Chemical Composition and Bioavailability of Zinc and Iron in Kunu-zaki, a Nigerian Traditional Beverage

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Abstract. The effect of processing (germination and fermentation) on the chemical composition (proximate, mineral and phytochemical contents) of a Nigerian traditional beverage, kunu-zaki made from sorghum (Sorghum bicolor) grains with the addition of sweet potato (Ipomoea batatas) paste were studied. The bioavailability of Zn and Fe in the beverage were also assessed using phytate, Zn, Ca and Fe molar ratios. Processing of sorghum into kunu-zaki significantly (p<0.05) increased its ash (26.32 %), Na (21.28 %), Ca (20.59 %), Fe (21.62 %), Zn (13.43 %), flavonoids (11.11 %) and alkaloids (30.00 %) contents, but decreased its protein (-43.75 %), fiber (-28.57 %), phenols (-43.80 %), saponin (-62.6 %), tannin (-43.80 %), oxalate (-33.33 %) and phytate (-60.27 %) contents. Addition of sweet potato paste to kunu-zaki apparently aided in enhancing its chemical composition. The combination of germination and fermentation as processing techniques were better than germination alone in significantly (p<0.05) improving the bioavailability of Fe and Zn, and reducing the anti-nutrient content of kunu-zaki, in comparison with both raw and germinated sorghum grains. In conclusion, the kunu-zaki had low contents of protein and fiber, adequate arrays of other functional nutrients and potentially high bioavailability of Fe and Zn.

1.0 Introduction

Kunu-zaki is an African traditional non-alcoholic beverage that has been popular in Nigeria for decades. It is produced by traditional fermentation of cereal grains by beneficial microbes, and has been widely known for its high acceptability, palatability and energy replenishing potentials, which is claimed to be related to its rich content of various functional components. Kunu-zaki, a product of lactic acid fermentation [1], is commonly prepared locally by women, and taken mainly by low and middle income earners for its thirst quenching and energy-giving properties. It is also used locally for entertainment at homes and during festivities such as Sallah and Christmas ceremonies [2]. Depending on the major cereal used in kunu-zaki production, the common types of the beverage include kunun zaki, kunun gyada, kunun akamu, kunun tsamiya and kunun baule [3]. Although, cereals can be used in composite form in the production of kunu-zaki, sorghum (Sorghum bicolor) and millet (Pennisetum americanum) grains are the most commonly used raw materials [2]. However, maize (Zea mays), rice, acha (Digitaria exilis), guinea corn and other grains may also be used [1].

The traditional production process involves steeping of the chosen cereal or mixture of the cereals in water for 24 – 72 hours and wet milling with the aid of a local grinding machine. The steeped and washed grains are usually ground with species such as ginger, clove, red and/or black pepper depending on the taste of the local producer. The slurry obtained is sieved and divided into two unequal portions. A potion (two-third volume) of the slurry is gelatinized with boiling hot water, while to the remaining portion (one-third volume), about an equal volume of sweet potato (Ipomonea batatas) tuber paste, malted rice paste or extract of Cadaba farinose stem is added and mixed. The latter mixture is mixed vigorously and thoroughly with the former, the gelatinized portion while still hot, and allowed to ferment for 24 hours at room temperature. After which, it is
filtered using a local sieve and the filtrate, kunu-zaki, is consumed as a beverage with or without the addition of a sweetener, usually sugar. The beverage can be consumed fresh or bottled and stored at refrigeration temperature [2,4,5].

Cereal grains and their different products form the major source of dietary nutrients for many people, especially those in the developing countries of the world. However, it has been reported that the nutritional quality of cereal grains and their products are inferior due to lower protein content, deficiency of certain essential amino acids, lower protein and starch availabilities, presence of certain anti-nutrients and the coarse nature of the grains [6]. In line with the above report, kunu-zaki has been noted to have a gross chemical composition of 87.85 – 89 % moisture, 9.84 – 12 % carbohydrate, 1.56 – 3 % protein, 0.10 – 0.30 % fat and 0.61 – 0.75 % ash, indicating that the drink is low in protein [2]. Furthermore, like other cereal based products, kunu-zaki may contain high amounts of anti-nutrient that may affect nutrient availability. Anti-nutrients found in cereals, such as phytate, can decrease the absorption of minerals such as Zn, Ca, Fe and Mn, and high intake of these anti-nutrients might lead to mineral deficiency. Zinc and iron are essential trace elements in human nutrition, whose deficiencies are among the common nutritional problems affecting the world. Their deficiencies are of major concern because of the serious health consequences they have, as well as the large number of people afflicted; especially in developing countries of the world. These deficiencies may be caused by the presence of certain anti-nutrients, especially the ability of phytate to reduce dietary Zn and Fe bioavailability by formation of insoluble mineral chelates at physiological pH. The formation and stability of these chelates depend on the relative concentrations of Zn, Fe and phytate as well as on the levels of dietary Ca present [7,8]. However, certain traditional food processing techniques, either individually or in combination, such as germination, steeping, fermentation, and boiling of grains for a limited period have been reported to cause increased activities of hydrolytic enzymes, improvement in the contents of certain essential amino acids, total sugars, B-group vitamins, and a decrease in dry matter, starch and anti-nutrients [9]. Thus, the present study is aimed at determining the effect of fermentation and germination, and the addition of sweet potato tuber paste on the chemical composition and bioavailability of Zn and Fe in kunu-zaki drink.

2.0 Materials and Methods

2.1 Collection of materials

The sorghum (Sorghum bicolor) grains and fresh sweet potato (Ipomoea batatas) tubers were purchased from Mami-Market, Obinze in Owerri-West Local Government Area of Imo State, and authenticated at the Department of Crop Science, Federal University of Technology, Owerri, Nigeria.

2.2 Laboratory Production of Kunu-zaki

The Kunu-zaki was produced by a modified method (Figure 1), based on the traditional method of production as described by [2,5]. In brief, 500 g of apparently healthy sorghum grains were washed and steeped in 1000 ml of water for 24 hours at room temperature (30°C). The steeped grains were drained of water, covered with moist paper-bag for 72 hours at room temperature for germination. Visible root portions of germinated grains were manually removed. After which, they were wet milled using Kenwood major blender (Model Titanium KMO20, Kenwood Chef Major, Giovanni, UK) and sieved with a muslin cloth to obtain a slurry. The paste was then divided into two unequal (two-third and one-third) portions. To the larger portion, 3 liters of boiling hot water was added and turned vigorously until a gelatinized paste was obtained. To the smaller portion, 10 g of washed and ground fresh sweet potato paste was added. The gelatinized sample was allowed to cool to about 50°C, and the later mixture was added to it and stirred vigorously but thoroughly for about 5 min. The combined mixture was covered with paper-bag, allowed to ferment for 12 hours and then filtered with the aid of a cloth sieve to obtain an un-sweetened kunu-zaki.
2.3 Chemical Analyses

For all chemical analyses, four different samples were collected during the course of production of the kunu-zaki:

i. Ground but unfermented sorghum paste (raw sorghum sample)
ii. Ground 72 hour germinated sorghum paste (germinated sorghum sample)
iii. Ground fresh sweet potato paste (raw potato sample)
iv. Complete, ready-to-drink un-sweetened kunu-zaki (kunu-zaki sample)

2.3.1 Proximate analyses

Crude fat was extracted by the soxhlet method with petroleum ether (40-60°C) for 8 hours and then determined along with moisture, total ash and crude fibre of the samples by the methods of [10]. Nitrogen was determined by the micro-Kjeldahl method and the crude protein content was calculated as N x 6.25. Carbohydrate was determined by difference. The energy values were calculated using the Atwater system as described by [11]. All the proximate results were reported as g/100g dry weight and based on triplicate determinations.

2.3.2 Mineral Analyses

A portion, 2.0 g of each sample was incinerated in a Carbolite furnace (Carbolite, Derbyshire, UK) at 600°C for 3 hours to constant weight. The ashed samples obtained were allowed to cool and each was transferred into a separate 50 ml beaker, with the crucible washed with 25ml of 6N HCl into the corresponding beaker. The beaker was then heated to boiling to break the ash. The solution was carefully filtered and transferred into a 50 ml standard flask and made up to the mark with distilled-deionized water. The resulting extract was used for the determination of sodium and potassium concentrations using a flame photometer (Model 405, Corning, Halstead Essex, UK), while
Calcium, magnesium, iron and zinc contents were determined by the use of an atomic absorption spectrophotometer (Alpha 4, Chem. Tech. Analytical, England). Phosphorous was determined as phosphate by the Vanadomolybdate colorimetric method [12]. All determinations were made in triplicates.

2.3.3 Phytochemical Analyses

The percentage compositions (g/100g) of saponins, tannins, alkaloids, flavonoids, oxalate and cyanogenic glycosides were determined according to the methods described by [13,14]. The amount of total phenols in the samples were determined using the Folin-Ciocalteau method [15]. Phytate was extracted according to the procedure described by [16]. Briefly, 1.0 g of each sample was extracted with 3% trichloroacetic acid (TCA) at 37°C for 45 min with simple shaking followed by centrifugation and extraction using anion exchange column. The extracted solution (0.2 ml) was mixed with 4.6 ml of distilled water and 0.2 ml of chromogenic solution and the tubes were heated in a water bath at 95°C for 30 min, and allowed to cool. The developed colour was read at 830 nm against blank. Standard phytate solution was prepared by dissolving sodium phytate in distilled water to prepare different phytate concentrations, which were treated as described above for the tested samples. The amount of phytate in the tested samples was expressed as mg phytate/100 g sample. All determinations were made in triplicates.

2.4 Calculated mineral and molar ratios

Ca/P, Na/K, Ca/Mg and the miliequivalent ratio of [K/(Ca+Mg)] were calculated as described by [17], while the molar ratios [Phy]:[Fe], [Phy]:[Zn], [Ca]:[Phy] and [Ca][Phy]:[Zn] were computed as described by [8].

2.5 Statistical Analysis

All determinations were carried out in triplicates and expressed as mean ± standard deviations. One-way analysis of variance (ANOVA) and post-hoc Tukey test were used to analyse data generated with the aid of GraphPad Prism version 5.3. Differences between means at p≤0.05 were considered statistically significant.

3.0 Results and Discussion

3.1 Changes in proximate composition of germinated sorghum and kunu-zaki.

Table 1 shows the proximate composition of the raw sorghum, raw potato, germinated sorghum and kunu-zaki beverage. Their carbohydrate content was generally high with values ranging from 65.10 g/100g (potato) to 69.02 g/100g (kunu-zaki), while the protein, lipid, ash and fiber contents were generally low. The energy values were high with a range of 1.67 x 10⁴ kJ/kg (germinated sorghum) to 1.78 x 10⁴ kJ/kg (potato). A further study of the Table 1 shows that addition of raw potato paste to the kunu-zaki did not significantly affect its proximate composition in comparison with the effect of the basic composite (raw sorghum), except for the energy values, where that of the raw potato was non-significantly (p>0.05) similar to that of kunu-zaki, but significantly (p<0.05) higher than those of both germinated and raw sorghum. This observation could be directly attributed to the significantly (p<0.05) higher lipid content of potato than those of both the raw and germinated sorghum.
Table 1: Proximate composition (g/100g) of raw sorghum, raw potato, germinated sorghum and kunu-zaki.

<table>
<thead>
<tr>
<th>Proximate</th>
<th>Raw sorghum</th>
<th>Raw potato</th>
<th>Germinated sorghum</th>
<th>Kunu-zaki</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carbohydrate</td>
<td>68.46 ± 1.38a</td>
<td>65.10 ± 2.01a</td>
<td>68.10 ± 1.23a</td>
<td>69.02 ± 2.00a</td>
</tr>
<tr>
<td>Protein</td>
<td>5.60 ± 0.81a</td>
<td>1.23 ± 0.12b</td>
<td>3.34 ± 0.24c</td>
<td>3.15 ± 0.15c</td>
</tr>
<tr>
<td>Lipid</td>
<td>11.75 ± 1.09a</td>
<td>17.66 ± 1.80b</td>
<td>12.37 ± 1.32a</td>
<td>14.20 ± 1.13ab</td>
</tr>
<tr>
<td>Moisture</td>
<td>8.27 ± 0.97a</td>
<td>10.09 ± 1.03ab</td>
<td>11.45 ± 0.98b</td>
<td>8.47 ± 0.87a</td>
</tr>
<tr>
<td>Ash</td>
<td>1.71 ± 0.33ab</td>
<td>2.22 ± 0.13a</td>
<td>1.54 ± 0.18b</td>
<td>2.16 ± 0.16c</td>
</tr>
<tr>
<td>Fiber</td>
<td>4.20 ± 0.51a</td>
<td>3.70 ± 0.32ac</td>
<td>3.20 ± 0.27bc</td>
<td>3.00 ± 0.17bc</td>
</tr>
<tr>
<td>Energy (x 10^4 kJ/kg)</td>
<td>1.69 ± 0.06ab</td>
<td>1.78 ± 0.02a</td>
<td>1.67 ± 0.03b</td>
<td>1.75 ± 0.01ab</td>
</tr>
</tbody>
</table>

Values are mean ± standard deviations of triplicate determinations. Values with similar superscripts per row are not significantly (p<0.05) different.

From Figure 2, it could be seen that germination and fermentation as traditional methods of processing raw sorghum grains into kunu-zaki non-significantly (p>0.05) increased its carbohydrate content (0.82 %), but significantly (p<0.05) reduced its protein (-43.75 %) and fiber (-28.57 %) contents. Although, this is in agreement with earlier reports of low protein content (1.56 – 3 %) in kunu-zaki [2,18], but the low levels of fiber and protein observed could be deleterious, since proteins are crucial for adequate body cells building and maintenance, while fiber is necessary for bowel movement and easy digestion of food.

![Figure 2: Percentage changes in the proximate compositions of germinated sorghum and Kunu-zaki drink in comparison with those of the raw sorghum sample.](image)

3.2 Changes in mineral composition of germinated sorghum and kunu-zaki.

The mineral composition of the samples studied is shown in Table 2. Potassium (K) and sodium (Na) are major macro-elements, while phosphorus (P) is the highest micro-element present in the samples. There were significantly (p<0.05) more minerals in the adjunct (raw sweet potato) than the main ingredient (sorghum) in kunu-zaki production. Sweet potato is one of the vegetable crops that is largely neglected but has been reported to contain essential mineral nutrients, vitamins and functional phytochemicals [19]. The presence of these essential minerals in potato may have contributed to the significantly (p<0.05) higher levels of these minerals, including zinc (Zn) and iron (Fe), in the kunu-zaki than the germinated sorghum; when compared with their baseline values.
in the raw sorghum (Figure 3). Furthermore, a combination of the two processing techniques of germination and fermentation in kunu-zaki production, unlike in the germinated sample only, may have aided in the significantly (p<0.05) high increase in the mineral content of kunu-zaki than in germinated sorghum. Germination and fermentation are traditional methods used in the processing of cereals and legumes, which have recently been found and documented to increase their palatability and nutritional value via increased activity of endogenous enzymes which breakdown large macromolecules to smaller units such as starch to sugars, and also degrade anti-nutrients, thereby liberating chelated minerals and increasing their availability [7,8].

Table 2: Mineral composition (mg/100g) of raw sorghum, raw potato, germinated sorghum and kunu-zaki.

<table>
<thead>
<tr>
<th>Mineral</th>
<th>Raw sorghum</th>
<th>Raw potato</th>
<th>Germinated sorghum</th>
<th>Kunu-zaki</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Macrominerals</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sodium</td>
<td>446.50 ± 30.30\textsuperscript{a}</td>
<td>484.50 ± 22.11\textsuperscript{bc}</td>
<td>522.50 ± 16.53\textsuperscript{bc}</td>
<td>541.50 ± 23.85\textsuperscript{b}</td>
</tr>
<tr>
<td>Potassium</td>
<td>750.50 ± 61.11\textsuperscript{a}</td>
<td>798.10 ± 50.72\textsuperscript{a}</td>
<td>722.00 ± 48.21\textsuperscript{a}</td>
<td>769.50 ± 50.56\textsuperscript{a}</td>
</tr>
<tr>
<td>Magnesium</td>
<td>2.55 ± 0.06\textsuperscript{ac}</td>
<td>2.80 ± 0.09\textsuperscript{b}</td>
<td>2.45 ± 0.04\textsuperscript{a}</td>
<td>2.65 ± 0.09\textsuperscript{bc}</td>
</tr>
<tr>
<td>Calcium</td>
<td>1.70 ± 0.08\textsuperscript{a}</td>
<td>1.95 ± 0.03\textsuperscript{bc}</td>
<td>1.80 ± 0.05\textsuperscript{ac}</td>
<td>2.05 ± 0.08\textsuperscript{b}</td>
</tr>
<tr>
<td><strong>Microminerals</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Phosphorus</td>
<td>66.60 ± 1.18\textsuperscript{a}</td>
<td>69.70 ± 2.02\textsuperscript{ab}</td>
<td>71.74 ± 1.71\textsuperscript{b}</td>
<td>70.72 ± 1.38\textsuperscript{ab}</td>
</tr>
<tr>
<td>Iron</td>
<td>1.85 ± 0.03\textsuperscript{a}</td>
<td>2.15 ± 0.07\textsuperscript{b}</td>
<td>1.95 ± 0.03\textsuperscript{a}</td>
<td>2.25 ± 0.05\textsuperscript{b}</td>
</tr>
<tr>
<td>Zinc</td>
<td>3.35 ± 0.11\textsuperscript{a}</td>
<td>3.70 ± 0.14\textsuperscript{bc}</td>
<td>3.45 ± 0.09\textsuperscript{ac}</td>
<td>3.80 ± 0.10\textsuperscript{b}</td>
</tr>
</tbody>
</table>

Values are mean ± standard deviations of triplicate determinations. Values with similar superscripts per row are not significant (p<0.05) different.

Figure 3: Percentage changes in the mineral compositions of germinated sorghum and kunu-zaki drink in comparison with those of the raw sorghum sample.
3.3 Changes in phytochemical composition of germinated sorghum and kunu-zaki.

As earlier inferred, anti-nutrients appreciably reduce palatability and nutritional value of plant-based foods that contain them. Although, most anti-nutrients are phytochemicals, but not all phytochemicals are anti-nutrients. Certain phytochemicals such as flavonoids are functional compounds that may act as antioxidants. Others like alkaloids and saponins are major sources of plant-based drugs, while oxalates and phytates are majorly described as anti-nutrients. The contents of these anti-nutrients in particular, and phytochemicals in general, have been reported to be significantly reduced by traditional methods of food processing such as soaking, boiling, fermentation, germination, dehydration, and cooking [7,8].

Table 3: Phytochemical composition (%) of raw sorghum, raw potato, germinated sorghum and kunu-zaki.

<table>
<thead>
<tr>
<th>Phytochemical</th>
<th>Raw sorghum</th>
<th>Raw potato</th>
<th>Germinated sorghum</th>
<th>Kunu-zaki</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flavonoid</td>
<td>4.50 ± 0.08a</td>
<td>3.00 ± 0.04b</td>
<td>4.50 ± 0.11a</td>
<td>4.05 ± 0.06c</td>
</tr>
<tr>
<td>Phenols</td>
<td>12.26 ± 1.03a</td>
<td>16.13 ± 1.00b</td>
<td>5.63 ± 0.80c</td>
<td>6.89 ± 1.07c</td>
</tr>
<tr>
<td>Alkaloid</td>
<td>5.00 ± 0.18a</td>
<td>3.00 ± 0.05b</td>
<td>4.50 ± 0.10c</td>
<td>6.50 ± 0.11d</td>
</tr>
<tr>
<td>Saponin</td>
<td>1.50 ± 0.05a</td>
<td>1.56 ± 0.01a</td>
<td>1.50 ± 0.01a</td>
<td>0.56 ± 0.00b</td>
</tr>
<tr>
<td>Tannin</td>
<td>1.76 ± 0.08a</td>
<td>1.58 ± 0.03b</td>
<td>1.60 ± 0.02b</td>
<td>1.23 ± 0.03c</td>
</tr>
<tr>
<td>Oxalate</td>
<td>3.00 ± 0.10a</td>
<td>4.60 ± 0.08b</td>
<td>2.40 ± 0.01c</td>
<td>2.00 ± 0.02d</td>
</tr>
<tr>
<td>Cyanogenic glycoside</td>
<td>2.72 ± 0.12ab</td>
<td>3.09 ± 0.11b</td>
<td>2.40 ± 0.03c</td>
<td>2.93 ± 0.06b</td>
</tr>
<tr>
<td>Phytate (mg/100g)</td>
<td>621.95 ± 3.18a</td>
<td>1479.08 ± 5.81b</td>
<td>553.79 ± 5.37c</td>
<td>247.07 ± 2.13d</td>
</tr>
</tbody>
</table>

Values are mean ± standard deviations of triplicate determinations. Values with similar superscripts per row are not significantly (p<0.05) different.

The results of the present study (Table 3) show that there were significantly (p<0.05) more array of phytochemicals in the sweet potato tuber than in the sorghum cereal. This observation corroborates the report that sweet potato is an excellent source of anti-oxidative polyphenolics, among them anthocyanins and phenolic acids; but that oxalic acid poses a problem when using sweet potato as food [20]. The food content of oxalate, along with other anti-nutrients are usually eliminated or drastically reduced during processing. This may explain the significant (p<0.05) reductions in the concentrations of almost all the phytochemicals in both the germinated sorghum and kunu-zaki. The reductions being more pronounced in the kunu-zaki than the germinated sorghum (Figure 4); notwithstanding the addition of the raw sweet potato paste. For instance, phytate and oxalate were reduced by 10.96 % and 20.00 % in germinated sorghum, and 60.27 % and 33.33 % in kunu-zaki, respectively (Figure 4). This may indicate a synergism between germination and fermentation in the activation of anti-nutrient-degrading enzymes. These process combinations may offer more practical advantage over germination or fermentation alone, since it was reported that long germination periods are needed to improve mineral bioavailability through germination of sorghum to enhance phytate-degrading enzyme activity [7]. The reason for the significant increases in the flavonoid and alkaloid contents of the kunu-zaki, which are apparently beneficial because of their functional potentials may not easily be deduced. However, they may have increased due to the addition of sweet potato paste, but because other phytochemicals did not, the reason may not be foolproof, and thus calls for more research for further clarification.
Flavonoids Phenols Alkaloids Saponin Tannins Oxalate Cyanogenic glycoside Phytate (mg/100g)
-70 -60 -50 -40 -30 -20 -10 0 10 20 30 40
Germinated sorghum
Kunun-zaki drink
Percentage Change in Concentration

Figure 4: Percentage change in the phytochemical compositions of fermented sorghum and kunu-
zaki drink in comparison with those of the raw sorghum sample.

3.4 Changes in computed mineral ratios, and Fe, Zn, Ca and Phytate molar ratios of
germinated sorghum and kunu-zaki.

The computed mineral ratio of the samples are shown in Table 4. The Ca/P ratio was generally
lower than 0.5 which was reported to be the minimum ratio required for favourable absorption of
Ca in the intestine [8]. Calcium is an important constituent of body fluids and is required along with
phosphorus for bone formation. Thus, the levels of Ca/P ratios in both the raw and processed
samples indicated that consumption of kunu-zaki as the sole sources of Ca and P may not enhance
strong bone development and maintenance, especially in children, since absorption under this
condition would not be adequate. The Na/K ratios, were greater than 0.60, a value reported not to
favour enhancement of high blood pressure in man [17], thus adequate for maintenance of
homeostatic localization of Na and K in the extracellular and intracellular compartments of the
body, respectively. Interestingly, the production of kunu-zaki significantly (p<0.05) increased this
ratio in comparison with those of the raw samples. On the other hand, the Ca/Mg ratio which ranged
from 0.67 ± 0.05 (raw sorghum) to 0.77 ± 0.06 (kunu-zaki) were lower than the recommended value
of 1.0 [17]. Similarly, the calculated milliequivalent ratios of [K/(Ca+Mg)] were all less than 2.2 as
reported by [17]. The results buttress the reported low mineral nutrient content of sorghum and
kunu-zaki, and calls for further fortification of kunu-zaki with other mineral nutrient rich sources.

Table 4 also shows the Fe, Zn, Ca and phytate (Phy) molar ratios of the samples. Phytate/Fe and
phytate/Zn molar ratios are associated with Fe and Zn bioavailability. The results show that the
phytate/Fe molar ratios ranged from 9.35 ± 0.05 in kunu-zaki to 31.06 ± 0.06 in raw potato, while
the phytate/Zn molar ratio ranged from 6.45 ± 0.05 in kunu-zaki to 20.70 ± 0.08 in raw potato. It
has been reported that ideal phytate/Fe molar ratio should be lower than 14, which is the critical
value above which Fe bioavailability is strongly impaired [21]. Similarly, it was showed that foods
with phytate/Zn molar ratios less than 10 encourage the bioavailability of Zn, but problems may
arise when the values are greater than 15 [22]. Our results proved that the combined processes of
germination and fermentation of sorghum into kunu-zaki significantly (p<0.05) reduced the
phytate/Fe and phytate/Zn molar ratios of sorghum to values (9.35 ± 0.05 and 6.45 ± 0.05,
respectively) within the acceptable limit. The above observations re-enforce previous results that
showed that the bioavailability of Zn and Fe in cereals and legumes are low but are generally
improved by various processing techniques [7,8,23].
Table 4: Computed mineral ratios, and Fe, Zn, Ca and Phytate molar ratios of raw sorghum, raw potato, germinated sorghum and kunu-zaki.

<table>
<thead>
<tr>
<th>Ratio</th>
<th>Raw sorghum</th>
<th>Raw potato</th>
<th>Germinated sorghum</th>
<th>Kunu-zaki</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Mineral ratios</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ca/P</td>
<td>0.03 ± 0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.03 ± 0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.03 ± 0.00&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.03 ± 0.00&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Na/K</td>
<td>0.60 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.61 ± 0.04&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.72 ± 0.02&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.70 ± 0.01&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ca/Mg</td>
<td>0.67 ± 0.05&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.70 ± 0.03&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.74 ± 0.08&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.77 ± 0.06&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>[K/(Ca+Mg)]&lt;sup&gt;a&lt;/sup&gt;</td>
<td>8.83 ± 0.12&lt;sup&gt;a&lt;/sup&gt;</td>
<td>8.40 ± 0.09&lt;sup&gt;bc&lt;/sup&gt;</td>
<td>8.49 ± 0.10&lt;sup&gt;b&lt;/sup&gt;</td>
<td>8.19 ± 0.08&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td><strong>Molar ratios</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>[Phy]:[Fe]</td>
<td>28.56 ± 0.09&lt;sup&gt;a&lt;/sup&gt;</td>
<td>31.06 ± 0.06&lt;sup&gt;b&lt;/sup&gt;</td>
<td>24.00 ± 0.10&lt;sup&gt;c&lt;/sup&gt;</td>
<td>9.35 ± 0.05&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>[Phy]:[Zn]</td>
<td>18.47 ± 0.07&lt;sup&gt;a&lt;/sup&gt;</td>
<td>20.70 ± 0.08&lt;sup&gt;b&lt;/sup&gt;</td>
<td>15.83 ± 0.04&lt;sup&gt;c&lt;/sup&gt;</td>
<td>6.45 ± 0.05&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>[Ca]:[Phy]</td>
<td>0.05 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.04 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.05 ± 0.02&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.14 ± 0.03&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>[Ca][Phy]/[Zn]</td>
<td>0.78 ± 0.06&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.01 ± 0.04&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.71 ± 0.09&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.34 ± 0.01&lt;sup&gt;c&lt;/sup&gt;</td>
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</table>

<sup>a</sup>Milliequivalent ratio. Values are mean ± standard deviations of triplicate determinations. Values with similar superscripts per row are not significantly (p<0.05) different.

Meanwhile, [24] suggested that the solubility of phytate and the proportion of Zn bound in a mineral complex with phytate in the intestine depend on the levels of Ca. Based on this assertion, phytate precipitation is not complete until dietary Ca/phytate molar ratio gets to a value of approximately 6:1. Thus, when Ca/phytate molar ratio is less than 6:1, phytate precipitation is incomplete and some of the dietary Zn remain in solution. By extension, the proportion remaining in solution increases with decreasing Ca/phytate molar ratio [17]. The Ca/phytate molar ratios of our samples are all below the critical ratio, indicating that their Ca content was sufficiently low to promote phytate-induced decreases in Zn bioavailability.

[25] suggested that [Ca][Phy]/[Zn] ratio may be a better predictor of Zn bioavailability than phytate/Zn and Ca/phytate; noting that if the value was greater than 0.50 mol/kg, there would be interferences with the availability of Zn. In our results, the [Ca][Phy]/[Zn] values for raw sorghum, raw potato and germinated sorghum were higher than the proposed cut-off point of 0.50 mol/kg; indicating that Zn would not be adequately available in the samples. However, the combined processes of germination and fermentation caused a significant (p<0.05) reduction in the [Ca][Phy]/[Zn] values of the raw and germinated sorghum to obtain a value of 0.34 ± 0.01 for kunu-zaki, which was lower than the recommended 0.5 mol/kg. The results underpin the synergistic advantage of combining germination and fermentation processes in kunu-zaki production in improving the nutritional quality of the beverage.

### 3.5 Conclusion

In conclusion, this study showed that kunu-zaki is a good source of carbohydrate, energy, minerals and functional phytochemicals, but requires fortification or intake with protein and fiber rich foods. Furthermore, the beverage showed potentially high Fe and Zn bioavailability; notwithstanding the poor bioavailability of these mineral in both raw and germinated sorghum. These were shown to have been significantly improved by the combined processes of germination and fermentation through significant reductions in anti-nutrient contents with concomitant liberation of the essential minerals.
References


