

Assessment of Metabolites Content of Native Alcoholic Beverages Obtained from Jos, Nigeria

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To cite this article:

Samuel Yusufu Gazuwa, Jonathan Dingkwoet Dabak, Kiri Hashimu Jaryum, Segun Afolabi Olomu. Assessment of Metabolites Content of Native Alcoholic Beverages Obtained from Jos, Nigeria. *International Journal of Food Science and Biotechnology*. Vol. 5, No. 4, 2020, pp. 97-101. doi: 10.11648/j.ijfsb.20200504.19

Received: November 3, 2020; **Accepted:** November 20, 2020; **Published:** December 16, 2020

Abstract: *Burukutu*, (BKT) *Pito* (PT) and *Goskolo* (GSK) are drinks consumed routinely within the area of study. Although consumption of GSK is illegal owing to its life threatening toxicity, people still consume it secretly. Whereas BKT and PT are locally prepared using unscientific methods in preparing the drinks, GSK is a mixture containing different compounds and chemicals. BKT and PT are more commonly ingested than GSK. Because methods applied during 'brewing' BKT and PT are not scientific, toxic metabolites might be generated. So, the drinkers are prone to their toxic effects. This work ascertained the presence and levels of unwanted metabolites in these drinks using Gas chromatographic-Mass spectrometric (GC-MS) technique. Metabolites could aggravate the toxicity of alcohol and alcohol-related disease conditions. Results, (% relative abundance), showed the significant presence in BKT of 6-Octadecanoic acid (38.51), Hexadecanoic acid (20.92), Isopentyl alcohol (12.27), 1,11,13-Octadecatriene (11.60) among others. In PT, 9-Octadecanoic acid (39.40), Hexadecanoic acid (15.98), Carbonic acid methylpentyl ester (14.22), 9,17-Octadecadienal (6.37) were the major components. In GSK, 6-Octadecanoic acid (42.43), n-Hexadecanoic (17.51), 1-Butanol (16.52) and 9,12-Octadecadienoic acid (10.72) were the dominant components. BKT, PT, and GSK are composed of compounds/metabolites that are organic in nature, with about 83% of them being organic acids. This leads to a decrease in pH thereby making the drinks acidic. It is concluded that the alcoholic beverages contain metabolites that predispose drinkers to metabolic or respiratory acidosis, due to drop in pH. Furthermore, these metabolites could aggravate the toxicity of alcohol. Other components of the samples include ketones, different types of alcohols, aldehydes, derivatives of benzene and esters.

Keywords: Organic, Metabolites, Acids, *Burukutu*, *Pito*, *Goskolo*, Toxic

1. Introduction

Native beers (BKT and PT) are widely consumed as food (because it is thick, heavy, and nutritious) in the rural areas of northern Nigeria and in poor urban neighborhoods because it is more affordable than commercially brewed beer [1]. Also, native beers find relevance during cultural activities such as payment of dowry during marriage ceremonies, naming of children and even for religious purposes [2, 3] Solange et al., 2014). But these drinks are prepared under unscientific conditions and techniques. This could result in metabolic shift from a precursor to undesirable and harmful

metabolites [4]. This work was designed to qualitatively and quantitatively assess the metabolites generated during preparation of native beers.

2. Materials and Methods

2.1. Materials

(GCMS-QP2010) SHIMADZU, Equipment. Stationary phase used was polyethylene glycol (PEG 400). Mobile phase used was helium gas.

Operating conditions (GC): Column oven temperature: 60°C; Injection Temperature: 250°C; Injection Mode: split;

Flow control Mode: Linear Velocity; Pressure: 100.20kPa; Total Flow: 6.2ml/min; Column Flow: 1.61 ml/min; Linear Velocity: 46.3 cm/sec; Purge Flow: 3.0ml/min; Split ratio: 1.0; High Pressure Injection: off, Carrier Gas Saver: off; Splinter Hold: off. Oven Temperature: At zero rate: temperature was 60°C, Hold Time (minute): 5.0. At 5.0 rate: Temperature was 140°C, Hold Time was 0.00. At 15 rate: Temp was 280°C whereas Hold Time was 10.00. Ion source Temperature: 200.00°C, Interface Temperature: 250°C, Solvent cut Time: 2.5/min; Detector Gain Mode: Relative; Detector Gain: 0.00Kv; Threshold: 4000.

Operating conditions (MS): Start Time: 3.0 min; End Time: 38.0 min; ACQ Mode: Scan; Event Time: 0.50 seconds; Scan Speed: 769; Start m/z: 30.00; End m/z: 400.00.

2.2. Methods

Preparation of samples

Native beer samples were purchased from local drinking joints scattered in area of study. They were kept in plastic containers and stored in a refrigerator. Prior to analysis, refrigerated samples were removed and kept on table in open laboratory to change to liquid form. They were filtered threefold using filter paper; spun at 3000 rpm for 2 hours in a refrigerated ultracentrifuge machine. The supernatant was separated from the pellet by careful decantation. Each supernatant was re-centrifuged using the same procedure above until a clear and transparent sample was obtained. Samples were then analysed using Gas chromatographic-Mass spectrometer (GCMS-QP2010) SHIMADZU Equipment.

3. Results and Discussion

Table 1. Gas Chromatographic-Mass Spectrometric analysis results of Burukutu.

Peak number	Retention time (minutes)	Chemical formula of Compound	Name of Compound	Area (%)
1	3.34	C ₅ H ₁₂ O	Isopentyl alcohol	12.27
2	3.67	C ₂ H ₄ O ₂	Ethyllic acid	0.09
3	4.27	C ₂ H ₈ O ₂ Si	Dihydroxydimethylsilane	1.64
4	5.45	C ₄ H ₁₀ O ₂	2,3- Dihydroxybutane	0.48
5	14.52	C ₈ H ₁₀ O	Benzeneethylalcohol	1.21
6	28.33	C ₁₆ H ₃₂ O ₂	Hexadecanoic acid	20.92
7	29.53	C ₁₈ H ₃₄ O ₂	6-Octadecanoic acid	38.51
8	29.79	C ₁₈ H ₃₂	1,11,13-Octadecatriene	11.6
9	30.38	C ₃₇ H ₇₄ NO ₈ P	*H1,2AHPOEE	2.45
10	30.78	C ₂₀ H ₄₀ O ₂	Arachic acid	4.36
11	31.51	C ₅₇ H ₁₀₄ O ₆	*2,3B9OP9O	6.46

H12AHPOEE = Hexadecanoic acid, 1-[[[(2 aminoethoxy)hydroxyphosphinyl]oxy]methyl]-1,2-ethanediy ester.

*2,3B9OP9O = 2,3-Bis[(9)-9-Octadecenoyloxy]propyl-9-octadecanoate.

Table 2. Gas Chromatographic-Mass Spectrometric analysis results of Pito.

Peak number	Retention time (minutes)	Chemical formula of Compound	Name of Compound	Area (%)
1	3.317	C ₇ H ₁₄ O ₃	Carbonic acid, methyl pentyl ester	14.22
2	4.52	C ₄ H ₁₀ O ₂	2,3-Butanediol	5.46
3	13.676	C ₇ H ₈ O	O-Hydroxytoluene	4.77
4	14.513	C ₈ H ₁₀ O	Benzene ethyl alcohol	3.64
5	28.283	C ₁₆ H ₃₂ O ₂	Hexadecanoic acid	15.98
6	29.48	C ₁₈ H ₃₄ O ₂	9-Octadecanoic acid	39.4
7	29.968	C ₁₈ H ₃₂ O	9,17-Octadecadienal	6.37
8	30.374	C ₂₀ H ₄₀ O ₂	Arachidic acid	2.05
9	30.756	C ₁₆ H ₃₀ O ₂	1,2-15,16-Diepoxyhexadecane	3.29
10	31.494	C ₃₇ H ₇₄ NO ₈ P	*H1,2AHPOEE	4.82

Table 3. Gas Chromatographic-Mass Spectrometric analysis results of Gskolo.

Peak number	Retention time (minutes)	Chemical formula of Compound	Name of Compound	Area (%)
1	3.42	C ₅ H ₁₂ O	1-Butanol	16.52
2	3.75	C ₂ H ₈ O ₂ Si	Dihydroxydimethylsilane	0.72
3	5.04	C ₅ H ₁₀ O ₃	Propanoic acid, 2-hydroxy-ethyl ester	0.05
4	11.09	C ₃ H ₈ O ₃	1,2,3-propanetriol	2.15
5	14.51	C ₉ H ₁₂ O	Benzene ethanol	0.14
6	28.29	C ₁₆ H ₃₂ O ₂	n-Hexadecanoic acid	17.51

Peak number	Retention time (minutes)	Chemical formula of Compound	Name of Compound	Area (%)
7	29.49	C ₁₈ H ₃₄ O ₂	6-Octadecenoic acid	42.43
8	29.75	C ₁₈ H ₃₂ O ₂	9,12-Octadecadienoic acid	10.72
9	30.38	C ₁₈ H ₃₆ O ₄	*PDAHHMEE	1.76
10	30.76	C ₂₀ H ₄₀ O ₂	Eicosanoic acid	3.56
11	31.5	C ₁₆ H ₃₀ O	cis-9-Hexadecenal	4.45

Pentadecanoic acid-2-hydroxy-1-(hydroxymethyl) ethyl ester.

4. Discussion

This work sought to evaluate the presence and levels of volatile organic metabolites and compounds in *Burukutu* (BKT) and *Pito* (PT) and also in *Goskolo* (GSK). GSK is a banned drink owing to its lethal attributes. Ethyl alcohol is the physiologically active component of alcoholic beverages. However, congeners do remain as fractions of brewing. Even if quantitatively small, congeners play a crucial role in the social use and of the alcohol abuse. They may be volatile compounds such as alcohols, acids, aldehydes, ketones and esters. Carbohydrates, tannins, phenols, metals, colouring agents, minerals, histamines and pharmacologically active components constitute other components. [5]. Ethyl acetate, ethyl formate, methyl alcohol, acetaldehyde, and the small aliphatic alcohols (n-propyl alcohol, isobutyl alcohol, n-butanol) constitute volatile congeners that make up the major volatile congener content of beers, wines and distilled spirits.

Consumption of native beers is common among residents of the area of study because they are relatively cheaper, culture friendly and readily available compared to branded factory-based lager beers. [4] reported that BKT samples obtained from B/Ladi contained trifluoroacetic acid, 2-methyl nonadecane and 2-tetradecen-1-ol. PT contained dichloroacetic, pentadecanoic acid, 1-hexadecanol, 1-tetracosanol, methyl tetradecanoate.

Results (% relative abundance), from this study showed the significant presence in BKT of 6-Octadecanoic acid (38.51), Hexadecanoic acid (20.92), Isopentyl alcohol (12.27), 1,11,13-Octadecatriene (11.60) among others. Isopentyl alcohol is a toxic compound. It targets the central nervous system (CNS), respiratory system, eye and skin. It is a product of gut microbial fermentation. It is the major higher chain alcohol in alcoholic beverages. It induces expression of CYP3A and CYT2E1 in human liver [6]. Another metabolite in BKT is Linolenic acid (Octadecatriene). It is particularly implicated in causing cardiovascular disease [7]. Although retention times of compounds may be similar or the same, it is not a fixed property of a component but a result of a system which contributes to the Retention Time (RT). RT does not guarantee a conclusive identification of a compound present in a sample. However, it gives a more reliable insight into the identity of metabolites when coupled with (GC-MS).

There were no metabolites in BKT with identical retention times, albeit some were very similar. For example, 6-Octadecanoic acid with retention time (RT) of 29.53 and 1,11,13-Octadecatriene with RT of 29.79. So also Arachidic

acid with RT of 30.78 and Hexadecanoic acid, 1-[[[(2-aminoethoxy) hydroxyphosphinyl]oxy]methyl]-1,2-ethanediyl ester with RT of 30.38.

In PT, retention times of most of the metabolites were generally similar, ranging between 29.48 to 30.76. Metabolites present in it included 9-Octadecanoic acid, 9,17-Octadecadienal, Arachidic acid, Carbonic acid methylpentyl ester, Benzene ethyl alcohol, 2,3-butanediol, O-Hydroxytoluene and 1,2-15,16-diepoxy hexadecane. Carbonic acid methylpentyl ester causes behavioral analgesia; attacks a segment of the central nervous system (CNS) including aquatic life with long side effects. PT is the major cause of illness among drinkers. It darkens the skin, and causes oedema and weakness. Its ethyl alcohol content is less than that in BKT [8].

As for GSK, the range of RT was between 29.29 and 31.38 (2.09). Some compounds present include 6-Octadecanoic acid, 9,12-Octadecanoic acid, Benzene alcohol, Pentadecanoic acid-2-hydroxy-1-(hydroxymethyl) ethyl ester, 1,2,3-propanetriol and Hexadecanoic acid. Hexadecanoic acid causes the generation of ketone bodies in excess of acetyl-coA, inhibits mouse pronuclear and 2-stage embryo development [9]. It raises the level of low density lipoprotein more than other saturated fats. High consumption of palmitate predisposes to cardiovascular disease. One unique metabolite in GSK is 1-Butanol. It is implicated in eye skin, central nervous system ailments [10]. Benzene ethyl alcohol is toxic and lethal to hepatocytes whose toxicity is not mitigated by ethyl alcohol [11]. Chronic ethanol ingestion exacerbates benzene myelotoxicity and reduces the urinary excretion of benzene metabolites in CD-1 mice [12].

BKT, PT and GSK are composed of diverse metabolites/compounds, most of which are organic acids. Among the major metabolites in all the drinks, some had identical RTs. In BKT, Hexadecanoic acid, 1-[[[(2-aminoethoxy) hydroxyphosphinyl]oxy]methyl]-1,2-ethanediyl ester had RT of 30.38, which is the same as Pentadecanoic acid-2-hydroxy-1-(hydroxymethyl) ethyl ester in GSK. Further, Eicosanoic acid in GSK had identical RT of 30.76 with 1,2-15,16-diepoxyhexadecane in PT. Arachidic acid in BKT had the same RT as cis-9-hexadecenal in GSK with RT value of 31.51.

In normal homeostasis, there is a need for a balanced state of equilibrium. The physiological pH for humans is 7.4. Since the living system is dynamic, there is a requirement to control the redox status (acidity and alkalinity) of cells. This is achieved by using appropriate buffers which help to

reasonably control the pH over a narrow range of changes compatible with life. Metabolic acidosis is a disturbance in the homeostasis of acidity of the plasma. If serum hydrogen ion concentration increases owing to processes, then it is a distinct acidosis. Acidosis is either respiratory which involves some changes in carbon iv oxide or metabolic which is influenced by bicarbonate. Metabolic acidosis increases the hydrogen ion concentration in the body particularly systemic circulation. Increased production of acid, acid ingestion, renal or gastrointestinal bicarbonate losses and decreased excretion of acid constitute etiologies of metabolic acidosis [13].

To be neutral, number of cations must balance anions. Sodium is the primary cation in the plasma. So, it is balanced by the sum of the bicarbonate and chloride ions which represents the anion gap. However, lactate and acetoacetate are unmeasured anions that are the main contributors to metabolic acidosis. Anion gap metabolic acidosis is frequently due to anaerobic metabolism and lactic acid accumulation. If the lungs are not able to remove carbon iv oxide produced, respiratory acidosis ensues. As a consequence, the pH of the blood and other fluids decrease making them too acidic and incompatible to homeostasis of the cell. In this work, at least 75% of compounds and metabolites present in samples of BKT, PT and GSK are organic acids. Hence, the pH of the blood, plasma, serum and other body fluids of drinkers reduce. As a result, drinkers are at risk of metabolic acidosis and its consequences. Mean amount of native beers consumed in a day by males and females drinkers in the area of study stands at 3,745dm³ and 2,946dm³ respectively [8].

Some compounds and metabolites in BKT and PT could be of benefit to the body. BKT contains Octadecanoic acid (Stearic acid) lowers the concentration of low density lipoprotein (LDL) cholesterol relative lauric and palmitic acids. LDL is harmful to the body because of its low density, which makes it to easily deposit on the inner surface of the blood vessels thereby reducing their diameter. This leads to artherosclerosis; a lethal disease condition that overworks the heart. BKT contains vitamins, iron, magnesium, manganese, phosphorus, calcium, 26.7 g starch, and 5.9 g of protein per liter [14]. Eicosanoic acid is a component of oily fruits such as coconut but in small quantity [15]. In addition, cis-9-hexadecanal has potential antimelanogenic antifungal attributes. It targets cell wall organisation, critical Growth Factor & virulence in *Aspergillus fumigatus* [16]. It contains almost all essential amino acids in required proportions except cystine and tryptophan which have been completely destroyed by heat during boiling [17].

PT is particularly enriched with nutrient elements such as Cl, Mg, and K. Apart from serving as an inebriating drink, it is important in fulfilling cultural norms and values including such as marriages, naming and burial ceremonies, parties, and other social gatherings [18]. The elemental composition in this native beverage is crucial due to its nutritional value especially mineral elements as well as estimation of toxic

levels [18]. PT can serve as functional food, owing to its antioxidant capabilities in addition to its gross energy content [19].

As for GSK, there has never been documented evidence about its beneficial attributes to the body. Consequently, its preparation and consumption has been banned by Plateau State Government, Nigeria.

5. Conclusion

It is concluded that BKT, PT, and GSK are composed of compounds and metabolites that are organic in nature. About 83% of them are organic acids, making the drinks acidic. This predisposes drinkers to metabolic or respiratory acidosis, due to drop in pH. Drinkers are exposed to the toxicities of metabolites albeit BKT and PT contain beneficial mineral elements and other nutrients. Other components of the samples include ketones, different types of alcohols, aldehydes, derivatives of benzene and esters.

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