

Original Research Article

Growth, Carbon Assimilation and Quality of Kesum (*Persicaria Minor*) as Exposed to Zinc Oxide Nanoparticles

ABSTRACT

Aims: This study was conducted to investigate the effect of zinc oxide nanoparticles towards the *Persicaria minor* that can be used as a guidance for further toxicity investigation of ZnO-NPs.

Study design: A Completely Randomized Block Design (RCBD) was used with three replication. Each unit was consisted with eight plants and the total of 96 plants were used in this study.

Place and Duration of Study: This study was conducted in plot 1, Vegetables Field plot for Teaching and Research, Taman Pertanian Universiti, Universiti Putra Malaysia (UPM) Selangor, Malaysia, from May 2018 until August 2018.

Methodology: *Persicaria minor* were exposed to four different concentration of zinc oxide nanoparticles (ZnO-NPs) which were (50,100 and 150 mg/L) and 0 mg/L as a control. The ZnO-NPs was dissolved in distilled water before being applied to plants. 40 mL of ZnO-NPs solution was applied to each plant. The growth, carbon assimilation and also secondary metabolites were measured in this experiment.

Results: The results showed that the treatment of zinc oxide nanoparticles enhanced growth of the *Persicaria minor* as the plant treated with zinc oxide nanoparticles ~~have had~~ higher plant height and total biomass when compared to control treatment. However, the analysis revealed that the treatment of zinc oxide nanoparticles highly and significantly influenced the carbon assimilation and quality of this plant as the treated plants showed reduction in chlorophyll content, photosynthesis rate, stomatal conductance and transpiration rate but increased in production of secondary metabolites. The increased in production of plant secondary metabolites may be attributed by the plant protection mechanism due to metabolic stress caused by high concentration of zinc oxide nanoparticles.

Conclusion: This research will progressively help in contributing some reliable and valid data on the effect of zinc oxide nanoparticles (ZnO-NPs), towards the *Persicaria minor* that can be used as guidance for further experimental investigation regarding this field.

Keywords: *Persicaria minor*, zinc- oxide nanoparticles, growth, carbon assimilation, secondary metabolites, toxicity

1. INTRODUCTION

According to National Health Portal [1], the World Health Organization (WHO) estimated that 80 percent of the world's population ~~consuming-consume~~ herbal medicine for their health care and there are about 21000 plants species ~~have-with the~~ potential to be utilized as medicinal plants. *Persicaria minor* is one of the plants that gained great attention in this field of study. According to Christopher et al. [2], *Persicaria minor* have gained great attention in scientific study due to its high content of antioxidant. This plant possesses variety of

23 pharmacological properties such as antioxidant activity, antiulcer activity, anti-inflammatory
24 activity, antimicrobial activity, anticancer activity and can enhance the digestive properties
25 and cytotoxic activity [2]. According to Vikram et al. [3], *Polygonum minus* has been used
26 traditionally in herbal medicine to treat digestive disorder, remove dandruff and the essential
27 oil that extracted from *Persicaria minor* leaves is used as aroma therapy and also in perfume
28 industry.

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30 Recently, nanoparticles (NPs) are widely studied because its beneficial properties in
31 agriculture and allied sector [4]. According to them, zinc oxide nanoparticles (ZnO-NPs) is
32 one of NPs ~~that~~ used widely as it ~~has~~ been utilized in variety of industrial sector including
33 medication, cosmetic materials, opposed microorganisms and textile industries. As it has
34 been commercially used, the toxicity effect of these ZnO-NPs to the environment and also
35 soil ecosystem are ~~of mainly concerned~~ [5]. Sabir et al. [6] stated that ZnO-NPs possess
36 significant characteristics which have antimicrobial, optical and physical properties therefore
37 it have great potential to enhance agriculture. The presence of ZnO-NPs may enhance the
38 antioxidant mechanism that helps to stabilize the plants and improve the photosynthetic
39 efficiency [7]. However, the effect ~~is~~ depends on the concentration of ZnO-NPs and ~~it is~~
40 varies from plant to plant [8].

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42 Secondary metabolites are the natural compound that produced by the plants and some of
43 the compound are utilized as medicines, flavoring and drugs. According to Biology
44 Reference [9], the simple classification of plants' secondary metabolites includes three main
45 groups which are terpenes, phenolics and nitrogen- containing compounds. Secondary
46 metabolites does not involve in the plant growth and development but required for plant to
47 survive in the environment because they give negatives impact on other organisms such as
48 pathogen and herbivores that can harm the plants [9]. Secondary metabolites possess
49 significant biological properties and also medicinal importance that can improve
50 pharmaceuticals field [10].

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52 There is no previous study ~~being~~ conducted on the effect of ZnO-NPs on physical and
53 biochemical response of *Persicaria minor* and ~~also~~ only ~~few limited~~ research ~~being has been~~
54 carried out ~~to discover~~ about the effect of ZnO-NPs on carbon assimilation and production of
55 secondary metabolite of this plant, ~~so~~ ~~So~~ this present study will reveal about the effect of
56 ZnO-NPs towards the *Persicaria minor* that can be ~~the of~~ guidance to improve the field of
57 agriculture in planting this important medicinal plant, ~~and also~~ other important herbs that
58 ~~being importantly have been~~ used recently in medicinal field and ~~also being~~ used as
59 guidance for further toxicity investigation of ZnO-NPs. Hence, the objectives of this study
60 were to study the growth, carbon assimilation and quality of *Persicaria minor* as affected by
61 zinc oxide nanoparticles, to determine the optimum concentration dose of zinc oxide
62 nanoparticles that can enhance the optimum growth and secondary metabolites of *Persicaria*
63 *minor* and to recognize the relationship between secondary metabolites and growth of
64 *Persicaria minor* as exposed by zinc oxide nanoparticles application.

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67 2. MATERIAL AND METHODS

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71 2.1 Experimental site

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73 This study was conducted in ~~plot-Plot~~ 1, Vegetables Field ~~plot-Plot~~ for Teaching and
74 Research, Taman Pertanian Universiti, Universiti Putra Malaysia (UPM) Selangor. The
75 research site was set up with net shading and black plastic to reduce the absorption of water
76 by sunlight since *Persicaria minor* require high amount of water and also to reduce the
77 competition with grasses and other plants. This experiment was conducted from the month
78 of May 2018 until August 2018.

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80 2.2 Planting material

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82 *Persicaria minor* was obtained from Jabatan Pertanian, Serdang. The first internodes at the
83 bottom of the shoots with five number of leaves was cut in 5_cm in height for the shoot stem
84 cutting preparation. The shoot stem cuttings were immersed in the tap water overnight to
85 increase their turgidity which increases the speed of the germination. Then, the propagation
86 step were done in the trays with peat moss as the medium and the shoots stem cutting were
87 left for two weeks for the development of the root. Then, the plants were transferred into
88 standard polybag (16_cm x 30_cm) which was filled with top soil as the medium [11].

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90 2.3 Soil preparation

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92 The soil medium was obtained from Unit Herba, Taman Pertanian Universiti and top soil was
93 used as the medium for the *Persicaria minor* planting. The top soil was transferred into the
94 polybag until it ~~occupied-filled~~ three- quarter of the polybag

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96 2.4 Synthesis and properties of Zinc-Oxide Nanoparticles (ZnO-NPs)

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98 The ZnO-NPs was synthesized chemically in the Laboratory of BioPhysics of Physics
99 Department, Faculty of Science UPM by using sol gel method. ZnO-NPs is characterized as
100 solid, white odorless powdery. ~~In-At~~ room temperature, 200ml of ethanol was added ~~to~~ 0.2M
101 of zinc acetate dehydrate. Then, the mixture was stirred for two hours to obtain clear
102 solution. Then, 1.0 M sodium hydroxide (NaOH) was titrated into the mixture until the pH 9 ~~is~~
103 ~~was reached~~. The milky white slurry was obtained from the titration. To allow homogenous
104 mixing, the white slurry was stirred for one ~~more~~ hour ~~more~~. The sample was left for 24
105 hours to allow the complete hydrolysis and gelation. The sample was separated from its
106 solution. Then, the sample was filtrated to obtain white precipitate. The sample ~~is-was~~ dried
107 in an oven for 48 hours at 100°C. The dried sample was ~~grinded-ground~~ by mortar and
108 pestle to yield ZnO powder.

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110 2.5 Experimental design

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112 A Completely Randomized Block Design (RCBD) was used in this study with ~~three~~
113 replications. Each unit ~~was~~ consisted ~~with~~ eight plants and ~~the_a~~ totals of 96 plants were
114 used in this study. After a month, *Persicaria minor* was exposed to four different
115 concentration of zinc oxide nanoparticles (ZnO-NPs) which were (50,100 and 150 mg/L) and
116 0 mg/L as a control. The ZnO-NPs was dissolved in distilled water before being applied to
plants. 40 mL of ZnO-NPs solution was applied to each plant.

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2.6 Plant maintenance

The maintenance steps are very crucial to ensure the plant develops healthy and to avoid the plants from wilting or attacked by any disease that can cause the plants to die. At the early phase of cultivation, *Persicaria minor* were watered two times daily. The watering was unnecessary only when the heavy rain occur. This was to avoid the over watering to the plants that can interfere with the plants growth. The common insects that can-could interfere with the plants growth were removed quickly from the planting area.

2.7 Collection of data

The growth data collection was conducted-done once a week after the application of treatment for the plants growth parameter. The destructive analysis and leaf gas exchange of the experiment were conducted at the end of the experiment.

2.7.1 Plant growth measurements

The plant growth measurements were conducted to obtain data about-on height, number of leaf and stem, diameter of stem, root to shoot ratio and the chlorophyll content.

2.7.1.1 Plant height

The plant height was measured starting from the stem on the soil-ground surface until-to the highest shoot growth using measuring tape.

2.7.1.2 Plant basal diameter

The plant basal diameter was measured by using vernier caliper at the tips of the plants.

Comment [O1]: base??

2.7.1.3 Plant leaves number

The leaves of the *Persicaria minor* were counted manually in-every three weeks.

2.7.1.4 Chlorophyll content measurement

The total chlorophyll content of the leaves were-was measured by using chlorophyll meter (SPAD 502). The leaves of the plants in each treatment for each replication were clipped by chlorophyll meter clipper to obtain the reading.

2.7.1.5 Plant fresh weight measurement

The plants were removed first from the soil and all the dirt were removed under the-flowing tap water. Then, the shoot and the root parts were separated for further analysis and all the plants parts were weighted separately using analytical balance.

2.7.1.6 Dry weight (biomass) measurement

The plants were dried in the oven at 60°C for 48 hours. Then, the measurements were recorded as observed by using electronic weighing scale.

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2.7.1.7 Root to shoot ratio

The root to shoot ratio was determined by dividing the weight of the roots part to the shoot part after the oven ~~dried~~-drying process.

3.7.1.8 Plant leaf temperature determination

The Infrared (IR) Thermometer was used to measure the plant leaves temperature.

2.7.2 Leaf gas exchange measurement

LI-6400XT (Li-COR Inc; Nebraska; USA) portable photosynthesis system was used to measure the leaf gas exchange. This equipment was warmed and was calibrated with ZERO IRGA mode for 30 minutes. The measurement was set at optimum condition which were 400 $\mu\text{mol mol}^{-1} \text{CO}_2$, 30°C cuvette temperature, 60% relative humidity with the rate of air flow set at 500 $\text{cm}^3 \text{min}^{-1}$ and then the cuvette condition was modified at 800 $\mu\text{molm}^{-2}\text{s}^{-1}$ photosynthetically photon flux density (PPFD). The measurement process of gas exchange was carried out between 9.00 am to 11.00 am by using fully expanded young leaves that give the measurement of net photosynthesis (A), stomata conductance (gs) and transpiration rate (E). Water use efficiency (WUE) was measured by using the formula of net photosynthesis dividing with transpiration rate. This is automatic operation and the results were saved in the the LI-6400XT console and Photosyn Assistant Software (Dundee Scientific, Dundee, UK) was used to analyze it. Precautions were taken to avoid mistakes during taking the measurements.

2.7.3 Total Phenolics and Flavonoids Quantification

Firstly, grounded plant tissue samples (0.1g) were extracted with 80% ethanol (10mL) on an orbital shaker for 120 minutes at 50°C. The mixture was filtered and the filtrate was used for the measurement of total phenolics and total flavonoids content. The total phenolic content in the leaves sample was measured by the Follin-Ciocalteu reagent (SigmaAldrich, Missouri, USA; diluted 10-fold). The absorbance was measured at 725 nm. The data were expressed as mg g^{-1} gallic acid equivalent (mg GAE g^{-1} dry sample). The determination process of total falvonoids content was measured by mixing a sample (1 mL) with NaNO_3 (Sigma Aldrich, Missouri, USA; 0.3 mL) in a test tube that covered with aluminium foil. The mixture then was left for 5 minutes. Then, 10% AlCl_3 (Wako Pure Chemical Industries Ltd; Tokyo, Japan; 0.3 mL) was added followed by the addition of 1.0 NaOH (Kanto Chemical Co. Inc.; Hokkaido, Japan; 2 mL). Then, the absorbance was measured at 510 nm using a spectrophotometer with rutting as a standard (data were expressed as mg g^{-1} rutting dry sample).

2.7.4 Chlorophyll fluorescence determination

The chlorophyll fluorometer was used to measure the chlorophyll florescence of the *Persicaria minor*. The mature leaf tissue was obtained from the *Persicaria minor* plant that cultivated at ~~20°C-20°C~~ in glasshouse exposed with artificial light to give minimum photon flux density of $550 \mu\text{mol} \cdot \text{m}^{-2} \cdot \text{S}^{-1}$ for 16 h photoperiod and phothosynthetically active radiation were supplied at $250 \mu\text{mol} \cdot \text{m}^{-2} \cdot \text{S}^{-1}$ during 16 h photoperiod.

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2.8 Statistical Analysis

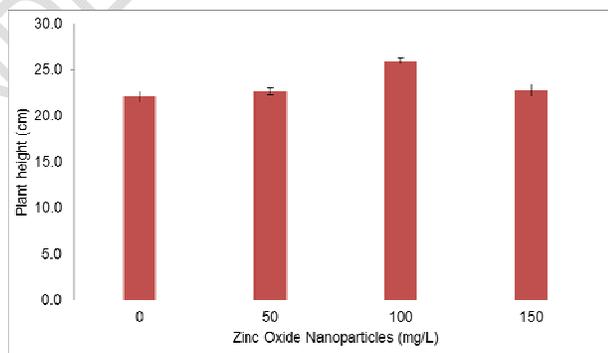
Statistical Package for Social Sciences (SPSS) version 24 was used to analyze the recorded data. A two-way ANOVA Test was conducted to analyze data for all the parameters used in the experiment. Results were significant if the p-value level ≤ 0.05 .

3. RESULTS AND DISCUSSION

3.1 Plant Height

Fig.1 ~~depicted~~ depicts the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the plant height of *Persicaria minor*. The result from analysis of variance showed that there was a significant effect between different concentration of zinc oxide nanoparticles treatment toward the height of *Persicaria minor* ($P=0.05$). From the figure, the increasing of concentration of zinc oxide nanoparticles treatment increased the plant height of *Persicaria minor*. The highest plant height was recorded in 100 mg/L on 12 weeks after treatment with mean 26.017 cm that might indicates the optimum concentration for the plant. Meanwhile, the control treatment recorded the shortest plant height with mean 22.083 cm.

The appropriate concentration of zinc oxide nanoparticles plays significant role in plant growth and promotion [12]. From the results, on twelfth week after harvesting, the plants treated with zinc oxide nanoparticles have higher plant height as compared to plants in control treatment. This finding indicates that the application of zinc oxide nanoparticles can induced the growth of the plants. Kouhi et al. [13] explained that the zinc oxide nanoparticles possess plant growth promoting effects and were used as micronutrient fertilizer, ~~as the~~ The presence of these nanoparticles triggered the physiological processes, acting as growth regulating compound that increased the plant growth parameters such as the plant height and biomass. In addition, Prasad et al. [14] also ~~supported~~ reported that zinc oxide nanoparticles possess beneficial effects in enhancing plant growth and development. The presence of zinc can enhance the biochemical, physiological and anatomical ~~responds~~ responses of the plants thus increased the plant growth such as plant height and biomass [15]. Therefore, it can be concluded that the zinc oxide nanoparticles treatment induced the plant growth and 100 mg/L can be considered as the best concentration, ~~among the~~ treatment rates used, in promoting the height of *Persicaria minor* plant.

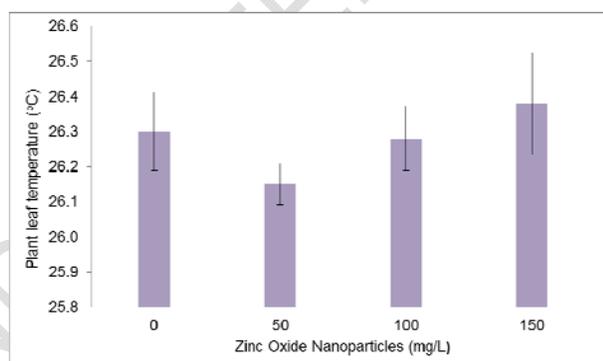


272 Fig.1. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on
273 plant height of *Persicaria minor*. Data are means with standard error of mean (SEM) of
274 24 replicates.
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278 3.2 Plant Leaf Temperature 279

280 Fig. 2 highlighted-highlights the effect of three months treatment of zinc oxide nanoparticles
281 (ZnO-NPs) on the total leaf temperature of *Persicaria minor*. The result from the analysis of
282 variance showed that there was a significant effect between different concentration of zinc
283 oxide nanoparticles treatment toward the total leaf temperature of *Persicaria minor* ($P=0.05$).
284 Based on the figure, the trend shows that the plant leaf temperature increased linearly with
285 the concentration of zinc oxide nanoparticles treatment, except for the control treatment.
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287 The plants maintaining their most important physiological process (photosynthesis) by
288 maintaining their average leaf temperature at around 21 degree celsius-Celsius [16]. The
289 plants leaf temperature depends on the stomatal conductance and transpiration rates of the
290 plants [17]. Transpiration is one of the best mechanisms used by plants to cool themselves
291 by 'pumping' out water from leaves through stomata [18]. From this study, the increasing of
292 plant leaves temperature can be explained through the reduction of the stomatal
293 conductance and transpiration rates of the plants due to the increasing in the concentration
294 of the treatment. This high temperature in turns will give negative effect to the
295 photosynthesis process thus affect the plant yields. Therefore, it can be concluded that the
296 zinc oxide nanoparticles treatment increased the plant leaf temperature due to the reduction
297 of stomatal conductance and also transpiration rate of *Persicaria minor*.
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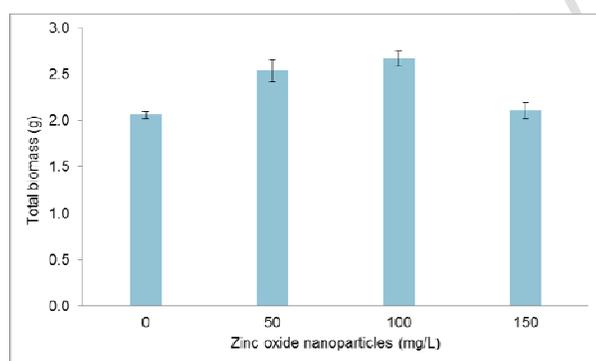
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314 Fig.2.The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on
315 plant leaf temperature of *Persicaria minor*. Data are means with standard error of
316 mean (SEM) of 24 replicates.
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318 3.3 Total Biomass 319

320 Fig.3 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on
321 the total biomass of *Persicaria minor*. The result from the analysis of variance showed that
322 there was a significant effect between different concentration of zinc oxide nanoparticles
323 treatment toward the total biomass of *Persicaria minor* ($P\leq 0.05$). From the results, it
324 showed that higher concentration of zinc oxide nanoparticles increased the total biomass of

325 *Persicaria minor* as plants treated with zinc oxide nanoparticles treatment have higher total
326 biomass as compared to the control treatment and 100 mg/L have the highest value of total
327 biomass with mean 2.675g.
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329 The highest value of total biomass indicates that 100 mg/L was the optimum concentration of
330 zinc oxide nanoparticles for *Persicaria minor*. This study showed that the treatment of zinc
331 oxide nanoparticles increased the plant biomass so the treatment might be effective in
332 boosting the plant growth and yield. Similar finding was observed from study conducted by
333 Venkatachalam et al. [19] that revealed total biomass significantly increased in the zinc oxide
334 nanoparticles treated plants as compared to control. This is also supported by findings of
335 Munir et al. [20] that stated the treatment of zinc oxide nanoparticles increased the shoot and
336 root dry weight thus increased the plant biomass. The presence of zinc can enhance the
337 biochemical, physiological and anatomical responds of the plants thus increased the plant
338 growth such as plant height and biomass [15]. Hence, it can be concluded that the presence
339 of zinc oxide nanoparticles can boost the *Persicaria minor* growth resulting in increasing of
340 the plant total biomass.
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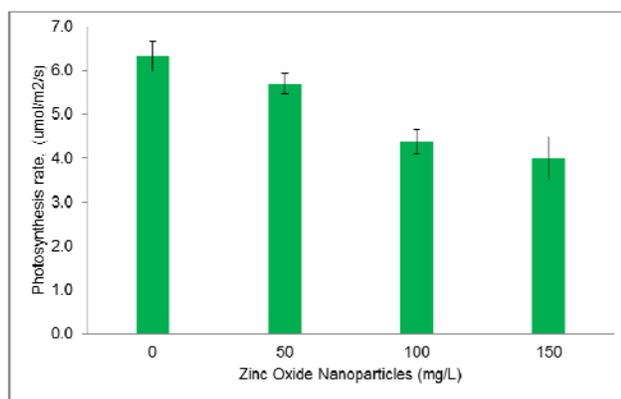
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357 **Fig.3.The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on**
358 **total biomass of *Persicaria minor*. Data are means with standard error of mean (SEM)**
359 **of 24 replicates.**
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361 3.4 Net photosynthesis rate (A)

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363 Fig.4 illustrated the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs)
364 on the net photosynthesis rate of *Persicaria minor*. The exposure of zinc oxide nanoparticles
365 toward the *Persicaria minor* was highly and significantly affected the net photosynthesis rate
366 of the plant ($P \leq 0.05$). From the result, the highest photosynthesis rate was recorded in
367 control treatment while the lowest photosynthesis rate was recorded in 150 mg/L of zinc
368 oxide nanoparticles treatment with mean 6.32 and 4 respectively. From the figure, the
369 photosynthesis rate of the plant reduced with the increasing concentration of zinc oxide
370 nanoparticles treatment.
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372 Photosynthesis is the perfect measurement to access plant performance. From this study,
373 the net photosynthesis of *Persicaria minor* was reduced with the increasing concentration of
374 zinc oxide nanoparticles treatment. Photosynthesis is highly affected in plants that exposed
375 to excess heavy metal where higher level of zinc oxide nanoparticles inhibits the
376 photosynthetic apparatus and caused critical changes to chlorophyll structure and amount
377 [21]. This finding is also similar with to that of Wang et al. [22] that revealed the presence of

378 zinc oxide nanoparticles reduced the chlorophyll content in leaves thus reduced the
379 photosynthetic efficiency in plants. In addition, plants exposed to high concentration of zinc
380 oxide nanoparticles have low photosynthetic efficiency due to the reduction of chlorophyll
381 content and also damaged to the photochemical system. Therefore, it can be concluded that
382 the presence of zinc oxide nanoparticles reduced the photosynthetic efficiency of *Persicaria*
383 *minor* plants. This finding appears to contradict that on biomass accumulation above (3.3).
384 Would the plant be storing more to make up for the reduced photosynthetic efficiency? An
385 adaptation perhaps?
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404 **Fig.4: The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L net**
405 **photosynthesis rate of *Persicaria minor*. Data are means with standard error of mean**
406 **(SEM) of 24 replicates.**

407 3.5 Transpiration rate (E)

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410 Fig.5 depicted depicts the effect of three months treatment of zinc oxide nanoparticles (ZnO-
411 NPs) on the transpiration rate of *Persicaria minor*. The exposure of different concentration of
412 zinc oxide nanoparticles toward the *Persicaria minor* was highly and significantly affected the
413 transpiration rate of the plant ($P \leq 0.05$). From the figure, the transpiration rate of the plant
414 reduced with the increasing concentration of zinc oxide nanoparticles treatment. The
415 transpiration rate of 50 mg/L treatment was significantly higher with mean 2.342 while the
416 lowest transpiration rate of *Persicaria minor* was observed in 150 mg/L of zinc oxide
417 nanoparticles treatment with mean 1.42.
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419 Transpiration is a process of the movement of water through plant and this process mainly
420 take place in leaves. Transpiration process is mainly controlled by the opening and closing of
421 the stomata. From the result, the transpiration rate reduced with the increasing concentration
422 of zinc oxide nanoparticles treatment. This finding is similar with Xiaoping et al. [23] that
423 revealed both transpiration rate and stomatal conductance reduced in plants treated with
424 zinc oxide nanoparticles. The stomatal closure reduced the transpiration rate of the plants.
425 According to Vankova et al. [24], the presence of zinc oxide nanoparticles induced the
426 production of plant stress hormone, abscisic acid and this hormone mainly accumulated in
427 leaves. The higher level of abscisic acid triggered the stomatal closure which in turn reduced
428 the transpiration rate in plants. Hence, the application of zinc oxide nanoparticles reduced
429 transpiration rate of *Persicaria minor* due to the high level of stress hormone that cause the
430 closure of stomata.

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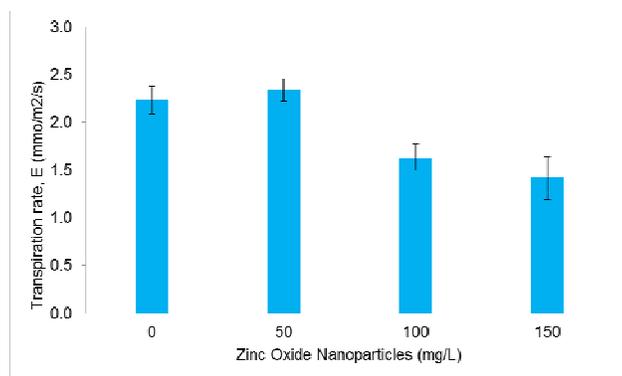


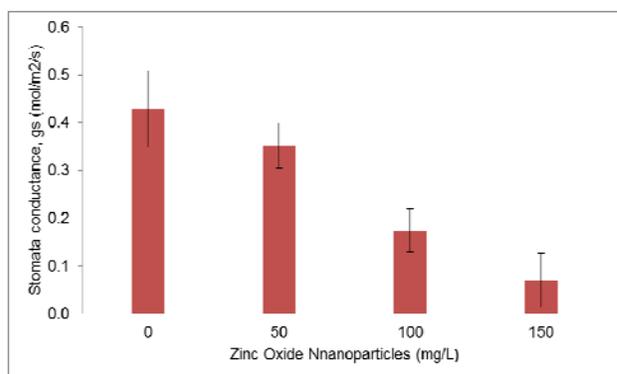
Fig.5. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L transpiration rate of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

3.6 Stomatal conductance (gs)

Fig.6 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the transpiration rate of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Polygonum minus* was highly and significantly affected the stomatal conductance of the plant ($P \leq 0.05$). From the figure, the stomatal conductance of the plant reduced as the concentration of the zinc oxide nanoparticles treatment increasing. The highest stomatal conductance was recorded in control treatment while the lowest stomatal conductance was observed in 150 mg/L of zinc oxide nanoparticles treatment with mean 0.428 and 0.07 respectively.

Stomatal conductance is a measure of the degree of stomatal opening and a good indicator in accessing plant water status [25]. The finding showed reduction in stomatal conductance due to the increasing of zinc oxide nanoparticles concentration is similar with Xiaoping et al. [23] that proved the higher concentration of zinc oxide nanoparticles reduced the stomatal conductance resulting in low photosynthetic efficiency of the plants. Singh and Bhati [26] also stated that high amounts of zinc oxide nanoparticles can restrict the stomatal conductance. This might due to the toxicity of the treatment disturbed the cell mechanism thus alters the stomatal function. Tsonev and Lidon [21] explained that the stomatal response to high concentration of zinc oxide nanoparticles is related to the changes in carbonic anhydrase (CA) activity. Carbonic anhydrase is an enzyme that is responsible for the stomatal activity and the presence of zinc oxide nanoparticles influenced the CA activity that triggered the stomatal closure thus reduced the stomatal conductance of the plants. Hence, it can be concluded that the presence of zinc oxide nanoparticles alter the stomatal mechanism thus reducing the stomatal conductance of *Persicaria minor* plants.

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Fig.6. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on stomatal conductance of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

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3.7 Maximum efficiency of photosystem II

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Fig.7 depicted the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the maximum efficiency of photosystem II of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Persicaria minor* was highly and significantly affected the maximum efficiency of photosystem II of the plant ($P \leq 0.05$). Based on the figure, increasing the concentration of zinc oxide nanoparticles caused the maximum efficiency of photosystem II to decrease. The highest value of maximum efficiency of photosystem II was observed in control treatment while the lowest value of maximum efficiency of photosystem II was recorded in 150 mg/L treatment with mean 0.758 and 0.522, respectively.

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The phytotoxicity of zinc oxide nanoparticles can be accessed through the efficiency of photosynthetic mechanism (chlorophyll fluorescence) that act as indicator in phytotoxicity assays. The finding of this study revealed that increasing the zinc oxide nanoparticles concentration resulting resulted in lower maximum efficiency of photosystem II of *Polygonum minus*. Wang et al. [22] stated that the treatment of zinc oxide nanoparticles reduced the chlorophyll fluorescence parameter and damaged the photochemical system. This finding can be explained further that the presence of zinc oxide nanoparticles induced the oxidative stress in plants and increase the production of reactive oxygen species (ROS) which alter the gene expression pathway thus reduced the chlorophyll fluorescence in plants. Therefore, it can be concluded that the zinc oxide nanoparticles treatment reduced the chlorophyll fluorescence parameters of *Persicaria minor* plants.

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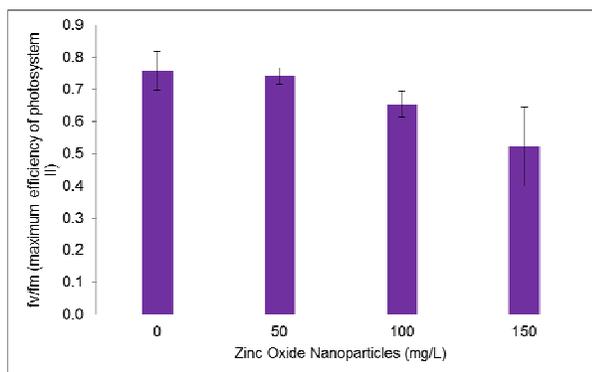


Fig.7. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on maximum efficiency of photosystem II of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

3.8 Maximum yield of photosystem II

Fig.8 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the maximum yield of photosystem II of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Persicaria minor* was highly and significantly affected the maximum yield of photosystem II of the plant ($P=0.05$). The trend shows that increasing the concentration of zinc oxide nanoparticles caused the maximum yield of photosystem II to decrease. Based on the figure, 50 mg/L of zinc oxide nanoparticles treatment shows the highest value of maximum yield of photosystem II with mean 3.166 when compared with other treatment.

The photosynthetic pigments (chlorophyll fluorescence) act as indicator in phytotoxicity assays in accessing the phytotoxicity of zinc oxide nanoparticles towards the plant. From this study, it was observed that the treatment of zinc oxide nanoparticles reduced the maximum efficiency of photosystem II which in turn reduced the maximum yield of photosystem II of *Persicaria minor*. According to Tsonev and Lidon [21], inside the chloroplast lamellae, the presence of zinc oxide nanoparticles caused the inhibition of photosynthetic electron transport and implicates the water evolving complex of photosystem II thus inhibits the photolysis and oxygen emission that disturb the conformation of photosystem II core complex. This mechanism explained how the zinc oxide nanoparticles treatment reduced the efficiency and yield of photosystem II in plants. Hence, the treatment of zinc oxide nanoparticles reduced the maximum yield and efficiency of photosystem II which in turn disturbed the photosynthetic process of *Persicaria minor* plant.

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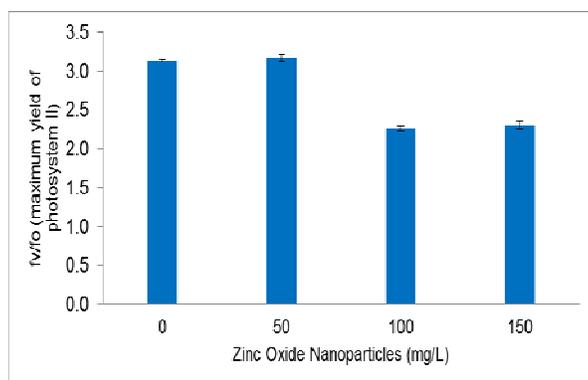


Fig.8. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on maximum yield of photosystem II of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

3.9 Minimal florescence

Fig.9 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the minimal florescence of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the Polygonum minus was highly and significantly affected the maximum yield of photosystem II of the plant ($P=0.05$). The result showed that higher concentration of zinc oxide nanoparticles treatment resulted in higher value of minimal florescence of *Persicaria minor*. The highest value of minimal inflorescence was recorded in 150 mg/L with mean 627.2 while the lowest value was observed in 50 mg/L with mean 462.6.

Higher minimal florescence indicates higher heat dissipation of plants. This might due to the presence of zinc oxide nanoparticles that induced stress in plants thus caused plants to produce high amount of heat. From this study, the treatment of zinc oxide nanoparticles reduced the transpiration rate of *Persicaria minor*. This reduction might be related with the increasing of minimal florescence of the plants. The high minimal florescence can cause heat stress to the plants. Heat stress is defined as the increase temperature beyond the threshold level that causes damage to plant growth and development [27]. Therefore, it can be deduced that the treatment of zinc oxide nanoparticles increased the minimal florescence of *Persicaria minor* due to the reduction in transpiration rate of the plants.

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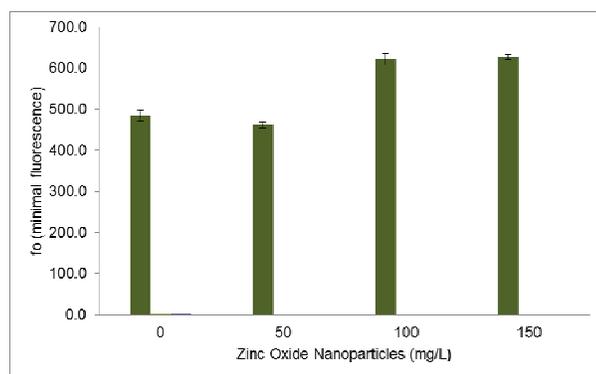


Fig.9. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on minimal fluorescence of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

3.10 Performance index (PI)

Fig.10 indicated the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the performance index of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Polygonum minus* was highly and significantly affected the performance index of the plant ($P=0.05$). Based on the figure, the highest and lowest performance index were observed in control treatment and 150 mg/L of zinc oxide nanoparticles treatment with mean 2.422 and 1.504, respectively. The performance index of *Persicaria minor* reduced when treated with higher concentration of zinc oxide nanoparticles that indicates zinc oxide nanoparticles increased the plant stress.

Define/explain PI here. Nanoparticles such as zinc oxide and silver were located on the surface of plants cells and induced the oxidative stress to the cells by the activation of oxidative stress signaling [28]. From this study, it was observed that the treatment of zinc oxide nanoparticles reduced the plants performance index. Zahed et al. [28] stated that the generation of reactive oxygen species (ROS) due to the zinc oxide nanoparticles treatment alter the gene expression and cell mechanism which in turn reduced the performance index of the plants. Wang et al. [22] explained that the toxicity of zinc oxide nanoparticles reduced chlorophyll content plants, resulted in low photosynthesis efficiency thus reduced the plants performance. Hence, it can be concluded that the presence of zinc oxide nanoparticles induced stress in *Persicaria minor* resulting in low performance index of the plants.

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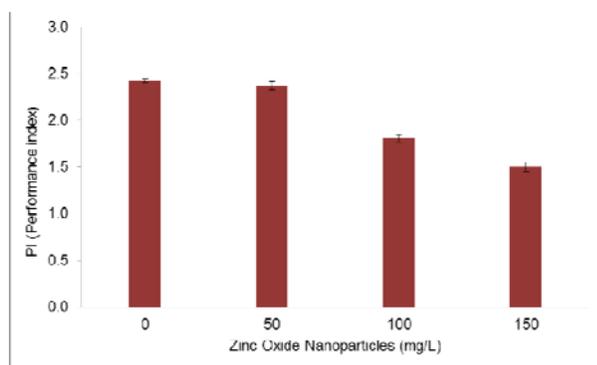


Fig.10. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on performance index of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates.

3.11 Total phenolics content

Fig.11 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the total phenolics production of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Polygonum minus* was highly and significantly affected the total phenolics production of the plant ($P \leq 0.05$). Based on the figure, the total phenolics production of the plant was directly proportional with the concentration of zinc oxide nanoparticles treatment. The lowest total phenolics production was recorded in control treatment while the highest total phenolics production was recorded in 150 mg/L treatment with mean 1.444 and 3.82 respectively. This result indicates that zinc oxide nanoparticles treatment induced stress and increased the secondary metabolites production of *Persicaria minor*.

Phenolics are compound that produced by plants to protect plants against stress. These compound play significant role in plant development (lignin and pigment biosynthesis) and also provided structural integrity for plant's support [29]. From this study, the greater production of total phenolics content in *Polygonum minus* with the increasing of concentration treatment revealed that the presence of zinc oxide nanoparticles induced stress towards the plants. This finding is supported by Rastogi et al. [30] that stated that zinc oxide nanoparticles treatment induced the Reactive Oxygen Species (ROS) production in plants thus increased plants stress. They also stated that higher concentration of zinc oxide nanoparticles lead to the damage of plant cell wall and plasma membrane thus induced the production of plant secondary metabolites for plants defense against disease and threat. Therefore, it can be concluded that higher concentration of zinc oxide nanoparticles lead to plant stress and boosted the plants secondary metabolites production which in turn enhanced the defense response of *Persicaria minor* plants.

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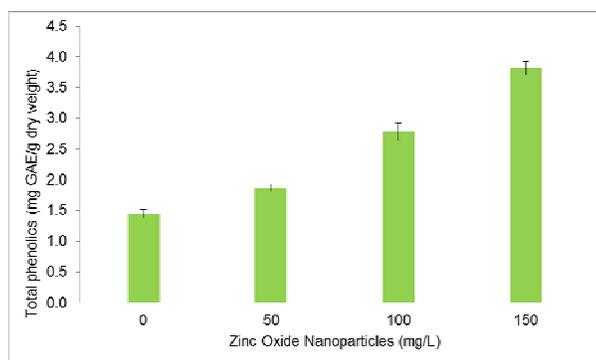


Fig.11. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on total phenolics production of *Persicaria minor*. Data are means with standard error of mean (SEM) of 24 replicates

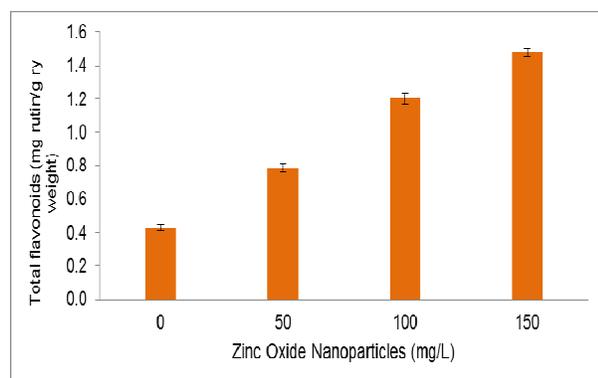
3.12 Total flavonoids content

Fig.12 shows the effect of three months treatment of zinc oxide nanoparticles (ZnO-NPs) on the total flavonoids production of *Persicaria minor*. The exposure of different concentration of zinc oxide nanoparticles toward the *Persicaria minor* was highly and significantly affected the total flavonoids production of the plant ($P=0.05$). Based on the figure, the total flavonoids production of the plant was directly proportional with the concentration of zinc oxide nanoparticles treatment. The lowest total flavonoids production was recorded in control treatment while the highest total flavonoids production was recorded in 150 mg/L treatment with mean 0.43 and 1.478 respectively. This result indicates that zinc oxide nanoparticles treatment induced stress and increased the secondary metabolites production of *Persicaria minor*.

Flavonoids are a wide group of plants chemicals (phytonutrients) that found mostly in fruits and vegetables. Flavonoids plays significant role in pharmacological field since this compound is a good source of antioxidant and anti-inflammatory, protect skin, enhanced brain function and also good for blood pressure regulation [31]. From this study, the greater production of total flavonoids content in *Polygonum minus* with the increasing of concentration treatment revealed that the presence of zinc oxide nanoparticles induced stress towards the plants. This finding is similar with that of Zafar et al. [32] who noted that the higher treatment concentration of zinc oxide nanoparticles generates oxidative stress of plants thus increasing the plant secondary metabolites production to protect plants against stress. The initial response of plants towards the presence of nanoparticles involved the increasing level of reactive oxygen species (ROS), cytoplasmic Ca^{2+} and up regulation of nitrogen activated protein kinase (MAPK) cascades thus activates the plants secondary metabolites that act against stress to protect the plants [33]. In addition, the presence of zinc oxide nanoparticles enhanced the expression of genes related to antioxidant capacity thus boost the defense mechanism of the plants by enhancing the production of plants secondary

Comment [O2]: M or N?

789 metabolites. Hence, it can be concluded that the presence of zinc oxide nanoparticles
790 enhanced the *Persicaria minor* secondary metabolites production by increasing the total
791 flavonoids production of the plants.
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810 **Fig.12. The effect of zinc oxide nanoparticles treatment (0, 50, 100 and 150) mg/L on**
811 **total flavonoids production of *Persicaria minor*. Data are means with standard error of**
812 **mean (SEM) of 24 replicates.**
813

814 4. CONCLUSION

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816 Overall, the treatment of zinc oxide nanoparticles increased the growth parameter of the
817 plants as the treated plants showed higher value of plant height and total biomass when
818 compared to plants in control treatment. However, the treatment of zinc oxide nanoparticles
819 reduced the photosynthesis rate, transpiration rate and stomatal conductance thus reduced
820 the performance index of *Persicaria minor* plants. The treatment also might have induced the
821 plants stress as it was significantly observed that the production of secondary metabolites
822 (total phenolics and flavonoids production) were directly proportional with the treatment
823 concentration that used mainly for plants production against stress. From this study, it can
824 be concluded that the optimum concentration of zinc oxide nanoparticles for enhancing the
825 *Persicaria minor* growth was 100 mg/L because it recorded the highest value for most of the
826 plants growth parameters.

827 Any recommendation/way forward?

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