

## Original Research Article

# Biochemical characteristics and Nutritional profile of the stem bark extracts from the red variety of *Byttneria catalpifolia*, an edible wild plant growing in the Western part of Côte d'Ivoire.

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### ABSTRACT

**Aims:** The objective of this work is to characterize the bark extracts of the red variety of *Byttneria catalpifolia*.

**Study design:** Dried bark powder and mucilage extracted from the fresh bark of the red variety of *Byttneria catalpifolia* were used to evaluate biochemical composition, minerals and nutritional profile.

**Place and Duration of Study:** The study was conducted in Laboratory of Biocatalysis and Bioprocesses, Food Sciences and Technology Unit, at Nangui Abrogoua University, between January 2015 and December 2017

**Methodology:** The AOAC method was used to evaluate the biochemical composition in bark and mucilage. The minerals were analysed by a variable pressure scanning electron microscope (SEM), amino acid and organic acid by HPLC and the nutritional profile was determined.

**Results:** The proximate analysis revealed high rates of ashes (11.51 %), crude fibre (50.33 %), reducing sugar (26.37 %), total sugar (44.88 %), caloric energy (135.18 Kcal /100g dw) in Bark powder while the amount of protein (6.01%) and carbohydrate (30.95%) were moderate and that of fat was low. Mucilage showed a content of ashes (3.82 %), total sugar (24.84 %), carbohydrate (87.79 %) and caloric energy (336.19 Kcal /100g dw) whereas the rate of crude fibre (0.5 %), reducing sugar (1.80 %) and fat (0.6 %) found to be low and that protein (7.29 %) was moderate. This study indicated that the both samples contained the amino acids and organic acids. The results also showed the both samples appeared to be good sources of minerals such as K, Ca, P, Mg and Fe.

**Conclusion:** The bark extracts of the red variety of *Byttneria catalpifolia* contain appreciable amounts of nutrients. Their nutritional profile by SAIN and LIM method showed that they belong to the group of foods recommended for health.

**Keywords:** *Byttneria catalpifolia*, mucilage, nutritional profile, vegetable, stem bark

## 1. INTRODUCTION

Many plants are consumed as vegetables [1]. Vegetables are the edible parts of plant that are consumed wholly or in parts, raw or cooked as part of main dish or salad. A vegetable includes leaves, stems, roots, flowers, seed, fruits, bulbs, tubers and fungi [2] [3]. It is worthwhile to note that consumption of numerous types of edible plants as sources of food could be beneficial to nutritionally marginal population especially in developing countries

22 where poverty and climate is causing havoc to the rural populace. Like other plants, Lianas,  
23 the woody climbing plants, have been intensively studied in the tropics [4] [5]. It is well  
24 known that they are an important component of tropical forests where they constitute up to  
25 32 % of the stems and up to 35 % of the woody species diversity [6]. Further, lianas are  
26 more diverse and typically more abundant in tropical than temperate forests, with subtropical  
27 forests being intermediate [7] [8]. Otherwise, the studies concerning the lianas were carried  
28 out on geographical, taxonomical and ecological aspects [9]. Among the liana species,  
29 *Byttneria catalpifolia* is a perennial plant that is widely distributed in the tropics of Africa, Asia  
30 and America. It belongs to the Family Sterculiaceae and it is the most abundant of the genus  
31 *Byttneria*. Based on the colour of the bark, there are two varieties of *Byttneria catalpifolia*.  
32 The white variety is characterized by the white colour of the bark when the skin is scraped  
33 while the red variety, by a purple colour of the bark. It is an edible wild plant whose bark is  
34 used to make a sticky sauce that is well appreciated by the populations of Western Côte  
35 d'Ivoire. However, information on the nutritional composition of this plant species is scarce or  
36 non-existent. Note that, proximate and nutrient analysis of wild edible plants plays a crucial  
37 role in assessing their nutritional significance [10]. The considerable use of wild edible plant  
38 species by the local people in their diet motivated us to carry out the present proximate and  
39 nutrients analysis. In spite of their importance as a food source, to the best of our  
40 knowledge, there are no published studies on the nutritional composition of the stem bark  
41 and the mucilage extracted from fresh bark of *Byttneria catalpifolia*. The mucilage of some  
42 plants is well known to science and has been studied by pharmacologists and found to  
43 possess biologically active principles. Therefore, the present study was designed to evaluate  
44 the biochemical composition and nutritional profile of the dried stem bark powder and the  
45 mucilage extracted from fresh bark of the red variety of *Byttneria catalpifolia* and provides  
46 consumers with the most appropriate mode of consumption which would give them health  
47 benefits.

## 48 49 **2. MATERIAL AND METHODS**

### 50 51 **2.1. Sample collection and preparation**

52 The stem samples of the red variety of *Byttneria catalpifolia* were collected from the western  
53 part of Côte d'Ivoire (7° 24'45 "North latitude and 7 ° 33'13" West longitude). These samples  
54 were immediately transported to the laboratory. After removing the epidermis with a kitchen  
55 knife, the barks were removed from the stems and chopped. The bark sample was divided  
56 into two parts. Part of sample was used for the extraction of mucilage. To do this, 20 g of  
57 fresh stem bark were ground in 200 mL of demineralized water using a crusher (Moulinex).  
58 The homogenate was centrifuged at 4000 rpm for 10 min. The pellet was ground again in  
59 200 mL and centrifuged under the same conditions as those previously described [11]. The  
60 two viscous supernatants were mixed and kept in the freezer for physicochemical analysis.  
61 The other part was dried in an oven at 65°C for 72 h. The dried samples were ground using  
62 a crusher (Moulinex) and the resulting sifted ( $\Phi$  2 mm) powder was stored in sealed plastic  
63 boxes for biochemical analysis.

### 64 65 **2.2. Proximate analysis**

66 Moisture, ash, crude fibre, crude protein, fat and carbohydrate contents were determined in  
67 accordance with the standard methods of the [12]. All determinations were run in triplicates.  
68 The moisture content was determined by the difference of weight before and after drying the  
69 sample (10 g) in an oven (memmert, germany) at 105°C until constant weight at least or 72  
70 h. Ash fraction was determined by the incineration of dried sample (5 g) in a muffle furnace

71 (Pyrolabo, France) at 550°C for 12 h. The residue weight percentage was expressed as ash  
72 content. Crude fibre content was obtained from the loss in weight of dried residue following  
73 the digestion for fat-free samples with 1.25% each of sulfuric acid and sodium hydroxide  
74 solutions. The crude protein content ( $N \times 6.25$ ) was estimated by the macrokjeldahl nitrogen  
75 assay method using a digestion apparatus. The fat content was determined by Soxhlet  
76 extraction using hexane as a solvent. Reducing sugar content was determined by extracting  
77 with 80% neutral aqueous ethanol followed by evaporation of the ethanol and subsequent  
78 measurement using the dinitrosalicylic acid method according to [13]. Total soluble sugar  
79 content in ethanolic extract was assessed using the phenol-sulfuric acid method according to  
80 Dubois et al. (1956). Carbohydrate and calorific values were calculated using the following  
81 formulas [14]:

82 Carbohydrates (dry weight basis):  $100 - (\% \text{ moisture} + \% \text{ proteins} + \% \text{ lipids} + \% \text{ ash} + \% \text{ fibre})$   
83

84 Calorific value (dry weight basis):  $(\% \text{ proteins} \times 2.44) + (\% \text{ carbohydrates} \times 3.57) + (\% \text{ lipids}$   
85  $\times 8.37)$ . The results of ash, crude fibre, crude protein, fat and carbohydrate contents were  
86 expressed on dry weight basis.

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### 88 **2.3. Minerals analysis**

89 The mineral elements were analysed after wet-ashing using the scanning electron  
90 microscope (SEM) with variable pressure (SEM FEG Zeiss Supra 40 VP). This SEM is  
91 equipped with an X-ray detector (Oxford Instruments) connected to an energy diffusion  
92 spectrometry (EDS) microanalyzer platform (Inca Cool Dry, without liquid nitrogen). About 10  
93 mg of the sample ash residue were applied evenly to a primed platform with double-sided  
94 adhesive carbon for analysis. To measure the content of chemical elements, the device  
95 performs a measurement of the transition energy of the electrons from electronic clouds of  
96 the K, L and M series of atoms of the sample.

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### 98 **2.4. Amino acids determination**

99 Amino acids were assayed by reverse phase high performance liquid chromatography (PTC  
100 column RP-18, 220 mm in length, 2.1 mm internal diameter) equipped with a pre-column  
101 (SHIMADZU SPD 20A). The sample was vacuum hydrolysed at 150 °C for 60 min in a  
102 Waters Pico-Tag work station (Waters' associates, Milford, MA, USA) in grade 6 N HCl at  
103 1% phenol. It was then taken up in ultra-pure water and derivatized automatically by means  
104 of an auto-derivator-analyzer-420 (SHIMADZU SPD 20A). The amino acid derivatives in the  
105 form of phenyl isothiocyanates (PITC-amino acids) were separated by chromatography  
106 system consisted of mobile phase A (45 M sodium acetate, pH 5.9) and mobile phase B  
107 (30% 105 mM sodium acetate pH 4.6; 70% acetonitrile) with a linear gradient of 7-36% at  
108 1.5 mL/min. The detection was done at a wavelength of 254 nm and the total duration of the  
109 analysis was 31 min. The acquisition and exploitation of the results were carried out using  
110 the Model 600 Data Analysis System (SHIMADZU SPD 20A) software.

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## 113 2.5. Organic acids determination

114 Organic acids were extracted from 1 g of sample with 50 mL of 80 % methanol saturated  
 115 with NaCl and were analysed according to the method of [15] using a HPLC system  
 116 (Shimadzu Corporation, Japan) consisting of a pump (Shimadzu LC-6A Liquid  
 117 Chromatograph), a UV detector (Shimadzu SPD-6A UV Spectrophotometric detector) and  
 118 an integrator (Shimadzu CR 6A Chromatopac). All separations were carried out in isocratic  
 119 mode with an ICsep ICE ORH-801 ion exclusion column (40 cm x 5 µm, Interchrom, France)  
 120 maintained at 35 °C using a Meta Therm™ furnace (Interchrom, France ). The HPLC  
 121 conditions are shown in the table 1. Organic acid standards (citric, fumaric, malic, oxalic,  
 122 tartaric acid, quinic acid, ascorbic acid, succinic acid and fumaric acid) were obtained from  
 123 Aldrich Co. (Sigma-Aldrich Chemie, Steinheim, Germany). The standard solutions were  
 124 prepared individually at different concentrations with double distilled water. The analysis was  
 125 carried out in triplicate. The levels of the organic acids in the samples is obtained by  
 126 comparing the retention times of the eluted compounds with the retention times of the  
 127 reference solutions.

128 Table 1: HPLC running conditions

Column	ICsep ICE ORH-801 ion exclusion column (40 cm x 5 µm, Interchrom, France)
Mobile phase	Isocratic mode: 0.004N Sulfuric acid, pH (HPLC-Grade)
Flow rate	0.6 mL/min
Oven Temperature	35 °C
UV Detection	Wavelength : 210 nm
Injection Volume	20 µl
Runtime	35 min + 10 min post-run

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## 130 2.6. Determination of B vitamins

131 A 5 g of liquid/powder sample was extracted with 20 mL of methanol (80%). The stock of  
 132 standard (Sigma Aldrich analytical grade reagent) was prepared by dissolving 0.01 g of each  
 133 standard in methanol (80%). A HPLC system (SHIMADZU SPD 20A) equipped with UV  
 134 detector (PAD) and C18 ODS column (250 x 4.6 de, Cluzeau France) was used in isocratic  
 135 mode for analysis. Mobile phase consisted of acetonitrile (55 mL), tetrahydrofurane (37 mL)  
 136 and water (8 mL) at 1.5 mL/min flow rate and 10 µl of each sample/standard were injected  
 137 and monitored at 325 nm wavelength.

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## 140 2.7. Nutritional profile of extracts of the red variety of *Byttneria catalpifolia* 141 according to the SAIN and LIM method

142 To establish the nutritional profiling of *Byttneria catalpifolia* extracts, several successive  
 143 steps have been carried out. First, it was necessary to establish the nutritional value of each  
 144 extract based on the nutrients that are needed to calculate SAIN and LIM. For this purpose,  
 145 it has been necessary beforehand, to establish for each of the extracts, a table of  
 146 composition which details the contents of these nutrients. The nutrient content is expressed

147 based on 100 g of raw material (RM). These data were collected from the French table  
148 Ciqual [16] and the FAO table.

149 The SAIN score of *Byttneria catalpifolia* extracts was calculated using proteins, crude fibre,  
150 calcium, iron and iodine. As for the LIM score, only sodium was used. The extracts of  
151 *Byttneria catalpifolia* contained either saturated fatty acids or added sugars.

$$SAIN = \frac{\frac{Prot}{65} + \frac{Fib}{30} + \frac{Ca}{900} + \frac{Vit.C}{110} + \frac{Fe}{12.5} + \frac{Iod}{150}}{6} \times 100$$

Energy

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$$LIM = \frac{\frac{Na}{3153} + \frac{SFA}{22} + \frac{Added\ sugar}{50}}{3} \times 100$$

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156 Prot : proteins, Fib : fiber, Ca : calcium, Vit. C : vitamin C, Fe : iron, Iod : Iodine, Na : sodium,  
157 SFA : saturated fatty acids

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## 159 2.8. Statistical analysis

160 All analyses were carried out in triplicates. Results were expressed by means of  $\pm$  SD.  
161 Means of proximate composition, amino acid, mineral, and organic acid contents of the dried  
162 bark powder and mucilage of *Byttneria catpifolia* were separated according to The Student's  
163 t-test ( $P < 0.05$ ), with the help of JMP® Pro software (version 12, SAS Institute Inc., Cary, NC,  
164 2007).

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## 166 3. RESULTS

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### 168 3.1. Proximate composition

169 Results of the composition of dried stem bark powder and mucilage extracted from the fresh  
170 bark of the red variety of *Byttneria catalpifolia* are presented in table 2. The pH value of dried  
171 stem bark powder ( $5.95 \pm 0.02$ ) and mucilage ( $5.95 \pm 0.01$ ) was similar. The dry matter  
172 content observed in fresh bark ( $45.03 \pm 0.34$  %) was significantly ( $p \leq 0.05$ ) higher than that  
173 obtained in the mucilage ( $6.21 \pm 1.15$  %). The ash contents obtained in dried stem bark  
174 powder and mucilage were  $11.51 \pm 0.45$  % and  $3.82 \pm 0.62$  % of dry weight (dw),  
175 respectively. There was meaningful difference ( $p \leq 0.05$ ) between ash contents of dried stem  
176 bark powder and mucilage. Our results showed that protein content ( $7.29 \pm 1.01$  % dw) and  
177 caloric energy ( $336.19 \pm 1.18$  kcal /100 g dw) were significantly ( $p \leq 0.05$ ) higher in mucilage  
178 than those observed in dried stem bark powder, with values of  $6.01 \pm 0.20$  % dw and  $135.18$   
179  $\pm 2.26$  kcal /100 g dw, respectively. Crude fibres observed in the dried stem bark ( $50.33 \pm$   
180  $0.22$  dw) was largely higher than that obtained in the mucilage ( $0.50 \pm 0.01$  dw).  
181 Furthermore, this study revealed the presence of parameters such as fat, reducing sugar,  
182 total soluble sugar, vitamin B9 and vitamin B2 in the both samples.

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185 Table 2: Proximate composition of the dried stem bark and mucilage from the fresh bark of  
 186 the red variety of *Byttneria catalpifolia*  
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Parameters	Dried bark powder	Mucilage
pH	5.95 ± 0.02 <sup>a</sup>	5.95 ± 0.01 <sup>a</sup>
Dry matter* (%)	45.03 ± 0.34 <sup>b</sup>	6.21 ± 1.15 <sup>a</sup>
Ashes (% dw)	11.51 ± 0.45 <sup>b</sup>	3.82 ± 0.62 <sup>a</sup>
Fat (% dw)	1.20 ± 0.01 <sup>b</sup>	0.60 ± 0.01 <sup>a</sup>
Proteins (% dw)	6.01 ± 0.20 <sup>a</sup>	7.29 ± 1.01 <sup>a</sup>
Carbohydrates (% dw)	30.95 ± 0.51 <sup>a</sup>	87.79 ± 1.01 <sup>b</sup>
Crude fibres (% dw)	50.33 ± 0.22 <sup>b</sup>	0.50 ± 0.01 <sup>a</sup>
Reducing sugar (mg/100g dw)	26.37 ± 0.05 <sup>b</sup>	1.80 ± 0.08 <sup>a</sup>
Total sugar (mg/100g dw)	44.88 ± 0.16 <sup>b</sup>	24.84 ± 0.16 <sup>a</sup>
Vitamin B2 (mg/100g dw)	694.83 ± 32.23 <sup>b</sup>	102.18 ± 0.61 <sup>a</sup>
Vitamin B9 (mg/100g dw)	31.97 ± 1.37 <sup>b</sup>	2.09 ± 0.01 <sup>a</sup>
Caloric Energy (Kcal/100g dw)	135.18 ± 2.26 <sup>a</sup>	336.19 ± 1.18 <sup>b</sup>

188 *Each value is an average of three replicate*

189 *Values are mean ± standard deviation*

190 *Means not sharing a similar letter in a line are significantly different (p ≤ 0.05) as assessed by the test*  
 191 *of student; dry weight: dw; \*Dry matter of the fresh bark of the red variety of *Byttneria catalpifolia**

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### 193 3.2 Amino acid contents

194 The amino acid compositions of the dried stem bark powder and mucilage from the fresh  
 195 bark of the red variety of *Byttneria catalpifolia* are illustrated in Table 3. These results  
 196 indicated that proline, valine, methionine, arginine, glycine, glutamic acid, tyrosine,  
 197 threonine, lysine and cysteine were found in the both analysed samples. The amino acid  
 198 contents of dried stem bark powder ranged from 0.11 ± 0.02 mg/ 100g dw (threonine) to 3.70  
 199 ± 0.01 mg/ 100g dw (arginine), while those of mucilage varied from 0.02 ± 0.01 mg/ 100g dw  
 200 (arginine) to 7.57 ± 0.31 mg/ 100g dw (cysteine). This study revealed the presence of  
 201 essential amino acids such as valine, methionine, threonine and lysine in both samples.  
 202 Their levels ranged from 0.11 ± 0.02 mg/ 100g dw (threonine) to 1.64 ± 0.01 mg/ 100g dw  
 203 (methionine) for dried stem bark powder, whereas those of mucilage varied from 1.09 ±  
 204 0.01mg/ 100g dw (lysine) to 4.77 ± 0.03mg/ 100g dw (methionine). The statistical analysis  
 205 showed that the essential amino acids contents of dried stem bark powder were lower  
 206 meaningfully (p ≤ 0.05) than those observed in mucilage. Besides, proline, arginine, glutamic  
 207 acid and glycine (non-essential amino acids) contents in dried bark powder were found also  
 208 to be lower significantly (p ≤ 0.05) than those obtained in mucilage, excepted for tyrosine and  
 209 cysteine contents.

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222 Table 3: Amino acid contents of the dried stem bark powder and mucilage from the fresh  
 223 bark of the red variety of *Byttneria catalpifolia*  
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Amino acids (mg/ 100g dw)	Dried stem bark powder	Mucilage
Proline	0.34 ± 0.11 <sup>a</sup>	2.33 ± 0.04 <sup>b</sup>
Valine	0.81 ± 0.28 <sup>a</sup>	1.11 ± 0.06 <sup>a</sup>
Methionine	1.64 ± 0.01 <sup>a</sup>	4.77 ± 0.03 <sup>b</sup>
Arginine	3.70 ± 0.01 <sup>a</sup>	7.57 ± 0.31 <sup>b</sup>
Glycine	0.60 ± 0.01 <sup>a</sup>	0.81 ± 0.06 <sup>b</sup>
Glutamic acid	0.43 ± 0.03 <sup>a</sup>	1.06 ± 0.29 <sup>b</sup>
Threonine	0.11 ± 0.02 <sup>a</sup>	2.28 ± 0.07 <sup>b</sup>
Tyrosine	0.15 ± 0.02 <sup>b</sup>	0.07 ± 0.00 <sup>a</sup>
Cysteine	0.43 ± 0.13 <sup>b</sup>	0.02 ± 0.01 <sup>a</sup>
Lysine	0.17 ± 0.02 <sup>a</sup>	1.09 ± 0.01 <sup>b</sup>

225 *Each value is an average of three replicates*

226 *Values are mean ± standard deviation*

227 *Means not sharing a similar letter in a line are significantly different  $p \leq 0.05$  as assessed by the test of*  
 228 *student; dry weight: dw.*

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### 230 3.3. Mineral composition

231 Table 4 summarizes some mineral composition of the dried stem bark powder and mucilage  
 232 from the fresh bark of *Byttneria catalpifolia*. The minerals detected were magnesium (Mg),  
 233 phosphorus (P), potassium (K), calcium (Ca), sodium (Na), iron (Fe), copper (Cu), Zinc (Zn)  
 234 and iodine (I). The amounts of these minerals in dried stem bark powder appeared  
 235 significantly higher ( $P \leq 0.05$ ) than those of mucilage. Ca ( $414.17 \pm 27.97 - 1104.36 \pm 223.30$   
 236 mg/ 100g dw) and K ( $440.00 \pm 20.93 - 990.69 \pm 227.31$  mg/100g dw) were the abundant  
 237 minerals. The least concentrated macro-elements in the mucilage and stem bark powder  
 238 was Na ( $8.02 \pm 1.27 - 28.72 \pm 17.92$  mg/ 100g dw, respectively). The micro-elements  
 239 observed in the dried stem bark powder were Fe, Cu, Zn and I. Cu and Zn were not detected  
 240 in mucilage. Iodine (I) content of the mucilage ( $6.99 \pm 1.10$  µg /100g dw) was meaningfully  
 241 lower ( $p \leq 0.05$ ) than that obtained in dried stem bark powder ( $14.61 \pm 1.10$  µg /100g dw).  
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243 Table 4: Minerals and mineral ratios of the dried stem bark and mucilage from the bark of the  
 244 red variety of *Byttneria catalpifolia*  
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Minerals (*mg/ 100 g dw)	Dried bark powder	Mucilage	Pellet
Mg	381.56 ± 162.27 <sup>c</sup>	198.47 ± 16.88 <sup>b</sup>	129.24 ± 18.18 <sup>a</sup>
P	38.50 ± 9.46 <sup>c</sup>	35.09 ± 7.36 <sup>b</sup>	8.01 ± 1.57 <sup>a</sup>
K	990.69 ± 227.31 <sup>c</sup>	440.00 ± 20.93 <sup>b</sup>	368.40 ± 3.78 <sup>a</sup>
Ca	1104.36 ± 223.30 <sup>c</sup>	414.17 ± 27.97 <sup>a</sup>	543.79 ± 85.62 <sup>b</sup>
Na	28.72 ± 17.92 <sup>c</sup>	8,02 ± 1.27 <sup>a</sup>	15.24 ± 0.62 <sup>b</sup>
Fe	11.05 ± 5.07 <sup>c</sup>	5.42 ± 2.92 <sup>a</sup>	6.85 ± 0.1 <sup>b</sup>
Cu	0.49 ± 0.02 <sup>b</sup>	ND	0.21 ± 0.01 <sup>a</sup>
Zn	0.46 ± 0.01 <sup>b</sup>	ND	0.14 ± 0.00 <sup>a</sup>
I (µg /100g dw)	14.61 ± 1.10 <sup>b</sup>	6.99 ± 1.10 <sup>a</sup>	13.55 ± 0.37 <sup>b</sup>
[Ca]/[P]	28.68	11.80	67.89
[Ca]/[Mg]	2.89	2.09	4.21

246 *Each value is an average of three replicates*

247 *Values are mean ± standard deviation*

248 *Means not sharing a similar letter in a line are significantly different  $p \leq 0.05$  as assessed by the test of*  
 249 *student; dry weight: dw. Not detected: ND*

250 \* the unit does not apply to ratios.

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### 252 3.4. Quantitative determination of organic acids

253 The organic acids were extracted from the powder of dried stem bark and mucilage of  
 254 *Byttneria catalpifolia*. They were identified and measured using HPLC system. Tannic,  
 255 oxalic, citric, tartaric, sulfanilic, salicylic, adipic, fumaric and benzoic acids were observed. All  
 256 of these organic acids were present in both samples. Table 5 clearly indicates that content  
 257 organic acids contents of the dried bark powder was found to be lower ( $p \leq 0.05$ ) than those  
 258 observed in the mucilage except tannic acid and citric acid. In both samples, levels of  
 259 fumaric and benzoic acids did not vary significantly ( $p > 0.05$ ). In addition, organic acids  
 260 varied from  $0.09 \pm 0.02$  mg /100 g dw to  $51.35 \pm 0.75$  mg /100 g dw and from  $0.11 \pm 0.02$  mg  
 261 /100 g dw to  $55.45 \pm 0.12$  mg /100 g dw for dry bark powder and mucilage, respectively.

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263 Table 5: Organics acids contents of the dried stem bark and mucilage extracted from the  
 264 fresh bark of red variety of *Byttneria catalpifolia*

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Organic acids (*mg/ 100g dw)	Dried bark powder	Mucilage
Tannic acid	$6.57 \pm 0.03^b$	$4.64 \pm 0.31^a$
Oxalic acid	$1.39 \pm 0.36^a$	$5.28 \pm 0.11^b$
Citric acid	$15.63 \pm 0.22^b$	$13.39 \pm 0.30^a$
Tartric acid	$51.35 \pm 0.75^a$	$55.45 \pm 0.12^b$
Sulfanilic acid	$0.15 \pm 0.03^a$	$0.31 \pm 0.01^b$
Salicylic acid	$0.32 \pm 0.03^a$	$2.86 \pm 0.23^b$
Adipic acid	$0.31 \pm 0.02^a$	$0.50 \pm 0.06^b$
Fumaric acid	$0.23 \pm 0.07^a$	$0.25 \pm 0.02^a$
Benzoic acid	$0.09 \pm 0.02^a$	$0.11 \pm 0.02^a$
[oxalic acid]/[Ca]	0.001	0.061

266 Each value is an average of three replicates

267 Values are mean  $\pm$  standard deviation

268 Means not sharing a similar letter in a line are significantly different  $p \leq 0.05$  as assessed by the test of  
 269 student; dry weight: dw. \* the unit does not apply to ratios.

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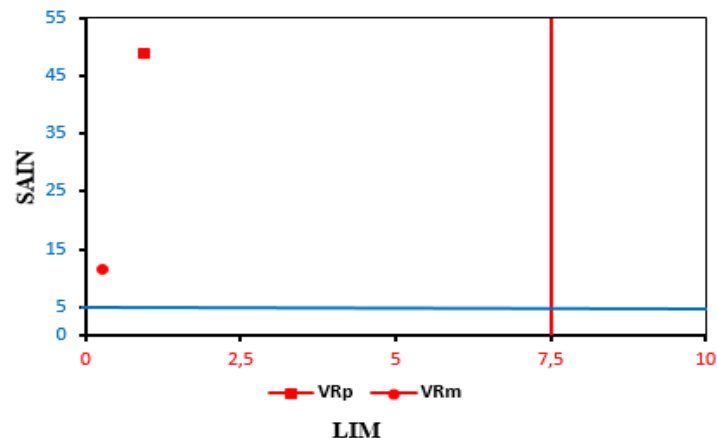
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### 3.5. Nutritional profile

The figure 1 shows the SAIN and LIM scores of *Byttneria catalpifolia* extracts (Dried bark and mucilage). The SAIN score for the dried bark was 49.05 and its LIM score was 0.91. As for mucilage, the SAIN score was 11.69 and its LIM score recorded was 0.25. The extracts of this stem vegetable had a high SAIN (SAIN > 5) and a low LIM (LIM < 7.5).



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**Fig. 1. SAIN and LIM profile of powder and mucilage extracted from the bark of *Byttneria catalpifolia***  
(VRp: powder of dried bark of red variety; VRm: mucilage extracted from fresh bark of red variety)

## 4. Discussion

### 4.1. Proximate composition

This work revealed that ash contents of the dried stem bark and mucilage from the red variety of *Byttneria catalpifolia* differed meaningfully ( $p \leq 0.05$ ). Note that, ash content is useful in assessing the quality grading of food materials and also gives an idea of the amount of their mineral composition [17]. This suggested that a significant part of the minerals was not extracted with the mucilage. The recorded values of ash were lower than that of the bark from *Bridelia thermifolia* ( $12.14 \pm 0.63$  %) and *Corchorus olitorius* (18.2 %) [18] [19]. By contrast, they were higher than that found by [20] in tubers of *Tacca leontopetaloides* ( $0.80 \pm 0.01$  %).

The fat contents of samples testified that the red variety of *Byttneria catalpifolia* is poor in lipids. Indeed, fat contents of dried stem bark and mucilage of this plant species were low compared with those of the seeds of *Daniella oliveri* (7.09 %) and *Olax subscorpoidea* (3.66%) [21]. Our results were also found to be slightly higher than those reported by [22] for the seed of *Parkia filicoidea* (0.43%).

Dietary fibre is an important component of food in human nutrition because it promotes intestinal mobility and reduces serum cholesterol, breast cancer and hypertension [23] [24]. In fact, there is evidence that dietary fibre is likely to reduce the absorption rate of glucose and fat [25], leading to health benefit. Our results revealed that the crude fibre of stem bark was high while it was low in mucilage. Moreover, the crude fibre of the dried bark powder of *Byttneria catalpifolia* was higher than that of *Melochia corchorifolia* ( $23.33 \pm 2.89$  %), a leafy vegetable [26]. According to [27], the recommend limit for fibre intake is from 1.4% to 3.5%. Based on this report, only the dried stem bark powder was good source of dietary fibre.

329

330 As concern reducing and total soluble sugars, their levels in the stem bark powder were  
331 higher than those observed in mucilage, whereas the carbohydrates content of mucilage  
332 was higher than that observed in dried bark powder. This showed that the mucilage could  
333 contain most complex sugars. The level of carbohydrates of the mucilage from the fresh bark  
334 of the red variety of *Byttneria catalpifolia* was high compared with the values of 52.6% in  
335 mucilage of *Ziziphus mauritiana* [28].

336 As for vitamin B2 and B9, the result showed that the both samples were a good source of  
337 the vitamin B9.

338 This work revealed that the calculated caloric energy of the both samples was low compared  
339 with the respective values of 1476.7KJ/ 100 g dw and 1513.5KJ/100 g dw in mushroom  
340 flours from *Ganoderma spp* and *Hebeloma mesophaeum* [29]. Therefore, the consumption  
341 of bark and mucilage of the red variety of *Byttneria catalpifolia* could be recommended to  
342 people suffering from obesity.

343

#### 344 **4.2. Amino acid contents**

345

346 Amino acid composition is a reliable indicator of the nutritional value of food. Free amino  
347 acids are among the main constituents of functionally essential compounds that are found in  
348 stem bark and mucilage of the red variety of *Byttneria catalpifolia*. The amino acid contents  
349 found in the mucilage was higher than those of the bark. These low levels of amino acids  
350 could be due to the effects of the processing (drying, grinding, sieving) undergone by the  
351 bark to obtain the powder. [30] in their work on *Moringa oleifera* showed that the quality of a  
352 dried product is often lower than the original food with an important impact on nutritional  
353 value. Otherwise, amino acids contents of bark powder or mucilage of *Byttneria catalpifolia*  
354 were found to be very lower than those of 11 wild edible mushrooms (from 153.09 mg/100 g  
355 dw in *F. hepatica* to 2267.32 mg/100 g dw in *B. edulis*) from northeastern Portugal [31].  
356 Besides, the essential amino acids contents of both samples studied were very low  
357 compared with the respective values of 154.3 mg/100 g in *B. craspedius* and 5232.5 mg/100  
358 g in *T. microcarpus* [32]. This suggested that both samples were poor in essential and non-  
359 essential amino acids. Moreover, levels of some of them were very lower than that  
360 recommended by [33]. In addition to that, protein contents were relatively low in the mucilage  
361 as well as in the dried bark powder and were not sufficient to meet protein requirements or  
362 the balance of the amino acids. Thus, we recommend that consumers eat *Byttneria*  
363 *catalpifolia* with meat or fish to balance the diet in protein.

364

#### 365 **4.3. Mineral composition**

366 The mineral composition indicated that the macro-elements such as Mg, P, K, and Ca had  
367 relatively high contents in *Byttneria catalpifolia* samples. These results showed a close  
368 agreement with those of [34] [35], who reported that Ca and K levels were highest in  
369 vegetables. Ca is the most abundant mineral in the body with 99 % in the teeth and bones. It  
370 is an essential nutrient that is needed for many functions in human health. Ca content of the  
371 dried bark of *Byttneria catalpifolia* was higher than that of the leaves of *Urtica urens* (830 mg  
372 / 100 g dw) [36], *Melochia corchorifolia* (750.37 mg / 100 g dw) [26] and that of mucilaginous  
373 vegetables such as *Iringia gabonensis* (452 mg / 100 g dw) and *Beilschmedia manii* (104  
374 mg / 100 g dw) [37]. Considering that the daily requirement of Ca is 1200 mg, one serving of  
375 the dried bark powder of *Byttneria catalpifolia* per day would help to meet daily Ca  
376 requirements [38].

377 The level of K in the dried bark of the red variety of *Byttneria catalpifolia* was higher than that  
378 of mucilage and both had high contents in K as well as leaves of *Urera trinervis* (1.25 %) and  
379 *Hippocratea myriantha* (1.29 %) [39]. K is known for its important role in the osmotic  
380 regulation of body fluids and in the acid-base balance. It also contributes to nerve and  
381 muscle excitability and carbohydrate metabolism [40].

382 Mg is closely associated with Ca and P and about 70% of the total Mg is found in the  
383 skeleton. Ca and Mg are essential elements of human physiology and are particularly  
384 important in the biological functions which are characteristics of the cardiovascular system  
385 [41]. Mg levels obtained in the bark and mucilage were greater than those reported by [42]  
386 and [43] in *Artocarpus altilis* flours (90.63 - 92.7 mg /100 g) and cassava (36.58 - 37.71 mg  
387 /100 g), respectively. Moreover, in the extracts of the bark of *Byttneria catalpifolia*, the [Ca] /  
388 [Mg] ratio was greater than 2. Note that, the absorption of Ca and Mg are interdependent,  
389 depending on their ratio. Diet is considered good if the ratio between these two minerals in  
390 the diet is from 2 to 1 in favour of Ca [44].

391 P has more functions than any other mineral element in the body. It forms a complex with Ca  
392 that gives stiffness to bones, teeth and muscles [45]. It acts as a cofactor for many enzymes  
393 and activates several of the B complex vitamins. It also affects the production of the  
394 adenosine triphosphate molecule, which is crucial for energy storage [46]. The P contents of  
395 powder and mucilage of *Byttneria catalpifolia* were higher than that of *O. gratissimum*, *T.*  
396 *occidentalis*, and *V. amygdalina* (13.8, 13.1 and 15.08 mg/100g dw, respectively) which are  
397 leafy vegetables eaten in South West of Nigeria [47]. It should be noted that, food is also  
398 considered "good" if the [Ca] / [P] ratio is greater than 1 and "mediocre" if less than 0.5 [44].  
399 The [Ca] / [P] ratio of dried bark powder and mucilage was 28.68 and 11.80, respectively.  
400 Thus, the levels of these minerals showed that the dried stem bark and mucilage of this plant  
401 may be used to complement the required macro-element needed for proper growth and  
402 development in human beings and other domesticated animals.

403 The high content of iodine (I) and other minerals in the pellet compared to the mucilage  
404 showed that the use of mucilage to prepare sauce does not allow access to all the minerals  
405 contained in this stem vegetable. The relative high concentration of I in dried stem bark  
406 powder compared with other vegetables such as eggplant leaves (7.6 µg / 100g dw) and  
407 melon leaves (6.13 µg / 100 g dw) [48] indicated that the bark could be a good source of I.  
408 Thus, its regular consumption would supply appreciable amount of I and would benefit  
409 consumers, particularly those in the western of Côte d'Ivoire, where endemic goitre is high  
410 (38.3%) [49]. Indeed, I is essential to produce thyroid hormones, thyroxine (T4) and  
411 triiodothyronine. Iodine deficiency can lead to goitre, hypothyroidism, poor growth and  
412 neurocognitive trouble [50].

413

#### 414 **4.4. Organic acids contents**

415

416 The observed organic acids were Tannic, oxalic, citric, tartaric, Sulfanilic, Salicylic, adipic,  
417 fumaric and benzoic acids. Citric and tartaric acids had the highest levels in both samples.  
418 Citric acid contents of samples were higher than that observed for *moringa* leaf (1.56 ± 0.45  
419 mg/100 g dw) [51]. However, they were lower than that observed in kale (*Brassica*  
420 *oleraceae* L.var. *acephala* DC.) leaf (2213 mg/100 g dw) [52]. As for tartaric acid contents of  
421 both samples, they were found to be lower than that obtained by [51] in Ceylon Spinach  
422 (1200 ± 11.56 mg/100g dw). In contrast, tartaric acid levels were higher than that found in  
423 *moringa* leaf by [51], who recorded the value of 0.45 ± 0.02 mg/100 g dw. Note that, the  
424 organic acid contents of foods do not only influence their flavour, but also their stability,  
425 nutrition, and acceptability [53]. The [oxalic acid] / [Ca] ratios of dried bark (0.001) and  
426 mucilage (0.061) were less than 2.25, which is the threshold value not to be exceeded.  
427 Indeed, oxalic acid interferes with Ca uptake. Thus, the [oxalic acid] / [Ca] ratios below 2.25  
428 indicated that the amount of oxalic acid in both samples will not interfere with Ca [54].

429

#### 430 **4.5. Nutritional profile**

431 The nutritional profile of extracts of the red variety of *Byttneria catalpifolia*, following the SAIN  
432 and LIM method showed that they belong to the group of foods recommended for health.  
433 These results corroborate those of [55] in their work on use of SAIN and LIM system for  
434 determination the nutritional profile of foods eaten by children under-five in the district of

435 Abidjan, Côte d'Ivoire. According to these authors, the different vegetable-based soups such  
436 as okra, tomato and eggplant have a good nutritional value because their SAIN > 5 and LIM  
437 < 7.5. [56] also showed that 80% of vegetables belong to the food group with a strong SAIN  
438 and a low LIM and therefore recommended for health. According to [57], non-starchy  
439 vegetables such as *Byttneria catalpifolia* are rich in fibres that improve intestinal transit by  
440 forming a mass, which leads to a more gradual absorption of nutrients, thus avoiding  
441 constipation. They are also very poor in energy and therefore can be consumed in relatively  
442 larger amounts to maintain a normal weight [58].

443

## 444 5. CONCLUSION

445

446 On the basis of the above results it can be concluded that the dried bark and the mucilage  
447 extracted from the fresh bark of the red variety of *Byttneria catalpifolia* contain appreciable  
448 amounts of nutrients such as carbohydrates, Proteins, essential amino acids, vitamins and  
449 minerals. Our findings indicated that the both samples were nutritionally important, since  
450 their nutritional profile by SAIN and LIM method showed that they belong to the group of  
451 foods recommended for health. The powder of the dried bark was very rich in iodine and  
452 other nutrients, therefore the consumption of this powder would be more beneficial to  
453 nutritional balance than the mucilage.

454

455

456

457

## 458 COMPETING INTERESTS

459

460 The authors declared that they have no conflict of interest.

461

462

463

## 464 CONSENT

465

466 It is not applicable.

467

## 468 ETHICAL APPROVAL

469

470 It is not applicable.

471

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