

RESILIENCE OF C₄ CROPS TO CLIMATE VAGARIES

ABSTRACT

Climate change poses significant challenges to agriculture worldwide, affecting productivity and threatening food security. Key drivers of climate change like altered water availability, temperature fluctuations and increased carbon dioxide concentrations, influence crop performance and ecosystem stability. Photosynthetic pathways in plants C₃, C₄, and CAM play a critical role in determining their adaptability to changing climatic conditions. Understanding the physiological responses of C₄ crops to various environmental stresses like water and temperature stress, highlights their potential for future climate resilience. Due to the efficient carbon concentrating mechanism, low stomatal conductance and high water use efficiency in C₄ plants, they are expected to show higher drought tolerance relative to C₃ plants. Studying the responses of C₄ crops to climate change is essential as they play a vital role in global food production, especially in tropical and subtropical regions prone to climate extremes. While C₄ crops are more resilient to heat and drought than their C₃ counterparts, their yield potential is still constrained by the increasing severity of abiotic stresses, such as prolonged droughts, heatwaves, and soil salinity. Understanding how C₄ crops respond to these challenges can provide insights into optimizing their growth and productivity in future climate scenarios.

Keywords: Climate change, Drought, Heat stress, C₄ photosynthesis, Carbon concentrating mechanism

1. INTRODUCTION

The climate of Earth has undergone alterations throughout history. There have been eight cycles of ice ages and warmer periods during the past 800,000 years, with the end of the last ice age approximately 11,700 years ago marking the beginning of the current climate era and the rise of human civilisation (NASA, 2024). Climate change occurs due to various factors, including atmospheric changes like El Niño, driven by winds and ocean currents. External forces, such as Earth's axial tilt and orbital shape, likely influence ice age cycles. Additionally, greenhouse gases like carbon dioxide (CO₂) trap heat, raising Earth's surface temperature (National Geographic Society, 2024). Human activities, such as burning fossil fuels for energy and transportation or using technology to boost meat production, release greenhouse gases (Garnett, 2009; Stavi & Lal, 2013). Deforestation for timber or industrial development reduces the carbon dioxide absorbed by trees, while factories contribute to the

greenhouse effect by emitting pollutants into the atmosphere (Rykowski, 2000; Aju et al., 2015; National Geographic Society, 2024). Global warming is linked to burning fossil fuels and tropical deforestation, with human activities increasing atmospheric CO₂ by about 30% over the past 150 years (National Geographic Society, 2024). Greenhouse gases like methane and chemicals such as chlorofluorocarbons, hydrofluorocarbons, and hydrochlorofluorocarbons contribute significantly to climate change (Sovacool et al., 2021). Methane levels are rising due to agriculture, industrial activities, and decomposing waste in landfills (Karakurt et al., 2012). Similarly, these gases, used in refrigeration and aerosol sprays, trap heat in the atmosphere. While many countries are phasing them out through laws and regulations, their impact remains a concern (National Geographic Society, 2024). The effects of climate change are becoming increasingly evident. Global temperatures are rising, and oceans are getting warmer, with much of the heat being absorbed by the top layers of the ocean (Levitus et al., 2017; NASA, 2024). Ice sheets are shrinking, particularly in Greenland and Antarctica, while glaciers are retreating in many regions around the world (Velicogna et al., 2020). Snow cover is decreasing, and sea levels are rising because of melting ice and thermal expansion of seawater. Global sea level rose about 8 inches in the last century (Nerem et al., 2018). Arctic sea ice is declining, and ocean acidification is increasing. The ocean has absorbed between 20% and 30% of total anthropogenic CO₂ emissions in recent decades (Sabine et al., 2004). Carbon dioxide in the atmosphere warms the planet, causing climate change. Human activities have raised the atmosphere's CO₂ content by 50% in less than 200 years, with the current level of 424 ppm (NASA, 2024).

Climate change impacts on agriculture are being witnessed all over the world in the recent years affecting farm level productivity and adversely impacting stability in food grain availability at the national level. Temperature, water, and CO₂ are three of the main causes of climate change. Adopting sustainable methods that lower greenhouse gas emissions, encourage reforestation, and boost the use of renewable energy sources is crucial to addressing the issues caused by climate change, particularly in agriculture. Under drought, high temperatures, and nitrogen or CO₂ limitation, C₄ plants are more competitive than C₃ plants. (Watson-Lazowski and Ghannoum, 2021). C₄ plants possess the carbon concentrating mechanism, and can have increased photosynthetic rate, yield, water-use efficiency (WUE), and nitrogen-use efficiency (NUE) compared with ecologically similar C₃ plants. Implementing agricultural practices that favor C₄ plants could increase crop resilience to climate stressors, leading to higher productivity and better food security.

2. IMPACT OF CLIMATE VAGARIES ON CROPS

Climate change may adversely impact the yield of irrigated crops throughout the world, attributable to rising temperatures and alterations in water supply (Lal, 2000; Mall et al., 2017). Crops may experience severe weather events such as drought, flooding, excessive heat, and cold during their life cycle, leading to significant yield reductions (Raza et al., 2019). The effects of these factors may differ based on area, crop, cropping systems, soil types, and management practices. Climate change affects crop production both directly and indirectly. The immediate consequences primarily result from alterations in crop length and influence reproductive processes, including pollination and fertilisation. The indirect effects of climate change are mostly attributable to variations in water availability, as well as changes in insect, disease, and weed dynamics. Weather aberrations can impact numerous factors, particularly in rainfed regions where over 80 percent of farmers are smallholders, hence possessing limited capacity to mitigate adverse effects (Maheswari et al., 2015).

3. CLIMATE RESILIENCE

The capacity to foresee, plan for, and react to potentially dangerous occurrences, patterns, or disruptions associated with the climate is known as climate resilience. Assessing how climate change will increase or change existing climate-related risks and acting to better manage them are key components of increasing climate resilience (C2ES, 2021).

3.1 Climate resilient agriculture

Climate-resilient agriculture is described as agriculture that alleviates poverty and hunger amidst climate change while enhancing the resources it relies upon for future generations. Climate resilient agriculture offers a broader view than just increasing output and aims to change the existing systems. It promotes socially, economically, and environmentally sustainable food production systems locally, regionally, and internationally (Cordaid, 2016).

3.2 Climate Smart Agriculture

Climate Smart Agriculture (CSA) is characterised as a methodology that directs the necessary actions to transform and realign agricultural systems to effectively promote development and guarantee food security in a changing climate. The initiative seeks to achieve three primary objectives: sustainably enhancing agricultural output and income; adapting to and fostering resilience against climate change; and minimising and/or eliminating greenhouse gas emissions, where feasible (Cordaid, 2016). CSA is frequently used interchangeably with climate-resilient agriculture (CRA); however, CRA is a subset of CSA that specifically addresses the impacts of climate change (Viswanathan et al., 2020).

4. CLIMATE CHANGE AND PHOTOSYNTHESIS

Through the process of photosynthesis, all plants absorb CO_2 from the atmosphere and transform it into sugars and starches, but they do it in different ways (Paul & Foyer, 2001; Long et al., 2004). Each class of plants uses a different type of photosynthesis, known as a pathway, which is a variant on a series of chemical reactions known as the Calvin Cycle. A plant's capacity to tolerate low carbon atmospheres, higher temperatures, and decreased water and nitrogen is crucial for understanding climate change (Dusenge et al., 2019). These reactions also affect the quantity and kind of carbon molecules a plant produces, the locations where those molecules are stored, and more.

The processes of photosynthesis designated by botanists as C_3 , C_4 , and CAM, are directly relevant to global climate change studies because C_3 and C_4 plants respond differently to changes in atmospheric CO_2 concentration and changes in temperature and water availability. Scientists have started looking into how plants might be able to adapt to the changing climate as the world continues to warm. Modifying the processes involved in photosynthesis could be one approach to do it (Lara and Andreo, 2011).

While the C_3 pathway is the most common, it is also inefficient due to the photorespiration, a process that wastes assimilated carbon when RuBisCO reacts not only with CO_2 but also O_2 . Under prevailing atmospheric conditions, potential photosynthesis in C_3 plants is suppressed by O_2 as much as 40%. The extent of that suppression increases under stress conditions such as drought, high temperatures and high light (Hirst, 2021). The C_4 plants are more efficient in photosynthesis than the C_3 plants (Schmitt & Edwards, 1981; Gowik & Westhoff, 2011). In C_4 plants, the photorespiration is suppressed by increasing the CO_2 concentration at the Rubisco site, hence suppressing the enzyme's oxygenase activity. C_4 photosynthesis relies on a biochemical CO_2 pump with spatial separation of CO_2 fixation and assimilation, typically involving Kranz anatomy where mesophyll and bundle sheath cells cooperate (Edwards et al., 2004; Lara and Andreo, 2011). The enzyme, phosphoenol pyruvate

carboxylase of the C₄ cycle is found to have more affinity for CO₂ than the ribulose diphosphate carboxylase of the C₃ cycle in fixing the molecular CO₂ in organic compound during Carboxylation (Hatch, 1987; Lara and Andreo, 2011).

The Crassulacean Acid Metabolism (CAM) cycle is a CO₂ fixation process occurring during the dark phase of photosynthesis in Crassulaceae plants, with malic acid as the first product (Black & Osmond, 2003; Osmond, 2007). Most CAM plants are succulents with fleshy leaves, abundant chloroplasts in mesophyll cells, and vascular bundles lacking well-defined bundle sheath cells. Their stomata open at night and close during the day. While less efficient in photosynthesis than C₄ plants, CAM plants are better adapted to extreme desiccation (Black & Osmond, 2003; Osmond, 2007; Schiller & Bräutigam, 2021).

5. CARBON CONCENTRATING MECHANISM IN C₄ PLANTS

C₄ plants achieve high productivity through a carbon concentrating mechanism (CCM) that increases CO₂ concentration around RuBisCO (ribulose-1,5-bisphosphate carboxylase/oxygenase), the primary CO₂-fixing enzyme in plants, algae, and cyanobacteria. This CCM operates between two cell types: mesophyll cells, where CO₂ is initially fixed, and bundle sheath cells, where it is ultimately fixed (Watson-Lazowski & Ghannoum, 2021). In C₄ plants, PEP carboxylation occurs in specialized mesophyll cells, which also perform the full light-dependent reactions of photosynthesis (Schlüter & Weber, 2020; Romanowska & Wasilewska-Dębowska, 2022). Malate or aspartate is then transported to bundle sheath cells, characteristic of Kranz anatomy, where pre-fixed carbon is released for final CO₂ fixation by RuBisCO. This process raises the CO₂ concentration in bundle sheath cells to about 10 times that of ambient air, significantly reducing photorespiration. Notably, most bundle sheath cells do not complete the full light-dependent reactions of photosynthesis (Zabaleta *et al.*, 2012; Watson-Lazowski & Ghannoum, 2021).

Table 1: C3 vs C4 cycle

	Calvin cycle or C3 cycle	Hatch and Slack pathway or C4 cycle
Enzyme	Ribulose bisphosphate carboxylase oxygenase (RuBisCO)	Phosphoenolpyruvate (PEP) carboxylase
Process	Convert CO ₂ into a 3-carbon compound 3-phosphoglyceric acid (or PGA)	Convert CO ₂ into 4-carbon intermediate
Steps	Carbon fixation, Reduction, Regeneration of RuBisCO	Carboxylation 2. Breakdown 3. Splitting 4. Phosphorylation
first stable product	Phosphor glyceric acid (PGA)	Oxaloacetate (OAA)
Where Carbon Is Fixed	All leaf mesophyll cells	The mesophyll cells (MC) and the bundle sheath cells (BSC). C4s have a ring of BSCs surrounding each vein and an outer

		ring of MCs surrounding the bundle sheath, known as the Kranz anatomy.
CO ₂ compensation point	50-150 ppm CO ₂	0-10 ppm CO ₂
Photorespiration	Present and easily detectable.	Present only to a slight degree or absent.
The CO ₂ concentration inside leaf	high (about 200 ppm).	low (about 100 ppm)
Net rate of photosynthesis in full sunlight	15-25 mg. of CO ₂ per dm ² of leaf area per hour.	40-80 mg. of CO ₂ per dm ² of leaf area per hour

(Source: Ehleringer & Cerling, 2002; Lara and Andreo, 2011; Hirst, 2021).

6. ADVANTAGES OF THE C₄ PHOTOSYNTHETIC PATHWAY

The C₄ pathway increases CO₂ concentration in bundle sheath cells to 70 μM, compared to 4 μM in mesophyll cells, reducing RuBisCO's oxygenase activity by over 80%, though this effect varies with temperature. Consequently, C₄ plants exhibit up to double the photosynthetic rate and yield, 1.5 to 3 times greater WUE, and 2.5 times higher NUE compared to ecologically similar C₃ plants (Lin et al., 2019). In addition, elevating the concentration of CO₂ within the bundle sheath allows RuBisCO to increase its in vivo catalytic activity two- to fivefold in warm climates. As a result, C₄ plants have only 50–80% of the RuBisCO content found in C₃ plants, allowing them to sustain a higher leaf area production rate with lower leaf nitrogen levels compared to C₃ species (Sage and Zhu, 2011; Lin et al., 2019). WUE is enhanced in C₄ plants because CO₂ is fixed more effectively and stomata remain less open, reducing transpiration rates (Way et al., 2014; Leakey et al., 2019). This improved WUE allows C₄ plants to have longer growing seasons and more adaptable allocation patterns, such as directing more biomass to shoots in moist conditions or to roots in dry conditions (Lopes et al., 2011; Leakey et al., 2019). The C₄ pathway is particularly advantageous in hot, high-light environments that encourage high photorespiration rates (Long, 1983; Sage & Kubien, 2007; Lara & Andreo, 2011).

7. SUBTYPES OF C₄ PHOTOSYNTHESIS

C₄ species not only exhibit superior physiological traits compared to C₃ species but also display considerable diversity within themselves. C₄ photosynthesis is broadly categorized into three subtypes: NADP-malic enzyme (NADP-ME), NAD-malic enzyme (NAD-ME), and phosphoenolpyruvate carboxykinase (PEP-CK), each adapted to specific environmental conditions (Ghannoum et al., 2011; Wang et al., 2014; Borghi, 2021).

In NADP-ME plants, malate serves as the main C₄ acid transported between mesophyll and bundle sheath cells, whereas aspartate is the primary transport acid in NAD-ME and PCK C₄ grasses. The C₄ subtypes are closely linked to specific grass subfamilies. Species with NADP-ME type anatomy are found in the Panicoideae subfamily within the Andropogoneae, Arundinelleae, and Paniceae tribes (Morrone et al., 2012; Venter, 2015). The NAD-ME and PEP-CK types are primarily associated with the Chloridoideae subfamily and have evolved only once in Panicoideae (Voznesenskaya et al., 2006; Koteyeva et al., 2023). Among major C₄ crops, sorghum exclusively uses NADP-ME, maize primarily employs NADP-ME with PEP-CK as a secondary decarboxylase, and millets exhibit a mix of NADP-ME, NAD-ME, and some PEP-CK species (Sonawane et al., 2018).

Research on C₄ grasses, categorized into three biochemical subtypes—NAD-ME, PCK, and NADP-ME—grown under ambient (400 $\mu\text{L L}^{-1}$) and interglacial (280 $\mu\text{L L}^{-1}$) CO₂ conditions revealed that the Chloridoideae/NAD-ME group had higher leaf mass per area and leaf nitrogen content. In contrast, NADP-ME and PCK grasses exhibited enhanced photosynthetic nitrogen use efficiency (Pinto et al., 2016).

8. C₄ CROPS

Approximately 60% of C₄ species are grasses, with around 40% of grasses utilizing the C₄ photosynthetic pathway. Most C₄ grasses belong to warm-origin taxa, particularly the PACMAD clade, and dominate warm-climate grasslands. These grasses include ecologically and economically significant species such as major staple food, fodder, and biofuel crops, as well as numerous prominent weeds (Sage et al., 2011; Watson-Lazowski & Ghannoum, 2021). C₄ crops are particularly prevalent in warm (Korres et al., 2016), drought-prone climates (Lopes et al., 2011; Korres et al., 2016) and are becoming increasingly crucial for food and bioenergy security (Watson-Lazowski & Ghannoum, 2021).

Maize, sorghum, and sugarcane are major C₄ crops. Maize, the most produced cereal crop globally, is mainly grown in North and South America, as well as Eastern and Southern Africa, with over 60% of production in temperate regions (Watson-Lazowski & Ghannoum, 2021). Sorghum, more drought-tolerant than maize, thrives in dry climates and is valuable in areas with limited rainfall and resources for fertilizers (Watson-Lazowski & Ghannoum, 2021; Khalifa & Eltahir, 2023; Liaqat et al., 2024). It is used for fodder, fuel, and construction materials. The drought resistance in sorghum is attributed to its root system, leaf rolling, osmotic adjustment, and ability to delay reproductive development (Nadew et al., 2021). Sugarcane, a key industrial crop for sugar and bioenergy, grows in tropical and subtropical regions (Raza et al., 2019). It benefits from high CO₂ levels, showing improvements in biomass, photosynthesis, and overall growth, and is capable of coping with rising CO₂ concentrations due to its low CO₂ compensation point and carbon sequestration abilities (Watson-Lazowski & Ghannoum, 2021). Marin et al. (2013) observed improved sugarcane WUE and yield in parts of Brazil due to climate change using crop simulation models. As a C₄ plant with a CO₂ compensation point of 0-10 ppm, sugarcane can deplete atmospheric CO₂ under certain conditions. High CO₂ levels partially close stomata, reducing transpiration and sap flow, enhancing xylem potential and water status. Sugarcane also sequesters carbon naturally, mitigating CO₂ emissions and global warming by forming phytoliths (PhytOC), storing ~300 Mt of CO₂ annually in soil for thousands of years (Misra et al., 2019).

Millet, including pearl millet (*Pennisetum glaucum*) and foxtail millet (*Setaria italica*), are C₄ crops vital for food and fodder, with over 95% produced in developing countries. Their drought and heat tolerance make them suitable for harsh climates, and their short life cycle (12-14 weeks) helps escape stress. Traits like small leaves, thickened cell walls, and dense roots enhance stress resilience. The C₄ mechanism concentrates CO₂ around RuBisCO, reducing photorespiration (~80%) and boosting photosynthesis, WUE, and NUE. This also improves growth, biomass allocation, and ecological performance in warm conditions (Lenka et al., 2020).

9. CLIMATE RESILIENCE IN C₄ GRASSES

Grass species are divided into two distinct clades: BOP (Bambusoideae, Oryzoideae, Pooideae) and PACMAD (Panicoideae, Arundinoideae, Chloridoideae, Micrairoideae, Aristidoideae, Danthonioideae) (Hodkinson, 2018; Pardo and VanBuren, 2021; Gallaher et al., 2022). BOP grasses, primarily cool-season species found in temperate climates, utilize C₃ photosynthesis, which outperforms C₄ in these regions. Frost tolerance has

independently evolved in many Pooideae grasses. PACMAD grasses, mainly warm-temperate and tropical species, include agriculturally significant crops like sugarcane, maize, sorghum, and various millets (Panicoideae) and underutilized grains like finger millet (Chloridoideae) (Pardo and VanBuren, 2021).

9.1 Climate resilience among PACMAD grasses

Grass stomatal anatomy enhances resilience with unique elongated, dumbbell-shaped guard cells and two subsidiary cells, enabling faster responses and higher WUE compared to kidney-shaped guard cells in eudicots and most non-grass monocots (Nunes et al., 2020; Pardo and VanBuren, 2021; Zahedi et al., 2024). Stomatal arrangement and density also influence drought tolerance (Mehri et al., 2009; Huang et al., 2020). Grasses typically have hypostomatic leaves with pores on the abaxial surface or amphistomatic leaves with pores on both surfaces, the latter promoting efficient CO₂ diffusion and higher photosynthetic rates. Unlike eudicots with dorsoventral leaves, grasses have isobilateral leaves oriented parallel to light, deeper veins, and vertical angles, minimizing WUE costs of amphistomaty (Pardo and VanBuren, 2021).

10. RESPONSE OF C₄ PLANTS TO CLIMATE VAGARIES

10.1 Effect of Water Stress on C₄ Photosynthesis

Water stress significantly limits global plant productivity (Kijne, 2006). Severe stress causes metabolic inhibition, including photoinhibition and damage to photosynthetic enzymes, leading to irreversible loss of photosynthetic activity (Goh et al., 2012). C₄ plants, with high WUE and low stomatal conductance, exhibit greater drought tolerance, reducing water stress development. However, high WUE often prioritizes biomass production over water conservation in areas with high evapotranspiration (Ghannoum, 2016; Watson-Lazowski & Ghannoum, 2021).

The C₄ pathway enhances WUE, enabling C₄ grasses to thrive in drier, more exposed habitats. The high substrate affinity of PEPcase and the carbon-concentrating mechanism allow C₄ plants to function at lower mesophyll CO₂ levels and stomatal conductance, achieving higher instantaneous WUE than C₃ plants (Ghannoum et al., 2011; Pardo & VanBuren, 2021). Under drought stress, leaf-level WUE increases as reduced water loss from stomatal closure outweighs the decline in CO₂ assimilation. WUE also varies by C₄ subtype, with NAD-me grasses showing higher WUE than NADP-me grasses under drought condition (Ghannoum et al., 2002; Pardo & VanBuren, 2021).

Sorghum genotypes exhibit varied responses and tolerance to drought, influenced by the interaction between genotype and water stress levels. Research by Tingting (2010) found that sweet sorghum exhibited the highest WUE under moderate drought stress during early and middle growth stages, and under severe drought stress in the late growth stage. Increasing drought stress raised the light compensation point but reduced the light saturation point, apparent quantum yield, and dark respiration rate. Severe drought stress caused photoinhibition, lowering WUE and stem biomass. In contrast, normal water conditions avoided photoinhibition and increased stem biomass but reduced WUE. Overall, moderate drought stress conditions maximized both WUE and stem biomass. Jabereldar et al. (2017) identified sorghum genotype Tagat 10 as the most drought-tolerant, followed by Tagat 14, while Tagat 9 and cv. Gadambalea were the most drought-sensitive. Withholding irrigation at the 3-leaf stage improved crop WUE, reflecting the crop's ability to convert water into grain. Tagat 10 demonstrated superior WUE due to its higher seed yield compared to other genotypes.

10.2 Effect of High Temperature on C₄ Photosynthesis

Understanding temperature effects on C₄ plants is crucial for predicting their performance in future climates. As temperature rises, the oxygenation reaction increases, reducing RuBisCO's CO₂ specificity and limiting carbon gain. C₄ plants overcome photorespiration by concentrating CO₂ around RuBisCO in bundle sheath cells, maximizing carboxylation (Watson-Lazowski and Ghannoum, 2021)

C₄ crops like maize and sorghum show varying responses to temperature, with sorghum having higher photosynthetic temperature optima and greater heat and drought tolerance (Watson-Lazowski and Ghannoum, 2021). In a study by Correia et al. (2021), two maize genotypes, B73 and P0023, with contrasting drought and heat tolerance levels, were acclimatized to high temperatures (38°C vs. 25°C) under well-watered and water deficit (WD) conditions. Both genotypes successfully acclimatized to high temperatures, employing different mechanisms: B73 maintained photosynthetic rates by increasing stomatal conductance (g_s), while P0023 preserved g_s and exhibited limited transpiration. The study concluded that key traits for drought and heat tolerance in maize include limited transpiration rates and synchronized regulation of carbon assimilation metabolism.

Table 2. Results of warming and heat stress studies in Maize

Location	Ambient temperature	Continual warming	Heatwave (°C / hours / growth stage)	Photosynthesis	Yield	Reference
Yucheng, China	~ 13.1	~2	N.A.	Increase	N.A.	Zheng <i>et al.</i> (2018)
Illinois, USA	~22.7	~2.64	6 / 72 / Vegetative	Decrease	ns	Siebers <i>et al.</i> (2017)
Illinois, USA	~22.7	N.A.	6 / 72 / Reproductive	Decrease	Decrease	Ruiz-Vera <i>et al.</i> (2015)

10.3 CO₂ levels

Elevated CO₂ concentration can influence the growth of C₄ plants through several mechanisms. One effect is the increase in intercellular CO₂ partial pressure, which enhances CO₂ assimilation rate. Another is the reduction in stomatal conductance, leading to lower leaf transpiration rate. This decrease in leaf transpiration rate can boost leaf CO₂ assimilation rate and growth by conserving soil water, improving shoot water relations, and raising leaf temperature. Additionally, elevated CO₂ may lower mitochondrial respiration, which reduces overall plant respiratory losses and contributes to increased biomass (Ghannoum *et al.*, 2000).

As CO₂ concentrations rise, some regions will also experience increased frequency and severity of droughts (Lara & Andreo, 2011). The potential for enhanced growth and yield of C₄ plants at elevated CO₂ concentrations is primarily attributed to reduced water use and decreased drought stress, rather than a direct increase in photosynthesis (Lara & Andreo,

2011; Pignon and Long, 2020). Pignon and Long (2020) found that C₄ species with CO₂ concentration in bundle sheath cells showed an indirect stimulation of photosynthesis when atmospheric CO₂ increased from 400 μmol mol⁻¹ to 550 μmol mol⁻¹. However, no yield gains were observed under elevated CO₂ without drought stress.

Elevated CO₂ reduced midday stomatal conductance of FACE-grown sorghum by 32% with irrigation and by 37% under drought stress (Wall et al., 2001). As atmospheric CO₂ continues to rise, sorghum yield is expected to increase in areas with limited water availability (Ottman et al., 2001). Some C₄ plants grown under Free-Air Carbon Dioxide Enrichment (FACE) showed enhanced photosynthetic rates only during drought or under conditions of high atmospheric vapor pressure deficits (Leakey et al., 2009). Additionally, cultivating sorghum under elevated CO₂ mitigated the loss in grain quality caused by drought during the grain-filling stage by delaying physiological and metabolic responses to drought (De Souza et al., 2015).

In a future high-CO₂ environment, water requirements for irrigated sorghum will decrease, while dry-land productivity is expected to rise, assuming minimal global warming (Conley et al., 2001). Elevated CO₂ in controlled environments has been shown to increase sugarcane photosynthesis, WUE, biomass, and productivity. The improved WUE of sugarcane under elevated CO₂ is mainly due to reduced stomatal conductance. Sugarcane grown in elevated CO₂ had lower leaf stomatal conductance and transpiration, leading to greater leaf WUE. This helped delay the adverse effects of drought, allowing the plants to continue photosynthesis for at least an additional day during episodic drought cycles (Vu and Allen, 2009).

10.3.1 General effects of elevated CO₂ on photosynthetic heat tolerance

In C₃ species, elevated CO₂ generally enhances heat tolerance for photosynthesis, although at supra-optimal temperatures, this benefit may be diminished or even result in a decrease in photosynthesis. In contrast, C₄ species often experience reduced photosynthetic thermotolerance at both near-optimal and supra-optimal growing temperatures with elevated CO₂. While both C₃ and C₄ plants show similar reductions in stomatal conductance with increasing CO₂, C₄ plants have lower stomatal conductance at any given CO₂ level. This leads to reduced transpiration and higher leaf temperatures in C₄ plants, which could make them more susceptible to heat-related damage compared to C₃ plants in the same environment (Lara and Andreo, 2011).

The growth of maize and pearl millet under elevated CO₂ and temperature improved their cellular tolerance to osmotic stress and heat shock. However, maize appeared to benefit more from increased CO₂, while pearl millet seemed to benefit more from higher temperatures. The effects of current and anticipated global climate changes are likely to vary between these two species and may similarly impact other C₄ plant species across different ecosystems, whether natural or managed (Bordignon et al., 2019).

Elevated CO₂ is expected to enhance carbon uptake and water-use efficiency, leading to increased productivity of broomcorn millet in semi-arid regions under future high-CO₂ climates (Zhang et al., 2021). Similarly, elevated CO₂ significantly boosted grain yield and the accumulation of Zn, K, and Mn over three years, as well as enhancing the concentration and accumulation of P in foxtail millet (Gong et al., 2021).

11. Role of C₄ crops in the future

Breeding crop varieties that can better withstand higher temperatures and extreme conditions is crucial for adapting to future climate challenges. Advances in technologies, particularly CRISPR-Cas9 gene editing, are significantly improving our ability to enhance germplasm. This technology allows for efficient overexpression, knockouts, and base pair edits within genetic sequences, making genetic improvements faster and more precise than traditional methods (Watson-Lazowski & Ghannoum, 2021). In addition to gene editing, another promising approach is the introduction of a C₄ carbon concentration mechanism (CCM) into C₃ crops, such as rice. The C₄ Rice project aims to achieve this, with predictions suggesting that incorporating a C₄ photosynthetic pathway into rice could increase yields by as much as 50 percent (Sheehy et al., 2008). The C₄ Rice Consortium is employing strategies like metabolic C₄ engineering and the identification of leaf anatomy determinants through mutant screens to develop C₄ rice (Kajala et al., 2011). These combined efforts offer great potential for improving crop resilience and productivity in the face of climate change.

12. CLIMATE RESILIENT MAIZE FOR ASIA (CRMA)

The "Climate Resilient Maize for Asia" project is a collaborative initiative aimed at addressing the challenges faced by resource-poor maize farming communities in South and Southeast Asia, especially considering the anticipated impacts of climate change. Supported by Germany's development agency GIZ and implemented through a public-private partnership, the project focuses on enhancing the resilience of maize crops by developing and distributing abiotic stress-tolerant maize hybrids. These hybrids are specifically designed to thrive in rain-fed, stress-prone production systems, thereby promoting crop diversification, intensification, and higher yields. Building on the successes of the GIZ-funded "Abiotic Stress-Tolerant Maize for Increasing Income and Food Security among the Poor in South and Southeast Asia" project, this initiative addresses critical challenges related to improving maize production, enhancing food security, and building economic stability for smallholder farmers in the region (CIMMYT, 2020).

13. CLIMATE RESILIENCE IN C₄ WEEDS

Climate change is expected to cause shifts in weed community composition, impacting their population dynamics, life cycles, phenology, and infestation levels. Some weed species may go extinct, while others may become more aggressive and invasive. While elevated CO₂ levels are likely to boost the productivity of major C₃ crops, many of the troublesome agricultural weeds are expected to respond more positively to the increase in CO₂ than the crops themselves, potentially leading to their dominance in agro-ecosystems. Rising temperatures will likely favor the growth of C₄ weeds, which could result in significant crop yield losses. As climatic factors shift, crop-weed interactions may change, with weeds gaining an advantage and some previously non-threatening species taking over cultivated land. Under conditions of elevated temperature and drought, C₄ weeds are expected to dominate over C₃ crops, while C₃ weeds may prevail under higher CO₂ concentrations. However, when both CO₂ and temperature levels rise, C₄ weeds are predicted to dominate, further impacting crop production (Anwar et al., 2021).

Climate change is expected to significantly impact weed demographics, leading to shifts in weed species within agroecosystems (Peters et al., 2014; Ramesh et al., 2017). These shifts are crucial for weed management strategies and agricultural productivity. For species to persist in a particular habitat, they must adapt to environmental changes that can result in the alteration of weed flora, range expansion, and migration to new areas. Climate change will likely create opportunities for weeds to invade new ecosystems (Clements & Ditommaso, 2011; Peters et al., 2014). In fact, climate change is predicted to enhance the ability of introduced weed species to adapt to new environments, increasing their potential for

invasion in both native and managed ecosystems. Weeds that are well-suited to altered environmental conditions, particularly with higher CO₂ concentrations, are expected to be more successful in utilizing available resources (Anwar et al., 2021). While C₃ crops may have a competitive edge over C₄ weeds under elevated CO₂ conditions alone, the simultaneous rise in both CO₂ and temperature could favor the growth of C₄ weeds (Alberto et al., 1996). For instance, soybean yields were reduced from 45% to 30% when grown alongside *Amaranthus retroflexus* under elevated CO₂ compared to ambient levels (Ziska, 2003).

Table 3. Climate Resilient Varieties crop varieties from different states in India

Crop	Varieties	State	Seed source
Bajra	GHB-538 and GHB-719	Gujarat	Pearl millet Research Station, JAU, Jamnagar
	WCC-75	Karnataka	GKVK, UAS, Bangalore/ KSSC/ NSC
	RBH-177, RBH-154, RBH-173	Rajasthan	RSSC, Rajasthan
Foxtail Millet	RS-118, K-211-1, PS-4, SIA-326	Karnataka	GKVK, UAS, Bangalore/ KSSC/ NSC
Finger Millet	VR-708,HR-374	Chhattisgarh	IGKV / NRC millets, Bangalore
	MR-1, MR-6, GPU-66	Karnataka	GKVK, UAS, Bangalore/ KSSC
	Phule Nachani	Maharashtra	MPKV, Rahuri; ZARS, Kolhapur
Sorghum	CSH-5, CSH-9, CSV-4	Karnataka	GKVK, UAS, Bangalore/ KSSC
	Pant Chari 5, Pant Chari 7	Uttarakhand	GBPUA&T, Pantnagar
	Phule Chitra, M-35-1 Phule Vasudha, CSV18	Maharashtra	MPKV, Rahuri

(Lenka et al., 2020)

14. CONCLUSION

The changing climate has the potential to exert considerable adverse effects on plant physiology, soil fertility, carbon sequestration, and microbial activity, hence inhibiting plant growth and productivity, which would ultimately influence food production. C₄ plants are affected by significant global change variables in ways that contrast with C₃ plants. Comprehending the responses of C₄ plants to factors such as temperature, CO₂, nutrients, and water is essential for forecasting the adaptability of agricultural and wild C₄ populations to climatic variations, particularly those projected with global climate change. This knowledge is crucial for formulating strategies to maintain plant yield and provide food security in a changing climate.

References

- Aju, P. C., Iwuchukwu, J. J., & Ibe, C. C. (2015). Our forests, our environment, our sustainable livelihoods. *Eur. J. Acad. Essays*, 2(4), 6-19.
- Alberto, A. M., Ziska, L.H., Cervancia, C. R. and Manalo, P. A. 1996. The influence of increasing carbon dioxide and temperature on competitive interactions between a C3 crop, rice (*Oryza sativa*) and a C4 weed (*Echinochloa glabrescens*). *Funct. Plant Biol.* 23(6): 795-802.
- Anwar, M., Islam, A.K.M., Yeasmin, S., Rashid, M., Juraimi, A.S., Ahmed, S. and Shrestha, A. 2021. weeds and their responses to management efforts in a changing climate. *Agron.* 11(10): 1921.
- Black, C. C., & Osmond, C. B. (2003). Crassulacean acid metabolism photosynthesis: working the night shift'. *Photosynthesis research*, 76, 329-341.
- Borghi, G. L. (2021). *Evolution and diversity of photosynthetic metabolism in C3, C3-C4 intermediate and C4 plants* Universität Potsdam].
- Bordignon, L., Faria, A. P., França, M. G., and Fernandes, G. W. 2019. Osmotic stress at membrane level and photosystem II activity in two C4 plants after growth in elevated CO2 and temperature. *Ann. Appl. Biol.* 174(2): 113-122.
- Borrell, A. K., Mullet, J.E., George-Jaeggli, B., van Oosterom, E. J., Hammer, G. L., Klein, P. E., and Jordan, D. R. 2014. Drought adaptation of stay-green sorghum is associated with canopy development, leaf anatomy, root growth, and water uptake. *J. Exp. Bot.* 65(21): 6251-6263.
- C2ES [Center for Climate and Energy Solutions]. 2021. C2ES home page [on line]. Available: <https://www.c2es.org/content/climate-resilience-overview/>. [03 Nov 2021].
- CIMMYT, 2020. *Climate-resilient maize for Asia (CRMA)*. International Maize and Wheat Improvement Center. Available at: <https://www.cimmyt.org/projects/climate-resilient-maize-for-asia-crma/> (Accessed: 23 December 2024).
- Clements, D., & Ditommaso, A. (2011). Climate change and weed adaptation: can evolution of invasive plants lead to greater range expansion than forecasted? *Weed research*, 51(3), 227-240.
- Conley, M. M., Kimball, B. A., Brooks, T. J., Pinter Jr, P. J., Hunsaker, D. J., Wall, G. W., Adam, N. R., LaMorte, R. L., Matthias, A. D., Thompson, T. L. and Leavitt, S. W. 2001. CO2 enrichment increases water use efficiency in sorghum. *New Phytol.* 151(2): 407-412.
- Cordaid. 2016. Promoting climate resilient agriculture for sustainable livelihoods [online]. Available: <https://www.cordaid.org/en/wp-content/uploads/sites/3/2016/11/2016-11-Cordaid-4P-lowres-Climature-Resilient-Agriculture.pdf> [03 Nov 2021].
- Correia, P. M., da Silva, A. B., Vaz, M., Carmo-Silva, E., and da Silva, J. M. 2021. Efficient Regulation of CO2 Assimilation Enables Greater Resilience to High Temperature and Drought in Maize. *Front. Plant Sci.* 12.

De Souza, A. P., Cocuron, J. C., Garcia, A. C., Alonso, A. P., and Buckeridge, M. S. 2015. Changes in whole-plant metabolism during the grain-filling stage in sorghum grown under elevated CO₂ and drought. *Plant Physiol.* 169(3): 1755-1765.

Dusenge, M. E., Duarte, A. G., & Way, D. A. (2019). Plant carbon metabolism and climate change: elevated CO₂ and temperature impacts on photosynthesis, photorespiration and respiration. *New Phytologist*, 221(1), 32-49.

Edwards, G.E., Franceschi, V.R. and Voznesenskaya, E.V., 2004. Single-cell C₄ photosynthesis versus the dual-cell (Kranz) paradigm. *Annu. Rev. Plant Biol.*, 55(1), pp.173-196.

Ehleringer, J. R., & Cerling, T. E. (2002). C₃ and C₄ photosynthesis. *Encyclopedia of global environmental change*, 2(4), 186-190.

Gallaher, T. J., Peterson, P. M., Soreng, R. J., Zuloaga, F. O., Li, D. Z., Clark, L. G., Tyrrell, C. D., Welker, C. A., Kellogg, E. A., & Teisher, J. K. (2022). Grasses through space and time: An overview of the biogeographical and macroevolutionary history of Poaceae. *Journal of Systematics and Evolution*, 60(3), 522-569.

Garnett, T. (2009). Livestock-related greenhouse gas emissions: impacts and options for policy makers. *Environmental science & policy*, 12(4), 491-503.

Ghannoum, O. 2016. How can we breed for more water use-efficient sugarcane?. *J. Exp. Bot.* 67(3): 557–559.

Ghannoum, O, Caemmerer, S. V., and Conroy, J. P. 2002. The effect of drought on plant water use efficiency of nine NAD-ME and nine NADP-ME Australian C₄ grasses. *Funct. Plant Biol.* 29:1337–1348.

Ghannoum, O., Caemmerer, S. V., Ziska, L. H., and Conroy, J.P. 2000. The growth response of C₄ plants to rising atmospheric CO₂ partial pressure: a reassessment. *Plant Cell Environ.* 23(9): 931-942.

Ghannoum, O., Evans, J. R., and Caemmerer, S. 2011. Nitrogen and water use efficiency of C₄ plants. In: Raghavendra, A. S. and Sage, R. F. (eds). *C₄ Photosynthesis and Related CO₂ Concentrating Mechanisms*. Springer, Dordrecht, pp 129–146.

Goh, C.-H., Ko, S.-M., Koh, S., Kim, Y.-J., & Bae, H.-J. (2012). Photosynthesis and environments: photoinhibition and repair mechanisms in plants. *Journal of Plant Biology*, 55, 93-101.

Gong, Z., Dong, L., Lam, S., Zhang, D., Zong, Y., Hao, X., and Li, P. 2021. Nutritional quality in response to elevated CO₂ concentration in foxtail millet (*Setaria italica*). *J. Cereal Sci.* 103318.

Gowik, U., & Westhoff, P. (2011). The path from C₃ to C₄ photosynthesis. *Plant Physiology*, 155(1), 56-63.

Hatch, M. D. (1987). C₄ photosynthesis: a unique blend of modified biochemistry, anatomy and ultrastructure. *Biochimica et Biophysica Acta (BBA)-Reviews on Bioenergetics*, 895(2), 81-106.

- Hirst, D., 2021. Aviation, decarbonisation and climate change. *Aviation*.
- Hodkinson, T. R. (2018). Evolution and taxonomy of the grasses (Poaceae): A model family for the study of species-rich groups. *Annual plant reviews Online*, 255-294.
- Huang, S., Knight, C. A., Hoover, B. K., & Ritter, M. (2020). Leaf functional traits as predictors of drought tolerance in urban trees. *Urban Forestry & Urban Greening*, 48, 126577.
- Jabereldar, A. A., El Naim, A. M., Abdalla, A. A., and Dagash, Y. M. 2017. Effect of water stress on yield and water use efficiency of sorghum (*Sorghum bicolor* L. Moench) in semi-arid environment. *Int. J. Agric. Forestry*. 7(1): 1-6.
- Kajala, K., Covshoff, S., Karki, S., Woodfield, H., Tolley, B. J., Dionora, M. J. A., Mogul, R.T., Mabilangan, A. E., Danila, F. R., Hibberd, J. M., and Quick, W. P. 2011. Strategies for engineering a two-celled C4 photosynthetic pathway into rice. *J. Exp. Bot.* 62(9): 3001-3010.
- Karakurt, I., Aydin, G., & Aydiner, K. (2012). Sources and mitigation of methane emissions by sectors: A critical review. *Renewable energy*, 39(1), 40-48.
- Khalifa, M., & Eltahir, E. A. (2023). Assessment of global sorghum production, tolerance, and climate risk. *Frontiers in Sustainable Food Systems*, 7, 1184373.
- Kijne, J. W. (2006). Abiotic stress and water scarcity: identifying and resolving conflicts from plant level to global level. *Field Crops Research*, 97(1), 3-18.
- Korres, N. E., Norsworthy, J. K., Tehranchian, P., Gitsopoulos, T. K., Loka, D. A., Oosterhuis, D. M., Gealy, D. R., Moss, S. R., Burgos, N. R., & Miller, M. R. (2016). Cultivars to face climate change effects on crops and weeds: a review. *Agronomy for Sustainable Development*, 36, 1-22.
- Koteyeva, N. K., Voznesenskaya, E. V., Pathare, V. S., Borisenko, T. A., Zhurbenko, P. M., Morozov, G. A., & Edwards, G. E. (2023). Biochemical and structural diversification of C4 photosynthesis in Tribe Zoysieae (Poaceae). *Plants*, 12(23), 4049.
- Lal, M. (2000). Climatic change-implications for India's water resources. *Journal of Social and Economic Development*, 3, 57-87.
- Lara, M. V., & Andreo, C. S. (2011). C4 plants adaptation to high levels of CO₂ and to drought environments. *Abiotic stress in plants-mechanisms and adaptations*, 415-428.
- Leakey, A. D., Ferguson, J. N., Pignon, C. P., Wu, A., Jin, Z., Hammer, G. L., & Lobell, D. B. (2019). Water use efficiency as a constraint and target for improving the resilience and productivity of C3 and C4 crops. *Annual review of plant biology*, 70, 781-808.
- Lenka, B., Kulkarni, G. U., Moharana, A., Singh, A. P., Pradhan, G. S., and Muduli, L. 2020. Millets: Promising Crops for Climate-Smart Agriculture. *Int. J. Curr. Microbiol. App. Sci*, 9(11): 656-668.
- Levitus, S., Antonov, J., Boyer, T., Baranova, O., Garcia, H., Locarnini, R., Mishonov, A., Reagan, J., Seidov, D., & Yarosh, E. (2017). NCEI ocean heat content, temperature anomalies, salinity anomalies, thermocline sea level anomalies, halosteric sea level

anomalies, and total steric sea level anomalies from 1955 to present calculated from in situ oceanographic subsurface profile data (NCEI Accession 0164586). *NOAA National Centers for Environmental Information*, 10, v53f54mvp.

Liaqat, W., Altaf, M. T., Barutçular, C., Mohamed, H. I., Ahmad, H., Jan, M. F., & Khan, E. H. (2024). Sorghum: a Star Crop to Combat Abiotic Stresses, Food Insecurity, and Hunger Under a Changing Climate: a Review. *Journal of Soil Science and Plant Nutrition*, 1-28.

Lin, H. C., Coe, R. A., Quick, W. P., & Bandyopadhyay, A. (2019). Climate-resilient future crop: Development of C 4 Rice. *Sustainable solutions for food Security: Combating climate change by adaptation*, 111-124.

Long, S. (1983). C4 photosynthesis at low temperatures. *Plant, Cell & Environment*, 6(4), 345-363.

Long, S. P., Ainsworth, E. A., Rogers, A., & Ort, D. R. (2004). Rising atmospheric carbon dioxide: plants FACE the future. *Annu. Rev. Plant Biol.*, 55(1), 591-628.

Lopes, M. S., Araus, J. L., Van Heerden, P. D., & Foyer, C. H. (2011). Enhancing drought tolerance in C4 crops. *Journal of experimental botany*, 62(9), 3135-3153.

Maheswari, M., Sarkar, B., Vanaja, M., Rao, S. M., Prasad, J. V. N. S., Prabhakar, M., Chary, R. G., Venkateswarlu, B., Choudhury, R. P., Yadava, D. K., and Bhaskar, S. 2015. *Climate resilient crop varieties for sustainable food production under aberrant weather conditions*. NICRA Bulletin No.4. ICAR-Central Research Institute for Dryland Agriculture, Hyderabad. 64p.

Mall, R. K., Gupta, A., & Sonkar, G. (2017). Effect of climate change on agricultural crops. In *Current developments in biotechnology and bioengineering* (pp. 23-46). Elsevier.

Marin, F. R., Jones, J. W., Singels, A., Royce, F., Assad, E. D., Pellegrino, G. Q., and Justino, F. 2013. Climate change impacts on sugarcane attainable yield in southern Brazil. *Clim. Change*. 117(1): 227-239.

Mehri, N., Fotovat, R., Saba, J., & Jabbari, F. (2009). Variation of stomata dimensions and densities in tolerant and susceptible wheat cultivars under drought stress. *Journal of Food Agriculture and Environment*, 7(1), 167-170.

Misra, V., Shrivastava, A. K., Mall, A. K., Solomon, S., Singh, A. K., and Ansari, M. I., 2019. Can sugarcane cope with increasing atmospheric CO₂ concentration. *Aust. J. Crop Sci.* 13(5): 780-784.

Morrone, O., Aagesen, L., Scataglini, M. A., Salariato, D. L., Denham, S. S., Chemisquy, M. A., Sede, S. M., Giussani, L. M., Kellogg, E. A., & Zuloaga, F. O. (2012). Phylogeny of the Paniceae (Poaceae: Panicoideae): integrating plastid DNA sequences and morphology into a new classification. *Cladistics*, 28(4), 333-356.

Nadew, D., Bejiga, T., & Teressa, T. (2021). Breeding of sorghum crop for resistance and tolerance to drought. *Int. Res. Plant Crops Sci*, 188-195.

NASA [National Aeronautics and Space Administration]. 2024. NASA home page [on line]. Available: <https://climate.nasa.gov/>. [12Dec 2024].

National Geographic Society, n.d. *Earth's Changing Climate*. National Geographic Education. Available at: <https://education.nationalgeographic.org/resource/earths-changing-climate/> (Accessed: 23 December 2024).

Nerem, R. S., Beckley, B. D., Fasullo, J. T., Hamlington, B. D., Masters, D., & Mitchum, G. T. (2018). Climate-change-driven accelerated sea-level rise detected in the altimeter era. *Proceedings of the National Academy of Sciences*, *115*(9), 2022-2025.

Nunes, T. D., Zhang, D., & Raissig, M. T. (2020). Form, development and function of grass stomata. *The Plant Journal*, *101*(4), 780-799.

Osmond, C. B. (2007). Crassulacean acid metabolism: now and then. In *Progress in Botany* (pp. 3-32). Springer.

Ottman, M. J., Kimball, B. A., Pinter, P. J., Wall, G. W., Vanderlip, R. L., Leavitt, S.W., LaMorte, R. L., Matthias, A. D., and Brooks, T. J. 2001. Elevated CO₂ increases sorghum biomass under drought conditions. *New Phytol.* *150*(2): 261-273.

Pardo, J. and VanBuren, R. 2021. Evolutionary innovations driving abiotic stress tolerance in C₄ grasses and cereals. *Plant Cell*. *33*(11): 3391-3401.

Paul, M. J., & Foyer, C. H. (2001). Sink regulation of photosynthesis. *Journal of experimental botany*, *52*(360), 1383-1400.

Peters, K., Breitsameter, L., & Gerowitt, B. (2014). Impact of climate change on weeds in agriculture: a review. *Agronomy for Sustainable Development*, *34*, 707-721.

Pignon, C. P. and Long, S. P. 2020. Retrospective analysis of biochemical limitations to photosynthesis in 49 species: C₄ crops appear still adapted to pre-industrial atmospheric [CO₂]. *Plant. Cell. Environ.* *43*(11): 2606-2622.

Pinto, H., Powell, J. R., Sharwood, R. E., Tissue, D. T., and Ghannoum, O. 2016. Variations in nitrogen use efficiency reflect the biochemical subtype while variations in water use efficiency reflect the evolutionary lineage of C₄ grasses at interglacial CO₂. *Plant. Cell. Environ.* *39*(3): 514-526.

Ramesh, K., Matloob, A., Aslam, F., Florentine, S. K., & Chauhan, B. S. (2017). Weeds in a changing climate: vulnerabilities, consequences, and implications for future weed management. *Frontiers in plant science*, *8*, 95.

Raza, A., Razzaq, A., Mehmood, S. S., Zou, X., Zhang, X., Lv, Y., & Xu, J. (2019). Impact of climate change on crops adaptation and strategies to tackle its outcome: A review. *Plants*, *8*(2), 34.

Raza, G., Ali, K., Hassan, M. A., Ashraf, M., Khan, M. T., & Khan, I. A. (2019). Sugarcane as a bioenergy source. *Sugarcane biofuels: status, potential, and prospects of the sweet crop to fuel the world*, 3-19.

Romanowska, E., & Wasilewska-Dębowska, W. (2022). Light-Dependent Reactions of Photosynthesis in Mesophyll and Bundle Sheath Chloroplasts of C₄ Plant Maize. How Our Views Have Changed in Recent Years. *Acta Societatis Botanicorum Poloniae*, *91*.

Ruiz-Vera, U. M., Siebers, M. H., Drag, D. W., Ort, D. R., and Bernacchi, C. J. 2015. Canopy warming caused photosynthetic acclimation and reduced seed yield in maize grown at ambient and elevated [CO₂]. *Glob. Change Biol.* 21(11): 4237-4249.

Rykowski, K. (2000). The role of forest ecosystems and wood in controlling the absorption and emission of carbon dioxide. *Geographia Polonica*, 73(2), 65-88.

Sabine, C. L., Feely, R. A., Gruber, N., Key, R. M., Lee, K., Bullister, J. L., Wanninkhof, R., Wong, C., Wallace, D. W., & Tilbrook, B. (2004). The oceanic sink for anthropogenic CO₂. *science*, 305(5682), 367-371.

Sage, R. F., & Kubien, D. S. (2007). The temperature response of C₃ and C₄ photosynthesis. *Plant, Cell & Environment*, 30(9), 1086-1106.

Sage, R. F., Christin, P. A., Edwards, E. J. (2011) The C₄ plant lineages of planet Earth. *J. Exp. Bot.* 62:3155–3169.

Sage, R. F. and Zhu, X. G. 2011. Exploiting the engine of C₄ photosynthesis. *J. Exp. Bot.* 62(9): 2989-3000.

Schiller, K., & Bräutigam, A. (2021). Engineering of crassulacean acid metabolism. *Annual review of plant biology*, 72(1), 77-103.

Schlüter, U., & Weber, A. P. (2020). Regulation and evolution of C₄ photosynthesis. *Annual review of plant biology*, 71(1), 183-215.

Schmitt, M. R., & Edwards, G. E. (1981). Photosynthetic capacity and nitrogen use efficiency of maize, wheat, and rice: a comparison between C₃ and C₄ photosynthesis. *Journal of experimental botany*, 32(3), 459-466.

Sheehy, J. E., Ferrer, A. B., Mitchell, P. L., Elmido-Mabilangan, A., Pablico, P., and Dionora, M. J. A. 2008. How the rice crop works and why it needs a new engine. In: *Charting new pathways to C₄ rice* .pp. 3-26.

Siebers, M. H., Slattery, R. A., Yendrek, C. R., Locke, A. M., Drag, D., Ainsworth, E. A., Bernacchi, C. J., and Ort, D. R. 2017. Simulated heat waves during maize reproductive stages alter reproductive growth but have no lasting effect when applied during vegetative stages. *Agric. Ecosyst. Environ.* 240: 162-170.

Sonawane, B. V., Sharwood, R. E., Whitney, S., Ghannoum, O. 2018. Shade compromises the photosynthetic efficiency of NADP-ME less than that of PEP-CK and NAD-ME C₄ grasses. *J. Exp Bot.* 69:3053–3068.

Sovacool, B. K., Griffiths, S., Kim, J., & Bazilian, M. (2021). Climate change and industrial F-gases: A critical and systematic review of developments, sociotechnical systems and policy options for reducing synthetic greenhouse gas emissions. *Renewable and Sustainable Energy Reviews*, 141, 110759.

Stavi, I., & Lal, R. (2013). Agriculture and greenhouse gases, a common tragedy. A review. *Agronomy for Sustainable Development*, 33, 275-289.

Tingting, X., Peixi, S. U., and Lishan, S. 2010. Photosynthetic characteristics and water use efficiency of sweet sorghum under different watering regimes. *Pak. J. Bot.* 42(6): 3981-3994.

Velicogna, I., Mohajerani, Y., Landerer, F., Mouginot, J., Noel, B., Rignot, E., Sutterley, T., van den Broeke, M., van Wessem, M., & Wiese, D. (2020). Continuity of ice sheet mass loss in Greenland and Antarctica from the GRACE and GRACE Follow-On missions. *Geophysical Research Letters*, 47(8), e2020GL087291.

Venter, N. (2015). *Drought responses of selected C4 photosynthetic NADP-Me and NAD-Me Panicoideae and Aristidoideae grasses* Rhodes University].

Viswanathan, P. K., Kavya, K., and Bahinipati, C. S., 2020. Global patterns of climate-resilient agriculture: A review of studies and imperatives for empirical research in India. *Review Dev. Change*. 25(2): 169-192.

Vu, J. C. and Allen Jr, L. H. 2009. Growth at elevated CO₂ delays the adverse effects of drought stress on leaf photosynthesis of the C₄ sugarcane. *J. Plant Physiol.* 166(2): 107-116.

Voznesenskaya, E. V., Franceschi, V. R., Chuong, S. D., & Edwards, G. E. (2006). Functional characterization of phosphoenolpyruvate carboxykinase-type C₄ leaf anatomy: immuno-, cytochemical and ultrastructural analyses. *Annals of Botany*, 98(1), 77-91.

Wall, G. W., T. J. Brooks, N. R. Adam, A. B., Cousins, B. A., Kimball, P. J., Pinter Jr, R. L., La Morte *et al.* 2001. Elevated atmospheric CO₂ improved sorghum plant water status by ameliorating the adverse effects of drought. *New Phytol.* 152(2): 231-248.

Wang, Y., Bräutigam, A., Weber, A. P., & Zhu, X.-G. (2014). Three distinct biochemical subtypes of C₄ photosynthesis? A modelling analysis. *Journal of experimental botany*, 65(13), 3567-3578.

Watson-Lazowski, A. and Ghannoum, O. 2021. The outlook for C₄ crops in future climate scenarios. In: Becklin, K. M., Ward, J. K., Way, D. A. (eds.), *Photosynthesis, Respiration, and Climate Change*. Springer, Cham, pp. 251-281.

Way, D. A., Katul, G. G., Manzoni, S., & Vico, G. (2014). Increasing water use efficiency along the C₃ to C₄ evolutionary pathway: a stomatal optimization perspective. *Journal of experimental botany*, 65(13), 3683-3693.

Zabaleta, E., Martin, M. V., and Braun, H. P. 2012. A basal carbon concentrating mechanism in plants. *Plant Sci.* 187:97-104.

Zahedi, S. M., Karimi, M., Venditti, A., Zahra, N., Siddique, K. H., & Farooq, M. (2024). Plant Adaptation to Drought Stress: The Role of Anatomical and Morphological Characteristics in Maintaining the Water Status. *Journal of Soil Science and Plant Nutrition*, 1-19.

Zhang, D., Li, A., Lam, S. K., Li, P., Zong, Y., Gao, Z., and Hao, X. 2021. Increased carbon uptake under elevated CO₂ concentration enhances water-use efficiency of C₄ broomcorn millet under drought. *Agric. Water Manag.* 245: 106631.

Zheng, Y. P., Li, R. Q., Guo, L. L., Hao, L. H., Zhou, H. R., Li, F., Peng, Z. P., Cheng, D. J., and Xu, M. 2018. Temperature responses of photosynthesis and respiration of maize (*Zea mays*) plants to experimental warming. *Russ. J. Plant Physiol.* 65(4): 524-531.

Ziska, L. W. 2003. Evaluation of yield losses in field sorghum from a C₃ and C₄ weeds with increasing CO₂. *Weed Sci.* 51: 914-918.

UNDER PEER REVIEW