerating the global problem (Cushnie and Lamb, 2005) [61]. Nowadays many antimicrobial drugs cannot destroy their pathogenic microorganisms as they becoming resistant. Therefore, researchers are finding some new drugs from mangrove species to control the pathogens due to the presence of antimicrobial compounds (Clercq, 2001; Poole, 2001) [62, 63].

The accompanying review will look at the antimicrobial action of green tea polyphenols, characteristic natural compounds, possessing an assorted scope of pharmacological properties. The antimicrobial activity of tea was first established almost 100 years ago by McNaught (1906) [64] (Taylor *et al.*, 2005) [65]. Many clashing reports of presumptive antimicrobial activity detect that the specific antibacterial spectrum of green tea is actually difficult to assess.

Tea exhibits antimicrobial properties, which are ascribed predominantly to its polyphenols (Dias *et al.*, 2013) [66]. The degree of animation depends on the bacterial species and the polyphenol structure (Campos *et al.*, 2003; Taguri *et al.*, 2004) [67, 68]. The antibacterial activity of green tea is being gradually documented and first reported from Japan by using Japanese tea against various diarrheal pathogens (Hossain and Mahmood, 2014) [57]. The revelation of potent plant and plant seed extracts can effectively avert bacterial growth. Therefore, This has prompted an expanding enthusiasm among the researchers to evaluate the capability of green tea seed to inhibit the growth of some species of fish pathogenic bacteria (*Y. Ruckeri*, *P. Putida*, *P. Luteola*, *A. Hydrophila* and *L. Anguillarum*) with the intention of assessing them as possible disease preventive measures in aquaculture (Halder *et al.*, 2005) [9]. Good evidence suggests that the catechin components of green tea are responsible for the observed antibacterial activity owing to the presence of EGC, EGCG, and ECG constituents (Tsai *et al.*, 2008; Yam *et al.* 1997; Hara 2001) [59, 69, 70]. Gram-negative bacteria seem to be more resistant to polyphenols than Gram-positive bacteria, due to differences in the exterior membrane (Negi *et al.*, 2003) [71]. Yang *et al.* (2011) [72] reported no antibacterial activity occurred against Gram-negative *E. Coli*, *S. Typhi*, and *P. Aeruginosa*. This study also disclosed that green tea extracts inhibited the growth of Gram-positive *M. Luteus*, *S. Aureus*, and *B. Cereus*, with *M. Luteus* being most delicate (Bansal *et al.*, 2013) [16]. Su *et al.* (2008) [6] also are of the view that the antimicrobial activity of the green tea extract against the pathogenic bacteria *Staphylococcus aureus*, *Streptococcus pyogenes*, and *E. Coli*. Hara-Kudo *et al.* (2005) [73] revealed the antibacterial effects of major green tea polyphenols using *Clostridium* and *Bacillus* spores where *C. Botulinum* and *C. Butyricum* spores were diminished in number while no effect was shown in *Bacillus cereus* spores (Wongkham *et al.*, 2001) [14]. Only 100 mg green tea/ml is able to inhibit *S. Typhimurium*, whereas twice the tea concentration is needed for the inhibitory effect to be observed for *E. Coli* (Yam *et al.*, 1997; Kim and Fung, 2004) [69, 74]. The main components responsible for the antimicrobial activity are EGCG and

EGC. EGCG at 10–100 μ M has shown to reduce *E. Coli* growth by approximately 50% (Gramza and Korczak, 2005) [75]. Due to the highest antioxidant activity (TEAC values), the antimicrobial activity of nonfermented tea is higher than that of semi-fermented or fermented tea (Nazer *et al.*, 2005) [76].

Attributable to the far-reaching capacity of green tea to repress spore germination of plant pathogens and proposed for use against contagious fungal pathogens. The antifungal impacts of EGCG were mainly studied against yeasts such as *Candida spp.* and molds such as dermatophytes (Steinmann *et al.*, 2013) [77]. As of late, data relates between green tea polyphenols and irresistible aspergilli or other human-pathogenic zygomycetes which cause dysfunction of the local defense system and infections of the skin, hair, and nails of humans and animals are lacking (Pappas *et al.*, 2009) [78]. In an *in vitro* study, it was revealed that EGCG, EGC, and ECG cause metabolic instability of *C. Albicans* (Evensen and Braun, 2009) [79].

The mechanism of antiviral action of polyphenolic compounds is based on various capacities to go about as antioxidant agents, to inhibit proteinaceous enzymes, to disrupt cell membranes, to avoid viral binding and penetration into cells, and to trigger the host cell. EGCG hinders infections by direct authoritative to biological molecules and persuades agglutination of the flu infection preventing their adsorption to target (Friedman, 2007) [80]. The antiviral mechanism of EGCG has been analyzed against endemic HBV (Hepatitis B virus) infection (Heet *et al.*, 2011; Wang *et al.*, 2007) [81, 82]. Many reports demonstrated that green tea catechin, EGCG is the most active compound against HIV infectious diseases (Hamza and Zhan, 2006) [83]. Furthermore, the evaluation has also been done with herpes simplex virus (HSV) and bovine coronavirus (BCV) to realize the resistance power of antiviral activity and therapeutic efficiency of catechin polyphenols (Matsumoto *et al.*, 2005) [84].

Anticarcinogenic activity

With time, Polyphenols are turning into the issue of medicinal research. They have been accounted for to have numerous helpful properties, including mitigating action, compound hindrance, antimicrobial activity (Harborne and Baxter, 1999) [85], antiallergic activity, antioxidant activity (Elliott and Chithan, 2017) [86], cell reinforcement action, vascular action, and cytotoxic antitumor action (Harborne and Williams, 2000) [87]. It isn't bewildering, in this way, leading pharmaceutical organizations tend to focus their efforts on improving antimicrobial agents in established classes in setting up classes due to antimicrobial obstruction (Taylor *et al.*, 2002) [88].

Be that as it may, the accessible chemotherapy portfolio, it has been recognized that researchers are approaching the end of the activity in terms of alterations in the structure of parents. Thusly, an intrigue was made for the improvement of new classes of drugs that work in various objective destinations for those as of now being utilized (Cushnie and Lamb, 2005) [61]. Ongoing studies and others have found that isolated green tea polyphenols have anticarcinogenic activity. The effects of green tea as a malignant growth chemopreventive agent in a wide range of animal models are of impressive intrigue. The oral administration of green tea inhibits the formation of chemically induced tumors in several models (Ho *et al.*, 1994) [46]. Green tea components can prevent the formation of carcinogens from forerunner substances by rummaging cancer-causing electrophiles. (Han

and Xu, 1990)^[89]. Researchers have investigated that the freeze-dried green tea containing EGCG turns away carcinogenesis in rodents (Tariq and Reyaz, 2013)^[55]. Various scientists have subsequently reported in this manner covered the preventive impacts of green tea, positive or negative, against human malignant growths, in view of epidemiological examinations. Specifically, esophageal cancer in urban Shanghai was averted with green tea consumption (over 150 g tea leaves per month) (Amarowicz *et al.*, 2005)^[90]. Dr Waun Ki Hong started his clinical investigations of green tea extricate with a part of U.S. population that had not recently been expending green tea. In 1997, the US Federal Drug Administration (FDA) conceded consent for a Phase I clinical preliminary with green tea containers, which were produced by a Japanese Tea Company, ITO EN (Molan, 2013)^[11]. Green tea is an exceptional cancer-preventive prescription with two features as refreshment and medication. Herbal remedies, including green tea, are the same old thing, yet by presenting the idea of malignant growth counteractive action that green tea spares individuals' lives.

Conclusion

It can be concluded that green tea has several health benefits. It is the reach source of phytonutrients like flavonoids, phenolic acids, polyphenols, and catechin tannins. Green tea also has several hydrophilic antioxidants properties as Trolox and free radical scavengers. Green tea is very good in consumption as a morning drink. It has a sufficient amount of caffeine to give you a good morning energy boost. Unlike coffee, tea contains an amino acid L-theanine, that prevents caffeine rush and gives you the energy to sustain throughout several hours instead.

References

- Molan AL. Antioxidant and prebiotic activities of selenium-containing green tea. *Nutrition*. 2013; 29(2):476.
- Namita P, Mukesh R, Vijay KJ. *Camellia sinensis* (green tea): A review. *Global Journal of Pharmacology*. 2012; 6(2):52-59.
- Narotzki B, Reznick AZ, Aizenbud D, Levy Y. Green tea: a promising natural product in oral health. *Archives of Oral Biology*. 2012; 57(5):429-435.
- Fujiki H, Suganuma M, Imai K, Nakachi K. Green tea: cancer preventive beverage and/or drug. *Cancer letters*. 2002; 188(1, 2): 9-13.
- Zou LQ, Liu W, Liu WL, Liang RH, Li T, Liu CM *et al.* Characterization and bioavailability of tea polyphenol nanoliposome prepared by combining an ethanol injection method with dynamic high-pressure microfluidization. *Journal of Agricultural and Food Chemistry*. 2014; 62(4):934-941.
- Su P, Henriksson A, Nilsson C, Mitchell H. Synergistic effect of green tea extract and probiotics on the pathogenic bacteria, *Staphylococcus aureus* and *Streptococcus pyogenes*. *World Journal of Microbiology and Biotechnology*. 2008; 24(9):1837-1842.
- Chen ZY, Wang S, Lee KMS, Huang Y, Ho WKK. Preparation of flavonol-rich green tea extract by precipitation with AlCl₃. *Journal of the Science of Food and Agriculture*. 2001; 81(10):1034-1038.
- Navas PB, Carrasquero-Durán A, Flores I. Effect of black tea, garlic and onion on corn oil stability and fatty acid composition under accelerated oxidation. *International Journal of Food Science & Technology*. 2006; 41(3):243-247.
- Halder B, Pramanick S, Mukhopadhyay S, Giri AK. Inhibition of Benzo [a] Pyrene induced mutagenicity and genotoxicity by black tea polyphenols theaflavins and thearubigins in multiple test systems. *Food and Chemical Toxicology*. 2005; 43(4):591-597.
- Zhu YX, Huang H, Tu YY. A review of recent studies in China on the possible beneficial health effects of tea. *International Journal of Food Science & Technology*. 2006; 41(4):333-340.
- An BJ, Kwak JH, Son JH, Park JM, Lee JY, Jo C *et al.* Biological and anti-microbial activity of irradiated green tea polyphenols. *Food Chemistry*. 2004; 88(4):549-555.
- Erol NT, Sari F, Polat G, Velioglu YS. Antioxidant and antibacterial activities of various extracts and fractions of fresh tea leaves and green tea. *Tarim Bilimleri Dergisi*. 2009; 15(4):371-378.
- Jin D, Hakamata H, Takahashi K, Kotani A, Kusu F. Determination of quercetin in human plasma after ingestion of commercial canned green tea by semi- micro HPLC with electrochemical detection. *Biomedical Chromatography*. 2004; 18(9):662-666.
- Wongkham S, Laupattarakasem P, Pienthaweechai K, Areejitranusorn P, Wongkham C, Techanitiswad T. Antimicrobial activity of *Streblus asper* leaf extract. *Phytotherapy Research*. 2001; 15(2):119-121.
- Almajano MP, Carbo R, Jiménez JAL, Gordon MH. Antioxidant and antimicrobial activities of tea infusions. *Food Chemistry*. 2008; 108(1):55-63.
- Bansal S, Choudhary S, Sharma M, Kumar SS, Lohan S, Bhardwaj V *et al.* Tea: a native source of antimicrobial agents. *Food Research International*. 2013; 53(2):568-584.
- Wheeler DS, Wheeler WJ. The medicinal chemistry of tea. *Drug Development Research*. 2004; 61(2):45-65.
- Boran H, Çiftçi C, Er A, Köse Ö, Kurtoğlu İZ, Kayış Ş. Evaluation of antibacterial activity of green tea (*Camellia sinensis* L.) seeds against some fish pathogens in rainbow trout (*Oncorhynchus mykiss*, Walbaum). *Turkish Journal of Fisheries and Aquatic Sciences*. 2015; 15(1):49-57.
- Barbosa DS. Green tea polyphenolic compounds and human health. *Journal für Verbraucherschutz und Lebensmittelsicherheit*. 2007; 2(4):407-413.
- Wanasundara UN, Shahidi F. Antioxidant and pro-oxidant activity of green tea extracts in marine oils. *Food Chemistry*. 1998; 63(3):335-342.
- Archana S, Abraham J. Comparative analysis of antimicrobial activity of leaf extracts from fresh green tea, commercial green tea and black tea on pathogens. *Journal of Applied Pharmaceutical Science*. 2011; 1(8):149.
- Mbata TI, Debiao LU, Saikia A. Antibacterial activity of the crude extract of Chinese green tea (*Camellia sinensis*) on *Listeria monocytogenes*. *African Journal of Biotechnology*, 2008, 7(10).
- Balentine DA, Wiseman SA, Bouwens LC. The chemistry of tea flavonoids. *Critical Reviews in Food Science & Nutrition*. 1997; 37(8):693-704.
- Frei B, Higdon JV. Antioxidant activity of tea polyphenols *in vivo*: evidence from animal studies. *The Journal of Nutrition*. 2003; 133(10):3275S-3284S.
- Zandi P, Gordon MH. Antioxidant activity of extracts

- from old tea leaves. *Food Chemistry*. 1999; 64(3):285-288.
26. Yang TTC, Koo MWL. Hypcholesterolemic effects of Chinese tea. *Pharmacological Research*. 1997; 35(6):505-512.
 27. Manzocco L, Anese M, Nicoli MC. Antioxidant properties of tea extracts as affected by processing. *LWT-Food Science and Technology*. 1998; 31(7, 8):694-698.
 28. Tang FY, Meydani M. Green tea catechins and vitamin E inhibit angiogenesis of human microvascular endothelial cells through suppression of IL-8 production. *Nutrition and Cancer*. 2001; 41(1, 2):119-125.
 29. Yang CS, Chung JY, Yang GY, Chhabra SK, Lee MJ. Tea and tea polyphenols in cancer prevention. *The Journal of Nutrition*. 2000; 130(2):472S-478S.
 30. da Silva Pinto M. Tea: A new perspective on health benefits. *Food Research International*. 2013; 53(2):558-567.
 31. Davidson PM, Sofos JN, Branen AL. *Antimicrobials in food*. Edn 3, CRC press, Boca Raton. 2005; 1:429.
 32. Huo C, Wan SB, Lam WH, Li L, Wang Z, Landis-Piwowar KR. The challenge of developing green tea polyphenols as therapeutic agents. *Inflammo pharmacology*. 2008; 16(5):248-252.
 33. Mukamal KJ, MacDermott K, Vinson JA, Oyama N, Manning WJ, Mittleman MA. A 6-month randomized pilot study of black tea and cardiovascular risk factors. *American Heart Journal*. 2007; 154(4):724-e1.
 34. Nagao T, Hase T, Tokimitsu I. A green tea extract high in catechins reduces body fat and cardiovascular risks in humans. *Obesity*. 2007; 15(6):1473-1483.
 35. Chang YY, Jiang QY. Medicinal Value of Tea Polyphenols [J] *Food and Drug*, 2007, 8.
 36. Rezaei-Zadeh K, Arendash GW, Hou H, Fernandez F, Jensen M, Runfeldt M *et al*. Green tea epigallocatechin-3-gallate (EGCG) reduces β -amyloid mediated cognitive impairment and modulates tau pathology in Alzheimer transgenic mice. *Brain Research*. 2008; 1214:177-187.
 37. Wei SQ, Xu H, Xiong X, Luo ZC, Audibert F, Fraser WD. Tea consumption during pregnancy and the risk of pre-eclampsia. *International Journal of Gynecology & Obstetrics*. 2009; 105(2):123-126.
 38. Dutta AK, Siddiquee MA, Hossain S, Kabir Y. Finlay green tea possesses the highest *in vitro* antioxidant activity among the 20 commercially available tea brands of Bangladesh. *Malaysian Journal of Pharmaceutical Sciences*. 2013; 11(2):11.
 39. Chan EW, Soh EY, Tie PP, Law YP. Antioxidant and antibacterial properties of green, black, and herbal teas of *Camellia sinensis*. *Pharmacognosy Research*. 2011; 3(4):266.
 40. Koech KR, Wachira FN, Ngure RM, Wanyoko JK, Bii CC, Karori SM *et al*. Antimicrobial, synergistic and antioxidant activities of tea polyphenols. 2017.
 41. Manning J & Roberts JC. Analysis of catechin content of commercial green tea products. *Journal of Herbal Pharmacotherapy*. 2013; 3(3):19-32.
 42. Senanayake SN. Green tea extract: Chemistry, antioxidant properties and food applications-A review. *Journal of Functional Foods*. 2013; 5(4):1529-1541.
 43. Kristanti RA, Punbusayakul N. Antioxidant and antimicrobial activity of commercial green tea in Chiang Rai. In *Asia Pacific Symposium on Assuring Quality and Safety of Agri-Foods*. 2008; 837:53-58.
 44. Chacko SM, Thambi PT, Kuttan R, Nishigaki I. Beneficial effects of green tea: A literature review. *Chinese Medicine*. 2010; 5(1):13.
 45. Jo YH, Yuk HG, Lee JH, Kim JC, Kim R, Lee SC. Antioxidant, tyrosinase inhibitory, and acetylcholinesterase inhibitory activities of green tea (*Camellia sinensis* L.) seed and its pericarp. *Food Science and Biotechnology*. 2012; 21(3):761-768.
 46. Ho CT, Ferraro T, Chen Q, Rosen RT, Huang MT. *Phytochemicals in teas and rosemary and their cancer-preventive properties*, 1994.
 47. Sharma M, Nagori K, Soni S, Swarnakar H, Vaishnav S, Khan N. Phytochemical constituents and pharmacological profile of green tea: An overview. *International Journal of Pharmaceutical and Chemical*. 2014; 3:110-117.
 48. Patil MP, Patil KT, Ngabire D, Seo YB, Kim GD. Phytochemical, antioxidant and antibacterial activity of black tea (*Camellia sinensis*). *International Journal of Pharmacognosy and Phytochemical Research*. 2016; 8(2):341-346.
 49. Bancirova M. Comparison of the antioxidant capacity and the antimicrobial activity of black and green tea. *Food Research International*. 2010; 43(5):1379-1382.
 50. Rice-Evans C. Implications of the mechanisms of action of tea polyphenols as antioxidants *in vitro* for chemoprevention in humans. *Proceedings of the Society for experimental Biology and Medicine*. 1999; 220(4):262-266.
 51. Guo Q, Zhao B, Shen S, Hou J, Hu J, Xin W. ESR study on the structure-antioxidant activity relationship of tea catechins and their epimers. *Biochimica et Biophysica Acta (BBA)-General Subjects*. 1999; 1427(1):13-23.
 52. Yokozawa T, Nakagawa T, Kitani K. Antioxidative activity of green tea polyphenol in cholesterol-fed rats. *Journal of Agricultural and Food Chemistry*. 2002; 50(12):3549-3552.
 53. Nanjo F, Goto K, Seto R, Suzuki M, Sakai M, Hara Y. Scavenging effects of tea catechins and their derivatives on 1, 1-diphenyl-2-picrylhydrazyl radical. *Free Radical Biology and Medicine*. 1996; 21(6):895-902.
 54. Pon Velayutham A, Liu D. Green Tea Catechins and Cardiovascular Health: An Update. *Current Medicinal Chemistry*. 2008; 15(18):1840-1850.
 55. Tariq AL, Reyaz AL. Antioxidant activity of *Camellia sinensis* leaves. *International Journal Current Microbiology and Applied Science*. 2013; 2(5):40-46.
 56. Frankel EN, Huang SW, Aeschbach R. Antioxidant activity of green teas in different lipid systems. *Journal of the American Oil Chemists' Society*. 1997; 74(10):1309-1315.
 57. Hossain MM, Mahmood S. *In vitro* studies on antibacterial, thrombolytic and antioxidant activities of green tea or *Camellia sinensis*. *American Journal of Phytomedicine and Clinical Therapeutics*. 2014; 2:1200-1.
 58. <http://www.greenteanutritionfacts.com>. 25 September, 2019.
 59. Cao G, Sofic E, Prior RL. Antioxidant capacity of tea and common vegetables. *Journal of Agricultural and Food Chemistry*. 1996; 44(11):3426-3431.
 60. Tsai TH, Tsai TH, Chien YC, Lee CW, Tsai PJ. *In vitro* antimicrobial activities against cariogenic streptococci and their antioxidant capacities: A comparative study of

- green tea versus different herbs. *Food Chemistry*. 2008; 110(4):859-864.
61. Cushnie TT, Lamb AJ. Antimicrobial activity of flavonoids. *International Journal of Antimicrobial Agents*. 2005; 26(5):343-356.
 62. Clercq E. New developments in anti-HIV chemotherapy. *Current Medicinal Chemistry*. 2001; 8(13):1543-1572.
 63. Poole K. Overcoming antimicrobial resistance by targeting resistance mechanisms. *Journal of Pharmacy and Pharmacology*. 2001; 53(3):283-294.
 64. McNaught JG. On the action of cold or lukewarm tea on *Bacillus typhosus*. *Journal of the Royal Army Medical Corps*. 1906; 7(4):372-373.
 65. Taylor PW, Hamilton-Miller JM, Stapleton PD. Antimicrobial properties of green tea catechins. *Food Science and Technology bulletin*. 2005; 2:71.
 66. Dias TR, Tomás G, Teixeira NF, Alves MG, Oliveira PF, Silva BM *et al.* White Tea (*Camellia sinensis* (L.)): Antioxidant Properties and Beneficial Health Effects. *International Journal of Food Science, Nutrition and Dietetics*. 2013; 2(2):19-26.
 67. Campos FM, Couto JA, Hogg TA. Influence of phenolic acids on growth and inactivation of *Oenococcus oeni* and *Lactobacillus hilgardii*. *Journal of Applied Microbiology*. 2003; 94(2):167-174.
 68. Taguri T, Tanaka T, Kouno I. Antimicrobial activity of 10 different plant polyphenols against bacteria causing food-borne disease. *Biological and Pharmaceutical Bulletin*. 2004; 27(12):1965-1969.
 69. Yam TS, Shah S, Hamilton-Miller JMT. Microbiological activity of whole and fractionated crude extracts of tea (*Camellia sinensis*), and of tea components. *FEMS microbiology letters*. 1997; 152(1):169-174.
 70. Hara Y. *Green tea: health benefits and applications*. edn 1, CRC press, New York, 2001, 139-148
 71. Negi PS, Jayaprakasha GK, Jena BS. Antioxidant and antimutagenic activities of pomegranate peel extracts. *Food Chemistry*. 2003; 80(3):393-397.
 72. Yang H, Landis-Piwowar K, Chan TH, Dou PQ. Green tea polyphenols as proteasome inhibitors: implication in chemoprevention. *Current Cancer Drug Targets* 2011; 11(3):296-306.
 73. Hara- Kudo Y, Yamasaki A, Sasaki M, Okubo T, Minai Y, Haga M *et al.* Antibacterial action on pathogenic bacterial spore by green tea catechins. *Journal of the Science of Food and Agriculture*. 2005; 85(14):2354-2361.
 74. Kim S, Fung DYC. Antibacterial effect of water-soluble arrowroot (*Puerariae radix*) tea extracts on foodborne pathogens in ground beef and mushroom soup. *Journal of Food Protection*. 2004; 67(9):1953-1956.
 75. Gramza A, Korczak J. Tea constituents (*Camellia sinensis* L.) as antioxidants in lipid systems. *Trends in Food Science & Technology*. 2005; 16(8):351-358.
 76. Nazer AI, Kobilinsky A, Tholozan JL, Dubois-Brissonnet F. Combinations of food antimicrobials at low levels to inhibit the growth of *Salmonella* sv. Typhimurium: a synergistic effect?. *Food Microbiology*. 2005; 22(5):391-398.
 77. Steinmann J, Buer J, Pietschmann T, Steinmann E. Anti- infective properties of epigallocatechin- 3- gallate (EGCG), a component of green tea. *British Journal of Pharmacology*. 2013; 168(5):1059-1073.
 78. Pappas PG, Kauffman CA, Andes D, Benjamin Jr DK, Calandra TF, Edwards Jr JE *et al.* Clinical practice guidelines for the management of candidiasis: 2009 update by the Infectious Diseases Society of America. *Clinical Infectious Diseases*. 2009, 503-535.
 79. Evensen NA, Braun PC. The effects of tea polyphenols on *Candida albicans*: inhibition of biofilm formation and proteasome inactivation. *Canadian Journal of Microbiology*. 2009; 55(9):1033-1039.
 80. Friedman M. Overview of antibacterial, antitoxin, antiviral, and antifungal activities of tea flavonoids and teas. *Molecular Nutrition & Food Research*. 2007; 51(1):116-134.
 81. He W, Li LX, Liao QJ, Liu CL, Chen XL. Epigallocatechin gallate inhibits HBV DNA synthesis in a viral replication-inducible cell line. *World Journal of Gastroenterology*. 2011; 17(11):1507.
 82. Wang H, Xu J, Deng F, Hu Z. Natural extract of green tea for inhibiting hepatitis B virus and its primary active ingredient. *Faming Zhuanli Shenqing Gongkai Shuomingshu*, CN 101028382, 2007, 9.
 83. Hamza A, Zhan CG. How can (-)-epigallocatechin gallate from green tea prevent HIV-1 infection? Mechanistic insights from computational modeling and the implication for rational design of anti-HIV-1 entry inhibitors. *The Journal of Physical Chemistry B*. 2006; 110(6):2910-2917.
 84. Matsumoto M, Mukai T, Furukawa S, Ohori H. Inhibitory effects of epigallocatechin gallate on the propagation of bovine coronavirus in Madin-Darby bovine kidney cells. *Animal Science Journal*. 2005; 76(5):507-512.
 85. Harborne JB, Baxter H. *The handbook of natural flavonoids*. Edn 1, John Wiley and Sons, Chichester, 1999, I-II
 86. Elliott M, Chithan K. The impact of plant flavonoids on mammalian biology: implications for immunity, inflammation and cancer. In *The flavonoids advances in research since 1986*. Edn 1, Routledge, 1994, 2017, 619-652.
 87. Harborne JB, Williams CA. *Advances in flavonoid research since 1992*. *Phytochemistry*. 2000; 55(6):481-504.
 88. Taylor PW, Stapleton PD, Luzio JP. New ways to treat bacterial infections. *Drug Discovery Today*. 2002; 7(21):1086-1091.
 89. Han C, Xu Y. The effect of Chinese tea on occurrence of esophagealtumor induced by N-nitrosomethylbenzylamine in rats. *Biomedical and Environmental Sciences*. 1990; 3(1):35-42.
 90. Amarowicz R, Pegg RB, Dykes GA, Troszynska A, Shahidi F. Antioxidant and antibacterial properties of extracts of green tea polyphenols. In *Phenolic compounds in foods and natural health products*. American Chemical Society. 2005; 9:94-106.