# Foliar Nutrient Concentrations and Antioxidant Activity of Tea (*Camellia sinensis* L. (O) Kuntze) Planted in Peninsular Malaysia and its Relation to Soil Edaphic Factors

# Wisnu Eko Murdiono<sup>1, 6</sup>, Nur Amirah Syafiqah Salman<sup>1</sup>, Uma Rani Sinniah<sup>1</sup>, Elisa Azura Azman<sup>1</sup>, Mohd Izuan Effendi Halmi<sup>3</sup>, Jean Wan Hong Yong<sup>4</sup>, Abbey Maleyki Mhd Jalil<sup>5</sup>, and Khairil Mahmud<sup>1, 2\*</sup>

#### 7 ABSTRACT

The global popularity of tea is due to its unique taste and health benefits, which are highly 8 linked with its nutrient and antioxidant activity (AOA). However, diverse growing habitats, 9 including distinct altitudes and soil edaphic factors, may regulate foliar nutrition and AOA of 10 tea. Thus, this study aimed to (1) compare the nutritional characteristics and AOA of clonal tea 11 grown in lowland and highland plantations and (2) investigate the influence of soil edaphic 12 factors on tea foliar nutrition and AOA. Tea leaves and soils of fourteen tea clones were sampled 13 between October 2021 to March 2022 from lowland and highland plantations in Peninsular 14 Malaysia. Leaves were analysed for nutritional content and antioxidant activity, while soil 15 samples underwent physical and nutritional analysis. Results showed significant variations in 16 most foliar nutrients, except for Ca in the lowlands and Fe in the highlands. While the highland-17 grown tea exhibited higher nutrient concentration, lowland-grown tea demonstrated superior 18 AOA. AT53 and 1248 were identified as promising among the clones, characterized by the 19 highest nutrients and AOA levels, respectively. Soil nutrient availability significantly 20 influenced foliar nutrient uptake, while soil pH was associated with the AOA. These findings 21 highlight the critical role of soil edaphic factors in shaping tea quality, providing valuable 22 insight for tea growers to optimize soil management strategies and maintain tea yield and 23 quality in the future. We found that soil nutrients have a significant association with nutrient 24

<sup>&</sup>lt;sup>1</sup> Department of Crop Science, Faculty of Agriculture, Putra University of Malaysia, 43400 Serdang, Selangor, Malaysia.

<sup>&</sup>lt;sup>2</sup> Biodiversity Unit, Institute of Bioscience (IBS), Putra University of Malaysia, 43400 Serdang, Selangor, Malaysia.

<sup>&</sup>lt;sup>3</sup> Department of Land Management, Faculty of Agriculture, Putra University of Malaysia, 43400 Serdang, Selangor, Malaysia.

<sup>&</sup>lt;sup>4</sup>Department of Biosystems and Technology, Swedish University of Agricultural Sciences, Alnarp, Sweden.

<sup>&</sup>lt;sup>5</sup> School of Nutrition & Dietetics, Faculty of Health Sciences, Sultan Zainal Abidin University (UniSZA), Terengganu 21300, Malaysia.

<sup>&</sup>lt;sup>6</sup> Department of Agronomy, Faculty of Agriculture, Brawijaya University, Malang, Indonesia.

<sup>\*</sup>Corresponding author; email: hairilmahmud@upm.edu.my

25 uptake, while soil pH is associated with the agronomic characteristics of tea. Investigating the

26 association between ecological variables and tea foliar properties (nutrients and AOA) is of

27 great importance for tea growers as they develop strategies to maintain the yield and quality of

tea in the future.

Keywords: Camellia sinensis, nutritional characteristics, antioxidant activity, altitude, soil
 edaphic.

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#### 32 INTRODUCTION

Tea (Camellia sinensis L. (O) Kuntze) is a popular caffeinated non-alcoholic beverage 33 around the globe and is currently consumed by 3 billion people (Pan et al. 2022). It is attributed 34 to several health benefits in addition to its unique taste and aroma. The health-promoting 35 properties of tea are due to the presence of polyphenols, particularly the flavan-3-ols, widely 36 37 known as catechins. Catechins are the main polyphenols in tea and have been reported to have a few derivatives (Lee et al. 2014). Epigallocatechin gallate (EGCG) is the catechin derivative 38 with the highest prevalence and pharmacological activity. EGCG is responsible for up to 70% 39 of total catechins and has been proven to have chemo-preventive/chemotherapeutic actions 40 against several malignancies (Steinmann et al. 2013) and other disorders such as obesity, 41 diabetes, neurological and cardiovascular diseases (Khan and Mukhtar 2019). In addition to 42 polyphenols, tea also contains minerals and trace elements which play an important role in 43 human metabolism. Regular consumption of tea may contribute to the daily dietary requirement 44 of certain elements, such as manganese. Furthermore, having more potassium than sodium may 45 benefit hypertensive patients (Fernandez et al., 2002). Hence, both nutrient elements and 46 polyphenols (particularly catechin, which has a strong antioxidant effect) are the main 47 compounds that have great contributions to human health benefits. 48

Variation in tea nutritional characteristics and AOA is associated with either internal (genetic 49 make-up and stage of leaf development) or external factors, including season, altitude as well 50 as soil physicochemical properties. For instance, earlier research revealed that certain nutrients, 51 such as N, P, K and Mg (Xiang et al., 2021) increase proportionally with altitude. Other 52 researchers addressed this phenomenon as a result of growth reduction experienced by the 53 plants grown in higher altitudes, which is highly associated with the highland's lower average 54 temperature. This result is quite the opposite of tea AOA. Previous studies conducted by Owuor 55 et al. (2011) and Martono et al. (2016) found that tea grown at lower altitudes tends to have 56 better antioxidant performance. This indicates that higher temperature is one of the factors 57 affecting catechin accumulation in plants grown at lower altitudes. Other environmental factors 58

influencing the accumulation of tea catechin are blue light, water stress, shading treatment as 59 well as soil physicochemicals which act by modulating the expression of biosynthetic genes 60 (Samynathan et al. 2021). An earlier study by Zhao et al. (2017) discovered that the contents of 61 Na, Mg, Ca, Cr, Fe, Ni, Sr and Cd in tea leaves were significantly and positively correlated with 62 those in topsoil or subsoil. It suggested a considerable rise of these elements in tea leaves when 63 soil contents increased. Another study by Tseng & Lai (2022) in Taiwan revealed that soil 64 characteristics, such as soil pH, exchangeable calcium and magnesium, significantly impact 65 tea's free amino acid content. 66

Previous studies on the influence of soil edaphic factors on nutritional characteristics as well 67 as AOA of tea have been conducted in Malaysia (Chan, et al. 2007; Izzreen and Fadzelly, 2013; 68 Amirah et al. 2023), mostly in lowland plantations. In Malaysia, tea has been planted in two 69 different altitudes for almost a century by BOH Plantation Sdn. Bhd., a Malaysian leading tea 70 company. However, research on the influence of soil physicochemical factors as well as altitude 71 on nutritional characteristics and the AOA of tea was less reported. Therefore, it is essential to 72 determine the characteristics of distinct altitudes and each soil edaphic factor on the capabilities 73 74 of tea clones in uptaking nutrients and accumulating catechin as this may benefit growers in designing future strategies for sustaining tea yield and quality. We hypothesized that soil 75 physicochemical properties from different altitudes might play a major role in influencing the 76 variation of foliar nutrient concentrations and the quality of tea. Therefore, in this study, we 77 aimed to 1) compare the nutritional characteristics and AOA among tea clones from lowland 78 and highland plantations; and 2) investigate the association between soil physicochemical 79 properties with nutritional characteristics and quality of clonal tea from both plantations. 80

#### 82 MATERIALS AND METHODS

#### 83 Study Sites

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Highland tea clones were planted in Cameron Highland, Pahang, while lowland clones were grown in Bukit Cheeding, Selangor. The Cameron Highland is located at approximately 1400 m above sea level (a.s.l), with an average temperature ranging from  $18 - 25^{\circ}$ C with an average humidity of 79 - 92% and 152.7 - 1077.8 mm rainfall. The Bukit Cheeding Plantation is at a lower elevation, approximately 20 m a.s.l, having a higher average temperature ranging from  $28 - 31^{\circ}$ C, lower humidity (74 - 86%), and rainfall (53.6 - 596.3 mm) compared to the highland location.

#### 92 Sample Collection and Analysis

Fourteen clones were sampled between October 2021 - March 2022 from lowland and 93 highland tea plantations. Leaf sampling sites were randomly selected (Table 1) and triplicated 94 for young tea leaves (bud and the first three fully expanded leaves). Fresh tea leaves were once 95 cleaned with tap water and distilled water to remove the adhering materials before oven-drying 96 at 60 °C for 4 days. The samples were then ground to get the fine powdered texture for further 97 antioxidant and foliar nutrient analysis. Total N was analyzed using a CNS analyzer 98 (CNSTruMax Determinator version 1.1x). For other foliar nutrients (total P, K, Ca, Mg, Al, and 99 Fe), tea leaves were digested using the dry-ashing method following Alarefee et al., (2021) and 100 subsequently analyzed by ICP-OES (Optima 8300, PerkinElmer, USA). 101

Tea leaves were extracted using 80% aqueous methanol by digital ultrasonic bath following 102 Bakht et al. (2019) with minor modifications using 40 °C for 30 min before AOA analysis. The 103 extract produced was evaluated for its total phenolic content (TPC), DPPH (2,2-diphenyl-1-104 picrylhydrazyl) radical scavenging assay and FRAP (Ferric Reducing Antioxidant Power) assay 105 according to Amirah et al. (2023). The TPC was evaluated employing the Folin-Ciocalteu 106 solution with gallic acid, which served as a reference. Ten-fold diluted tea extract (15 µL) was 107 combined with distilled water (240 µL) and 0.25N Folin-Ciocalteu reagent (15 µL), and then 108 mixed thoroughly. After 3 minutes of dark incubation at room temperature, 30 µL of 1N Na<sub>2</sub>CO<sub>3</sub> 109 was added. Following 2 hours of dark incubation at room temperature, absorbance was 110 measured at 765 nm. In the DPPH assay, 20 µL tea extract was added to 180 µL of DPPH 111 solution (150 µmol L<sup>-1</sup>) and incubated for 40 minutes in darkness. Absorbance was measured 112 at 517 nm, with 80% methanol used as blank. L-ascorbic acid served as the positive control. 113 The antioxidant activity was expressed as an IC<sub>50</sub> value, calculated using GraphPad Prism 8 114 115 software (GraphPad Software, San Diego, CA, USA). In the FRAP assay, the FRAP reagent needs to be freshly prepared. The reagent was made by mixing acetate buffer (0.3M, pH 3.6), 116 117 10 mM TPTZ in 40 mM HCl and 20 mM FeCl<sub>3</sub> in a 10:1:1 ratio, then warmed at 37°C. A mixture of FRAP reagent (280 µL) and tea extract (20 µL) was incubated at 37°C for 30 min 118 and measured the absorbance at 593 nm. 119

The soils were sampled at 0-20 cm depth using an auger within 1-2 meters from each clone for soil physicochemical analyses. The soil samples were air-dried for several days and filtered through a 2-mm sieve before chemical and physical analysis (Alarefee et al., 2021; Khairil & Burslem, 2018). Soil pH was measured at a soil: water ratio of 1:2.5. Soil total N was analysed using CNS analyser (CNSTruMax Determinator version 1.1x), while other soil nutrients (total

P, K, Ca, Mg, Fe and Al) were analysed using an ICP–OES (Optima 8300, PerkinElmer, USA) 125

after the aqua regia extraction method (Alarefee et al. 2021). 126

Clones	Lowland	Plantation	Highland	Plantation
Clones	North latitude (°)	East longitude (°)	North latitude (°)	East longitude (°)
AT53	2.92450	101.57789	4.52340	101.39976
TV9	2.91870	101.57658	4.52110	101.40415
1248	2.92172	101.58327	4.52091	101.40018
2024	2.92813	101.58134	4.52322	101.39959
663	2.92231	101.58356	_	_
1294	2.92201	101.58292	_	_
2026	2.92368	101.58454	_	_
TBR2020	_	_	4.51790	101.41153
196	_	_	4.52321	101.39954
664	_	_	4.52281	101.39865

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#### **Statistical Analysis** 129

All analytical results were performed as the average of three replicates using R-studio 130 version 4.1 (R Core Team, 2021). Data were subjected to One-way ANOVA continued by 131 Tukey's HSD to examine the variation of nutritional characteristics, AOA and soil 132 physicochemical properties. The significance level was set at a 95% confidence level ( $\alpha = 0.05$ ). 133 In combination with Pearson's correlation, Principal Component Analysis (PCA) was 134 performed to determine the influence of soil edaphic factors (nutrient concentrations and pH) 135 on nutritional characteristics and the AOA of tea from both plantations. These were conducted 136 137 using R-studio version 4.1 (R Core Team, 2021). PCA helps to identify patterns and relationships between variables by transforming the data into Principal Components (PCs) that 138 summarize the dataset. Principal components were selected based on eigenvalues > 1 and 139 cumulative variance explained. 140

#### 141 **RESULTS** 142

#### Nutritional Characteristics and AOA of Tea 143

The concentration of foliar nutrients differed significantly (p<0.05) between lowland tea 144 clones, except for Ca (Table 2). Among seven clones evaluated, AT53 had the highest foliar K 145  $(1.84\pm0.38 \text{ mg g}^{-1})$ , Mg  $(0.80\pm0.16 \text{ mg g}^{-1})$ , Fe  $(13.00\pm0.79 \text{ mg g}^{-1})$  and Al concentration 146  $(16.60 \pm 0.78 \text{ mg g}^{-1})$ . Meanwhile, clones 2026 and 663 had the highest foliar N  $(4.39 \pm 0.2\%)$ 147 and P concentration ( $13.76 \pm 1.06 \text{ mg g}^{-1}$ ), respectively. In highland tea plantations, the majority 148 of the foliar nutrition concentrations varied significantly (p<0.05) among clones, except for Fe 149 (Table 3). Of the seven clones evaluated, TBR2020 had the highest foliar Ca (3.89 mg g<sup>-1</sup>), Mg 150

151 (1.78 mg g<sup>-1</sup>) and Al content (0.69 mg g<sup>-1</sup>). Meanwhile, clones AT53, 1248 and 196 had the 152 highest foliar N (5.18 %), P (8.35 mg g<sup>-1</sup>) and K content (13.4 mg g<sup>-1</sup>), respectively.

In terms of the AOA of tea, DPPH assay from lowland (Table 2) and highland (Table 3) plantations varied significantly (p<0.05) among clones. Clone 1248 (50.66  $\mu$ g/mL) and TBR2020 (127  $\mu$ g/mL) had the highest value for lowland and highland plantations, respectively. In addition, FRAP (Ferric Reducing Antioxidant Power) value significantly varied among the highland population only, with clone 1248 (1.19 mM Fe (II)/g) having the highest value. However, total phenolic content (TPC) displayed insignificant variations among tea clones from both plantations.

A PCA based on population mean values of seven nutrient elements and three antioxidative 160 assays for tea grown in both plantations is displayed first axis explaining 76.97% of the 161 variation in the data (Figure 1). This axis reflected a positive correlation with most of the 162 variables (loadings 0.83 - 1.02), except N. The Second PC axis which explained 8.45% 163 variations, captured variation in N concentration (loadings 0.83) of tea. Following the PCA, tea 164 clones were clustered into two distinct groups. Clones with numbers 1 - 7 were from the 165 lowland plantation and had a significant association with all the antioxidative assays, as well as 166 167 foliar P, Fe and Al concentrations. Another cluster consists of clones with numbers 8 - 14 originating from the highland plantation and found to be highly associated with foliar K, Ca 168 169 and Mg.

Four clones out of the ten clones evaluated were planted in both locations, namely AT53, TV9, 1248 and 2024. Even though they have similar genetic make-up, different geographical areas (altitude) and their microclimate influence their physiology and metabolism. Based on our foliar analysis and PCA result, we found that four clones grown in both locations had similar clustered patterns as shown in Figure 1. Tea clones planted in the highlands tend to have higher foliar nutrients, particularly K, Ca and Mg. On the other hand, lowland-grown clones were associated with better antioxidant performance, as well as foliar P, Fe and Al.

#### 177 Soil Edaphic Factor

Soil P, K, Mg, Fe, and Al varied significantly among the seven lowland clones evaluated (Table 4). Soil obtained around clone 663 had the highest P ( $0.37 \text{ mg g}^{-1}$ ), Mg ( $0.39 \text{ mg g}^{-1}$ ) and Fe ( $10.20 \text{ mg g}^{-1}$ ) concentration. The highest soil K ( $1.17 \text{ mg g}^{-1}$ ) and Al ( $23.8 \text{ mg g}^{-1}$ ) concentrations were found from soil-derived near clones TV9 and 2026, respectively. In highland plantations, most of the soil nutrient content varied significantly, except for Ca (Table

5). Soil derived near clone 196 had the highest soil N (0.49 %) and P (7.22 mg g<sup>-1</sup>) concentration. The highest soil K (2.83 mg g<sup>-1</sup>), Mg (3.10 mg g<sup>-1</sup>), Fe (29.9 mg g<sup>-1</sup>) and Al (24.1 mg g<sup>-1</sup>) concentration was obtained around clone TBR2020.

In terms of pH, both lowland (Table 4) and highland (Table 5) tea plantations had acidic pH, with averages of 4.23 and 3.58, respectively. The pH of lowland plantations differed significantly among clones with soil near clones AT53 and TV9 as the lowest and clones 1248 and 1294 as the highest.

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#### 191 Association of Soil Edaphic with Foliar Nutrient and AOA of Tea

We found that soil physicochemicals had a significant association with foliar nutrients and 192 AOA of tea (Table 6). Certain soil nutrients displayed direct correlation as well as 193 intercorrelation with foliar nutrients. For instance, soil N with foliar N (r=0.55), soil Ca with 194 foliar Ca (r=0.77), and soil Mg with foliar Mg (r=0.59). In addition, soil N is also significantly 195 intercorrelated with foliar K (r=0.83), Ca (r=0.64), and Mg (r=0.76). Soil P is significantly 196 intercorrelated with foliar K (r=0.80), Ca (r=0.61), and Mg (r=0.69). Both Ca and Mg 197 demonstrated substantial intercorrelation with foliar K (r=0.90; r=0.57), while they showed an 198 intercorrelation. Soil pH demonstrated a positive and significant correlation with foliar P 199 (r=0.81), Fe (r=0.53), and Al (0.55). Furthermore, pH also had a positive and significant 200 correlation with TPC (r=0.59) and AOA of tea, represented by DPPH (r=0.84) and FRAP (0.78). 201

#### 202 DISCUSSION

#### 203 Foliar Nutrient and AOA of Lowland and Highland Tea Plantation

Our result demonstrated that clones grown in a highland plantation have a higher foliar 204 nutrient content than those grown at a lower elevation. However, not all nutrients would 205 increase proportionally with altitude. Our findings indicate that the foliar concentrations of N, 206 207 K, Ca, and Mg increase with altitude. This is similar to the result of the previous study. For instance, a recent study by Xiang et al. (2021) also found that foliar N and P content as well as 208 leaf C: N ratio increase significantly with altitude. Several researchers assumed that the increase 209 in leaf nutrient content of tea with altitude could be considered a consequence of biomass 210 production decreasing with altitude, mainly due to cold limitation in trees in mountainous 211 regions (Jeyakumar et al. 2020). 212

Highland plants tend to grow at a slower pace than those grown at lower altitudes. Decreasing temperatures, lower nutrient availability, and slower rates of photosynthesis are among the factors that influence the poor growth of highland plants (Jeyakumar et al. 2020).

Generally, every 1 km increase of elevation results in a  $6.5^{\circ}$ C decrease in temperature. Low temperatures reduce soil microbial and enzymatic activity, thus limiting nutrient availability, hence high-altitude soils are less fertile (Xu et al. 2015). Furthermore, lower air density and atmospheric pressure occurring at higher altitudes produced lower CO<sub>2</sub> levels and slower transpiration rates, which eventually led to lower rates of photosynthesis (Wang et al., 2017). Therefore, previous researchers assumed that the increase in leaf nutrient content of plants with

altitude could be considered as the plants' inability to use the absorbed resources for growth.

# Journal of Agricultural Science and Technology (JAST), 28(2)

# In Press, Pre-Proof Version

			Antioxidant activity							
Clone	Ν	Р	Κ	Ca	Mg	Fe	Al	TPC	DPPH	FRAP
	% mg/g								µg/mL	mM Fe(II)/g
1248	$4.22 \pm 0.7ab$	$11.11 \pm 0.4bc$	0.26±0.06b	$0.61 \pm 0.11$	0.22±0.07b	$12.77 \pm 0.8a$	$4.53 \pm 1.2b$	19.64±0.15	50.66±1.86c	2.10±0.2
2024	3.79 ± 0.2ab	$9.45 \pm 0.5c$	0.31±0.12b	$0.38 \pm 0.04$	0.14±0.06b	$6.01 \pm 1.2b$	$16.19 \pm 0.2a$	19.29±0.30	74.25±2.79a	$1.99 \pm 0.1$
AT53	$3.55 \pm 0.4ab$	$10.22 \pm 1.0$ bc	1.84±0.38a	$0.70\pm0.08$	0.80±0.16a	$12.97 \pm 1.4a$	$16.61 \pm 1.4a$	19.29±0.26	74.36±4.99a	1.55±0.3
TV9	$3.99 \pm 0.6ab$	$11.85 \pm 0.9ab$	0.15±0.04b	$0.68 \pm 0.09$	0.11±0.03b	$10.26 \pm 0.6a$	$3.66 \pm 0.7b$	19.30±0.04	73.89±4.31a	$1.86\pm0.3$
663	$2.97 \pm 0.2b$	$13.76 \pm 1.06a$	0.55±0.09b	$0.58\pm0.13$	0.27±0.06b	$3.32 \pm 0.8 bc$	$4.43\pm0.9b$	19.03±0.21	62.90±4.93abc	$1.90\pm0.5$
2026	$4.39 \pm 0.2a$	$9.19 \pm 1.2c$	0.47±0.11b	$0.64\pm0.12$	0.22±0.003b	$3.43 \pm 1.3 bc$	$3.72 \pm 0.8b$	19.23±0.20	72.49±1.97ab	$1.65 \pm 0.3$
1294	$4.57 \pm 0.7ab$	$11.06 \pm 0.5 bc$	0.24±0.03b	$0.38 \pm 0.03$	0.23±0.04b	$2.67 \pm 0.4c$	$3.91 \pm 1.7b$	19.04±0.10	54.61±3.64b	$1.76\pm0.5$
Mean	$3.93\pm0.2$	$10.95 \pm 1.6$	0.54±0.13	$0.57 \pm 0.04$	$0.29 \pm 0.05$	$7.35\pm4.4$	$7.58\pm5.8$	19.26±0.08	66.17±2.40	1.83±0.4
p-value	0.018*	0.000***	0.000***	0.116	0.000***	0.000***	0.000***	0.428	0.000***	0.551

Table 2. Foliar nutrient concentration and antioxidant activity of lowland tea plantation

The significance of the values is indicated as follows: \* P<0.05; \*\* P<0.01; \*\*\* P<0.001.

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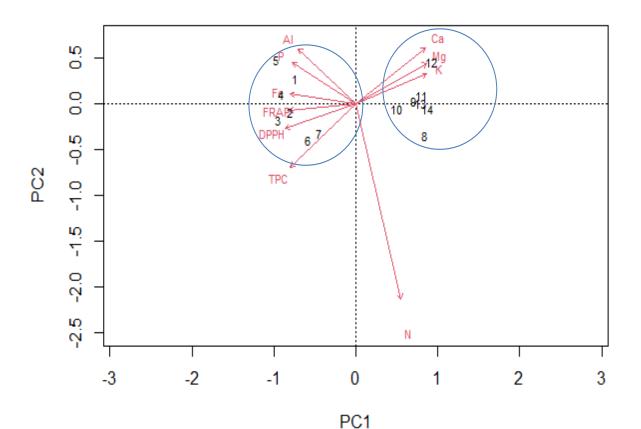
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Table 3. Foliar nutrient concentration and antioxidant activity of highland tea plantation.

				Antioxidant activity						
Clone	Ν	Р	Κ	Ca	Mg	Fe	Al	TPC	DPPH	FRAP
	% mg/g							mg GAE/g	µg/mL	mM Fe(II)/g
1248	4.46±0.1ab	8.35±0.8a	11.1±1.0ab	2.20±0.1b	1.28±0.1b	$0.057 \pm 0.003$	0.38±0.1ab	$18.80 \pm 0.94$	140±1.15a	1.190±0.04a
2024	4.47±0.3ab	7.98±0.2ab	12.2±0.5ab	2.42±0.1b	1.59±0.1ab	$0.057 \pm 0.01$	0.30±0.1b	17.34±0.16	141±2.38a	0.975±0.05ab
AT53	5.18±0.1a	6.74±1.0ab	10.3±1.4ab	2.63±0.2b	1.74±0.2ab	$0.057 \pm 0.01$	0.36±0.1ab	$18.80 \pm 0.49$	133±3.53ab	0.851±0.05b
TV9	4.31±0.6ab	5.78±0.5b	10.8±0.5ab	2.52±0.3b	$1.46\pm0.02$	$0.050 \pm 0.004$	0.44±0.1ab	18.35±0.56	132±2.34ab	0.989±0.11ab
TBR2020	3.84±0.1b	6.22±0.3ab	10.9±1.0ab	3.89±0.3a	1.78±0.1a	$0.053 \pm 0.01$	0.69±0.1a	16.97±0.16	127±2.99b	0.932±0.05ab
196	4.59±0.1ab	6.71±0.3ab	13.4±0.3a	2.62±0.2b	1.91±0.1a	$0.060 \pm 0.01$	0.51±0.1ab	$17.80 \pm 0.50$	131±0.58ab	1.180±0.10a
664	4.74±0.1ab	6.73±0.1ab	8.55±0.6b	2.25±0.1b	1.47±0.1ab	$0.047 \pm 0.003$	0.22±0.02b	$16.58 \pm 0.03$	129±3.43ab	0.801±0.01b
Mean	4.51±0.1	6.93±0.3	11.03±0.4	2.65±0.1	$1.60\pm0.1$	$0.054 \pm 0.003$	$0.41\pm0.04$	16.81±0.31	133±1.35	1.12±0.07
p-value	0.049*	0.0454*	0.0347*	0.000***	0.009**	0.741	0.022*	0.051	0.014*	0.006**

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The significance of the values is indicated as follows: \* P<0.05; \*\* P<0.01; \*\*\* P<0.001.



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Figure 1. Biplots showing the distribution of 14 populations of tea along principal component
 axes 1 and 2 from PCAs summarizing variation in foliar nutrient elements and antioxidant
 activity assay (Note: 1–7 = lowland clones, 8–14 = highland clones).

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Table 4. Soil physicochemical properties of lowland tea plantation.

Clone	Ν	Р	Κ	Ca	Mg	Fe	Al	Soil pH
Cione	%			mş	g/g			
1248	$0.11 \pm 0.009$	0.20±0.030ab	1.04±0.076ab	$0.37 \pm 0.018$	0.35±0.029a	9.45±0.31abc	14.2±1.78b	4.35±0.25ab
2024	$0.12\pm0.012$	0.12±0.038b	0.83±0.118bc	$0.23 \pm 0.023$	$0.20\pm 0.006b$	6.16±0.61c	20.2±1.68ab	4.27±0.03ab
AT53	$0.08\pm0.028$	0.19±0.034ab	0.51±0.026cd	$0.33 \pm 0.003$	$0.18 \pm 0.009 b$	6.52±0.80bc	16.2±1.75ab	3.83±0.10b
TV9	$0.10\pm0.028$	0.30±0.047ab	1.17±0.030a	$0.28\pm0.200$	0.22±0.009b	8.69±0.96abc	20.8±1.93ab	3.86±0.15b
663	$0.11 \pm 0.003$	0.37±0.037a	1.08±0.094ab	$1.04\pm0.401$	0.39±0.027a	10.20±0.83a	22.1±1.78a	4.80±0.21a
2026	$0.16\pm0.021$	0.09±0.012b	0.39±0.021d	$0.29 \pm 0.009$	$0.18 \pm 0.010 b$	9.78±0.98ab	23.8±1.41a	4.18±0.10ab
1294	$0.11 \pm 0.030$	0.37±0.090a	0.77±0.030bc	$0.54 \pm 0.074$	0.19±0.012b	7.27±0.51abc	18.1±1.22ab	4.35±0.25ab
Mean	$0.11 \pm 0.007$	$0.24 \pm 0.029$	$0.83 \pm 0.065$	$0.44\pm0.08$	$0.24 \pm 0.019$	8.30±0.41	19.33±0.87	4.23±0.09
p-	0.166	0.003**	0.000***	0.058	0.000***	0.007**	0.012*	0.022*
value								

The significance of the values is indicated as follows: \* P<0.05; \*\* P<0.01; \*\*\* P<0.001.

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	Table 5. Soli physicochemical properties of highland tea plantation.										
Clone	Ν	Р	Κ	Ca	Mg	Fe	Al	Soil pH			
Clotte	% mg/g										
1248	0.23±0.04ab	6.63±0.8a	0.83±0.03ab	$1.44\pm0.13$	1.03±0.07ab	15.5±2.1b	20.8±3.4ab	3.35±0.2			
2024	0.42±0.05ab	2.16±1.8ab	0.29±0.19b	$1.56\pm0.15$	0.59±0.22ab	8.70±4.0b	8.64±4.2bc	3.77±0.2			
AT53	0.27±0.02ab	1.50±0.1b	0.16±0.03b	$1.58\pm0.02$	0.41±0.05b	3.56±0.2b	7.68±0.4c	3.41±0.5			
TV9	0.41±0.08ab	4.89±2.3a	0.51±0.12b	$2.16\pm0.56$	0.62±0.13ab	28.4±0.5a	21.3±1.1ab	3.57±0.2			
663	0.14±0.06b	1.67±0.2b	2.83±1.15a	$1.23\pm0.16$	3.10±1.27a	29.9±4.5a	24.1±3.8a	$3.73 \pm 0.01$			
2026	0.49±0.04a	7.22±1.8a	0.69±0.12ab	$2.62 \pm 0.52$	1.28±0.46ab	12.3±1.6b	11.4±1.3abc	3.79±0.3			
1294	0.32±0.12ab	2.57±0.8ab	0.23±0.09b	$1.50\pm0.40$	0.40±0.15b	7.90±2.5b	4.81±1.4c	3.46±0.3			
Mean	0.32±0.03	3.80±0.6	0.79±0.24	1.73±0.15	$1.06\pm0.26$	15.19±2.3	14.12±1.8	3.58±0.1			
p-	0.032*	0.039*	0.011*	0.128	0.029*	0.000***	0.000***	0.857			
value											

Table 5. Soil physicochemical properties of highland tea plantation.

241 The significance of the values is indicated as follows: \* P<0.05; \*\* P<0.01; \*\*\* P<0.001.

242

240

Table 6. Pearson correlation coefficient (r) between soil nutrients and foliar nutrientconcentration and AOA from both plantations.

Pearson (	Correlation								
Foliar	Foliar P	Foliar	Foliar	Foliar	Foliar	Foliar	TPC	DPPH	FRAP
Ν		Κ	Ca	Mg	Fe	Al			
0.55*	-0.67**	0.83**	0.64*	0.76**	-0.73**	-0.62*	-0.67**	-0.79**	-0.60*
0.38	-0.56*	0.80**	0.61*	0.69**	-0.65*	-0.56*	-0.50	-0.74**	-0.38
-0.46	0.07	-0.02	0.27	-0.004	0.08	0.01	-0.12	0.09	0.13
0.41	-0.61*	0.90**	0.77**	0.86**	-0.80**	-0.69**	-0.71**	-0.82**	-0.64*
-0.05	-0.45	0.57*	0.77**	0.59*	-0.47	-0.40	-0.60*	-0.49	-0.39
-0.15	-0.42	0.45	0.61*	0.41	-0.40	-0.39	-0.42	-0.41	-0.30
060*	0.34	-0.40	-0.26	-0.47	0.26	0.25	0.44	0.38	0.49
-0.62*	0.81**	-0.76**	-0.72**	-0.76**	0.53*	0.55*	0.59*	0.84**	0.78**
-0.51	$0.78^{**}$	-0.96**	-0.86**	-0.91**	$0.80^{**}$	$0.71^{**}$	$0.77^{**}$	$0.92^{**}$	$0.71^{**}$
0.47	0.01	-0.04	-0.30	-0.02	0.04	0.05	0.10	-0.02	-0.09
	Foliar N 0.55* 0.38 -0.46 0.41 -0.05 -0.15 060* -0.62* -0.51	N           0.55*         -0.67**           0.38         -0.56*           -0.46         0.07           0.41         -0.61*           -0.05         -0.45           -0.15         -0.42          060*         0.34           -0.62*         0.81**           -0.51         0.78**	$\begin{array}{c ccccc} Foliar & Foliar P & Foliar \\ N & K \\ \hline 0.55^{*} & -0.67^{**} & 0.83^{**} \\ 0.38 & -0.56^{*} & 0.80^{**} \\ -0.46 & 0.07 & -0.02 \\ 0.41 & -0.61^{*} & 0.90^{**} \\ -0.05 & -0.45 & 0.57^{*} \\ -0.15 & -0.42 & 0.45 \\060^{*} & 0.34 & -0.40 \\ -0.62^{*} & 0.81^{**} & -0.76^{**} \\ \hline \end{array}$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	$\begin{array}{c c c c c c c c c c c c c c c c c c c $	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$

245 The significance of the values is indicated as follows: \* P<0.05; \*\* P<0.01; \*\*\* P<0.001.

Nitrogen is the most abundant nutrient available in tea foliar from both plantations, followed 246 by K. However, in some cases, tea may absorb more Ca than K (Hawkesford et al. 2011). This 247 is similar to our finding, especially in lowland plantations, where K ranked as the fourth 248 macronutrient after Ca. Sufficient K greatly boosts the yield and quality of tea, as it speeds 249 250 metabolism, triggers catechin synthesis, and promotes biotic and abiotic resistance by activating and governing several enzymes (Huang et al., 2022; Ruan et al., 2013). Meanwhile, Ca is 251 beneficial for improving plant resistance, enhancing photosynthesis capacity, and promoting 252 plant growth (Huang et al., 2022). Magnesium is also involved in activating and governing 253 several other physiological processes, such as photosynthesis, respiration and nucleic acid 254 metabolism (Pongrac et al. 2020). 255

Our study found that lowland plantations had a higher foliar Fe and Al content than clones derived from highland plantations. This is suggested due to the higher foliar P in lowland over highland plantation, which is supported by the PCA result (Figure 1), with both the arrow of P and Fe going in the same direction. Iron (Fe) is an important micronutrient for plants since it is involved in metabolic activities such as DNA synthesis, respiration, and photosynthesis. In contrast, Al is toxic to most plants since a micromolar dose of it may inhibit root growth

(Mahmud et al. 2024). However, tea plants are recognised as Al-hyperaccumulators, which was
initially coined for plants containing Al exceeding 1.0 mg per gram dry mass (Chenery, 1955).
However, distinguishing Al accumulators requires different thresholds depending on
geographic origin: tropical plants need higher levels (2.3 – 3.9 mg Al per gram of leaf dry mass
(Metali et al., 2012).

In tea plants, Al concentration varies between young and mature leaves, whereas young 267 leaves contain lower Al than mature leaves. Zhang et al. (2018) reported that the Al 268 concentrations in young leaves range from 0.25 - 0.66 mg g<sup>-1</sup>, while in mature tea leaves, they 269 range from  $4.3 - 10.4 \text{ mg g}^{-1}$ . A recent study revealed that Al concentrations in young leaves 270 were between 0.67 and 2.21 mg  $g^{-1}$  and in mature leaves between 2.63 and 7.83 mg  $g^{-1}$  (Zaman 271 et al. 2024). In our study, we found that young tea leaves are even able to accumulate up to 272 16.61 mg g<sup>-1</sup> or 8-fold higher than the result of Zaman et al. (2024). Al primarily enters the 273 plant root from acidic soils (pH < 5.0), where Al is solubilized into its toxic form (Al<sup>3+</sup>). Organic 274 acids like malate, citrate and oxalate are critical in Al detoxification. The formation of Al-275 organic acid complexes (such as Al-malate) is essential for their transport within the plant. 276 These complexes are less toxic and can be transported through the symplastic pathway and 277 loaded into the xylem for translocation to the shoots (Wang et al., 2017). Thus, we assume that 278 high Al accumulation of tea was due to the efficiency of detoxification and/or compartmentation 279 mechanisms. Otherwise, such high Al concentrations would be extremely toxic. In the leaves, 280 Hajiboland et al. (2013) revealed that up to 60% of Al in tea plants is stored in cell walls, 281 primarily by binding to pectin and hemicellulose components. A significant portion of Al is 282 sequestered in the vacuoles of tea leaves, reducing its toxicity to other cellular organelles (Gao 283 et al. 2014). In addition, Al can be deposited in vacuoles that exist as complexes with phenolic 284 substances, such as catechin (Barceló and Poschenrieder, 2002). P is among its components. 285 Therefore, high Al uptake is generally coupled with high uptake of P, which was supported by 286 our PCA result (Figure 1). 287

In terms of AOA, clones grown at a lower elevation tend to have a superior antioxidant, as demonstrated by greater TPC and FRAP values and lower DPPH IC50 values. This result was similar to a previous study conducted in Africa (Owuor et al., 2011), Taiwan (Chen et al. 2014), Indonesia (Martono et al., 2016), and China (Wang et al., 2022). Altitude increases are generally linked with decreases in tea polyphenol content (Ahmed et al. 2019), whereas catechin is its major constituent. Catechin was reported to be inversely correlated to cultivation altitude as the EGCG (major catechin derivative) declined when the cultivation altitude was elevated (Chen

et al. 2014). Wang et al. (2022) confirmed the inhibition of catechin biosynthesis in high 295 altitudes by stimulating the changes induced by temperature and light at different altitudes. 296 Light intensity plays a major role in influencing the catechin content of tea. Xiang et al. (2021) 297 proved that the catechin content and photosynthetic capacity of tea plants increased under 298 optimum light intensities (250  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup> – 350  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>). However, it will decrease under 299 shading treatment (150 µmol m<sup>-2</sup> s<sup>-1</sup>) or extreme high light intensity in highland plantation (550 300  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>). Lowland-grown tea may produce up to 28% higher polyphenols compared to 301 highland (Zhang et al., 2018). 302

303

#### 304 Association of Nutritional Characteristics and AOA with Soil Edaphic

The Principal Component 1 (PC1) of the soil properties (nutrient concentrations and pH) 305 displayed a positive and significant correlation with foliar P, Fe, Al, and AOA (TPC, DPPH, 306 FRAP). In PCA, PC1 is the first principal component that captures the maximum variance in 307 the data. This suggested that PC1 represent soil fertility and nutrient availability gradient that 308 influences tea nutritional composition and antioxidant activity. A strong correlation between 309 PC1 and foliar P or Fe suggested that PC1 reflects a soil axis related to P and Fe availability, 310 essential for plant metabolism and enzymatic functions (Bhat et al. 2024). The presence of Al 311 in the correlation might indicate soil acidity, which is common in tropical soil, further affects P 312 availability and metal uptake (Ur Rahman et al., 2024). Higher foliar TPC, DPPH, and FRAP 313 linked to PC1 suggested that increasing soil P, Fe, and Al may enhance the synthesis of phenolic 314 compounds, boosting antioxidant activity. This could be due to plant stress responses to metal 315 presence (Fe and Al), increasing secondary metabolites such as phenolics. This suggests 316 nutrient availability and potential soil stressors (such as Al) are key to plant biochemical 317 composition. 318

319 Based on our analysis, total soil N was the most important element for tea growth and development, as it had a positive association not just with foliar N but also with foliar K, Ca, 320 and Mg, as well as a negative correlation with foliar Fe and Al. In addition, total soil P, Ca, and 321 Mg showed a substantial association with foliar K, Ca, and Mg, indicating that variations in 322 323 tea's nutritional characteristics were influenced by the availability of nutrients. The results were similar to those of previous studies conducted in China (Zhao et al., 2017) and Taiwan (Tseng 324 and Lai, 2022). Nitrogen is an essential nutrient for tea production and accounts for 4-5% of tea 325 leaf dry weight (Hamid et al. 2014). Along with Mg and Mn, it up-regulated the expression of 326

key genes for chlorophyll synthesis and promoted its synthesis (Chen et al. 2021), thus resultingin a proportional increase in economic yield (Sitienei et al. 2013).

Soil pH was the only soil parameter significantly correlated with foliar nutrients (P, Fe, and 329 Al) and AOA of tea (TPC, DPPH, and FRAP assay). Soil pH from both plantations was highly 330 acidic, with an average value of 4.23 and 3.58 for lowland and highland, respectively. Tea has 331 also been planted on acidic soil in Vietnam (pH 3.7-3.9) (Huu Chien et al., 2018), Taiwan (pH 332 3.5 - 5.21) (Tseng and Lai, 2022) and China (pH 3.96-5.48) (Yan et al., 2020). Soil pH 333 significantly affects the availability of foliar P concentration. P tends to form insoluble 334 complexes with Fe and Al in acidic soils, reducing availability. In contrast, as pH increases 335 towards neutral, P becomes more available for plant uptake (Baguy et al., 2024). 336

The increased Fe and Al uptake in tea plantations with increased soil pH was rare. Typically, 337 Fe and Al solubility and availability decrease as soil pH increases (Ruan et al., 2006; Alekseeva 338 et al., 2011). However, several mechanisms could explain this phenomenon. Tea plants have 339 adapted to acidic soils and may release organic acids and chelating compounds from their roots, 340 such as malate, citrate and oxalate. These compounds can solubilize Fe and Al even at higher 341 pH by forming metal-organic complexes that remain plant-available. For instance, the 342 formation of al-malate is essential for transport within plants since these complexes are less 343 toxic. Subsequently, they can be transported via the symplastic pathway and loaded into the 344 xylem for translocation to the leaves (Wang et al., 2017). Another mechanism includes 345 microbial activity changes. An increase in soil pH could shift the microbial community 346 structure. The shift had the potential to enhance siderophore-producing organisms that make Fe 347 more available to plants (Choi et al., 2024). 348

Higher soil pH has been associated with increased enzymatic activities in tea plants, which 349 350 could enhance antioxidant defences. For instance, increasing soil pH from 3.3 to 5.3 enhances the nutrient availability in the rhizosphere. Subsequently, improving pH facilitates the uptake 351 352 of essential elements, including C, K, Ca, Mg, Mn, P and S, which play a vital role in phenolic biosynthesis (Jia et al. 2024). The antioxidant activity, measured by DPPH and FRAP assay, 353 354 tends to increase with higher soil pH. The improved nutrient uptake and enhanced photosynthetic capacity under high pH conditions contribute to accumulating phenolic 355 compounds, potent reducing agents. This results in higher FRAP values, indicating better 356 357 antioxidant potential (Jahan et al., 2022; Jia et al., 2023).

358

#### 359 CONCLUSIONS

We conclude that altitude and soil physicochemical properties are among the factors that 360 influenced the variation of nutritional characteristics and the AOA of tea. Most foliar tea 361 nutrients varied significantly among clonal teas, except Ca in the lowlands and Fe in the 362 highlands. The highland tea population tended to have higher foliar nutrient concentrations, 363 while the lowland population had better AOA performance. AT53 and 1248 were considered 364 promising clones for having higher foliar nutrients and better AOA performance, respectively. 365 Regarding soil edaphic, we found that soil nutrients and pH displayed a significant correlation 366 with foliar nutrients, while soil pH was also significantly associated with the AOA of tea. 367

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#### 369 ACKNOWLEDGEMENTS

The authors are grateful to BOH Plantation Sdn. Bhd. for their invaluable cooperation and support in providing the sample subject for the research study. This study was funded by Geran Putra Universiti Putra Malaysia (grant number GP/IPM/2020/9690400) and Tadom Ecoliving Research Grant (grant number 6300421-12038).

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