The tolerance of oil palm (*Elaeis guineensis*) seedlings to Al stress is enhanced by citric acid and natural peat water

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Abstract. Hidayah AN, Yahya S, Sopandie D. 2020. The tolerance of oil palm (*Elaeis guineensis*) seedlings to Al stress is enhanced by citric acid and natural peat water. *Biodiversitas* 21: 4850-4858. Management technology on soil containing high levels of Aluminum (Al) toxicity is still needed to be developed so that the growth and development of plants will be optimum. The aims of the research were to investigate the response of oil palm seedlings (*Elaeis guineensis* Jacq.) toward aluminum stress, and to evaluate the effects of several exogenous compounds to improve the tolerance of oil palm. The research was conducted from September 2018 to March 2019 at PT Gunung Sejahtera Ibu Pertiwi, Central Kalimantan. This research consisted of two nutrient culture experiments, namely: Al toxicity on oil palms seedlings and the role of various exogenous compounds to improve plant tolerances. The results revealed that the solution at concentrations of 400 μM, 800 μM, and 1600 μM of Al significantly inhibited root growth, increased MDA levels, decreased the photosynthesis rate, activity of CAT and APX. Therefore, a solution at concentration of 400 μM of Al can be used as the selection level of Al tolerant oil palm varieties on nutrient culture. Ethephon at concentrations of 25 ppm, 50 ppm, and 100 ppm inhibited root and shoot growth, increased MDA levels but reduced the photosynthesis rate, chlorophyll content, APX, and CAT activity. Addition of 25 ppm and 50 ppm of citric acid, 200 ppm and 300 ppm of peat water significantly increased root length, root dry weight, photosynthesis rate, chlorophyll content, carotenoids, CAT, and APX activities as MDA levels decreased. Addition of citric acid and peat water enabled seedlings of oil palm to improve their tolerance to Al stress on nutrient culture.

Keywords: CAT activity, chlorophyll content, MDA level, photosynthesis rate, root growth

Abbreviations: APX: ascorbate peroxidase, CA: citric acid, CAT: catalase, DAT: day after treatment, Et: ethephon, HAT: hour after treatment, MDA: malondialdehyde, PW: peat water

INTRODUCTION

One of the constraints in oil palm (*Elaeis guineensis* Jacq.) cultivation is the poor management technology of plant cultivation in marginal land, thus the plants productivity is under their potential. The upland soil in Indonesia is dominated by acid mineral soil approximately 107.36 million ha (74.31%) (Ritung et al. 2015). This type of soil has the characteristics i.e low pH and cation exchange capacity, low base saturation and C organic, high Al saturation, high Fe and Mn content, subject to erosion, and deficient on macronutrients (especially P, K, Mg, and Ca) (Sopandie 2014). Efforts to improve cultivation on acid soil, however, still use soil ameliorant such as agricultural lime and dolomite fertilizer (Lubis 2008), which needs a lot of input per acre of land, and it is costly.

The diversity of oil palm in Indonesia is quite high, it is proven that 53 varieties have been registered with the Indonesian Ministry of Agriculture (Direktorat Perbenihan Perkebunan 2020). In addition, Setiawan (2017) stated that the Indonesian oil palm consortium in collaboration with the Government had introduced them from Cameroon in 2008 and from Angola in 2010 to increase their source of diversity. The tolerance level of oil palm varieties to Al stress and acid soils varies widely. Although Corley and Thinker (2015) pointed out that oil palm grows well in acid soils and able to accumulate of Al around 200 ppm on leaves and 600 ppm on roots, but lateral root growth is reduced at high aluminum levels and low pH (Cristancho et al. 2011). From those upland soil characteristics, high Al concentrations in soils are the main limiting factors for plant growth and development. Al toxicity reduced leaf size, plant height, and total dry weight of oil palm seedlings (Ramasari et al. 2016), primary root length, and sugar content in shoots and roots (Supena et al. 2014). In fact, one of the varieties planted in oil palm land with an Al saturation above 75% shows high flower and fruit abortion so that the productivity achievement is only about 70% of its potential. Therefore, it is necessary to develop more adaptive varieties through rapid and precise selection methods.

Plants treated with Al have a natural tolerance mechanism but their expression is often slow, so it is necessary to accelerate their internal mechanism through induction of signal transduction. One of the Al-tolerant varieties is Simalungun DxP (Supena et al. 2014), but its tolerance could possibly be increased by adding exogenous compounds. Several studies revealed that some chemical compounds could induce the signal transduction in plant which is thought to have a contribution to the change of...
gene expression, that leads to the increase of activity of several antioxidant enzymes such as APX, CAT, and GPX. These are very important enzymes in protecting the cell from oxidative damage by reactive oxygen species (ROS) resulted from Al toxicity. There are several chemical compounds that have induced the tolerance to Al stress, such as calcium on pepper plant (Yang 2014), citric acid on wheat plant (Sidak and Orabi 2015), ethephon on Kentucky bluegrass seedlings (Zhang et al. 2018), and peat water on Aloe vera (Chotimah et al. 2013). Sopandie (2014) pointed out that a short-term problem-solving approach can be through tolerance induction by using plant growth regulators (PGRs), hormone, osmoprotectant, Ca2+, and Si. Induction of tolerance can be defined as a complex change, involving a variety of different metabolic pathways until cellular and molecular level when it is responding to certain stress (Onaga and Wydra 2016).

Levitt (1980) mentioned that exogenous compounds (PGRs, hormone, osmoprotectant, Ca2+) could influence the transduction signal process that is involved in the gene expression change when the plants are stressed. This change will induce the activities of antioxidant enzymes, such as CAT, APX, and GPX that play a role in ROS scavenging, so that it can reduce MDA accumulation (Apel and Hirt 2004). The concept of making use of exogenous compounds in oil palm cultivation is expected to become an initial step to overcome aluminum stress. The aims of the research were to obtain a quick selection method for the aluminum stress-tolerant on oil palm at the seeding phase, oil palm growth, and physiological responses to stress and to evaluate the effects of several exogenous compounds to improve the tolerance of oil palm.

**MATERIALS AND METHODS**

**Plant materials**

Simalungun DxP variety seedlings obtained from the Indonesian Palm Oil Research Institute (IOPRI, Medan, Indonesia) were selected for their shape, color, and size. Planting was done by inserting a shell sprout into the planting hole where the radicle tip facing down and plumula to the top and then covered again with sand. After planting, media was watered with water in advance as much as 100 mL/polybag. Ten-week-old seedlings with similar growth were selected and transplanted into the hydroponic system with nutrient solutions (Sopandie, 1990). Nutrient culture composition as follows: 1.5 mM Ca (NO3)2·4H2O, 1.0 mM NH4NO3, 1.0 mM KCl, 0.4 mM MgSO4·7H2O, 1.0 mM KH2PO4, 0.068 mM Fe (C6H8O7), 0.50 ppm MnSO4·H2O, 0.02 ppm CuSO4·5H2O, 0.05 ppm ZnSO4·7H2O, 0.5 ppm H2BO3 dan 0.01 ppm NH4MoO4·2H2O. Nutrient solution was renewed every 5 days and the pH was kept constant (4.0 ± 0.1) every 2 days though addition of 0.3 M sodium hydroxide (NaOH) or 0.05 M hydrochloric acid (HCl), as required. This research was conducted at the Experimental Farm, PT Gunung Sejahtera Ibu Pertiwi, Central Kalimantan, Indonesia in shade house for 6 months (September 2019 - March 2020).

**Procedures**

**Aluminum toxicity to the growth of oil palm seedling stage on nutrient culture**

This experiment was arranged in a completely randomized design with various aluminum concentrations (0 μM, 100 μM, 200 μM, 400 μM, 800 μM and 1600 μM) with 3 replications. The addition of Al was done when transplanted and renewed of nutrient solution every 5 days. Determination of photosynthesis rate, malondialdehyde (MDA) level, ascorbate peroxidase (APX) enzyme activity, catalase (CAT) enzyme activity were carried out at 3 HAT (hours after treatment), 7 HAT, 24 HAT, and 72 HAT while growth measurements (root length, plant height, number of leaves, diameter of stem, root and shoot dry weight) at 30 days after treatment (DAT).

**The effects of various exogenous compounds in alleviating the adverse effect of Al on oil palm seedlings**

This experiment used a completely randomized design with the following treatments: 0 μM Al (control), 400 μM Al, Al + Ethephon (Et) 25 ppm, Al + Et 50 ppm, Al + Et 100 ppm, Al + citric acid (CA) 25 ppm, Al + CA 50 ppm, Al + CA 100 ppm, Al + peat water (PW) 100 ppm, Al + PW 200 ppm, Al + PW 300 ppm with 3 replications. Natural peat water is peat water from the peatlands of Bereng Bengkel, Central Kalimantan, Indonesia. The Al concentration used 400 μM (based on the results of Al toxicity experiments) and all treatments were given during transplanting and renewal of nutrient solutions every 5 days for new ten-weeks-old seedlings. Determination of MDA levels, APX, and CAT enzyme activity was done at 1 DAT and 3 DAT, while growth measurements (root length, plant height, number of leaves, diameter of stem, root and shoot dry weight, photosynthesis rate, contents of chlorophyll, carotenoid, and anthocyanin) at 30 DAT.

**Growth measurements**

Root length and plant height are recorded before transplanting into nutrient solution. Then, the root length, plant height, number of leaves, diameter of stems was measured at 30 DAT. Diameter of stem was measured using a caliper at the base of the stem. Roots and shoots were separated for fresh weight determination. All plant parts were washed in double-distilled water and dried in an oven at 70 °C for 10 h and the dry weights recorded with an electronic balance. The rate of photosynthesis was measured on the third leaf using CI-340 Handheld Photosynthesis System in open system mode according to schedule. Whereas the contents of chlorophyll (Chl), carotenoid, and anthocyanin in the third leaf were determined following the Sims and Gamon (2002) method. The third leaf sample was weighed 0.02 g ± 0.005 g and recorded. Mashed fresh leaf samples, added with acetone tris 2 mL 85:15 (%), and centrifuged (14000 rpm, for 10 minutes). Then, the supernatant (1 mL) was added with acetone tris (3 mL) and mixed evenly. Absorbance of the mixture is measured at wavelength 470, 537, 647, and 663 nm, measured with UV-VIS spectrophotometer. The result expressed as mol per 100 g wet weight (mol 100 g⁻¹).
**MDA level and CAT activity**

Determination of MDA levels and CAT activity used the method of Chen and Zang (2016). The first step, prepare the crude protein/ enzyme extract. As much as 0.2 g of the fresh leaf was weighed and ground them with a mortar and pestle in liquid nitrogen. The leaf powder homogenized by adding 3 ml of 100 mM PBS buffer. The homogenate assigned to two 1.5 ml centrifuge tubes and centrifuge at 10,000 x g for 20 min at 4 °C. The supernatant transferred to new centrifuge tubes for further analysis. The concentration of crude protein in the supernatant measured with Analytik Jena ScanDrop 250 by a spectrophotometric method with Formula Warburg-Christian (protein); protein concentration (mg/ml) = 1.55 x A280 - 0.76 x A260. For MDA assay, add 100 µL extract and 1 mL 0.25% thiobarbituric acid (TBA) were mixed and heated at 100 °C for 15 min, then quickly cooled in an ice bath (± 5 min). 1 mL 0.25% TBA solution with 100 µL 100 mM PBS (pH 7.8) serves as a reference. The absorbance of 200 µL supernatant was measured at 532 nm and 600 nm with Analytik Jena ScanDrop 250 respectively. The MDA level was expressed in nmol mg⁻¹ protein. Then for CAT assay, the reaction solution prepared by adding 77.5 µL 30% H₂O₂ in 50 mL 100 mM PBS (pH 7.0) (for 50 reactions). Add 50 µL crude enzyme and 1 mL reaction solution into the cuvette, and immediately record the dynamic absorbance at 240 nm with Analytik Jena ScanDrop 250 at every 15 sec for 1 min, looking for the steady average alteration. Reaction solution with 50 µL 100 mM PBS (pH 7.8) serves as a reference. The CAT activity was expressed in unit mg⁻¹ protein.

**APX activity**

Activity of APX was determined by the method of Nakano and Asada (1981) by recording the decrease in absorbance of ascorbate at 290 nm. The assay mixture contained phosphate buffer (50 mM, pH 7.0), 0.1 mM EDTA, 0.5 mM ascorbate, 0.1 mM H₂O₂, and the enzyme extract. APX activity was calculated by using the extinction coefficient 2.8 mM⁻¹ cm⁻¹. One unit of the enzyme is the amount necessary to decompose 1 µmol of substrate per min at 25°C. The APX activity was expressed in umol min⁻¹ mg⁻¹ protein.

**Data analysis**

The data were subjected to analysis of variance (ANOVA) and for mean value comparisons by Duncan Multiple Range Test were evaluated where P<0.05 was considered significant. Data analysis used the statistical program Genstat v19.1.0.21390 and the graph reproduced by software Microsoft excel 2010.

**RESULTS AND DISCUSSION**

**Toxicity and tolerance of aluminum in growth of oil palm seedling**

Al stress inhibited the seedlings growth, including root length, plant height, dry weight of root and shoot, except the leave number, and stem diameter at 30 DAT (Table 1). Application of 100 and 200 µM Al did not affect root length, but 400 µM Al treatment reduced the root length by about 33.9 %, and at higher concentration, the obstacle on root length becomes bigger. It can be seen that with 200 µM Al treatment, plant height was decreased, and with 400 µM Al treatment the decrease was the biggest, about 31.4 %. Dry weight of root was not affected by 100 and 200 µM Al treatments, but 400 µM Al decreased root dry weight about 32.75 %. All Al treatments reduced shoot dry weight, the higher Al concentration the bigger the decrease (Table 1). Based on these results, 400 µM concentration of Al was determined as the appropriate concentration to be used in selection of oil palm seedlings tolerance to Al toxicity in nutrient culture. Sopandie (2014) pointed out that the best indicator to differentiate plant resistance to Al toxic is the value of root growth resistance. The root apex is the critical site for Al toxicity and genes for Al tolerance are likely to be expressed (Dellaire and Ryan 1995). The root apex (root cap, meristem, and elongation zone) accumulates more Al and attracts greater physical damage than the mature root tissues. The characteristic of root length has a positive correlation with the result of Al stress (Panda et al. 2009). Some studies on oil palm seedlings concluded that Al stress has inhibited root growth (Mendez et al. 2014; Christiansen et al. 2011; Marlina et al. 2017) and reduced total dry weight (Christiansen et al. 2011; Marlina et al. 2017). Rout et al. (2001) mentioned that Al interferes with cell division in roots, decreases root respiration and uptake and use of water and nutrients, particularly calcium and phosphorous and metabolic pathway. Inhibition of root division was due to the binding of P at DNA by Al that enters the cell nucleus of root tip so that the DNA activities decline, and cell division is hampered (Rout et al. 2001). The selection of oil palm seedlings tolerant Al can be performed under 400 µM Al condition since this concentration showed inhibition in root length, root and shoot dry weight.

**Effects of aluminum on photosynthetic rate, organic compound and enzyme activity of oil palm seedlings**

Figure 1 shows the effect of Al on photosynthetic rate, MDA content, CAT, and APX activities in oil palm seedlings. Al stress has brought about the decrease in the rate of photosynthesis, the effect of which increased with increasing Al concentration, and the length of time of exposure (Figure 1A). Proklamasiningish et al. (2012) and Marlina et al. (2017) have also shown the inhibition effect of Al on photosynthesis rate in oil palm. They have conveyed that the decrease in photosynthesis rate was thought to be due to declines in chlorophyll content (Yang et al. 2015). The inhibition of photosynthetic rate could also be associated with the inhibition of electron transport in photosystem II (Jiang et al. 2009).

Al stress significantly raised MDA content in all Al concentrations, compared to control, and it keeps increasing as the Al concentration increases (Fig 1B). The activity of CAT enzyme increased significantly in all Al treatments starting at 3 HAT and 7 HAT then decreased at 24 HAT and 72 HAT (Figure 1C). The same pattern has also occurred in APX, the activity of this enzyme increased
following the increase in Al concentration applied, starting at 3 HAT and 7 HAT then decreased dramatically at 24 HAT and 72 HAT (Figure 1D). Panda et al. (2009) mentioned that Al stress has caused lipid peroxide in plasma membrane so the production of ROS rises. Yamamoto et al. (2001) pointed out that MDA is one of the end products of lipid peroxide that accumulates when the plant is exposed to oxidative stress; the accumulation of MDA in tissue is widely used to estimate cell damage. However, plants have a complex antioxidant system for ROS scavenging, in which specific enzymes act to neutralize the action of free radicals (Miller et al. 2010). When plants are exposed to Al stress, they will produce antioxidants such as superoxide dismutase (SOD), glutathione peroxide (GPX), CAT, and APX (Shahnawaz and Sanadhya 2017). The magnitude of enzyme antioxidation production and the degree of activity of these enzymes can be used in the selection of genotypes that are susceptible and tolerant to abiotic stress (Kusvuran et al. 2016). Our current results showed that when the seedlings were exposed to Al stress, the increase of enzymes antioxidation activity occurred at 3 HAT and 7 HAT, then decreased dramatically at 24 HAT and 72 HAT, indicating that the increase in enzyme activity can be attributed to the quick response to Al stress at 3 until 7 hours after exposurement of Al, while at 24 and 72 hours a steady-state condition has achieved. Exposure oil palm seedling to Al stress caused the decrease in rate of photosynthesis, increase MDA levels, followed by the increase of CAT and APX enzyme activity.

Table 1. Effect of Al treatment to root length, plant height, number of leaves, diameter of stem, roots and shoots dry weight of oil palm seedlings at 30 DAT

<table>
<thead>
<tr>
<th>Concentrations of Al (µM)</th>
<th>RL (cm)</th>
<th>PH (cm)</th>
<th>NL</th>
<th>DS (cm)</th>
<th>Dry weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>34.63 a</td>
<td>36.90 a</td>
<td>4.33 a</td>
<td>1.03 a</td>
<td>0.58 a</td>
</tr>
<tr>
<td>100</td>
<td>33.00 a</td>
<td>34.27 a</td>
<td>4.67 a</td>
<td>1.10 a</td>
<td>0.57 a</td>
</tr>
<tr>
<td>200</td>
<td>34.63 a</td>
<td>30.30 b</td>
<td>4.00 a</td>
<td>1.00 a</td>
<td>0.51 ab</td>
</tr>
<tr>
<td>400</td>
<td>24.70 b</td>
<td>25.30 c</td>
<td>4.33 a</td>
<td>0.97 a</td>
<td>0.39 bc</td>
</tr>
<tr>
<td>800</td>
<td>20.37 c</td>
<td>26.93 c</td>
<td>4.67 a</td>
<td>0.97 a</td>
<td>0.34 c</td>
</tr>
<tr>
<td>1600</td>
<td>19.73 c</td>
<td>26.20 c</td>
<td>4.33 a</td>
<td>0.93 a</td>
<td>0.31 c</td>
</tr>
</tbody>
</table>

Note: The numbers followed by the same letters in the same column show no significant difference based on DMRT at the level α: 5%; DAT= day after treatment; RL= root length; PH= plant height; NL= number of leaves; DS= diameter of stem

Figure 1. The effect of Al on photosynthetic rate (A), MDA content (B), CAT (C), and APX (D) activities; HAT = hour after treatment. The bars represent the standard error of the mean.
Effects of various exogenous compounds in growth of oil palm seedlings exposed to Aluminum stress

Table 2 showed that 400 µM Al inhibited root length, dry weight of root, and shoot drastically at 30 DAT. There was no adverse effect of Al on plant height, number of leaves, and diameter of stem. In general, except ethephon, the application of citric acid and peat water in some extents could reduce the adverse effects of Al, so that seedlings were able to obtain recovery, although not fully recovered. The results of the experiment showed that ethephon treatment actually worsened the growth of palm oil seedlings after exposure to Al, as indicated by a greater decrease in root length, root dry weight, and shoot dry weight. There was relatively no change in plant height, number of leaves and stem diameter of seedlings after exposure to high concentration of Al and application of ethephon. According to Sharma et al. (2019), ethylene, plays a crucial role in ameliorating the harmful effects of these abiotic stress conditions like temperature, drought, heavy metals, and salt. Exposure of tobacco protoplasts to ethephon and ACC led to activation of a plasma membrane cation channel that was permeable to Ba(2+), Mg(2+), and Ca(2+), and inhibited by Al(3+) (Zhao et al. 2007). Koppittke (2016) pointed out that the production of ethylene and auxin seems to be a component of a plant-response to toxic Al, resulting in cell wall modification or regulation of organic acid release. Yu et al. (2019) revealed that 0.1 µM ethrel (ethylene donor) treatment has a maximum biological effect on promoting the adventitious rooting in cucumber under salt stress. However, Sun et al. (2016) pointed out that Al-triggered ethylene acts as a signaling molecule with impacts on auxin biosynthesis and distribution in roots, especially in the root transition zone, and thereby inhibits root growth.

There was an alleviating effect of Al toxicity on root growth due to the application of 25 and 50 ppm of citric acid, in which citric acid tremendously increased the root length and root dry weight, although root growth recovery was not fully. The effect of citric acid on those concentrations did not occur in shoot dry weight. Citric acid at a concentration of 100 ppm did not have a positive effect on the recovery of seedlings growth after exposure to Al. The research results of Hongyu et al. (2018) also revealed exogenous citric acid could effectively alleviate the toxicity of aluminum on P. massoniana seedlings, and it had a better effect on aluminum-sensitive variety than aluminum-resistant variety. Hu et al. (2016) revealed that exogenous citric acid application may alleviate growth and physiological damage caused by high temperature. Li et al. (2009) pointed out that citrate is considerably more effective as an Al chelator than is malate. Chotimah et al. (2013) pointed out that application of peat water increased frond number, frond width, and decreased Al-frond.

Table 2 indicates that not all concentrations of peat water have a positive effect on increasing the tolerance of oil palm seedlings to the adverse effects of Al, i.e. only 200 ppm peat water showed a beneficial impact on seedlings growth. Adding 200 ppm peat water into nutrient culture containing high concentration of Al enabled seedlings to grow normally under Al stress conditions, as indicated by root length, root dry weight, and shoot dry weight which were relatively similar to the control condition without Al. There was no so much information obtained about the effect of peat water on increasing tolerance to Al toxicity in oil palm seedlings. The peat water there was 24.56 ppm of the total carboxylic acids and 12.72 ppm of total phenolic acids (Chotimah et al. 2007). Chotimah et al. (2013) revealed that application of carboxylic acid derivatives group increased Al root and P root accumulation, while at the top, carboxylic acid derivatives group decreased Al shoot Aloe vera. Peat water ameliorant application was able to lead to more efficient chalk usage for 75% in black soybean adaptation toward Al stress (Pujiwati et al. 2015). Mimmo et al. (2013) revealed that the release of organic acids plays an important role in alleviating Al toxicity. A. virginicus showed an Al-stress-induced synthesis and secretion of citrate and malate in roots, this mechanism may help to suppress an increase of toxic Al ions in the root region (Ezaki et al. 2013). Wang et al. (2012) revealed that construction of a new citrate synthesis pathway by simultaneous overexpression of CS and PEPC in the cytoplasm of transgenic plant leaves could enhance Al resistance in plants. Approximately 60-80% of synthesized organic acids were secreted to the medium by each plant, independent of Al stress (Ezaki et al. 2013).

Table 2. Effect of exogenous compounds on root length, plant height, number of leaves, diameter of stem, root dry weight, shoot dry weight of oil palm seedlings at 30 DAT

<table>
<thead>
<tr>
<th>Treatments</th>
<th>RL (cm)</th>
<th>PH (cm)</th>
<th>NL</th>
<th>DS (cm)</th>
<th>Dry Weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Roots</td>
</tr>
<tr>
<td>0 Al</td>
<td>47.09 a</td>
<td>33.80 b</td>
<td>6.33 a</td>
<td>1.80 a</td>
<td>1.08 ab</td>
</tr>
<tr>
<td>400 µM Al</td>
<td>37.10 c</td>
<td>37.73 ab</td>
<td>6.33 a</td>
<td>1.80 a</td>
<td>0.88 d</td>
</tr>
<tr>
<td>Al + Et 25</td>
<td>32.50 d</td>
<td>36.50 ab</td>
<td>6.00 a</td>
<td>1.80 a</td>
<td>0.67 e</td>
</tr>
<tr>
<td>Al + Et 50</td>
<td>26.93 e</td>
<td>36.03 ab</td>
<td>6.00 a</td>
<td>1.83 a</td>
<td>0.64 e</td>
</tr>
<tr>
<td>Al + Et 100</td>
<td>18.74 f</td>
<td>33.60 b</td>
<td>6.33 a</td>
<td>1.80 a</td>
<td>0.33 f</td>
</tr>
<tr>
<td>Al + CA 25</td>
<td>44.26 ab</td>
<td>35.43 ab</td>
<td>6.00 a</td>
<td>1.77 a</td>
<td>1.10 a</td>
</tr>
<tr>
<td>Al + CA 50</td>
<td>46.99 a</td>
<td>38.70 a</td>
<td>6.33 a</td>
<td>1.80 a</td>
<td>0.98 bc</td>
</tr>
<tr>
<td>Al + CA 100</td>
<td>40.02 bc</td>
<td>34.03 b</td>
<td>6.00 a</td>
<td>1.77 a</td>
<td>0.89 d</td>
</tr>
<tr>
<td>Al + PW 100</td>
<td>41.28 bc</td>
<td>35.77 ab</td>
<td>6.33 a</td>
<td>1.70 a</td>
<td>1.03 bc</td>
</tr>
<tr>
<td>Al + PW 200</td>
<td>47.80 a</td>
<td>36.60 ab</td>
<td>7.00 a</td>
<td>1.70 a</td>
<td>1.16 a</td>
</tr>
<tr>
<td>Al + PW 300</td>
<td>37.59 e</td>
<td>36.87 ab</td>
<td>7.00 a</td>
<td>1.87 a</td>
<td>1.08 ab</td>
</tr>
</tbody>
</table>

Note: The numbers followed by the same letters in the same column show no significant difference based on DMRT at the level a: 5%; DAT: day after treatment; RL: root length; PH: plant height; NL: number of leaves; DS: diameter of stem; Et: ethephon; CA: citric acid; PW: peat water
Exposing seedlings of oil palm to high concentration of Al (400 uM) brought about a decrease of all physiological characters except carotenoids and anthocyanin (Table 3), the physiological characters of which were photosynthesis rate, chlorophyll a and chlorophyll b content at 30 DAT. Application of ethephon did not have a significant effect on photosynthesis rate, chlorophyll a and chlorophyll b content, and carotenoids, but it enhanced extremely anthocyanins more than three times. The results of this experiment have justified why ethylene was not able to increase the growth performance of seedlings under Al stress conditions (Table 2), it was clearly proven that ethylene actually reduced photosynthesis and both chlorophyll a and chlorophyll b content (Table 3). The increase of anthocyanin when ethylene was applied is an interesting phenomenon, as it is known that anthocyanin is a pigment that often increases when abiotic stress occurs. Ezaki et al. (2013) revealed that Al-induced synthesis of polyphenolic compounds including anthocyanin also occurred in roots as a long-term response to Al toxicity and anthocyanin production did not co-localize with either Al accumulation, nitric oxide (NO) production or lipid peroxides production in the roots. The concentration of the blue Al(3+)-anthocyanin complex reached a maximum when a sufficient excess of aluminum was present (Schreiber et al. 2011).

Application of 200 ppm peat water elevated the rate of photosynthesis, as well as content of chlorophyll a, chlorophyll b, and carotenoid. According to Panda et al. (2009), Al stress could reduce the content of chlorophyll, but giving exogenous compounds such as citric acid (Hu et al. 2016; Song et al. 2018) and humic acid (Mirdad, 2016) enabled seedlings to alleviate chlorophyll damage, thus improving the photosynthesis rate. Chotimah et al. (2007) explained that peat water contains 0.104% humic acid. This increase in photosynthetic rate is expected to be associated with the effect of the humic acid compound contained in peat water. There are other compounds that have the same effect, which able to act as antioxidants when plants are exposed to Al stress, i.e. carotenoids and anthocyanin (Apel and Hirt 2004; Sopandie 2014). Li et al. (2017) mentioned that ethephon was able to induce the synthesis of anthocyanin in root hairs which were thought to have important role in the mechanism of tolerance when plants were subjected to high concentration of Al.

A high concentration of Al (400 uM) elevated tremendously MDA content compares with control at both 1 DAT and 3 DAT (Fig 2A). An increase in MDA levels in the tissue indicates that there has been damage due to lipid peroxidation of the membrane after exposure of seedlings to high concentration of Al. Xu et al. (2012) pointed out that the addition of Al(3+) significantly increased MDA. Meanwhile, the activity of CAT (Fig. 2B) and APX (Fig. 2C) enzymes in the addition of Al was lower than that of the control, presumably, the plant’s recovery ability was not able to compensate for cell damage. Nasr et al. (2011) pointed out that increasing Al concentration in root medium, APX activity was significantly decreased. Addition some exogenous compounds can reduce stress levels with different magnitudes as shown in Fig.2, the alleviating effect of Al stress was observed more clearly at 1 DAT as shown by lower accumulation of MDA at that time. The experimental results revealed that ethylene has a contribution to the response pathway of the signal transduction as shown that addition of 25 ppm ethylene has returned MDA levels back almost to their original level. There was a little effect of ethephon at concentrations of 50 and 100 ppm in reducing MDA levels at 1 DAT, their effect was not observed at 3 DAT, even MDA elevated much higher. These results corresponded to the increase in the activity of CAT (Fig 2B) and APX (Fig 2C) enzymes under the presence of ethylene at both 1 DAT and 3 DAT. Zhang et al. (2018) showed that ethephon treatment on tobacco plants increased the enzyme activity of APX, POD, and CAT under PEG-induced drought conditions.

### Table 3. Effect of exogenous compound in photosynthesis rate, chlorophyll a, chlorophyll b, carotenoids and anthocyanin in oil palm seedlings at 30 DAT

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Photosynthesis (mol·m⁻²·s⁻¹)</th>
<th>Chlorophyll a (mol·100 g⁻¹)</th>
<th>Chlorophyll b (mol·100 g⁻¹)</th>
<th>Carotenoids (mol·100 g⁻¹)</th>
<th>Anthocyanin (mol·100 g⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 Al</td>
<td>6.34 bc</td>
<td>3.66 ab</td>
<td>1.14 ab</td>
<td>1.57 abcd</td>
<td>0.13 d</td>
</tr>
<tr>
<td>400 µM Al</td>
<td>3.00 de</td>
<td>2.60 b</td>
<td>0.84 b</td>
<td>1.18 d</td>
<td>0.23 cd</td>
</tr>
<tr>
<td>Al + Et 25 ppm</td>
<td>4.31 cd</td>
<td>2.56 b</td>
<td>0.91 b</td>
<td>1.26 cd</td>
<td>0.29 bc</td>
</tr>
<tr>
<td>Al + Et 50 ppm</td>
<td>1.57 e</td>
<td>2.66 b</td>
<td>0.93 b</td>
<td>1.56 abcd</td>
<td>0.39 ab</td>
</tr>
<tr>
<td>Al + Et 100 ppm</td>
<td>2.01 e</td>
<td>2.48 b</td>
<td>0.85 b</td>
<td>1.38 bcd</td>
<td>0.42 a</td>
</tr>
<tr>
<td>Al + CA 25 ppm</td>
<td>8.78 a</td>
<td>4.53 a</td>
<td>1.50 a</td>
<td>2.00 a</td>
<td>0.28 bc</td>
</tr>
<tr>
<td>Al + CA 50 ppm</td>
<td>6.08 bc</td>
<td>4.25 a</td>
<td>1.43 a</td>
<td>1.79 abc</td>
<td>0.19 cd</td>
</tr>
<tr>
<td>Al + CA 100 ppm</td>
<td>9.38 a</td>
<td>4.23 a</td>
<td>1.45 a</td>
<td>1.88 ab</td>
<td>0.16 cd</td>
</tr>
<tr>
<td>Al + PW 100 ppm</td>
<td>5.44 c</td>
<td>2.57 b</td>
<td>0.85 b</td>
<td>1.25 cd</td>
<td>0.18 cd</td>
</tr>
<tr>
<td>Al + PW 200 ppm</td>
<td>6.85 b</td>
<td>4.25 a</td>
<td>1.40 a</td>
<td>1.83 abc</td>
<td>0.17 cd</td>
</tr>
<tr>
<td>Al + PW 300 ppm</td>
<td>5.98 bc</td>
<td>3.65 ab</td>
<td>1.15 ab</td>
<td>1.69 abc</td>
<td>0.21 cd</td>
</tr>
</tbody>
</table>

Note: The numbers followed by the same letters in the same column show no significant difference based on DMRT at the level α: 5%; DAT: day after treatment; Et: ethephon; CA: citric acid; PW: peat water
Citic acid appears to be more effective in reducing MDA levels, especially at concentrations of 100 ppm at 1 DAT and 3 DAT (Figure 2.A). There was an increase in CAT activity (Figure 2.B) and APX (Figure 2.C) when the oil palm seedlings were added with citric acid at 1 DAT and 3 DAT. This fact can answer the relationship between citric acid application with decreased MDA levels and increased activity of CAT and APX enzymes as well as increased tolerance of seeds to Al side effects (Table 2). The application of citric acid can increase the activity of CAT and APX enzymes which play a role in ROS scavenging so that MDA accumulation is reduced drastically. Hongyu et al. (2018) showed that after giving exogenous citric acid, the activity of antioxidant enzymes in the leaves increased, and the content of H2O2, MDA, and osmotic regulatory substances in the leaves decreased. Similar results were also seen in the addition of 200 ppm or 300 ppm of peat water which was able to reduce MDA levels. The addition of 200 ppm and 300 ppm of peat water also increased the enzyme activity of CAT (Fig. 2B) and APX (Fig. 2C). These results related to the increased activity of CAT and APX enzymes with the addition of peat water which indicates the involvement of peat water in the signal transduction response. This response is thought to play a role in the humic acid contained in peat water. Zykova et al. (2018) revealed that all humic acid fractions from peat showed antioxidant activity in radical scavenging. Application of humic acid under stress reduces H2O2 levels through activation of superoxide dismutase (SOD) and CAT (Yildiztugay et al. 2019). The addition of citric acid and peat water exogenously was able to increase the tolerance of oil palm seedlings to Al stress, as indicated by the increase in root length, root dry weight, photosynthesis rate, chlorophyll content, activity of CAT and APX enzymes, and decreased levels of MDA.

ACKNOWLEDGEMENTS

The authors would like to thank the PT Astra Agro Lestari, Tbk., Indonesia for all of the supports.

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