Physiological Effects of γ-Aminobutyric Acid Application on Improving Heat and Drought Tolerance in Creeping Bentgrass

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ABSTRACT. Gama-aminobutyric acid (GABA) is a nonprotein amino acid in plant cells, which responds to changes in environmental factors. The objectives of this study were to evaluate the effects of foliar spray of GABA on drought and heat tolerance in creeping bentgrass (Agrostis stolonifera), and to investigate physiological factors altered by GABA application that contribute to improved drought tolerance and heat tolerance. GABA-treated plants (cv. Penncross) or non-GABA-treated control plants were then subjected to the following three treatments in growth chambers: 1) nonstress control [plants irrigated every 2 days to maintain soil water content at the pot capacity and maintained at 21/19 °C (day/night) for 35 days], 2) heat stress [plants exposed to 35/30 °C (day/night) and well-watered conditions for 35 days], and 3) drought stress [plants unirrigated for 9 days and maintained at 21/19 °C (day/night), and then rewatered for 2 days]. As compared with untreated plants, GABA-treated plants showed 22% to 39% and 8% to 21% significantly lower leaf electrolyte leakage (EL) and 35% to 143% and 21% to 24% significantly higher turf quality (TQ), 8% to 17% and 17% to 24% relative water content (RWC), 22% to 39% and 25% to 27% chlorophyll content, 7% to 11% and 6% to 17% photochemical efficiency, and an 84% to 683% and 57% to 76% osmotic adjustment (OA) exposed to heat or drought stress across days of treatment, respectively. GABA-treated plants accumulated 7% to 10% more water-soluble carbohydrates (WSC) and 11% to 43% more free proline than nontreated plants under heat stress, and 12% to 30% higher accumulation of WSC under drought stress. After 2 days of rewatering, a significantly better recovery also was observed in GABA-treated plants than that in nontreated plants previously exposed to drought stress. The results suggest that foliar application of GABA significantly improved heat and drought tolerance of creeping bentgrass, which was associated with maintenance of cell membrane stability, delaying in leaf senescence, and enhancing OA. The effectiveness of exogenous GABA application was more pronounced under heat stress than under drought stress.

Drought and high temperature are the most common and detrimental abiotic stresses for growth of cool-season plant species in the summer, including heat- and drought-sensitive cool-season turfgrass species, such as creeping bentgrass (Fry and Huang, 2004; Pessarakli, 2007). Drought and heat stress injury are typically characterized by leaf dehydration, reflected as a decline in leaf water content or accelerated leaf senescence due to loss of chlorophyll and photosynthetic activities, as well as decline in membrane stability (Pessarakli, 2007; Rivero et al., 2007; Veerasamy et al., 2007). Leaf wilting and senescence contributes to TQ decline, which negatively affects the playability and functionality of turfgrasses. Therefore, it is important to develop effective strategies to alleviate those physiological damages from drought or heat stress to maintain turfgrass quality in environmental conditions with elevated temperature and limited water resource for irrigation.

Plant growth regulators (PGR) or biostimulants have been used to improve turfgrass tolerance to drought or heat stress, such as trinexapac-ethyl [TE (Bian et al., 2009; Etemadi et al., 2015; Krishnan and Merewitz, 2015)], abscisic acid [ABA (Lu et al., 2009)], cytokinins [CK (Wang et al., 2012; Xu and Huang, 2009; Zhang et al., 2010)], and CK-containing bio-stimulants (Zhang and Ervin, 2004). Effectiveness and mode of actions of different PGR or biostimulants vary in different turfgrass species and stresses, with TE mainly affecting shoot growth rate and water demand (McCann and Huang, 2008), ABA regulating stomatal closure during drought stress (Kholova et al., 2010), and CK mediating leaf senescence under heat stress (Veerasamy et al., 2007). In recent years, a nonprotein amino acid, GABA, has been found to exhibit PGR effects due to its central roles in maintaining carbon and nitrogen balance and its involvement in carbohydrate and amino acid metabolism (Barbosa et al., 2010; Bouché and Fromm, 2004). In general, GABA is typically produced at a low level under...
favorable environmental conditions, but increases in response to abiotic stresses such as salinity, cold, and drought stress (Kinnersley and Turano, 2000; Serraj et al., 1998; Shi et al., 2010). There is some evidence indicating that GABA is involved in regulation of cytosolic pH (Bouché and Fromm, 2004), protection against oxidative stress (Bouché et al., 2003), and plant defense against insects (McLean et al., 2003). A recent study on functional genomics and metabolomics in Arabidopsis thaliana suggested that GABA may act as a signaling molecule interacting with phytohormones and other metabolites (Batushansky et al., 2014; Renault et al., 2011). Exogenous application of GABA has been found to enhance plant tolerance to abiotic stresses, such as chilling tolerance in peach fruit [Amygdalus persica (Shang et al., 2011)], heat tolerance in rice seedlings [Oryza sativa (Nayar et al., 2014)], and drought tolerance in black pepper [Piper nigrum (Vijayakumari and Puthur, 2015)]. Krishnan et al. (2013) reported that spraying with GABA improved membrane stability through upregulated peroxidase activity in leaves of perennial ryegrass (Lolium perenne) under drought stress, but no further data supported whether GABA was involved in the regulation of osmolytes or OA since leaf RWC was significantly increased in GABA-treated plants. Hence, limited information is available on the effectiveness of GABA on improving plant tolerance to drought and heat stress in turfgrass. Furthermore, physiological effects of GABA in regulating plant responses to drought or heat stress are not well understood in turfgrass species and other plant species.

Based on the potential biological functions of GABA (Barbosa et al., 2010; Bouché and Fromm, 2004; Bouché et al., 2003; Kinnersley and Turano, 2000; Serraj et al., 1998; Shi et al., 2010), we hypothesize that GABA may regulate drought or heat tolerance in creeping bentgrass through alteration in OA for water retention and suppressing stress-accelerated leaf senescence. The objectives of this study were to evaluate the effectiveness of foliar spray of GABA on drought and heat tolerance in creeping bentgrass, and to investigate physiological factors altered by GABA application for improving drought or heat tolerance.

**Materials and Methods**

**Plant material and treatment.** Creeping bentgrass (cv. Penncross) sod was collected from Horticultural Farm II at Rutgers University, North Brunswick, NJ. The sod pieces were planted in 24 polyvinyl chloride tubes (30 cm length and 10 cm diameter) filled with fritted clay in a greenhouse for 2 months (Sept.–Oct. 2014). The greenhouse had average temperatures of 23/16 °C (day/night) and 790 μmol·m⁻²·s⁻¹ photosynthetically active radiation (PAR) with natural sunlight and supplemental sodium lights when lack of natural sunlight. Plants were fertilized weekly with half-strength Hoagland’s nutrient solution (Hoagland and Arnon, 1950) and trimmed twice per week to maintain a canopy height of ~4 cm. Plants were transferred to controlled growth chambers after 2-month establishment in the greenhouse. The environmental conditions of growth chambers were maintained at day/night temperatures of 21/19 °C, 70% relative humidity, and 12-h photoperiod at PAR of 660 μmol·m⁻²·s⁻¹ at the canopy level. Plants were maintained in those conditions for 7 d to allow plant acclimation to the growth chamber conditions before the imposition of heat and drought treatment.

For the GABA treatment, all plants were sprayed twice at 2-d intervals before exposed to heat or drought stress with 5–10 mL of 0.5 mm GABA solution or water (non-GABA-treated control) until dripping. The concentration of GABA was chosen based on a preliminary test with a range of concentrations (0.5, 1, 2, 4 mm) for the most effective concentration on phenotypic changes based on TQ and growth. GABA-treated plants or non-GABA-treated control plants were then subjected to the following three treatments in growth chambers: 1) nonstress control (20 May, 27 May, and 3 June 2015)—plants were irrigated every 2 d to maintain soil water content at pot capacity and maintained at 21/19 °C (day/night) for 35 d; 2) heat stress (21 May, 28 May, and 4 June 2015)—plants were exposed to 35/30 °C (day/night) and well-watered conditions for 35 d; and 3) drought stress (5 May, 9 May, and 12 May 2015)—plants were unirrigated for 9 d and maintained at 21/19 °C (day/night), and then rewatered for 2 d. The duration of heat and drought stress was determined based on when stress symptoms were observed (mainly based on TQ, the degree of leaf wilt, and the inhibition of growth under stresses).

The experimental design was completely randomized block design with three stress treatments and GABA application under each stress treatment. Each temperature treatment (21/19 °C or 35/30 °C day/night) was replicated in four growth chambers, which served as experimental systems or units. Four pots or replicates of drought-stressed or nonstressed plants were placed in four growth chambers set at 21/19 °C (day/night). GABA treatment was replicated in four pots for each stress treatment (nonstress control, drought, or heat) that were replicated in four growth chambers.

**Physiological analysis.** Turf quality was evaluated based on color, density, and uniformity of the grass using a scale of 1 to 9 [9 = fully turgid, dense green canopy; 6 = minimal acceptable level; 1 = completely desiccated and brown plants (Beard, 2001)]. For leaf RWC measurement, leaves were detached from the plant and immediately weighed for fresh weight (FW) and then immersed in deionized water for 12 h at 4 °C. Turgid leaves were blotted dry and weighed for turgid weight (TW), then leaf tissues were dried in an oven at 80 °C for at least 72 h to get a dry weight (DW). Leaf RWC was calculated using the formula RWC (%) = [(FW–DW)/(TW–DW)] × 100 (Barrs and Weatherley, 1962). EL was used for estimating cell membrane stability of leaves. Fresh leaves (0.1 g) were washed three times with deionized water and immersed in 35 mL of deionized water and shaken for 24 h. The conductance of the solution was measured as initial conductivity (C_initial) using a conductivity meter (model 32; Yellow Springs Instrument Co., Yellow Spring, OH). Leaves then were autoclaved at 120 °C for 20 min. Once at room temperature, the conductivity of tissues was measured as maximum conductance (C_max). EL was calculated as the percentage of C_initial/C_max (Blum and Ebercon, 1981).

Chlorophyll was immerged in 10 mL of dimethyl sulfoxide and left in the dark for 48 h, and then the leaf extract was measured at 663 and 645 nm with a spectrophotometer (Spectronic Instruments, Rochester, NY). Chlorophyll content was calculated using the formula described by Arnon (1949). A chlorophyll fluorescence meter (Fim 1500; Dynamax, Houston, TX) was used for measuring photosynthetic efficiency (F_v/F_m). Individual leaves were adapted to darkness for 30 min using leaf clips and the F_v/F_m ratio was recorded with the fluorescence meter. Three measurements of F_v/F_m ratio were taken per replicate at each sampling day.
The content of WSC was determined according to the method of Buysse and Merckx (1993) with some modification. Leaves (20 mg DW) were added into 10 mL 80% ethanol and then extracted in boiled water (100 °C) for 15 min. After centrifugation at 3600 g for 10 min, supernatants were collected for further analysis. The reaction mixture contained 1 mL of supernatant, 1 mL of 18% phenol solution, and 5 mL concentrated sulfuric acid. The mixture was shaken, and absorbance was read at 490 nm using a spectrophotometer (Spectronic Instruments).

Free proline content was quantified spectrophotometrically using the ninhydrin method according to Bates et al. (1973). Fresh leaves (0.1 g) were immersed in 5 mL 3% aqueous sulfosalicylic acid and then heated in a water bath (100 °C) for 20 min. After centrifugation at 12,000 g, the supernatant was used for the estimation of the proline concentration. The reaction mixture consisted of 1 mL of supernatant, 1.5 mL of 2.5% acid ninhydrin, and 1 mL of glacial acetic acid, which was heated at 100 °C for 40 min. The reaction mixture was immediately cooled down in an ice bath, and extracted with 2.5 mL of toluene then absorbance was read at 520 nm.

For determination of OA, fresh leaf tissues were submerged in deionized water for 8 h at 4 °C to fully hydrate leaves. Tissues were blotted dry and frozen in liquid nitrogen for further analysis. Following thawing in ice bath, leaves were ground with a micropestle to extract leaf sap and 10 mL sap from a 1.5-mL tube was inserted into an osmometer (Wescor, Logan, UT) to determine osmolality (millimoles per kilogram). According to the formula ψS (megapascals) = [(osmolality) (0.001) (2.58)], osmolality was converted to ψS. OA was then calculated as the difference in ψS at full turgor between stressed leaves and well-watered control leaves (Blum, 1989).

Statistical Analysis. The General Linear Model procedure of SAS (version 9.1; SAS Institute, Cary, NC) was used to determine the significance of main treatment effects and interactions between GABA and heat or drought stress for all measured parameters. The significance of differences between GABA treatments under nonstress control, drought, or heat stress were tested using the least significance test with $P = 0.05$ at a given day of stress treatment.

Results

Effects of foliar GABA application on turf growth and physiological factors under nonstress condition. All parameters evaluated in this study, including TQ (Fig. 1A), RWC (Fig. 2A), EL (Fig. 3A), chlorophyll content (Fig. 4A), $F_v/F_m$ (Fig. 5A), WSC content (Fig. 6A), and free proline content (Fig. 7A), remained relative constant during the course (35 d) of nonstress treatment. Foliar application of GABA had no significant effects on those parameters under nonstress conditions.

Effects of GABA on TQ and EL under drought or heat stress. TQ decreased gradually with prolonged heat stress duration for GABA-treated or nontreated plants, but GABA-treated plants maintained a significantly higher TQ than nontreated plants during 35 d of heat stress; TQ was 35%, 90%, and 143% higher from 21 to 35 d of heat stress, respectively (Fig. 1B). TQ declined with drought duration, but GABA-treated plants showed a 21% higher TQ at 5 d of drought and 124% higher at 9 d of drought when compared with nontreated plants (Fig. 1C).

GABA-treated plants previously exposed to drought stress
Fig. 2. The effects of γ-aminobutyric acid (GABA) on relative water content (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d; control + GABA, GABA-treated plants maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants exposed to 35/30 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants were exposed to 35/30 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.

Fig. 3. The effects of γ-aminobutyric acid (GABA) on electrolyte leakage (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d; control + GABA, GABA-treated plants maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants exposed to 35/30 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants were exposed to 35/30 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.
Fig. 4. The effects of γ-aminobutyric acid (GABA) on chlorophyll content (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) for 35 d under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d; control + GABA, GABA-treated plants maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought + GABA, GABA-treated plants unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.

Fig. 5. The effects of γ-aminobutyric acid (GABA) on photochemical efficiency ($F_v/F_m$) (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d; control + GABA, GABA-treated plants maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat + GABA, GABA-treated plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought + GABA, GABA-treated plants unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.
Fig. 6. The effects of γ-aminobutyric acid (GABA) on water-soluble carbohydrate (WSC) (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.

Fig. 7. The effects of γ-aminobutyric acid (GABA) on free proline (A) under well-watered condition, (B) under heat stress, and (C) under drought stress in creeping bentgrass under normal water condition, heat stress, and drought stress. Means of four independent samples are presented. Bars represent standard errors. The same letters above columns indicate no significant difference at $P = 0.05$ based on least significant difference. Control, plants were maintained at 21/19 °C (day/night) for 35 d under well-watered conditions; heat, plants were exposed to 35/30 °C (day/night) under well-watered conditions for 35 d; drought, plants were unirrigated for 9 d and maintained at 21/19 °C (day/night) and then rewatered for 2 d.
also recovered to a higher TQ than nontreated control plants after 2 d of rewatering. Leaf EL increased during 35 d of heat stress (Fig. 3B) and 9 d of drought stress (Fig. 3C) regardless of GABA treatment. GABA application resulted in significant decrease in EL under heat or drought stress, which were 18%, 24%, and 32% lower than nontreated plants at 21, 28, and 35 d of heat stress and 21% and 8% lower at 5 and 9 d of drought stress, respectively (Fig. 3B and C).

**Effects of GABA on chlorophyll and photochemical efficiency under drought and heat stress.** Leaf chlorophyll content declined with prolonged duration during heat (Fig. 4B) and drought stress (Fig. 4C) in GABA-treated and nontreated plants. Compared with nontreated plants, GABA-treated plants had 22%, 34%, and 39% higher chlorophyll content at 21, 28, and 35 d of heat stress, and 25% and 27% higher chlorophyll content at 5 and 9 d of drought stress, respectively (Fig. 4B and C). GABA application also enhanced chlorophyll content at 2 d of rewatering following 9 d of drought stress (Fig. 4C). Leaf Fv/Fm declined significantly from 21 to 35 d of heat stress in plants without GABA treatment, whereas Fv/Fm did not change in GABA-treated plants during 35 d of heat stress (Fig. 5B). Under drought stress, Fv/Fm in both GABA-treated and nontreated plants declined significantly with stress duration from 5 to 9 d; exogenous GABA significantly increased Fv/Fm, compared with nontreated plants at both 5 and 9 d of drought. In addition, a significantly higher Fv/Fm ratio also was observed in GABA-treated plants than that in nontreated plants after 2 d of rewatering (Fig. 5C).

**Effects of GABA on water-soluble carbohydrate and free proline accumulation.** The content of WSC and proline did not change significantly under nonstress conditions (Figs. 6A and 7A). The content of WSC decreased in leaves of creeping bentgrass during 21 to 35 d of heat stress, whereas it was unchanged during 5 to 9 d of drought stress (Fig. 6). WSC content was 7%, 19%, and 10% significantly higher in plants treated with GABA compared with nontreated ones at 21, 28, and 35 d of heat stress (Fig. 6B). The application of GABA also resulted in 12% and 30% higher WSC content in GABA-treated plants than nontreated plants at 5 and 9 d of drought stress. Significant difference of WSC content between two treatments was not observed after 2 d of rewatering (Fig. 6C). Proline content decreased significantly from 21 to 35 d of heat or 5 to 9 d of drought stress (Fig. 7B and C). Exogenous GABA increased proline accumulation in plants under heat stress, but had no effects on proline accumulation under drought stress, although GABA-treated plants had greater proline content at rewatering (Fig. 7C).

**Effects of foliar GABA application on leaf RWC and OA under drought or heat stress.** Leaf RWC declined with stress duration for both drought and heat stress in GABA-treated or nontreated plants, but exogenous application of GABA effectively suppressed the decline in RWC (Fig. 2B and C). GABA-treated plants exhibited significantly higher RWC compared with nontreated plants during 35 d of heat stress and 9 d of drought stress. Application of GABA significantly increased OA in leaves under heat or drought stress. The OA of GABA-treated plants increased by 2-, 8-, and 5-fold compared with that of nontreated plants at 21, 28, and 35 d of heat stress, respectively (Fig. 8A). Under drought stress, there was an increase in OA for the GABA-treated plants, which was two times higher at 5 d. This increase was also found in the treated plants at 9 d with a 2-fold increase. GABA application had no effects on OA as plants were rewatered (Fig. 8B).

**Discussion**

Enhanced stress tolerance induced by application of GABA has been reported in different species and stress conditions, such as in *Caragana intermedia* under NaCl stress (Shi et al.,
heat stress, suggesting that both WSC and proline could contribute to GABA-induced OA. Nayyar et al. (2014) also found that GABA application raised the endogenous levels of proline under heat stress associated with improved heat tolerance in rice seedlings. However, under drought stress, the increased OA was mainly related to greater WSC content, not proline, in GABA-treated plants.

Overall, foliar application of GABA was effective in improving heat and drought tolerance of creeping bentgrass associated with maintenance of cell membrane stability, suppressing leaf senescence, and enhanced OA associated with the accumulation of WSC or proline. Additional studies are required regarding metabolic and molecular mechanisms of GABA regulation of drought or heat tolerance in cool-season turfgrass and other plant species.

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