Response of morphological and biochemical traits of maize genotypes under waterlogging stress

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ABSTRACT

Maize (Zea mays L.) is one of the most important cereal crops cultivated around the world. Waterlogging stress is a major production constraint of maize production in rain-fed agricultural systems. The main objective of this experiment was to investigate the effect of continuous waterlogging on morphological and biochemical traits of maize genotypes at the vegetative stage. Ten maize genotypes were treated under no waterlogging (control) and continuous waterlogging of five centimeters depth for 10 days. The treatments were applied to the plants at their 45 days of age. Visual leaf injury scores from Leaf 4 (youngest leaf is the reference point) to Leaf 7 separated tolerant and susceptible genotypes. Waterlogging stress significantly reduced the total number of live leaves and chlorophyll content in leaf tissues in susceptible genotypes. The anatomical study revealed that tolerant maize genotypes produce a large number of aerenchyma cells under waterlogging stress compared to susceptible genotypes. The enzymatic activities of ascorbate peroxidase (APX) and peroxidase (POD) exhibited a greater increase in tolerant genotypes than susceptible genotypes whereas the contents of reactive oxygen species (H$_2$O$_2$) greatly increased in susceptible genotypes than tolerant genotypes under waterlogging stress compared to control. Principal component 2 (PC2) indicated that increasing plant height in the genotypes BHM-14, BHM-13 and BHM-9 was associated with waterlogging tolerance. The findings of this experiment will add value to maize breeding to screen out maize genotypes for waterlogging stress tolerance.

KEYWORDS: Zea mays L., waterlogging, morphology, biochemical traits

INTRODUCTION

Maize (Zea mays L.) is called the queen of the cereals due to its high productivity, wider adaptability in the various agro-ecological regions and high genetic potential compared to other cereals (Mahesh et al., 2013). maize is one of the most widely produced and consumed crops in the world by producing about 1116.2 million metric tons in 2019-2020 (USDA, www.fas.usda.gov/data/grain-world-markets-and-trade). In recent years, abiotic stresses such as drought, waterlogging, submergence, salinity and extreme temperature are increasingly affecting crop production and productivity (Bray et al., 2000).

Waterlogging has become one of the main constraints of maize production worldwide by affecting its yield performance (Du et al., 2017). The main causes of waterlogging in maize production are—continuous rainfall with inadequate drainage, contingent flooding, and high water-table. Sometimes maize is grown in converted paddy fields or in poorly drained soil that help create waterlogging during the rainy season (Amin et al., 2014). In the Asian monsoon regions, the main source of waterlogging stress of the summer maize is flooding of fields during late spring and summer (Mano et al., 2006). Over 18% of total maize growing areas in the South and South-East Asia are frequently affected by waterlogging stress (Zaidi et al., 2009). The agronomic and yield performance of maize is severely affected by waterlogging stress (Osman et al., 2013). Extinction of plant species and alteration of plant distribution can be caused by extreme or gradual waterlogging (Bailey-Serres & Voosenek, 2008).

Waterlogging causes a reduction of gas exchange between the atmosphere and root tissue by 100 times lowering the diffusion rate of gases in the flooded soil (Zaidi et al., 2009). Waterlogging condition decreases the available O$_2$ for plants (Capon et al., 2009). With the gradual decrease of oxygen, plant root suffers from hypoxia (low oxygen) followed by anoxia (no oxygen) when faced with excess moisture for more than three days (Dennis et al., 2000; Zaidi and Singh, 2002). Some physiological processes of plants that require oxygen— including growth and cell division, uptake and

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transpiration of nutrients and respiration—become affected under waterlogging. Plants growth, development and survival are dramatically affected by waterlogging conditions (Parent et al., 2008). However, the level of damage significantly depends on the developmental stages of the crop. From the early seedling stage to tasseling stage are the susceptible stages of maize under waterlogging (Zaidi et al., 2004). Plants suffer from chlorosis, necrosis, decreased growth, defoliation, yield loss and eventually plants die due to waterlogging stress (Hasanuzzaman et al., 2016).

Plants under waterlogging conditions survive through various morphological, physiological, anatomical and biochemical adaptations. Production of ethylene and formation of adventitious roots are some of the major adaptive responses under waterlogging stress due to O₂ deficiency (Pezeshki, 2001; Suralta and Yamauchi, 2008). Under waterlogging conditions, the plant exhibits several anatomical changes by the formation of gas space or aerenchyma. The presence of the aerenchyma or gas space is the most common adaptive feature for the distribution and flow of oxygen from root to shoot under waterlogging stress (Colmer & Voosenek, 2009; Shiono et al., 2011). Reduction of soil redox potential and enhancement of generation of toxic compounds occur due to waterlogging (Fiedler et al., 2007). Production of active oxygen species (AOS) such as superoxide (O₂⁻), singlet oxygen (¹O₂), hydrogen peroxide (H₂O₂) and hydroxyl radicals (OH•) are the results of oxidative stress due to waterlogging (Subbaiah & Sachs, 2003; Liao & Lin, 2003; Zhang et al., 2005; Jackson & Colmer, 2005). Due to normal aerobic metabolism, AOS is present in plants at different levels and contribute to waterlogging damage vitally (Kuk et al., 2003). AOS is highly reactive and therefore any oxidative damage to lipids, proteins and nucleic acids can change normal cellular metabolism (McKersie & Leshem, 1994; Alschler et al., 1997; Imlay, 2003; de Azevedo Neto et al., 2006). However, plants have the potential to detoxify the adverse effects of reactive oxygen species (ROS) by generating various forms of antioxidants such as ascorbate peroxidase (APX), superoxide dismutase (SOD), peroxidase (POD), catalase (CAT), glutathione reductase (GR), ascorbic acid, glutathione, tocopherols and carotenoids (Biswas &Kalra, 2018). In addition, significant genetic variability in the tolerance of maize to water-logging stress has been reported (Zaidi et al., 2002, 2003, 2007).

Response of morpho-physiological traits associated with tolerance and identification of promising genotypes of maize are the prerequisite for the success of breeding under waterlogging stress. Identification and development of genotypes capable of withstanding water logging conditions could be an ideal and affordable approach, which could be suitable for maize-growing farmers lacking sufficient resources in the sub-tropics. The major bottlenecks are the lack of appropriate screening techniques, morpho-physiological traits associated with tolerance and the identification of promising genotypes (Zaidi & Singh, 2001). Therefore, it is important to run an experiment to assess the response of morphological and biochemical traits of maize genotypes under water-logging. The objective of this study was to find out traits of maize genotypes responsive to waterlogging stress at morphological and biochemical levels that may offer tolerance under stress conditions.

**MATERIALS AND METHODS**

### Plant Materials and Growth Conditions

Seeds of ten selected maize genotypes were collected from Bangladesh Agricultural Research Institute (BARI), Gazipur-1700 (Table 1). The experiment was laid out following a Randomized Complete Block Design (RCBD) with three replications. There were two treatments—no water (control) and 5.0 cm water treated plot (Figure 1). A waterlogging treatment was applied in the field continuously for 10 days with an average depth of 5.0 ± 1.0 cm at the vegetative stage of plant in order to introduce stress when the plants were 45 days old. Water level of the treated field was maintained continuously at the same level throughout the experiment from one end to another end of the plots.

### Measurements and Data Collection

**Morphological traits analysis**

**Leaf scoring**

Leaves of ten genotypes from both control and waterlogging-treated plants were scored based on injury level using 1-9 scores at 10 days after application of waterlogging stress (Figure 2, Table 2, Ari et al., 2019).

**Plant height**

Plant height of individual plants was measured at 3, 6 and 9 days after treatment imposition. The measurement of plant height was carried out in centimeters (cm) from the soil surface to the highest point of the arch of the uppermost leaf whose tip is pointing down.

**Leaf area**

The length and width of each leaf was measured individually at 3, 6 and 9 days after treatment and the area of individual leaf laminae was estimated using the following formula: LA = 0.75 × Leaf Length × Leaf Width (Ren et al., 2013). Leaf area of the individual leaves was added to estimate the total leaf area of each plant.

**Number of live leaves (NLL)**

At 3, 6 and 9 days after applying waterlogging stress, the live leaves per plant were counted. A leaf was considered as an alive leaf when more than 50% of its leaf area was green.

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**Figure 1:** Experimental plot (A) control plot (no waterlogging) (B) treated plot (waterlogging)
Percent leaf (%) mortality rate
Percent leaf mortality rate was estimated by counting total number of dead leaves (less than 50% alive) and the total number of live leaves (more than 50% alive) per plant at 9 days after applying the waterlogging stress and percent mortality was counted by using the following formula:

\[
\text{Leaf mortality rate (\%) = \frac{\text{Number of dead leaves/plant}}{\text{Total number of leaves/plant}}} \times 100
\]

Chlorophyll content
At 9 days after applying waterlogging stress, the chlorophyll content of the 3rd leaf of each plant was measured with the help of a chlorophyll meter (SPAD-502 PLUS, 3V=200 Mw, Konica Minolta, Osaka, Japan). This was done similarly for each plant under control and waterlogging treatment. It was a non-destructive method. There is a small sensor inside the meter and leaves were placed between them. For each leaf three positions (proximal, distal and middle) were brought under measurement and the measurements were averaged (Abbasi et al., 2015).

Anatomical study
During the experimentation, 0.5% acetocarmine solution was used for staining roots. To prepare the solution 0.5 g carmine was dissolved in 100 L of 45% glacial acetic acid, and refluxed for 24 hours (KSU, 2018). The dissected roots were stained with 0.5% acetocarmine well enough to visualize under a microscope. The cross-sections of roots were visualized at 100x magnification and the root anatomical structure was captured.

Biochemical trait analysis
Leaf samples from the 4th youngest leaf of four selected genotypes (BHM-9, BHM-13, BM-6 and Popcorn) were collected at 10 days after waterlogging stress and were subjected for biochemical analysis. The ascorbate peroxidase (APX) and peroxidase (POD) activities were determined by following the method of Nakano and Asada (1981). Hydrogen peroxide was dissolved in 100 L of 45% glacial acetic acid, and refluxed for 24 hours (KSU, 2018). The dissected roots were stained with 0.5% acetocarmine well enough to visualize under a microscope. The cross-sections of roots were visualized at 100x magnification and the root anatomical structure was captured.

Table 1: List of ten maize genotypes used in this study with their characteristics (DHCP, 2017, BARI, http://baritechnology.org/en/home/tech_commodity#result). BARI, Bangladesh Agricultural Research Institute

| Genotypes | Given identity | Developed by | Growing season | Average yield (t ha\(^{-1}\)) | Year of release | Characteristics |
|-----------|----------------|--------------|----------------|-----------------------------|----------------|----------------|
| BHM-14    | G1             | BARI         | Rabi, kharif   | Rabi 10.84, Kharif 10.52    | 2017           | Early maturing, high temperature tolerant, root and stalk lodging tolerant |
| BHM-9     | G2             | BARI         | Rabi, kharif   | Rabi 10.0-11.0, Kharif 7-7.5 | 2007           | Higher yield potential, resistant to disease and pest |
| Popcorn   | G3             | BARI         | Rabi, kharif   | Rabi 5.5, Kharif 4.0-4.5    | 1986           | High popping quality (95%) |
| BHM-13    | G4             | BARI         | Rabi, kharif   | Rabi 10.0-11.0, Kharif 7-7.5 | 2016           | Higher yield potential, resistant to disease and pest |
| BHM-7     | G5             | BARI         | Rabi, kharif   | Rabi 10.0-11.0, Kharif 7-7.5 | 2006           | Higher yield potential, resistant to disease and pest |
| BHM-12    | G6             | BARI         | -              | -                           | 2016           | -              |
| BM-6      | G7             | BARI         | Rabi, kharif   | Rabi 6.5-7.5, Kharif 5-6.0  | 1998           | High yield potential, resistant to disease and pest |
| BM-7      | G8             | BARI         | Rabi, kharif   | Rabi 6.5-7.5, Kharif 5-6.0  | 2002           | Open pollinated variety, high yield, bold grain |
| Mohor     | G9             | BARI         | Rabi, kharif   | Rabi 5.5-6.0, Kharif 3.5-4.5| 1991           | High yield |
| Barnali   | G10            | BARI         | Rabi, kharif   | Rabi 5.5-6.0, Kharif 4.0-4.5| 1986           | High yield, resistant to pest and disease |

Table 2: Scoring criteria for the visual injury scoring under waterlogging stress at the vegetative stage for ten maize genotypes

| Score | Leaves                                      |
|-------|---------------------------------------------|
| 1     | Normal color and growth                     |
| 3     | Nearly normal condition, but leaf tip become discolored and wilting started |
| 5     | Leaf become rolled, most part of the leaf become discolored and started drying |
| 7     | Leaf become mostly dry, totally discolored  |
| 9     | Leaf dead or near to die                    |

Percent leaf (%) mortality rate
Percent leaf mortality rate was estimated by counting total number of dead leaves (less than 50% alive) and the total number of live leaves (more than 50% alive) per plant at 9 days after applying the waterlogging stress and percent mortality was counted by using the following formula:

\[
\text{Leaf mortality rate (\%) = \frac{\text{Number of dead leaves/plant}}{\text{Total number of leaves/plant}}} \times 100
\]
(H$_2$O$_2$) activity was determined by following the protocol of Velikova et al. (2000).

**Statistical Analysis**

Data were analyzed using MINITAB 17 statistical software packages (Minitab Inc., State College, Pennsylvania, USA). A two-way analysis of variance (ANOVA) was executed for different morphological and biochemical traits following a general linear model (GLM) to explore treatment, genotype and treatment genotype interaction. Principal Component Analysis (PCA) of the morphological traits was carried out to investigate the association between morphological traits and tolerance of genotypes under treatment. The Principal Component (PC) scores were stored and ANOVA of the PC scores was performed following a one way ANOVA to explore the statistical significance between treatment and genotypes and treatment-variety interaction.

**RESULTS**

**Treatment Effect**

Waterlogging stress significantly affected the survivability of older leaves exhibiting genotypic variations (Table 2, Figure 3). The number of leaves per plant ranged between 8 and 11 in maize genotypes under both control and waterlogging treatment during data collection. The first three young leaves were fully alive (score 1) showing no sign of injury under both control and waterlogging treatments (Table 2). On the other hand, leaf injury scores from the leaf position 7 to leaf position 9 were nine for all genotypes under waterlogging treatment although that ranged between 1 and 3 under control treatment. Notably, the leaf injury scores of leaf position 4–6 of 10 maize genotypes under waterlogging treatment showed notable variations among genotypes (Figure 3). The genotypes BHM-14, BHM-13 and Barnali accounted for leaf injury scores lower than 3 up to leaf position 5 and that of lower than 5 up to leaf position 6 (Figure 3). These three genotypes were therefore graded as tolerant. The genotypes BHM-9, BHM-12 and Mohor accounted for leaf injury scores lower than 3 up to leaf positions 5 and 6 (Figure 3). These three genotypes were graded as moderately tolerant (Figure 3). The four other genotypes, Popcorn, BHM-7, BM-6 and BM-7 accounted for leaf injury scores higher than 5 at the leaf position 6 (Figure 3). These four genotypes were graded as susceptible (Figure 3).

Waterlogging stress significantly altered the morphological traits of maize plants including plant height, number of live leaves, mortality rate of leaves, leaf area and chlorophyll content (Figure 4, Figure S1-S7). Plant height was significantly increased by 1.35 folds ($P<0.001$) for BHM-9 at 3 days after waterlogging stress compared to control (Figure S1). Plant height at 6 days after waterlogging increased by 1.31 folds ($P<0.001$) in genotype BHM-9 under waterlogging stress compared to control (Figure S2). The total number of live leaves was decreased by 1.26 folds ($P<0.001$) in BM-6 at 6 days after waterlogging treatment compared to control (Figure S5). The total number of live leaves was decreased from 1.3 to 1.4 folds ($P<0.001$) in genotypes BHM-13, BHM-7, BHM-12, BM-6, BM-7, Mohor and Barnali at 9 days after waterlogging treatment compared to control (Figure 4A). Percent leaf mortality rate (%MR) was increased by 5.12 folds in genotype BHM-12 followed by 4.69 folds in genotype BHM-13 ($P<0.001$) under waterlogging treatment compared to control (Figure 4A). The genotypes BHM-14, Popcorn, BHM-7 showed a reduction in chlorophyll content (CC) by 1.15, 1.58, 2.27 folds ($P<0.001$), respectively, under waterlogging stress compared to control (Figure 4C).

**Response of Biochemical Traits Under Waterlogging Stress**

The contents of APX and POD were increased in both tolerant and susceptible genotypes under waterlogging stress compared to control (Figure 5A) but their contents were greatly increased in two tolerant genotypes BHM-9 and BHM-13 compared to two susceptible genotypes, Popcorn and BM-6 (Figure 5A). The content of APX was increased by 3.67 folds ($P<0.001$) in the moderately tolerant genotype BHM-9 followed by 2.33 folds in the tolerant genotype BHM-13 (Figure 5A) whereas the content of POD was increased by 2.46 folds ($P<0.001$) in tolerant genotype BHM-13 under waterlogging treatment compared to control (Figure 5B).

The content of hydrogen peroxide (H$_2$O$_2$) as a measure of oxidative stress greatly increased in both susceptible genotypes Popcorn and BM-6 but that was decreased on tolerant genotype BHM-9 but slightly increased in BHM-13 (Figure 5C). The highest increase in the content of H$_2$O$_2$ by 9.05 folds was observed in susceptible genotype BM-6 under waterlogging stress compared to control (Figure 5C).
A remarkable number of morphological traits exhibited significant genotypic variation including plant height at 3, 6 and 9 days after waterlogging, number of live leaves at 3, 6 and 9 days after waterlogging, percent leaf mortality rate (\%MR), chlorophyll content (CC), leaf area at 3, 6 and 9 days after waterlogging (Table S1). The content of hydrogen peroxide (H$_2$O$_2$), peroxidase (POD) and ascorbate peroxidase (APX) also showed a significant varietal difference (Suppl. Table 1).

**Anatomical Difference**

Maize genotypes under control condition (no waterlogging) produce no aerenchyma cells in epidermal regions (Figure 6A). Anatomical study showed that roots of tolerant maize genotypes such as BHM-13, BHM-14 and Barnali under waterlogging stress produced large aerenchyma cells (Figure 6B). By contrast, the susceptible maize genotypes such as BHM-7, BM-6 and BM-7 produced aerenchyma cells either none or comparatively much lower in number compared to tolerant genotypes under waterlogging stress (Figure 6B vs. 6C).

**Genotype Difference**

A remarkable number of morphological traits exhibited significant genotypic variation including plant height at 3, 6 and 9 days after waterlogging, number of live leaves at 3, 6 and 9 days after waterlogging, percent leaf mortality rate (\%MR), chlorophyll content (CC), leaf area at 3, 6 and 9 days after waterlogging (Table S1). The content of hydrogen peroxide (H$_2$O$_2$), peroxidase (POD) and ascorbate peroxidase (APX) also showed a significant varietal difference (Suppl. Table 1).

**Trait Association**

**Principal component analysis (PCA)**

The first three principal components (PC) explained 72% of the total data variation that showed variation due to the effect of waterlogging and genotypes (Table 3). The first principal component (PC1) explained 40.9% variation, second principal component (PC2) explained 19.6% and third principal component (PC3) explained 12% variation (Table 3). Scores of PC1 were highly significant for treatment (waterlogging), genotype and treatment-genotype interaction (Table 3). PC2 showed that PC scores were highly significant for treatment, genotype and treatment-genotype interaction. PC3 scores were highly significant for genotype and genotype treatment interaction (Table 3).

The PC1 separated the genotype BHM14 (G1) from the genotype BM7 (G8) for their highest positive and negative PC scores, respectively (Figure 7). PC1 also accounted for higher coefficients for a number of live leaves per plant and plant height (Figure 7). PC2 separated genotypes based on control versus waterlogging stress for their positive and negative
scores, respectively (Figure 7). Plant height at 3 days after waterlogging, plant height at 6 days after waterlogging and plant height at 9 days after waterlogging treatments also had negative coefficients for PC2 indicating that the genotypes BHM14 (G1), BHM 9 (G2) and popcorn (G3) were the least affected under waterlogging stress and thus exhibited higher stress tolerance (Figure 7).

DISCUSSION

This study was conducted to investigate the response of morpho-physiological traits of maize genotypes under waterlogging conditions. The increasing trend of plant height of a few maize genotypes along with the response of biochemical traits under waterlogging are discussed herein.

Impact of Waterlogging on Morphological Traits

Plant height

According to Parent et al. (2008), the height of the plant and the height of the ear in maize were severely affected in almost all genotypes. In contrast, Li et al. (2011) concluded that waterlogging did not significantly affect plant height. In the present study, significant differences in plant height under waterlogging and significant varietal interaction indicated that selection for waterlogging tolerance could be effective (Figure S1- S3).

Table 3: Principal components and their coefficients from principal component analysis

| Variable                        | PC1     | PC2     | PC3     |
|---------------------------------|---------|---------|---------|
| Plant height (cm) 3DAT          | 0.361   | -0.381  | 0.032   |
| Plant height (cm) 6DAT          | 0.383   | -0.353  | 0.062   |
| Plant height (cm) 9DAT          | 0.404   | -0.291  | 0.086   |
| No. of live leaves 3DAT         | 0.369   | -0.11   | -0.211  |
| No. of live leaves 6DAT         | 0.39    | 0.19    | -0.243  |
| No. of live leaves 9DAT         | 0.361   | 0.329   | -0.238  |
| Mortality rate                  | -0.179  | -0.551  | 0.102   |
| Chlorophyll Content             | 0.264   | 0.381   | 0.13    |
| Leaf area 3DAT                  | 0.134   | 0.117   | 0.535   |
| Leaf area 6DAT                  | 0.085   | 0.064   | 0.548   |
| Leaf area 9DAT                  | 0.116   | 0.151   | 0.462   |
| % variation explained           | 40.9    | 19.6    | 12.1    |

$P$—Probability of statistical significance; PC—principal component; DAT—days after transplanting

Figure 5: Treatment effect, genotypic variation and treatment × genotype interaction for (A) APX (B) POD and (C) $\text{H}_2\text{O}_2$ of four selected maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction.

Figure 6: Cross sections of roots of maize plants (A) cross-section of roots of BHM-14 grown without waterlogging (control), (B) cross-section of roots of tolerant BHM-14 grown under waterlogging condition showing presence of large number of aerenchyma cells, (C) cross-section of roots of susceptible BM-7 genotype grown under waterlogging condition showing presence of a few aerenchyma cells.
Chlorophyll Content

Chlorophyll is a major component of the chloroplast that has a positive relationship with photosynthesis (Anjum et al., 2011). One of the first stress symptoms, which may be related to nitrogen deficiency caused by leaching and de-nitrification of soil nitrogen, has been identified as a significant decrease in leaf chlorophyll content under water-logging stress (Zaidi & Singh, 2001). In this experiment, the highest decrease of chlorophyll was found in the susceptible genotype BHM-7 (Figure 4C). The same agreement was found in the experiment of Lone et al. (2009) that the relative greenness of leaf was also affected due to flooding treatment as there was fading of leaf color in most of the cases as reflected by their corresponding SPAD values (relative greenness).

Impact on Anatomical Structure

Aerenchyma formation under waterlogging stress is a remarkable anatomical modification. Aerenchyma is the oxygen storage area that facilitates the movement of gas in the root cortex and thus facilitates aerobic respiration in submerged organs (Mano & Omori, 2013). Once the tubes form between the roots and shoots, cell metabolism can be oxygenated and maintained. A large number of species including maize can form aerenchyma in the shoots and roots (He et al., 1996a). In this experiment, we found differences in root anatomical structure between two treatments and the tolerant genotypes produced a large number of aerenchyma cells compared to susceptible genotypes (Figure 6). Aerenchyma is related to hydraulic conductivity in roots, where the absorption of water and nutrients can occur even at a lower rate (Irfan et al., 2010). The role of antioxidant and cell wall loosening enzymes in the formation of aerenchyma in Saracura maize (Zea mays L.) roots with contrasting tolerance to waterlogging revealed that long-term stress improves the activity of enzymes involved in loosening the cell wall, linked to a more effective defense of antioxidants (De Souza et al., 2017). Plants in their roots showed constitutive aerenchyma, and it is known that these structures are associated with greater tolerance to excess water (Imaz et al., 2013; Mano & Omori, 2013).

Impact of Waterlogging on Biochemical Traits

Naturally, plants face various kinds of biotic and abiotic stresses due to changes in biochemical components that cause changes in their metabolic system. According to Jaiswal (2018), under waterlogging conditions increased APX content has been found. In order to protect the cells from detrimental effects, an enzyme such as APX is essentially needed to scavenge ROS or regenerate antioxidants. To detoxify $H_2O_2$, APX as an isoenzyme is more effective as it is widely distributed inside the cells of higher plants (Shigeoka et al., 2002). In the present study, APX content was greatly increased in tolerant genotypes in treated condition compared to control (Figure 5A).

In response to biotic/abiotic stresses, peroxidase synthesis and accumulation are mostly stimulated in plants (Giorgi et al., 2009). Yadav et al. (2017) found that with the increase in the
duration of waterlogging, POD activity gradually increased and resistant genotype reported higher POD activity in response to waterlogging stress compared to sensitive genotype. POD is responsible for the scavenging of H$_2$O$_2$ generated under oxidative stress. Thus, in resistant genotype, these enzymes eliminate the excess H$_2$O$_2$ more effectively and confer resistance to waterlogging stress. The genotypes that either have better ability to detoxify reactive oxygen species enzymatically (POD) or non-enzymatically or produce less reactive oxygen species are reported to be resistant to waterlogging stress (Lin et al., 2004). In the present investigation, the highest increase of POD was observed in tolerant genotype BHM-13 (Figure 5B) which indicates that this genotype was relatively tolerant to waterlogging stress.

Abiotic stresses have led to excessive generation of ROS, such as H$_2$O$_2$ (Polle, 2001). These ROS are highly reactive in nature that causes harm to a variety of cellular molecules and metabolites (Ashraf, 2009). According to Yadav et al. (2017), after imposing waterlogging stress, the level of hydrogen peroxide (H$_2$O$_2$) increased in both resistant and sensitive maize genotypes, but the sensitive genotype revealed relatively higher H$_2$O$_2$ content under waterlogging condition. In the present investigation, the highest increase of H$_2$O$_2$ was observed in BM-6 (Figure 5C) that indicates it was more susceptible to waterlogging stress because it has a lower scavenging capacity.

**CONCLUSION**

In the rice-based cropping system, maize (Z. mays L.) has become an important cereal crop due to its high productivity and suitability for cultivation in most areas. This study investigated traits related to waterlogging tolerance and their response due to treatment at the vegetative stage. Plant height was increased in BHM-14 and BHM-9 compared to control under waterlogging conditions. The total number of live leaves was decreased mostly in the susceptible genotype BM-7 and BM-6 and chlorophyll content was decreased in the susceptible genotype BHM-7 under waterlogging stress compared to control. The biochemical analysis showed that the tolerant genotypes exhibited greater enzymatic activities of APX and POD and accumulated lower content of H$_2$O$_2$ under waterlogging stress compared to control. The result suggested that the genotypes BHM-9, BMH-13 and BHM-14 exhibited higher waterlogging tolerance whereas BM-6, BM-7 and BMH-7 exhibited susceptibility to waterlogging stress. However, the experiment was conducted with only 10 maize genotypes. Further studies with more genetically diverse populations can elucidate and confirm the results of this study.

**FUNDING**

This research was supported by the Ministry of Science and Technology, Government of the People’s Republic of Bangladesh (Project no. 2020/6/MoST).

**ACKNOWLEDGEMENTS**

We acknowledge Bangladesh Agricultural Research Institute for providing seeds of maize.

**AUTHOR CONTRIBUTIONS**

AHKR and LH conceived and designed the study. SNA executed all experiments, analyzed the data and wrote the manuscript. SA and NS assisted with biochemical analysis. AHKR critically revised the manuscript.

**CONFLICTS OF INTEREST**

The authors declare no conflicts of interest for this study.

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### Table S1: Analysis of variance for morphological and biochemical traits of ten maize genotypes under control and waterlogging treatments

| Traits             | df  | Treatment (T) | Genotype (G) | T x G | F value | P value |
|--------------------|-----|---------------|--------------|-------|---------|---------|
| PH 3DAT            | 1   | 9             | 9            | 7.73  | 10.55   | 3.28    | 0.006  | <0.001 | <0.001 |
| PH 6DAT            | 1   | 9             | 9            | 4.8   | 10.01   | 3.12    | 0.029  | <0.001 | <0.001 |
| PH 9DAT            | 1   | 9             | 9            | 1.1   | 9.01    | 2.15    | 0.295  | <0.001 | 0.026  |
| NL 3DAT            | 1   | 9             | 9            | 0.2   | 12.89   | 1.54    | 0.657  | <0.001 | 0.135  |
| NL 6DAT            | 1   | 9             | 9            | 52.72 | 9.39    | 0.75    | <0.001 | <0.001 | 0.663  |
| NL 9DAT            | 1   | 9             | 9            | 143.62| 5.63    | 1.52    | <0.001 | <0.001 | 0.141  |
| %MR                | 1   | 9             | 9            | 571.13| 3.12    | 1.59    | <0.001 | <0.001 | 0.12   |
| LA 3DAT            | 1   | 9             | 9            | 119.14| 3.14    | 3.28    | <0.001 | <0.001 | <0.001 |
| LA 6DAT            | 1   | 9             | 9            | 0.89  | 3.76    | 0.61    | 0.352  | 0.002  | 0.78   |
| LA 9DAT            | 1   | 9             | 9            | 1.26  | 2.96    | 1.35    | 0.268  | <0.001 | 0.245  |
| H$_2$O$_2$         | 1   | 3             | 3            | 55568.96| 12475.5| 17505.52| <0.001 | <0.001 | <0.001 |
| POD                | 1   | 3             | 3            | 8117.81| 798.66  | 971.81  | <0.001 | <0.001 | <0.001 |
| APX                | 1   | 3             | 3            | 4225.55| 3693.58 | 1559.52| <0.001 | <0.001 | <0.001 |

df = Degrees of freedom, P = Probability of statistical significance, Tx G = Treatment genotype interaction, PH 3DAT = plant height at 3 days, PH 6DAT = plant height at 6 days, PH 9DAT = plant height at 9 days, NL3DAT = total number of live leaves at 3 days, NL 6DAT = total number of live leaves at 6 days, NL 9DAT = total number of live leaves at 9 days, %MR = percent (%) mortality rate, CC = chlorophyll rate, LA 3DAT = total leaf area at 3 days, LA 6DAT = total leaf area at 6 days, LA 9DAT = total leaf area at 9 days, H$_2$O$_2$ = hydrogen peroxide, POD = peroxidase, APX = ascorbate peroxidase

**Figure S1:** Treatment effect, genotype variation and treatment × genotype interaction for plant height at three days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali

**Figure S2:** Treatment effect, genotype variation and treatment × genotype interaction for plant height at six days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali
Figure S3: Treatment effect, genotype variation and treatment × genotype interaction for plant height at nine days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali.

Figure S4: Treatment effect, genotype variation and treatment × genotype interaction for total number of live leaves at three days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali.
Figure S5: Treatment effect, genotype variation and treatment × genotype interaction for total number of live leaves at six days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali.

Figure S6: Treatment effect, genotype variation and treatment × genotype interaction for total leaf area per plant at three days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali.
Figure S7: Treatment effect, genotype variation and treatment × genotype interaction for total leaf area per plant at six days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali

Figure S8: Treatment effect, genotype variation and treatment × genotype interaction for total leaf area per plant at nine days after waterlogging of ten maize genotypes under control and waterlogging treatments. Vertical bars indicate standard error of mean; different letters indicate significant difference among the treatment × genotype interaction. In the graph, G1= BHM-14, G2= BHM-9, G3= Popcorn, G4= BHM-13, G5= BHM-7, G6= BHM-12, G7= BM-6, G8= BM-7, G9= Mohor, G10= Barnali