

Phytochemical Screening of *Etlingera elatior* Cultivated on Different Dosage of Biochar

Abstract

This study was to investigate the impact of biochar on phytochemical composition in plant. The phytochemical composition was extract from different dosage application of biochar (0%,5% and 20% by weight). These treatments found significantly increased in the phytochemical composition. It is recommended that application of biochar not only improve the growth rate but also enhance the phytochemical of the plants. The Fourier Transform Infrared spectra confirmed that the biochar rich with humus like-compounds which increase the nutrient content of the soil.

Keywords: phytochemical, biochar, ethanolic extract, Cultivated

Introduction

Recently, food safety is becoming a major issue. Eventhough the advances in technology have been applied to agricultural industry, the global food supply is yet insufficient to meet the markets demands. A prominent environmental issues such as climate change, desertification, and soil pollution are still remain to be resolved for the agriculture sector [1]. Soil quality and agricultural productivity become deteriorated due to the impact of industrialization. The pollutants like heavy metals, engineered nanomaterials, polycyclic aromatic hydrocarbons (PAHs), and persistent organic and inorganic chemicals as derivatize from the industrial process acceleratedd the soil contamination [2]. These anthropogenic sources have constantly affected plant environments, thereby causing harm to the terrestrial food. Many studies have witnessed the negative impacts of contaminants on soil composition and biodiversity.

Pyrolysis and composting technologies had introduced biochar into global agriculture attention due to its benefits [3,4]. Biochar produced by biomass pyrolysis [5]. Its elemental composition consists of carbon, nitrogen, hydrogen, potassium, and magnesium all of which can serve as major nutrients in plant growth. The addition of biochar increases the amount of organic matter in the soil (e.g, organic carbon), thereby improving soil physicochemical and

32 biological properties. Biochar can positively or negatively affect the soil microbial growth to
33 alter the agricultural environment [7].

34 Biochar is a very stable, carbon-based material obtained from the pyrolysis of biomass under
35 anaerobic conditions, and is highly recalcitrant in soils. The parent material or biomass can be
36 obtained from agricultural, municipal, animal, or industrial sources. The pyrolysis
37 temperatures generally employed range from 300 to 1000 °C. Biomass is largely composed of
38 organic compounds such as cellulose, hemicellulose, and lignin. Lignin is the most stable of
39 these compounds, and is resistant to degradation at even higher temperatures. In contrast,
40 temperatures above 300 °C can decrease the cellulose and hemicellulose contents. The
41 temperature and duration of pyrolysis are determined based on the target purpose. In some
42 cases, catalytic additives such as K_3PO_4 and clinoptilolite are used to reduce the pyrolysis
43 temperature. During pyrolysis, water and volatile organics from the biomass may evaporate,
44 thereby increasing the aromatic content. The parent biomass source and pyrolysis temperature
45 affect the physiochemical properties of the biochar obtained. For instance, pyrolysis
46 temperature is the main control on atomic ratio and structural composition. Although raw
47 biomass is slightly acidic, pyrolysis at high temperatures increases its alkalinity. This is due to
48 the partial detachment of the functional groups leading to the formation of unpaired negative
49 charges such as carboxyl (COO^-) and hydroxyl groups (OH^-) that have the ability to attract
50 positive charges [9]. High-temperature pyrolysis also causes release of hydrogen- and
51 oxygen-containing groups, contributing to increased carbon content. The tendency of the
52 surface functional groups to attract positive charges enhances the cation exchange capacity,
53 which is an important property of biochars for remediation of metal-contaminated soils.
54 Furthermore, the porosity, pore size, and surface area of biochars depend on pyrolysis
55 temperature as the high temperature of pyrolysis leads to formation of pores via the release of
56 volatile organics. Thus biochar's properties can be targeted to a range of different purposes by
57 adjusting the pyrolysis temperature.

58 Application of biochar has been shown to yield a wide range of benefits to plant growth and
59 stress management. Studies have documented the role of biochar in improving agronomical
60 parameters and environments for various plant species. Addition of less than 5% biochar
61 enhanced the germination, yield, and root development of the halophytes such as sesbania and
62 seashore mallow. Rice hull-based biochar applied at a rate of 2% to a sandy soil increased the
63 biomass yield and sucrose content of sugarcane plants.

64 Application of biochar also can improve nutrient cycles. In wheat culture, biochar addition at
65 10% in combination with urea improved agronomic efficiency of N by 63%; biochar mixed
66 with 10% KOH significantly increased Si uptake by plants. It is noted that Si plays a vital role
67 in both biotic and abiotic stress management in plants. P-loaded biochar at 30 g kg⁻¹ also
68 improved the bioavailability of P. Sewage sludge biochar at 50% acts as a soil conditioner and
69 was able to stimulate turf grass growth, even in urban soils. Other studies have found that
70 biochar can expand plant diversity by benefitting reforestation activities in low-carbon soils.
71 The reforestation capacity is due to increased total soil carbon in the reforested sites.

72 In this study, it was hypothesized that application of biochar to *E. elatior* with different
73 dosages of biochar might affect the type of secondary metabolite. Therefore, this present
74 study aimed to evaluate the effect of biochar at two dosages (10 and 20% of fresh mixture
75 weight) on the type of secondary metabolite extracted from the *E. elatior*.

76 **Materials and Method**

77 Cultivation of *Etilingera elatior*

78 *Etilingera elatior* was cultivate on the pot with 20 cm diameter and 35 cm height. 3 replicates
79 for pots of *Etilingera elatior* was cultivated and label as 0%, 5% and 20%. The cultivation soil
80 used was non-fertilized soil(0%) fertilized soil with biochar; 5% and 20%. The pots was
81 placed in the netting house and the growth of *Etilingera elatior* was monitored. Watering
82 process was done twice a day. The biochar used in this study were purchased from Black Owl
83 Biochar Products. The dosage of biochar used was calculated over the weight of soil. The soil
84 treatments are as follows:

85 a) Non-fertilized soil (0%), b) non-fertilized + biochar (5% and 20%)

86 Sample extraction

87 Phytochemical was extracted according to [12,13,14] with some adjustments. 100 gm of the
88 dry leaf powder of *E. elatior* were weighted, transferred to a flask, soaked with ethanol until
89 the powder was fully immersed and incubated overnight. The extracts were then filtered
90 through Whatman filter paper No.1 along with 2 gm sodium sulfate anhydrous to remove the
91 traces of water in the filtrate. Before filtering, the filter paper along with sodium sulphate was
92 wetted with 95% ethanol. The filtrates were then air dried and subjected to gas
93 chromatography-mass spectrometry analysis.

95 **Fourier Transform Infrared (FTIR)**

96 The biochar samples were analysed using an ATR-FTIR equipped with diamond crystal,
97 controlled by OMNIC software (Thermo Nicolet Analytical Instruments, Madison, WI). A
98 flat tip powder press was used to achieve even distribution and contact. All spectra were
99 collected at 12 scans with a resolution of 4 cm⁻¹ in the range 4000–670 cm⁻¹. The spectrum
100 of each sample was ratioed against a fresh background spectrum recorded from the bare ATR
101 crystal. The ATR crystal was cleaned with ethanol.

102 **Gas Chromatography Mass Spectrometry (GCMS) analysis**

103 The gas chromatography mass spectrometry analysis was performed by using a non-polar
104 BPX-5 capillary column with an initial temperature of 50°C hold for five minutes and then
105 increased to 300°C at a rate of 5.0°C per minutes and hold 10 minutes. The temperature of the
106 injector and detector were set at 320°C respectively. 1µl of the fractions was diluted in 100µl
107 hexane was introduced into the gas chromatography. The gas used as the carrier was Helium.
108 Interpretation of mass-spectrum was conducted using the database of National Institute
109 Standard and Technology (NIST17). The spectrum of the secondary metabolites components
110 was compared with the spectrum of known components stored in the NIST library. The name,
111 molecular mass and structure of the components of the test materials were ascertained.

112 **Results and discussions**

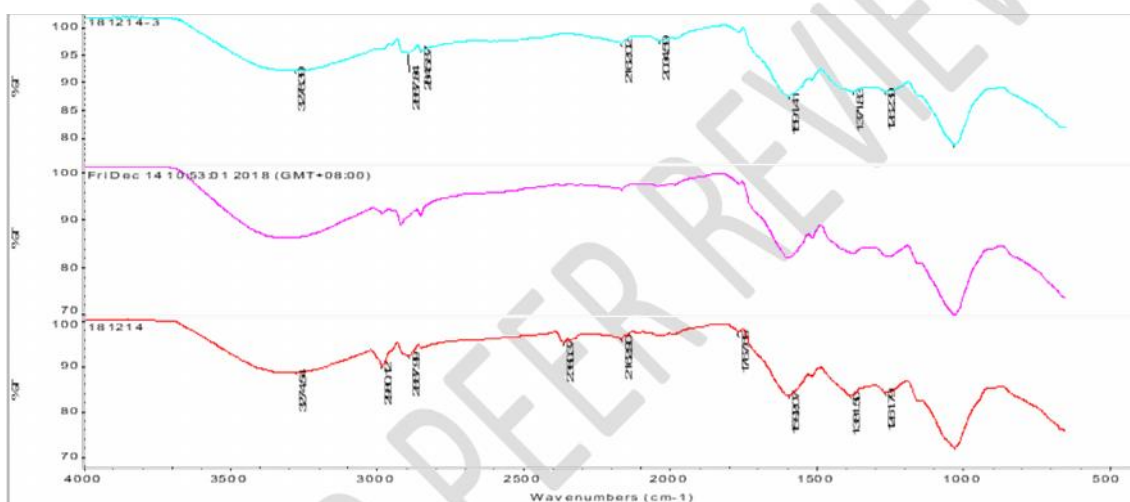
113 The Fourier Transform Infrared spectra shown in Figure 1. The spectra with pronounced
114 bands at 1591 cm⁻¹ corresponding to carboxyl group of protein compounds. Broad bands at
115 1700 cm⁻¹ due to stretching of C-O bond in polysaccharide. However, polysaccharide would
116 have degraded during the process of biochar, whilst the phosphate content in biochar increase.
117 The phosphate band can be observed at 1029 cm⁻¹. These findings confirm that the biochar
118 was rich in humus compounds which is probably from mixture of poultry materials.

119 The gas chromatography's chromatograms of *Etlingera elatior* obtained from ethanolic
120 extract are shown in Figure 2. The chromatograms demonstrate different peaks for different
121 dosage of biochar (Figure 2(a)-(c)). Different pattern of chromatogram was observed on
122 different dosage of biochar applied. The major compounds with their percentage area (PA%)
123 are summarize in Figure 3. The results revealed that phytol (13%), Hexadecanoic acid
124 (9.76%), Neophytadiene (6.51%), coumarin (5.65%), precocene (5.27%) and caryophyllene
125 (4.59%) were among the major compounds identified from *Etlingera elatior* ethanolic extract
126 on 0% biochar (Figure 3(a)). Increasing biochar dosage to 5%, the identified compounds

127 shows that, the 6 major compounds are Dihydrocucurbitacin (13.69%), Niacinamide
128 (11.02%), α -Limonene (10.01%), Phyrachen (9.23%), Phytol (7.24%) and Neophytadiene
129 (5.75%) shown in Figure 3(b). In 20% biochar the major compounds of *Etlingera elatior* are
130 Linoleic acid (39.98%), 2-pinen-4-ol (12.32%), Hexatriacontyl pentafluoropropionate
131 (6.89%), Benzofuran (5.12%), Acethophenon (4.41%) and furfural (4.03%) shown in Figure
132 3(c). Most of the compounds known to be secondary metabolite which are rich in medicinal
133 values.

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Figure 1. Stack spectra of biochar.

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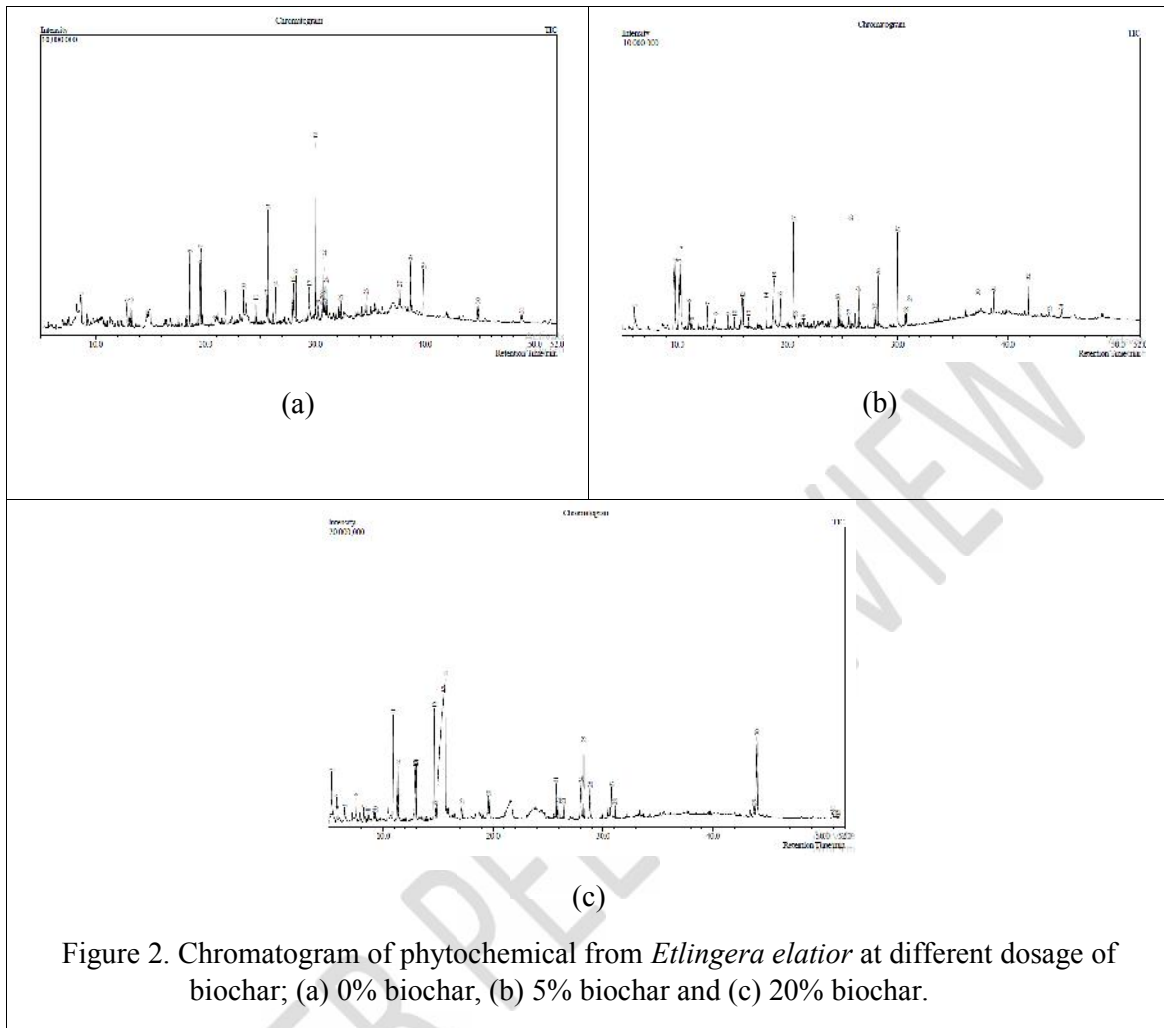
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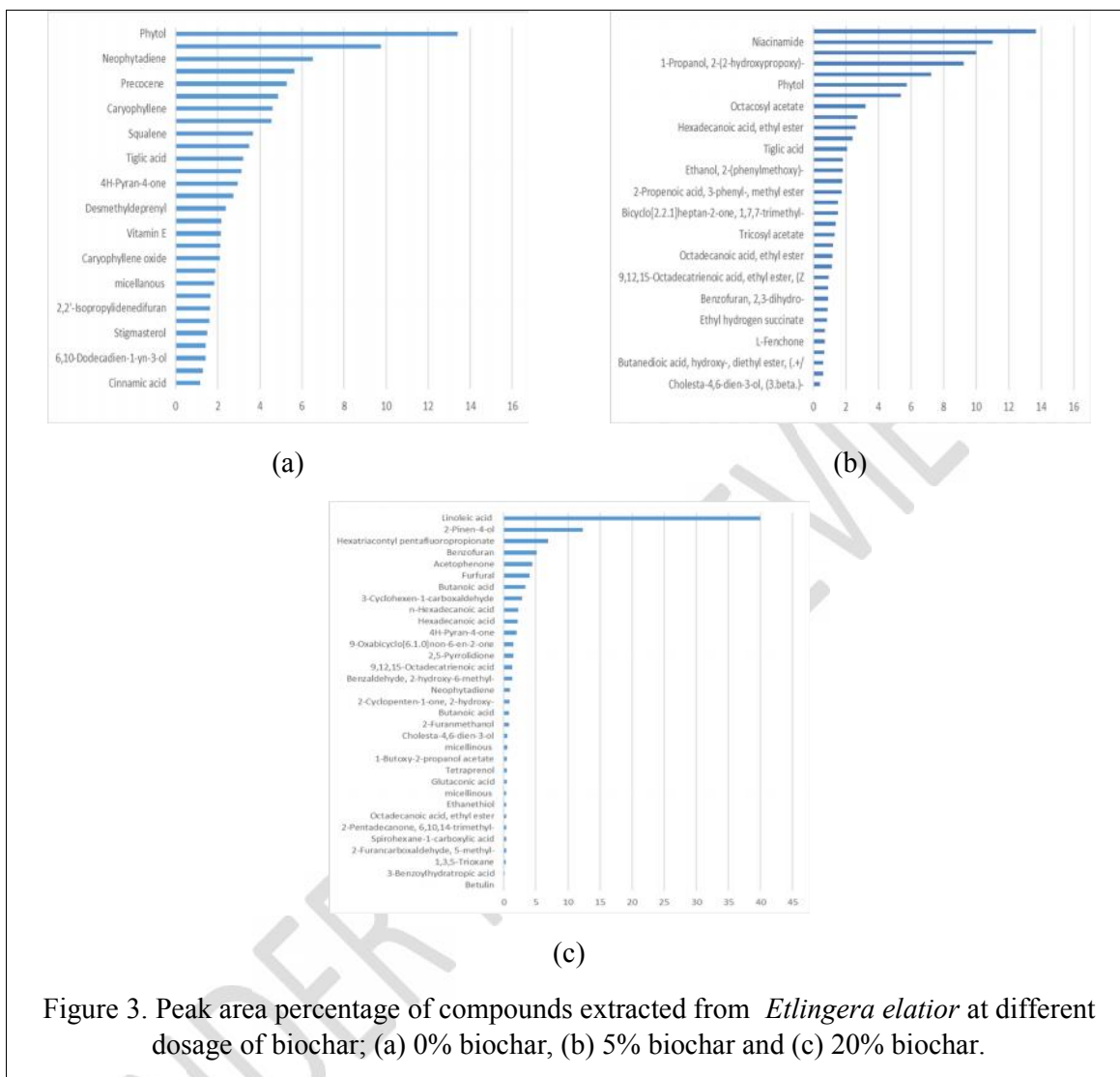


Figure 3. Peak area percentage of compounds extracted from *Etilingera elatior* at different dosage of biochar; (a) 0% biochar, (b) 5% biochar and (c) 20% biochar.

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167 Conclusion

168 Application of biochar on soil can increase nutrient availability and enhance the development
169 of phytochemical composition in plants. Without biochar, the chemical composition *Etilingera*
170 *elatior* extract was slightly low. At 5% and 20% biochar, some compounds are increasing and
171 new compounds are develop compared to 0% biochar. This suggest that the biochar not only
172 able to increase the growth rate of plants but also the nutrients of the plants.

173 **Conflicts of Interest**

174 The authors declare that there is no conflict of interests regarding the publication of this
175 paper.

176 **References**

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