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A review on health benefits of kombucha nutritional compounds and metabolites

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**ABSTRACT**
Kombucha is a beverage made by fermenting sugared tea using a symbiotic culture of bacteria and yeasts. Kombucha consumption has been associated with some health effects such as: the reduction of cholesterol levels and blood pressure, reduction of cancer propagation, the improvement of liver, the immune system, and gastrointestinal functions. The beneficial effects of kombucha are attributed to the presence of bioactive compounds that act synergistically. Bacteria contained in kombucha beverage belong to the genus *Acetobacter*, *Glucobacter*, and the yeasts of the genus *Saccharomyces* along with glucuronic acid, contribute to health protection. This review focuses on recent findings regarding beneficial effects of kombucha and discusses its chemical compounds, as well as the metabolites resulted by the fermentation process. Besides, some contraindications of kombucha consumption are also reviewed.

1. Introduction

All foods are essentially functional to a certain level because they provide the energy and nutrients needed to maintain life. However, there is evidence of the existence of bioactive food components that are not considered nutrients but can provide beneficial health effects (Crowe & Francis, 2013; Kaur & Singh, 2017; Shimizu, 2012; Tur & Bibiloni, 2016). Functional foods are those that have scientifically proven their beneficial effects in the organism, in one or more of its functions, providing optimal health and well-being, regardless of their nutritional value (Kaur & Singh, 2017).

These foods have a preventive function, due to the fact that they reduce risk factors that cause diseases. Shimizu (2012) refers to functional foods called FOSHU (Food for Specified Health Uses), which were approved by Japan’s Consumer Affairs Agency, a leader in functional foods regulations. Some of these foods improve the intestinal microbiota, regulate nutrient absorption and/or reduce the risk of chronic non-communicable diseases (Crowe & Francis, 2013).

It has been reported that certain dietary factors, such as lactic acid bacteria, oligosaccharides, amino acids, and polyphenols, may be promising ingredients for the future development of functional foods (Shimizu, 2012). However, the bioavailability and efficacy of these compounds at levels that are scientifically achievable in typical eating patterns should be revised (Kaur & Singh, 2017). For a product to be considered as a functional food, it must meet the following requirements: being a food product, having scientific evidence that supports the benefit of the product, having measurable physiological effects and being consumed daily as part of a normal diet (Tur & Bibiloni, 2016).

2. Kombucha definition

Kombucha is the name of the beverage obtained from the fermentation of tea, mainly black tea (there are also other varieties that can be used as a base for its preparation, such as green and oolong tea, also known as blue tea); with added sugar as a substrate for fermentation. Although this beverage has originally been prepared using tea, it is possible to find variations made with infusions like mint, lemon balm or jasmine. The taste of the beverage is slightly acidic...
and slightly carbonated, which provides greater acceptance among consumers. Some metabolic products of Symbiotic Culture of Bacteria and Yeast (SCOBY), like acetic acid and other organic acids, posses antibacterial activity and prevents contamination of the drink by pathogenic bacteria (Watawana, Jayawardena, Gunawardhana, & Waisundara, 2015).

3. Preparation of kombucha

This beverage is prepared by fermenting sugared tea with a SCOBY (Jayabanal, Malbaša, Lončar, Vitas, & Sathishkumar, 2014). Its flavour is slightly sweet and sour at the same time, plus it may contain traces of carbon dioxide (Nummer, 2013).

The typical production of kombucha beverage is based on black, green or oolong tea. For its production, 5 g of tea leaves per litre of water may be used, then, sugar is added, which will serve as a substrate for tea fermenting bacteria and yeasts. Approximately 50 g of sugar per litre of water is enough. Before adding the SCOBY or a bit of prepared kombucha, the beverage should be at a temperature close to 20°C. It is extremely important to use sanitized utensils and work in clean areas while making kombucha, in order to have control over the growth of microorganisms and to prevent unwanted contamination (Watawana et al., 2015). Likewise, it is important to control pH levels during fermentation of kombucha, and preferably stop this process when a pH level of 4.2 is reached, since the overproduction of acetic acid may be counterproductive (Kovacevic et al., 2014). Other food safety methods include pasteurizing the final product to prevent overproduction of alcohol and carbon dioxide, as well as the addition of 0.1% of sodium benzoate and 0.1% of potassium sorbate as food preservatives, and finally, keeping it refrigerated (Watawana et al., 2015).

3.1. Fermentation

Kombucha fermentation period is typically known to require a minimum of 3 days to a maximum of 60 days, depending on cultural practices (Watawana et al., 2015). The fermentation of kombucha is carried out at room temperature, optimizing fermentation time. Sucrose is used as the main carbon source in a concentration of 5–20%, providing the media and nutrients necessary for microorganism development. A SCOBY or the resulting liquid at a 10% concentration from a previous fermentation may be used as starter culture for fermentation (Vina, Semjonov, Linde, & Petatko, 2013).

According to Vina et al. (2013), the variables of the fermentation process, such as time, temperature and sucrose concentration, will determine the final concentration of organic substances such as acids and pH. Organic acids produced during this fermentation process diminished the tea’s pH value, which leads to a lack of oxygen induced by the acidity. Due to this, the number of possible pathogenic microbial cells, if any, diminishes, resulting in a safe beverage for consumption, despite having a microbial origin (Watawana et al., 2015).

3.2. SCOBY growth

The culture used for the kombucha fermentation has a variable microbiological composition according to its origin, the weather, geographical location and medium used for the fermentation process (Watawana et al., 2015). In kombucha beverage, “the most abundant prokaryotes in the symbiotic culture belong to the Acetobacter and Gluconobacter genus” (Jayabalan et al., 2014). These genus belong to the Acetobacteraceae family (Table 1), the bacteria are Gram-negative aerobic bacilli (Stasiak & Blazejak, 2009). They can be told apart by their ability to oxidize the acetate anion in carbon dioxide. The strains of the genus Acetobacter produce acetic acid from ethanol. This process is carried out by alcohol dehydrogenase and aldehyde dehydrogenase, enzymes that produce acetic acid, which enters the Krebs cycle obtaining water and carbon dioxide as end products (Teyssier & Hamdouche, 2016). The genus Gluconobacter is not capable of oxidizing acetate through the Krebs cycle, since it lacks the enzymes necessary for the oxidation process, like succinate dehydrogenase and alpha-ketoglutarate dehydrogenase, leading to accumulation of products, like gluconate, in the medium (Zoecklein, Fugeslang, Gump, & Nury, 1999).

Moreover, different yeast species can be found in kombucha, which outnumbers the acetic acid bacteria (AAB) (Jayabanal, Malini, Sathishkumar, Swaminathan, & Yun, 2010). The enzyme invervates, derived from yeasts, catalyses the hydrolysis of sucrose to glucose and fructose, producing ethanol through the glycolysis pathway. On the other hand, Gluconobacter and Acetobacter bacteria use glucose to produce gluconic acid and ethanol to produce acetic acid (Jayabalan et al., 2014). The production of ethanol and acetic acid inhibits the growth of pathogenic bacteria in the kombucha (Dufrense & Farnworth, 1999).

The osmophilic yeast and bacteria that are inoculated in the beverage for fermentation are the ones responsible for the growth of what is known as tea fungus, which has the scientific name of Medusomyces gisevii. Using sucrose as a carbon source, the acetic acid bacteria of the tea produce a network of cellulose as a secondary metabolite of fermentation, mainly the bacteria Acetobacter xilinus. The symbiotic mass of bacteria and yeast adheres to the biofilm, forming a thick jelly-like membrane also called zoogleal biofilm (Jayabanal et al., 2014).

The biofilm of microorganisms remains floating on the surface of the tea with an appearance very similar to a mushroom cap, which is why it usually receives that name (Watawana et al., 2015). Illana (2007) mentioned that the

| Bacteria                        | Yeasts                                      |
|--------------------------------|--------------------------------------------|
| Acetobacter xylinum, Acetobacter | Saccharomyces cerevisiae,                  |
| xilinaeides, Bacterium gluconicum, | Zygosaccharomyces bailii,                  |
| Acetobacter aceti, Acetobacter | Schisosaccharomyces pombe,                  |
| pasteurianus and Gluconobacter | Saccharomyces ludwigii,                     |
| oxydans, Lactobacillus sp.,     | Zygosaccharomyces rouxii,                   |
| Lactococcus sp., Leuconostoc sp., | Torulaspora delbrueckii,                   |
| Bifidobacterium sp., Thermus sp., | Brettanomyces bruxellensis,                |
| Allobaculum sp.,               | Brettanomyces lambicus,                    |
| Ruminococcaceae Incerate Sedis, | Brettanomyces custeri,                     |
| Propionibacterium sp.,         | Pichia membranaefaciens,                   |
| Enterococcus sp.               | Roekero apiculata and Torulopsis sp.       |

References: Marsh et al. (2014); Vina, Semjonov, Linde, & Petatko (2013); Battikh, Chaieb, Bakhrouf and Ammar (2011).
growth of this consortium of bacteria and yeasts induces the addition of new thicker membranes that take the shape of their container and heightens the symbiotic effect between bacteria and yeast (Table 1). The cellulose membrane keeps the microorganisms on the surface, allowing enough oxygen availability for its development and protecting the microorganisms from UV rays (Suhartatik, Karyantina, Marsono, Rahayu, & Kuswanto, 2011).

Several factors play an important role in the concentration of kombucha constituents, one of them is temperature. According to the investigation by Fu, Yan, Cao, Xie, and Lin (2014), keeping kombucha tea refrigerated at 4°C mildly decreases the content of acetic acid bacteria, from $9.3 \times 10^6$ CFU/mL to $3.4 \times 10^6$ CFU/mL during 14 days of storage; while the content of lactic acid bacteria decreases significantly, from approximately $23.5 \times 10^6$ CFU/mL to $2.7 \times 10^3$ CFU/mL during 8 days of storage. It has been reported that yeast has a positive impact on the survival of lactic acid bacteria at 30°C, but not at 12°C (Suharja, Henriksson, & Liu, 2012), which could mean that the low cooling temperature of 4°C may have limited the positive effect of yeasts over lactic acid bacteria, reducing its survival rate (Fu et al., 2014).

Marsh, O’Sullivan, Hill, Ross, and Cotter (2014) reported a sequence analysis of multiple samples of kombucha, in order to provide the most in depth study of microflora to date and to observe the changes occurred in the microbial population during kombucha production. They extracted DNA from cellulosic pellicles from 5 different geographic locations at two fermentation times. Different profiles were detected among samples, however, the major bacteria genus present was *Glucanacetobacter* (>85%) and a prominent *Lactobacillus* population was also identified (up to 30%) with a number of sub-dominant genera that have not been detected previously on kombucha. *Zygosaccharomyces* genus was the yeast found at >95% in the fermented tea and other greater fungal diversity not previously identified.

Jayabalan et al. (2010), analysed the microbiological and chemical composition of the kombucha fungus. Three samples of the tea fungus were used to evaluate its composition in different stages of fermentation, at 7, 14 and 21 days. Fibre and protein were the main components of the SCOBY. In regards to the proteins, a significantly large amount of amino acids was determined, the highest in concentration being the essential amino acids leucine and isoleucine (Table 2). In addition, an increase in all the components was observed over fermentation time. Likewise, minerals like sodium, potassium, and magnesium were found.

### 4. Chemical components of black tea and green tea

Tea comes from a leafy perennial crop, from the family Theaceae, known as *Camellia sinensis*, which was originally harvested in China. The young and tender leaves are used to make different varieties of tea, depending on the process to which it is subjected, resulting in black tea, green tea or oolong tea (González, 2003). For the production of black tea, the leaves are crushed and left exposed to high humidity, which causes an enzymatic oxidation by polyphenol oxidases (Valenzuela, 2004). To produce green tea, heating methods that inactivate enzymes using steam are utilized, which prevents fermentation (González, 2003). The oolong tea is produced by a partially fermented Chinese tea that is oxidized in the range from 10 to 70% (Chen et al., 2011) and is made by wilting fresh leaves by sun, then slightly bruising (Weerawatanakorn et al., 2015).

Tea has various components, like caffeine, alkaloids, amino acids, carbohydrates, proteins, chlorophyll, fluoride, aluminium, minerals and trace elements (National Cancer Institute, 2010). When the leaves are fresh, its flavonol or catechin content is very high. These flavonols, flavonoid derivatives, are characterized by their monomeric structure, and the ones commonly found in tea are epicatechin (EC), epigallocatechin (EGC), epicatechin gallate (ECG) and epigallocatechin gallate (EGCG) (Valenzuela, 2004).

It is important to consider that the oxidation process to which black tea is subjected induces a change in the monomeric structure of catechins, resulting in dimeric and polymeric flavonols, known as theaflavins and thearubigins. Therefore, green tea has a lower content of theaflavins and thearubigins due to the fact that it does not go through a fermentation process. For this very reason, teas have a different composition and thus its effect on health varies from one to another (Valenzuela, 2004). Table 3 presents the flavonoid components of tea, both black and green tea. The benefits that have been attributed to tea, both black and green, are mainly due to its catechin content, which is polyphenol derivative. These substances act as potent antioxidants and protect against the development of diseases. The beneficial effects of tea mentioned below have been studied mainly *in vitro*, while others are based on clinical and epidemiological evidence (Valenzuela, 2004).

| Table 2. Amino acid content (mg/g dry weight) of tea fungus (SCOBY) at different times of fermentation. | Table 3. Contenido de aminoácidos (mg/g peso seco) de hongos de té (SCOBY) en distintos momentos de la fermentación. Referencia: Jayabalan et al. (2010). |
|-----------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| **Amino acids**           | **Fermentation time**                                                                                                            |
|                            | 7th day                        | 14th day                      | 21st day                      |
| Essential amino acids     |                                |                                |                              |
| Isoleucine                | 28.1 ± 0.51                   | 35.2 ± 0.40                   | 44.2 ± 0.36                   |
| Leucine                   | 27.2 ± 0.75                   | 35.9 ± 0.64                   | 45.1 ± 0.60                   |
| Lysine                    | 39.5 ± 0.50                   | 48.0 ± 0.36                   | 53.1 ± 0.40                   |
| Methionine                | 6.3 ± 0.55                    | 11.3 ± 0.80                   | 20.2 ± 0.50                   |
| Phenylalanine             | 13.6 ± 0.55                   | 22.3 ± 0.70                   | 30.2 ± 0.60                   |
| Threonine                 | 7.7 ± 0.20                    | 13.2 ± 0.58                   | 20.1 ± 0.65                   |
| Valine                    | 15.1 ± 0.26                   | 22.3 ± 0.45                   | 30.2 ± 0.61                   |
| Tryptophan                | 7.4 ± 0.26                    | 12.3 ± 0.51                   | 21.1 ± 0.45                   |
| Non-essential amino acids |                                |                                |                              |
| Alanine                   | 30.9 ± 0.55                   | 41.9 ± 0.40                   | 53.0 ± 0.50                   |
| Arginine                  | 14.5 ± 0.60                   | 30.8 ± 0.60                   | 42.2 ± 0.60                   |
| Aspartic acid             | 30.3 ± 0.70                   | 42.0 ± 0.65                   | 53.2 ± 0.60                   |
| Cysteine                  | 10.3 ± 0.70                   | 15.2 ± 0.35                   | 24.4 ± 0.47                   |
| Glutamic acid             | 32.2 ± 0.61                   | 42.3 ± 0.50                   | 50.1 ± 0.47                   |
| Glycine                   | 9.5 ± 0.55                    | 17.2 ± 0.30                   | 26.6 ± 0.81                   |
| Histidine                 | 6.0 ± 0.50                    | 10.6 ± 0.55                   | 18.5 ± 0.60                   |
| Proline                   | 28.5 ± 0.45                   | 35.2 ± 0.60                   | 43.4 ± 0.55                   |
| Serine                    | 11.2 ± 0.36                   | 22.2 ± 0.51                   | 31.7 ± 0.61                   |
| Tyrosine                  | 9.9 ± 0.45                    | 18.6 ± 0.40                   | 27.0 ± 0.55                   |

*Values are mean ± SD; n = 3*
Protective effect of polyphenols on cancer initiation, promotion, and progression. Adapted from: Valenzuela (2003).

Theaflavin-3-gallate, theaflavin-3,3′-digallate
Quercetin, kaempferol, rutin
Caffeic acid, quinic acid, gallic acid.

Another relevant effect of tea polyphenols is their protection against the development of cardiovascular diseases (CVDs). By inhibiting the oxidation of low-density lipoproteins (LDL), they assist in the prevention of the development of atheroma. They may also be involved in cholesterol metabolism by inhibiting pancreatic lipase, thus decreasing cholesterol and triacylglycerol absorption. Finally, they promote smooth muscle relaxation, preventing high blood pressure induced by vasoconstrictors, such as thromboxanes (González, 2003).

5. Chemical components of kombucha and their beneficial effects

Chemical assays of kombucha beverage have indicated the presence of a variety of compounds, including organic acids, mainly acetic, gluconic, and glucuronic acid (GlcUA), although citric, L-lactic, malic, tartaric, malonic, oxalic, succinic, pyruvic, and usnic acids may also be found; sugars (sucrose, glucose, and fructose), water soluble vitamins (B₁, B₂, B₆, B₁₂, C), amino acids, biogenic amines, purines, pigments, lipids, proteins, hydrolytic enzymes, ethanol, acetic acid bacteria and lactic acid bacteria, carbon dioxide, polyphenols, minerals (manganese, iron, nickel, copper, zinc, plumb, cobalt, chromium, and cadmium), anions (fluoride, chloride, bromide, iodide, nitrate, phosphate, and sulphate), D-saccharic acid-1,4-lactone (DSL), and metabolic products of yeasts and bacteria (Jayabal, Malbaša, et al., 2014, Jayabal, Malini, et al. 2010).

The presence and quantity of the chemical components are variable, mainly depending on the microorganisms of the symbiotic culture used for fermentation of kombucha, as well as fermentation time and temperature, sucrose content,
and type of tea used, in addition to the analysis methods used for quantification. Beneficial metabolites produced in kombucha are synthetized in Figure 2.

5.1. Vitamins

Regarding the vitamin content of this beverage, Bauer-Petrovska and Petrushevska-Tozi (2000) analysed a brew of kombucha beverage prepared with 70 g of sucrose and 5 g/L of black tea, finding the following values of B vitamins: 74 mg/100 mL of vitamin $B_1$, 52 mg/100 mL of vitamin $B_6$ and 84 mg/100 mL of vitamin $B_{12}$. Meanwhile, Malbaša, Lončar, Vitas, and Canadanović-Brunet (2011) reported that the content of vitamin $B_2$ was 8.3 mg/100 mL, while the concentration of vitamin C constantly increased, reaching 28.98 mg/L on the tenth day of fermentation.

5.2. Minerals

Minerals are inorganic substances needed in small amounts for normal body functions and growth, as well as for the maintenance of its tissues. According to Bauer-Petrovska and Petrushevska-Tozi (2000), copper, iron, manganese, nickel, and zinc are minerals that increased due to metabolic activity of kombucha. Mineral concentration was in a range of 0.004 μg/mL for cobalt and 0.462 μg/mL for manganese. Besides, traces of lead (0.005 μg/mL) were detected. It is worth noting that, according to the Agency for Toxic Substances and Disease Registry (ATSDR, 2007), toxic blood lead levels are 20 μg/dL for adults and 10 μg/dL for children, which is equivalent to 0.2 and 0.1 μg/mL, respectively. Kombucha tea has much lower concentrations, thus not representing a potential health risk. Additionally, Markowitz (2011) mentioned that small amounts of lead in the blood of adults are not harmful.

Nevertheless, it is important to consider that because children are more susceptible to the effects of lead than adults, it would be advisable for children not to drink this beverage on a regular basis, to prevent a chronic exposure that could cause them lead poisoning. Meanwhile, Kumar, Narayan, and Hassarajani (2008) established the presence of fluoride, chloride, bromide, iodide, nitrate, phosphate, and sulphate after seven days of fermentation of kombucha prepared with 100 g of sucrose and 5 g/L of black tea; being fluoride the anion with the highest concentration (3.2 mg/g).

5.3. Polyphenols

Polyphenols are active substances with more than one phenol structural unit per molecule. They represent the largest group of phytochemicals and they are the most abundant antioxidants present in the diet. Total intake of polyphenols can be up to 1 g/day (Scalbert, Johnson, & Saltmarsh, 2005). Moreover, they play a role in preventing several diseases related to oxidative stress, such as cancer, CVDs, and neurodegenerative diseases (Manach, Scalbert, Morand, Rémésy, & Jiménez, 2004). They modulate the activity of a variety of enzymes and cell receptors as a means of defence against oxidative stress caused by reactive oxygen species (Tsao, 2010). Main dietary polyphenol sources include fruits, vegetables, cereals, legumes, natural fruit juices, tea, coffee, and red wine (Scalbert et al., 2005).

The protective effect of kombucha beverage is mainly due to polyphenol activity, compounds produced during fermentation, and the synergistic effect of the different compounds found in the tea (Jayabal, Subathradevi, Marimuthu, Sathishkumar, & Swaminathan, 2008). Total polyphenol content in kombucha tea shows a linear increase during fermentation time (Chu & Chen, 2006). As an example, both epicatechin (EC) and epigallocatechin (EGC) are found predominantly in the tea (Manach et al., 2004). A higher level of EC (~150%) was found on day 12 of fermentation of kombucha made with green tea, and of EGC (~115%) on the same day of one made with black tea (Jayabal, Marimuthu, & Swaminathan, 2007).

A research conducted by Fu et al. (2014) used different types of kombucha to compare free-radical scavenging abilities against 2,2-diphenyl-picrylhydrazyl (DPPH), hydroxyl radicals, and superoxide radicals. Different types of tea were used to prepare kombucha: low-cost green tea, black
teas, and tea powder and fermentation process was 90 h. The results showed that kombucha made with green tea had the highest free-radical scavenging ability against DPPH, hydroxyl radicals, and superoxide anions. According to Chu and Chen (2006), black tea reaches its maximum activity against DPPH radicals until day 15 of fermentation. Tea powder had greater antioxidant activity than black tea against DPPH and hydroxyl radicals, but not against superoxide radicals, upon which black tea had a greater activity than tea powder.

5.4. D-saccharic acid-1,4-lactone (DSL)

D-saccharic acid-1,4-lactone (DSL) is a component derived from D-glucaric acid (product of the GlcUA pathway), which has detoxifying and antioxidant properties (Bhattacharya, Gachhui, & Sil, 2012; Ziółtaszek, Hanausek, Kiliarńska, & Walaszek, 2008). DSL content in kombucha has been found to range between 57.99 and 132.72 μg/mL, depending on the origin of the product. The highest DSL value was found on the eighth day of fermentation and diminished afterward. It was established that lactic acid bacteria had a positive effect on DSL production, in symbiosis with Gluconobacter genus bacteria (Yang et al., 2010).

5.5. Ethanol

According to Chen and Liu (2000), ethanol concentration in kombucha increases with fermentation time, reaching an approximate maximum value of 5.5 g/L on the 20th day of fermentation, followed by a slow reduction.

The entire chemical composition of kombucha beverage, including residual sugar concentration, carbon dioxide, and organic acids, is what finally determines its flavour and depending on the fermentation time, different flavours will be obtained. It has been observed that kombucha tea with higher acetic acid concentration produces a more acid and astringent flavour, meanwhile, another with more gluconic acid produces a milder flavour. Therefore, by controlling fermentation conditions it is possible to obtain the desired quality in kombucha tea (Chen & Liu, 2000).

The French paradox is the observation of low coronary heart disease (CHD) death rates despite the high intake of dietary cholesterol and saturated fat. It was proposed by French epidemiologists in the 1980s, particularly for wine drinking in France, where a high intake of dietary cholesterol and saturated fat, daily consumption of wine, but low CHD death rates were observed. The French paradox states that moderate alcohol consumption has a protective effect against CHD, since it raises high-density lipoprotein (HDL) cholesterol concentrations (Ferrières, 2004). Since the daily consumption of ethanol in a lower concentration can protect the human body from CVDs, the consumption of kombucha, which has a low concentration of ethanol, could also have a role in preventing CVDs.

5.6. Acetic acid

Acetic acid bacteria in kombucha produce acetic acid when these act on ethanol from sucrose, which is metabolized into glucose and fructose (Specking, 2015). Acetic acid is the chemical compound responsible for the acidic smell and taste of vinegar. Its name comes from the Latin acetum, which means vinegar (Bramforth, 2014). Acetic acid tends to slowly increase reaching 11 g/L at 30 days of fermentation, gradually diminishing until ending at 8 g/L at 60 days of fermentation. This decrease is due to its later utilization as a carbon source for bacteria when sugars in the tea are used up, or because of the decrease in ethanol metabolism by yeast due to low pH (Chen & Liu, 2000). If a different carbon source is used, such as molasses, the amount of acetic acid produced is considerably lower (Jayabalan et al., 2014).

Other important organic acids are contained in kombucha depending on the tea basis. Profiles of organic acids change during fermentation of green and black tea. Besides of acetic acid, lactic and citric acid are produced (Jayabalan et al., 2007). Gluconic acid, ethyl-glucoluate, oxalic acid, saccharic acid, keto-gluconic, succinic and carbonic acids were considered to be present in kombucha. Some of those acids were proved to have in vitro antimicrobial activity and improve sleep (Greenwald, Steinkraus, & Ledford, 2000; Sreearamulu, Zhu, & Knol, 2000). Kombucha is a high source of glucuronic acid, which has a detoxifying effect against drug, bilirubin, and chemicals, as well as pollutants and excess of steroid hormones (Nguyen, Nguyen, Nguyen, & Le, 2015). Knowing that glucuronic acid has specific benefits some of its specific properties are described below.

5.7. Glucuronic acid (GlcUA)

GlcUA plays a role in xenobiotic liver detoxification, as it has the ability to combine with toxin molecules to favour their elimination from the organism, which makes it very important as an auxiliary on liver functions. It is as well involved in endobiotic elimination. One of this endobiotics is bilirubin, which is how GlcUA (by means of glucuronidation) prevents these pigment’s toxic effects and impedes inhibition of a variety of enzymes involved in protein and carbohydrate metabolism. Most of the bilirubin is excreted through bile and only a small portion of conjugated bilirubin is excreted through urine, which is why a high level of bilirubin in urine is an indicator of damage somewhere along the process of glucuronidation (Vina, Linde, Patetko, & Semjonovs, 2013).

GlcUA also takes part in polyphenols’ increased bioavailability. Phenols conjugate with GlcUA, improving its transport and bioavailability. The UGT1A isoform of the glucuronosyltransferase family, located in the bowels, is the one that takes part in polyphenol glucuronidation. Polyphenols are secreted via bile into the duodenum, where they are subjected to the activity of β-glucuronidase (which promotes deglucuronidation, separating in this case the polyphenol from GlcUA by hydrolysing its glycosidic bond) and are then reabsorbed. This enterohepatic circulation can lead to a longer presence of polyphenols in the body, where they carry out its antioxidant activity preventing different diseases related to oxidative stress (Vina et al., 2013).

Several steroid hormones and vitamin D derivatives, such as estrogens, androgens, glucocorticoids, mineralocorticoids, progestogens and cholecalciferol are essential for health. They regulate immune functions, decrease inflammatory response, balance extracellular fluid volume, among others (Cutolo et al., 2014). Deficiencies and/or excesses of steroid hormones have undesirable effects on health. Glucuronidation can prevent both scenarios: it prevents deficiencies by increasing steroid water solubility, therefore improving its transport and bioavailability; and prevents toxic effects and impedes inhibition of a variety of enzymes involved in protein and carbohydrate metabolism.
excesses by facilitating the elimination of excess steroids (Vina, Linde, et al., 2013, Vina, Semjonovs, et al., 2013).

As a constituent of glucuronidation, GlcUA is also important for biotransformation and protection of fatty acids from lipid peroxidation. Especially, polyunsaturated fatty acids, which are essential compounds for the body, essential components of cell membranes and precursors of eicosanoids. Polyunsaturated fatty acids are susceptible to react with reactive oxygen species, triggering chain reactions that damage the fatty acid molecule. Peroxidation has been considered a risk factor for the development of certain pathologies, such as atherosclerosis, kidney damage and Parkinson's disease (Mylonas & Kouretas, 1999).

Kombucha consumption prevents polyunsaturated fatty acid oxidation in the human body, due to its GlcUA content that takes part in glucuronidation, increasing polyphenol bioavailability, which neutralizes free radicals that promote lipid peroxidation (Jayabalal et al., 2008).

GlcUA is necessary for many body functions since it is a constituent of various essential polysaccharides in the body, glycosaminoglycans (GAGs). GAGs are a series of compounds formed by dimers consisting of an amino sugar (D-Glucosamine or D-Galactosamine) and a uronic acid (D-glucuronic acid or L-iduronic acid), except for keratan sulphate which contains a galactose molecule instead of an acid. They are bound to sulphate groups in variable proportions and covalently bind to proteins, forming proteoglycans, except for hyaluronic acid which is not sulphated (Mylonas & Kouretas, 1999). Different GAGs are formed depending on the constituents of the dimer.

Glycosaminoglycans molecules are part of the extracellular matrix of all the body organs and they have multiple functions. GlcUA is part of the following GAGs: hyaluronic acid, chondroitin sulphate, heparin and dermatan sulphate. They all have structural functions, except for heparin, which is a non-structural GAG. Hyaluronic acid serves as a lubricant and shock absorber, and it is present in higher concentrations in the vitreous fluid in the eye, conjunctive tissue, synovial fluid of joints and cartilage (Frati-Munari, 2012; Vina et al., 2013).

Chondroitin sulphate is mostly present in bones and cartilage, in the latter it binds to collagen and keeps fibres in a strong network; it also helps to prevent joint problems, being helpful in the relief of osteoarthritis. Heparin is an intracellular compound and a potent anticoagulant produced by mast cells. Dermatan sulphate is present in higher concentrations in the vascular endothelium, connective tissue, cartilage, skin, cornea, and bones. L-iduronic acid is the predominant uronic acid present in dermatan sulphate, and it is an epimer of D-glucuronic acid, although a variable amount of β-D-glucuronic acid is also present (Frati-Munari, 2012; Vina et al., 2013).

Moreover, GlcUA is a precursor of L-ascorbic acid (vitamin C) in kombucha beverage, since it is synthesized from L-gulonic acid, which is involved in the metabolic pathway of GlcUA. Thus, GlcUA concentration in kombucha increases its antioxidant activity as well (Vina et al., 2013).

Actually, there are published data describing the similarity of GlcUA contained in kombucha and the acid produced in the human body by metabolic pathways. Preliminary studies carried out by Jayabalal et al. (personal communication) did not find similarity. As previously discussed, the concentration and presence of GlcUA in kombucha tea is basically determined by the strains and species that acts by symbiosis. Meaning that the GlcUA concentration in kombucha is exhibited due to the specific SCOBY microorganisms.

6. Other reported beneficial effects of kombucha

Researchers and testimonials of individuals, which mention having consumed this drink, declare beneficial effects for human health. Aloulou et al. (2012) evaluated the suppressing effect of α-amylase enzyme (secreted by intestinal epithelium and necessary for carbohydrate digestion) in diabetic rats (aloxan induced), which were administered 5 mL/kg of kombucha or black tea daily during 30 days. The results showed that the rats which drank kombucha had a better suppressing effect of α-amylase enzyme in pancreas and plasma, as well as postprandial glucose compared to those of the rats which drank black tea. Besides glucose metabolism disorders, pancreatic and plasma enzymatic changes were also evaluated.

These enzymes “act on triacylglycerol to metabolize it into free fatty acids and monoaoylglycerol”, an abnormal increase can be caused by pancreatic damage (Sastre, Sabater, & Aparisi, 2005). In the study, the rats administered aloxan, presented damage in pancreatic structure, more than that of the rats from the control group or those treated with kombucha. Aloxan increases reactive oxygen species (ROS), producing toxicity in pancreatic cells. An increase in pancreatic and plasmatic lipase concentration causes an increase in lipid absorption, which leads to an increase of triacylglycerol and low density proteins. The group treated with kombucha had a significant reduction of pancreatic and plasmatic lipase. The group treated with black tea had a decrease on both enzymes as well, but the decrease was lower than in the group consuming kombucha.

One more scientific support carried out by Kabiri, Setorki, and Ahangar (2013), a study which determines the protective effects of kombucha beverage and silymarin (milk thistle) in rats with liver damage induced by thioacetamide (hepatic fibrosis related toxin). In the study, 36 rats were divided into 6 groups, where group 1 was designated as control group. Group 2 was integrated by rats injected with thioacetamide; group 3 included the rats injected with thioacetamide and later treated with kombucha (50 mL/during 3 weeks); group 4 included rats treated with kombucha (50 mL/during 3 weeks) and later injected with thioacetamide; group 5 included rats injected with thioacetamide and later treated with silymarin (200 mg/kg during 3 weeks); and group 6 included rats injected with thioacetamide and later treated with kombucha (50 mL/per rat) and silymarin (400 mg/kg) during 3 weeks.

The results showed that the group treated with silymarin had a significant descent in the previously mentioned parameters with exempt of bilirubin. This same situation happened with the group treated with kombucha tea and silymarin. The protective action of both foods is ought to their polyphenol component, which protect the liver against free radical formation which can produce hepatocyte mal-function and liver damage.

Deghrique, Chriaa, Battikh, Abid, and Bakhrouf (2013), evaluated kombucha’s anti-proliferative properties, prepared with black or green tea, fermented during 12 days, on two human cancer cell lines (A549, lung cell carcinoma and Hep-2, epidermoid cell carcinoma). The cells were incubated in 96
7. Contraindications of kombucha

Toxicity caused by kombucha has been suspected in several cases, reporting dizziness and nausea after consumption. Lead poisoning and gastrointestinal toxicity was found on cases, reporting dizziness and nausea after consumption.

Regarding on the increase in phenolic compounds and antioxidant activities of kombucha, Sun, Li, and Chen (2015) elaborated the beverage with mixes in various ratios of sweetened black tea and wheatgrass juice. The highest antioxidant activity was obtained using a 1:1 (v/v) black tea decoction to wheat grass juice ratio and 3 days of fermentation. Under those processing conditions, this beverage produced various types of complementary phenolic acids with antioxidant effect, fact that was considered by the authors as an advantage over traditional kombucha beverage.

On the other hand, it has been proved the tea has potential to revert hepatic toxicity induced by CCl₄ (carbon tetrachloride), a liquid which transforms into gas at room temperature. It comes from aerosol and certain refrigerators production and can cause adverse health effect and hepatic toxicity if recommended dose is exceeded (Agency for Toxic Substances and Disease Registry [ATSDR], 2005; Kovacevic et al., 2014). Four HIV positive patients reported secondary effects related to its consumption, such as allergic reaction, ictericia, nausea, vomit, neck pain and headache. However, kombucha cannot be declared toxic for human health due to the presented evidence not being substantial for the affirmation of its toxicity or the disease occurrence in previous studies (Jayabal et al., 2014).

Some harmful effects of kombucha consumption have been described by several authors (Greenwalt et al., 2000). Organs internal lesions on rats after 12 weeks of kombucha consumption were reported and it was concluded that the susceptibility to toxicity depends on the specie (Ibrahim, Kwanashie, Njoku, & Olurinola, 1993). Developing of acidosis was described in individuals having severe pre-existing conditions. One of them increased the fermentation time from 7 to 14 days which produces a very acidic beverage (CDC, 1995). High acidity and microbial contamination of kombucha was reported as a warning for possible illness (Perry, 1995), and mycotoxigenic substances (as secondary metabolites) have health repercussions including toxic and carcinogenic effects.

6. Conclusions

Kombucha beverage is a source of bioactive components, such as polyphenols and glucuronic acid. The beneficial outcomes of kombucha consumption are attributed to the synergistic effect between these components, making it a drink with potential beneficial health properties (when elaborated under adequate sterile conditions). It is apparent that its consumption can protect against the development of CVDs, mainly due to its polyphenol content that inhibits the oxidation of LDL, regulates cholesterol metabolism, and prevents high blood pressure by promoting smooth muscle relaxation. GlcUA, one of its main components, plays a role in xenobiotic liver detoxification and endobiotic elimination, thus potentially enhancing liver functions. It must be emphasized that concentration of the drink’s active components will vary depending on the scoby and elaboration methods. Health effects on humans under controlled research are merited, because some contraindications have been reported.

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