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Comprehensive review on the consequences and omics approaches for boosting wheat production under drought stress

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Abstract

Plants are subjected to a number of abiotic stresses in the field, many of which are detrimental to plant development and output. Drought is one of the most detrimental environmental conditions for crop cultivation. Wheat, a major crop is severely affected by drought stress. Wheat production declines due to drought are likely to have significant losses from all other factors. There are big consequences of drought stress on wheat's physiological, biochemical, and morphological properties, as well as the effects on growth, water relations, and photosynthesis. Drought stress causes changes in leaf size, stem length, and root multiplication, as well as an imbalance in plant-water relationships and a decrease in water efficiency. Drought stress causes alterations in the wheat plant, which are being investigated through physiological investigations. One of the most effective strategies to make wheat plants drought tolerant is to boost their antioxidant defense system to minimise oxidative stress. Plant biologists are developing novel techniques to increase the antioxidant defense system in order to mitigate the impacts of droughtinduced plant damage. Drought tolerance in plants is conferred via a number of genes and their overexpression. Recent advances in wheat genomic, transcriptomic, proteomic, and metabolomic research under varying levels of drought give critical information for generating drought-tolerant wheat cultivars. This overview aims to summarise and update understanding on drought stress's impacts on wheat and tolerance mechanisms, as well as the omics approach to wheat plant drought stress adaptation.

Keywords

Wheat; Drought stress; Consequences; Omics approaches; Agriculture productivity

1. Introduction

Wheat (Triticum aestivum) is the world's most frequently farmed food crop, and it is endangered by climate change in the future (Pequeno et al., 2021). By mid-century, climate change is expected to reduce world wheat production by 1.9% (Pequeno et al., 2021). The poor nations in tropical regions are expected to bear the brunt of the negative consequences. African countries' productivity are expected to reduce 15% by 2050, while yields in Southern Asia are expected to fall 16% (Pequeno et al., 2021). Plants are subjected to a variety of abiotic stresses in the field, all of which are detrimental to plant growth and development. Because of the extreme and rapid worldwide climatic changes, the simultaneous occurrence of abiotic stress, particularly drought stress in plants, might alter the morphological, physiological, and molecular responses in plants (Hussain et al., 2018). As a result of the drought, maize, peanut, wheat, sunflower, sugarcane, and cotton yields and yield components have all decreased significantly (Furlan et al., 2012; Tahir et al., 2002; Barnabas et al., 2008; Vasantha et al., 2005; Kamara et al., 2003).

Wheat production has always been subject to several constraints, resulting in the formation of biotic and abiotic limitations, among them, drought is a major concern. Due to global climate change, drought stress is becoming increasingly severe and frequent in the Indo-Gangetic floodplain, particularly during the winter wheat growing season (Shekhar and Paul, 2012).

Drought stress is characterised by a shortage of water that results in morphological, biochemical, physiological, and molecular changes (Sallam et al., 2019). Photosynthesis, chlorophyll production, nutrient metabolism, ion absorption and translocation, respiration, and carbohydrates metabolism are all affected by drought stress in plants (Li et al., 2011; Farooq et al., 2009; Oyiga et al., 2020). The immediate result is decreased organ output, greater flower abortion, and a shorter grain filling time, all of which have an impact on crop yield (Koua et al., 2021).Drought stress, along with high temperatures during the reproductive stage (ultimate development phase), is a major contributor to low wheat production in the tropics and subtropics (Sattar *et al.*, 2020).

Wheat must adapt to drought conditions in order to survive, and numerous resistant genotypes have been developed to assist preserve soluble sugars, proline content, amino acids, chlorophyll content, enzymatic and nonenzymatic well as as antioxidant activities (Abid et al., 2016). Wheat has enhanced its drought tolerance mechanisms; nevertheless, these mechanisms differ and are dependent on the crop kinds and cultivars (Tefera et al., 2021). Durum wheat was discovered to be able to survive water stress better than bread wheat (Tefera et al., 2021). Drought stress reduced the grain filling period by 15-24% and grain production by 11-34% (Islam et al., 2021). When compared to irrigated circumstances, yield decreased by 3.09t ha⁻¹ (-46.8%) under semiarid conditions in winter wheat (Neha et al., 2021). Yield and kernel per meter square decreased by 68.71% and 66.05% respectively (Koua et al., 2021). Moreover, plant height, the number of spikelets spike⁻¹, spike length, chlorophyll content, and relative water were all reduced by 26%, 23%, 9%, 11%, and 16%, respectively under drought stress (Tefera et al., 2021). Thus, to maintain food security under the changing climatic conditions, the development of stress tolerance in wheat might be a promising approach.

An overview of omics technologies, such as transcriptomics, metabolomics, and proteomics, is currently used to gain an in-depth, precise, and systemic understanding of key biological and cellular pathways engaged by agricultural plants during stress (Zargar *et al.*, 2022). Mutant libraries, cDNAs, expression profiles, sequence data sets, and quantitative trait loci (QTLs) are all valuable resources in structural and genomics research (Jiang *et al.*, 2011). Cloning of many genes and discovered QTLs were significant for drought tolerance (Vikram *et al.*, 2012). To address the complicated nature of drought stress and mitigating mechanisms for tolerating drought

stress, a large number of QTLs were found (Fleury et al., 2010). In addition, numerous QTLs responsible for increased grain output and other secondary traits that allow rainfed uplands and lowlands to tolerate drought stress have been found (Bernier et al., 2007; Venuprasad et al., 2009). Under drought conditions, different OTL performance has been seen in upland and lowland habitats. Under drought conditions, different QTL performance has been seen in upland and lowland habitats. Finally, the best QTLs for the ecosystems, genetic background, and environmental conditions are chosen. Furthermore, marker assisted back crossing (MABC) has been successfully used in crop plants to generate high yielding cultivars. MABC has been used to boost the yield of crop varieties in Thailand's north-eastern provinces (Kanjoo et al., 2012). Applications of molecular breeding techniques such as marker-assisted selection (MAS), SNP marker applications, and genome wide assisted selection (GWAS) are expected to pave the way for understanding the molecular mechanisms underlying crop plant resilience to a wide range of environmental challenges, such as drought stress, in the near future.

1.1. Consequences of drought on wheat

Drought causes plants to undergo a number of physiological and molecular changes, the majority of which aid in their adaptation to the harsh environment. Drought stress has both direct and indirect effects on plant metabolism. Droughtinduced stress changes the morpho-anatomical, physiological, and biochemical composition of plants, reducing transpiration and improving the efficiency with which the plants use the water they have stored. Leaf water deficits come from constant water loss through transpiration. Drought stress, on the other hand, has a variety of effects, ranging from lesions to confusion (Bijalwan *et al.*, 2022). Some consequences are discussed below-

1.1.1. Morphological effect

As a response to drought stress, wheat undergoes morphological changes such as reduced plant size, early maturity, decreased leaf area, reduced yield, limited leaf extension, small leaf size, reduced number of tillers, reduced leaf longevity, reduced total shoot length, decreased plant height, increased leaf rolling, and reduced plant biomass (Rijal et al., 2021). Drought stress has a substantial impact on seed germination, seedling development, dry matter partitioning, root growth, root depth, and extension (Lonbani and Arzani, 2011). The root is the first organ to be influenced by drought stress (Shimazaki et al., 2005). Under water stress, the root grows in order to find water, while the growth of the shooting component is stunted (Ahmad et al., 2018). The fall in biomass (Wang et al., 2005), grain number (Dolferus et al., 2011), is mostly caused by terminal drought, which results in a large reduction in wheat output. One of the most visible symptoms of leaf senescence is chlorosis, which causes a reduction in photosynthesis (Ali et al., 2020). Wheat grown in extreme drought circumstances can bring the entire plant to senescence, but it also improves the mobilisation of stored carbohydrates from the stem and leaves to forming grains during parenthesis, so compensating the yield loss incurred by senescence during drought stress (Nawaz et al., 2013; Faroog et al., 2014).

Drought stress requires specific physiological changes in the plant in order to mitigate the effects of drought stress (Vinocur and Altman, 2005). Adaptation mechanisms in wheat for dealing with drought stress include osmotic root adjustment, greater root penetration into the soil, increased root density, and increased root to shoot ratio (Ali *et al.*, 2020). The relevance of root angle in irrigated settings and roots per shoot in drought situations for boosting grain production might be crucial for designing drought-tolerant cultivars was concluded by Neha *et al.*, (2021).

Canopy stay green traits aided in the selection of drought-tolerant genotypes, since they may confer longer photosynthesis, nutrient, and water absorption under stress (Lopes and Reynolds, 2012). The use of an RGB-based vegetation index to characterize canopy green area dynamics might reduce selection time and improve precision (Neha *et al.*, 2021).

1.1.2. Physiological effect

Drought stress has been linked to a wide range of physiological responses. Drought stress on wheat crops is mitigated by a variety of physiological characteristics (Rijal et al., 2021). The availability of water has a direct link with the functioning of many physiological processes in plants. These physiological processes are disrupted when water availability is reduced, and plants are unable to create appropriate amounts of dry matter (Barbeta et al., 2015). Water deficit causes a decrease in water status during crop growth, soil water potential, and plant osmotic potential for water and nutrient absorption, which reduces leaf turgor pressure, causing plant metabolic processes to be disrupted (Mehraban and Miri, 2017). Drought significantly impacts water relations, nutrient absorption, growth, and yield in wheat at later stages (6 weeks after emergence) than at earlier stages (3 weeks after seedling emergence) (Nawaz et al., 2014). In comparison to their individual impacts, combined drought and heat shocks had the most devastating consequences on plant relative water relations except turgor potential (Sattar et al., 2020). Plants restrict their stomata (perhaps via ABA signalling) when the volume of available water decreases, reducing CO_2 input. CO_2 reduction not only reduces carboxylation but also directs additional electrons to generate reactive O₂ species (Ahmad et al., 2018). Closing of stomata, a reduction in photosynthetic activity, the onset of oxidative stress, changes in cell wall integrity, and the formation of toxic compounds that lead to plant death are some of the physiological changes (Alghabari and Ihsan, 2018). Different abiotic stressors attack biological membranes first and foremost. The durability of maintenance

membranes under water stress is thought to be a key component of drought resistance in plants. Drought stress damages membrane integrity (Almeselmani et al., 2012). Drought stress also affects the electron transport chain, resulting in the generation of reactive oxygen species (ROS) that are detrimental to plant cells and organelles mitochondria, such chloroplasts, as and peroxisomes (Farooqi et al., 2020).Under drought conditions, RuBisCO (ribulose-1, 5-bisphosphate carboxylase/oxygenase) enzyme activity is inhibited, resulting in a decrease in photosynthetic amount and the generation of ATP (Dulai et al., 2006).

Plants must adapt to drought conditions in order to survive, and several tolerant genotypes have been developed to assist preserve soluble sugars, proline content, amino acids, chlorophyll content, and non-enzymatic enzymatic antioxidant activities (Abid et al., 2016). When compared to leaf water potential, relative water content (RWC) is an effective predictor of water status in drought conditions (Lugojan and Ciulca, 2011), as it decreases when leaf exposed to drought (Nayyar & Gupta, 2006) and it has also been used to find drought-tolerant cultivars (Bayoumi et al., 2008). These several forms of plant drought tolerance mechanisms aid in the understanding of the response that physiological aids in the maintenance of growth and productivity during times of stress (Rijal et al., 2021). Similarly, these qualities are important in breeding projects aimed at developing drought-tolerant cultivars that can thrive in areas where water is scarce.

1.1.3. Biochemical effect

Reduced efficiency of Rubisco, photochemical, accumulation of stress metabolites (glybet, glutathione. and polyamines), antioxidant enzymes [peroxidase (POD), superoxide dismutase (SOD), ascorbate peroxidase (APX), catalase (CAT)], and reduced ROS accumulation are some of the biochemical effects in plants due to water stress (Nezhadahmadi et al., 2013). Drought stress has a deleterious impact on total soluble sugar (TSS), total carbohydrate (TC), and

enzyme activity in wheat (Hammad and Ali, 2014). The culm reserves, or water-soluble carbohydrates (WSCs), contributed less to the grains as a result of the stress (Islam *et al.*, 2021). When the generation and build-up of ROS increases, it affects cellular processes such as protein degradation, enzyme inhibition, oxidative damage to DNA and RNA, and membrane lipid peroxidation, all of which lead to cell death (Ishikawa *et al.*, 2010; Huseynova, 2012).

Wheat plants mostly in the heading stage, accumulate proline content to a greater extent than any other osmoregulators due to a shortage of water (Maralian et al., 2010). Tefera et al., (2021) stated that wheat in response to drought increases proline and total soluble sugar by 28% and 6% respectively and biochemical parameters need to be considered as a better option for the selection process under drought stress. Todorova et al., (2021) conclude that Serrate (selective herbicide) affects the biochemical reactions of drought-stressed wheat seedlings and after the combined herbicide+drought treatment, stress indicators, the enzymatic and non-enzymatic antioxidant defense were also elevated throughout the stress phase.

As the production of ROS increases in stress (Farooqi et al., 2020), enzymatic and nonenzymatic systems get activated to detoxify the toxic level of ROS (Caverzan et al., 2016). This reaction is also influenced by other aspects such as tissue type, stress duration, and intensity, and developmental stage, demonstrating the intricacy of the process of ROS formation and detoxification, as well as the impact of ROS on the antioxidant mechanism (Caverzan et al., 2016). Hussain et al., (2021) remarked that the increase in output was attributed to the wheat genotype's improved drought tolerance, as ABA (a form of abscisic acid) aids the plant in watersaving because an increase in the concentration of ABA near the guard cell causes the closer of stomata (Sourour et al., 2017). Silicon application (anthesis stage- foliar spray, tillering stageenhances plant fertigation) biochemical characteristics, nutrient absorption, and growth

rate, all of which aid the plant in mitigating drought-stressing impacts (Bukhari *et al.*, 2021). The latest findings by Gyugos *et al.*, (2021) of far-red light's adaptive adjustment of glutathione and amino acid levels during drought, which can be used as a foundation for spectrum-dependent modification of crops' defensive metabolites (glutathione, proline) to prevent stress-induced damage.

2. Omics approaches for increasing productivity of wheat under drought stress

Omics, or the study of an organism's genes, transcripts, proteins, and metabolites, decodes the complete genome in order to get a better knowledge of plant molecular reactions and develop precise crop development tactics (Jain et al., 2019). The key requirement for demonstrating the exact link between genotype and phenotype is the integration of crop functional genomics and phenomics data (Yang et al., 2020). One of the reasons for the limited contribution of genomicsassisted selection-based production of droughttolerant wheat cultivars might be the polygenic nature of drought tolerance and the huge genome size of wheat (Mwadzingeni et al., 2016). Koobaz et al., (2020) used metabolomics and proteomics to investigate the mechanism behind wheat desiccation tolerance, as well as the function of several stress-responsive proteins and metabolites. Figure 1 represents an overview of the integrated multi-omics approach for crop improvement. Combining multi-omics and systems biology to construct a top-down (phenotype to genotype) and bottom-up (genotype to phenotype) model for crop breeding improvement under environmental difficulties could be beneficial (Yang et al. 2021). The drought signalling route in wheat is depicted in Figure 2.



Figure 2: Drought signalling pathway in wheat.

2.1. Genomics approach

Genomics focuses on the structure, function, evolution, mapping, epigenomic, metagenomic, and genome editing aspects of genes and genomes (Muthamilarasan et al., 2019). To enhance drought tolerance. several methods like quantitative trait locus (QTL) mapping, marker aided breeding, and introgression of a gene from a wild gene pool have been used (Merchuk-Ovnat et al., 2016; Mwadzingeni et al., 2016). Pour-Aboughadareh et al., (2017) suggest that wild cousins of wheat might be used to increase stress Khanna-Chopra tolerance. et al.. (2020)discovered notable OTLs on chromosome 3B regarding cell membrane stability as well as on chromosome 2D for flag leaf influencing factors (area, breadth, and length). A key yield QTL (Qyld.csdh.7AL) was recently incorporated in four elite Indian wheat lines to create droughttolerant genotypes with good yielding qualities under stressed conditions (Gautam et al., 2020). Zandipour et al., (2020), found a QTL hotspot for nine key characters (grain number per spike, spike length, spike weight, plant height, biological yield, grain yield, stem weight, and thousandgrain weight) on wheat chromosome 1B under terminal drought stress. Table 1 shows all recently

developed QTLs and related traits for drought tolerance.

Molecular markers have greatly aided us in improving desired qualities, such as drought tolerance, with more efficacy and dependability (Budak et al., 2015). The development of sequence-based simple sequence repeats (SSRs) and single nucleotide polymorphisms (SNPs) markers has been aided by recent advances in DNA sequencing and genotyping techniques (Pascual et al., 2020), mostly SNPs due to their abundance in nature (Winfield et al., 2016). Hua et al., (2019) found target genes in wheat that might be used to increase drought resistance through genetic engineering. By studying the expression of wheat genes P5CS and P5CR during drought stress, Karolina et al., (2019) discovered that they play a significant role in determining tolerance to water deficiencies. According to Neha et al., (2021), TaCwi.4A genetic marker for drought tolerance was highly associated with grain yield in wheat under water stress and also suggested it for use in future breeding programs. Figure 3 provides a percentage of QTLs with related trait and their presence in chromosome responsible for drought tolerance in wheat.



Figure 3: Graphical representation of percentage of drought responsible QTLs in chromosomes of wheat according to the work of previous researchers (Gahlaut *et al.*, 2017; Xu *et al.*, 2017; Ayalew *et al.*, 2018; Condorelli *et al.*, 2018; Fatima *et al.*, 2018; Sukumaran *et al.*, 2018; Dolferus *et al.*, 2019; Liu *et al.*, 2019; Goel *et al.*, 2019; Li *et al.*, 2019; Touzy *et al.*, 2019; Tura *et al.*, 2020; Ballesta *et al.*, 2020; Koua *et al.*, 2021).

Table 1.	Genes/Transgene	s responsible fo	or drought to	olerance in wheat
Lable 1.	Oches/ Hansgene	s responsible ru	n urought o	on and m wheat

S. No	Genes/Tran sgenes	Source	Putative function/Improved traits	Reference
1.	ERF1-V	Wheat	Induce biochemical and molecular changes in transgenic wheat plants during salt and drought stress	(Xing <i>et al.</i> , 2017)
2.	SeCspA	E. coli	More grain weight, yield, proline content, less reduction in chlorophyll content	(Yu <i>et al.</i> , 2017)
3.	TaODORA NT1	Wheat	Under salt and drought stress, an R2R3-MYB gene increases the expression of ROS and stress-related genes in transgenic tobacco	(Wei <i>et al.,</i> 2017)
4.	TaFER-5B	Wheat	Enhance temperature and drought tolerance, improve ROS and leaf iron content	(Zang <i>et al.</i> , 2017)
5.	TaDrSR1 and TaDrSR2	Wheat	Wheat drought tolerance proteins with differing levels of abundance	(Wang <i>et al.</i> , 2018)
6.	TaPEPKR2	Wheat	Enhance root length and drought tolerance	(Zang <i>et al.</i> , 2018)
7.	TaSHN1	Wheat	Enhance recovery from stress and lowers leaf water loss and stomatal density	(Bi <i>et al.</i> , 2018)
8.	TaWRKY2	Wheat	Higher soluble sugar, proline, chlorophyll and survival rate	(Gao <i>et al.</i> , 2018)
9.	CIPK23	Wheat	Enhanced ABA sensitivity, higher survival rate, increased osmolyte, stomata closure	(Cui <i>et al.,</i> 2018)
10.	TabZIP2	Wheat	More single seed weight, less spikes and seed	(Luang <i>et al.</i> , 2018)
11.	TaCML20	Wheat	In transgenic wheat, a CaM-like gene boosts water-soluble carbohydrate levels and yield	(Kalaipandian <i>et al.</i> , 2019)
12.	TaH2B-7D	Wheat	Knock-down wheat plants were used to investigate its involvement in drought tolerance transmission	(Wang <i>et al.</i> , 2019)
13.	TaCOMT	Wheat	In transgenic <i>Arabidopsis</i> , increases melatonin content and enhances drought tolerance	(Yang <i>et al.</i> , 2019)
14.	TaDr1	Wheat	A transcriptional repressor that is co-expressed in wheat with TaFT1 and TaVrn1 in response to drought stress	(Zotova <i>et al.</i> , 2019)

15.	TaSIM	Wheat	Reduces water loss and improves soluble sugar, proline, and stress-sensitive gene expression in transgenic <i>Arabidopsis</i> under drought stress	(Yu <i>et al.,</i> 2019)
16.	TdPIP2;1	Wheat	In transgenic wheat plants, an aquaporin gene engaged in salt and drought tolerance is found	(Ayadi <i>et al.,</i> 2019)
17.	TaMYBsm3	Wheat	In transgenic <i>Arabidopsis</i> plants, a MYB-CC transcription factor increases DT by upregulating the expression of stress-related genes (P5CS1, DREB2A, and RD29A)	(Li <i>et al.</i> , 2019)
18.	OTS1	Arabidopsis	More RWC, photosynthesis, and antioxidants, delayed senescence	(Li <i>et al.</i> , 2019)
19.	AtWRK30	Arabidopsis	Induces gas-exchange properties, osmolytes biosynthesis, antioxidant system, and stress-responsive gene expression in transgenic wheat plants, as well as heat and drought tolerance	(El-Esawi <i>et</i> <i>al.</i> , 2019)
20.	TdSHN1	Wheat	In transgenic tobacco plants, a transcription factor that increases salt and drought tolerance	(Djemal and Khoudi, 2019)
21.	SBPase	Brachypodium	Fully driven GUS expression promoted	(Alotaibi <i>et al.</i> , 2019)
22.	HaHB4	Sunflower	More water use efficiency and yield	(González <i>et al.</i> , 2019)
23.	GmDREB1	Soybean	Drought tolerance in transgenic wheat plants is improved.	(Zhou <i>et al.</i> , 2020)

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2.2. Transcriptomics approach

Transcriptomics is concerned with the transcriptome, which is the total collection of RNA transcripts generated by an organism's genome in a cell or tissue (Raza et al., 2021). With more whole-genome transcriptomic studies the genes linked to downstream signalling, production of stress response molecules, are undermined al., 2020). (Zang et The transcriptome analysis based on microarray and RNA sequencing (RNA-seq) has been used to deconstruct the wheat drought tolerance (Liu et al., 2015).

Drought-stressed wheat at the reproductive stage was studied using RNA-seq, and 309 droughtresponsive differentially expressed genes (DEGs) linked to photosynthetic activity, floral development, and stomatal movement were discovered (Ma *et al.*, 2017). Liu *et al.*, (2017) investigated the transcriptome of wheat during the early grain-filling stage of drought and discovered 566 DEGs related to signal transduction, metabolism, and transcription. They also looked at the involvement of phenylalanine metabolism and flavonoid production in wheat spikes under drought tolerance. Plant development has been found to be aided by the use of laser irradiation under various abiotic stress situations and with the use of RNA-seq based transcriptome analysis Qui et al., (2017) explored He-Ne radiation's role in wheat drought tolerance and found that the irradiation promoted 820 DEGs in wheat under water stress. Transcriptome profiling was done in roots of wheat under water stress using RNA-seq based transcriptome analysis by Hu et al., (2018). discovery of 45139 DEGs, The 13820 transcription factors, and 288 miRNAs by RNAseq of drought-stressed roots of tolerant (NI543941) and sensitive (WL711) wheat genotypes further highlighted the applicability of RNA-seq data in the dissection of particular QTL (Iquebal et al., 2019).

2.3. Proteomics approach

Proteomics is a technique for profiling the total expressed protein in an organism and it is separated into four parts: sequence, structural, functional, and expression proteomics (Mosa et al., 2017; Aizat and Hassan, 2018). The amino acid sequences are determined using sequence proteomics High-Performance by Liquid Chromatography (HPLC; Twyman, 2013). The structure of proteins is studied in structural proteomics to better understand their possible activities using computer-based modelling, and experimental methods including crystallization, nuclear magnetic resonance (NMR), X-rav diffraction of protein crystals, and electron microscopy (Woolfson, 2018).Functional proteomics identifies a protein's activities, which are then investigated using a variety of techniques such as yeast-one or two hybrids and protein microarray profiling (Lueong et al., 2014).

Wheat's different organs' proteomic responses to drought stress have been extensively studied, including seedling leaves and roots (Cheng *et al.*, 2015; Hao *et al.*, 2015), glume and awn development (Deng *et al.*, 2019), flag leaves (Deng *et al.*, 2018; Zhu *et al.*, 2020), and grain development (Ge *et al.*, 2012; Deng *et al.*, 2018; Duan *et al.*, 2020). Subcellular proteomics has recently been established to decode several organelle proteins involved in abiotic stress defense, such as those found in the plasma membrane, endoplasmic reticulum, mitochondria, and chloroplast (Zhang *et al.*, 2021; Zhu *et al.*, 2021).

Drought stress had a negative impact on the Bahar leaf proteome, resulting in a significant drop in total protein content, mostly due to deficiencies in photosynthetic proteins and enzymes associated with sugar and nitrogen metabolism, as well as the ability to detoxify toxic compounds, according to a study conducted on two spring wheat cultivars (Kavir, drought-tolerant; Bahar, droughtsusceptible). On the other hand, relatively little protein alterations were seen in Kavir stressed leaves (Michaletti *et al.*, 2018).

Comparative proteomics to evaluate and analyse the wheat types Chinese Spring and *D. villosum* (a species of *Triticeae*) discovered a total of 883 distinct abundant proteins (DAPs) (Wang *et al.*, 2021). Many proteins associated in defense/stress change radically between *D. villosum* and the wheat variety Chinese Spring, including the G subunit, zinc finger protein family, PR1, HSP family, LEA protein, and serpin family (Wang *et al.*, 2021). This discovery has paved the way for more research into the molecular mechanisms of wheat resilience and quality, as well as the development of wheat varieties based on wild relatives of wheat (Wang *et al.*, 2021).

In the cell nucleus subproteome of wheat growing grains, using label-free quantitative proteomic analysis Li *et al.*, (2021) found 398 water-deficit sensitive DAPs out of which 146 up-regulated DAPs were mostly engaged in oxidation-reduction and stress response, whereas 252 down-regulated DAPs were mostly involved in translation, oxidation-reduction, and cellular amino metabolic pathway. The wheat grain nuclear sub proteome's metabolic route in response to water shortage was hypothesised, providing new knowledge from a subcellular amine were for understanding the molecular

processes of plant drought stress response (Li et al., 2021).

2.4. Metabolomics approach

Metabolomics is a powerful method for obtaining detailed data on metabolite profiling and metabolic network analysis (Kumar et al., 2021). It also provides information on metabolites that have been identified as well as those that have yet to be identified (Kumar et al., 2021). Metabolite profiling techniques have been frequently utilized to analyse plant molecular responses to drought stress and assess metabolite levels in a specific metabolite class or pathway (Kusano et al., 2007). Proteomics simply detects gene products, but metabolomics may reveal how proteins are expressed metabolically and uncover biochemical processes that are critical for gene function (Lindon and Nicholson, 2008). Genome sequencing techniques like next-generation sequencing (NGS) and metabolite measurement through mass-spectrometry (MS) are used collectively to generate crop-improvement strategies (Pandey et al., 2016). In order to correctly identify and evaluate metabolites, accurate annotation reporting and of metabolomics data are critical. Alseekh et al., published standards (2021).recently for annotation and quantification of gas/liquid chromatography-mass spectrometry (GC/LC-MS) based metabolomics data reporting to maintain transparency and avoid misinterpretation of metabolite data.

Drought stress responses of spring wheat leaf tissue are revealed by metabolomics and proteomics, which provide a better framework for understanding the mechanisms that drive plant cell responses to drought stress and knowledge of chemicals that can be exploited in crop improvement programmes (Michaletti et al., 2018). Drought stress had a significantly higher negative impact on the distribution and accumulation of metabolites in JD8 than in JD17. according to a study conducted on shoots of two wheat genotypes (JD17- drought tolerant, JD8drought sensitive), and GC-MS was recommended as an effective approach for

understanding plant biochemistry under water stress (Guo et al., 2018).

Using GC/MS, researchers discovered that lysine, asparagine, methionine, serine, and glutamine are the major metabolites involved in drought resistance in wheat leaves (Yadav et al., 2019). Beside this GC-MS analysis of wheat roots and leaves under drought stress revealed that tryptophan and valine, as well as other metabolites such fumaric acid, malic acid, and citric acid, are the primary metabolites connected to drought tolerance (Kang et al., 2019). Guo et al., (2020) examined the metabolome of drought tolerant (HX10) and sensitive (YN211) wheat genotypes ultra-performance using liquid chromatography-mass spectrometry (UPLC-MS). In response to drought stress, there was a difference in the levels of 56 metabolites between these genotypes. Future QTL or Genome-Wide Association Studies (GWAS) research might leverage this metabolome data to discover locus (loci) or gene(s) linked with these metabolic traits (s), and therefore offer gene(s) or trait-specific marker(s) for crop improvement was further remarked.

2.5. Ionomics approach

Ionomics is a method for studying the molecular systems underlying critical mineral nutrient and trace element composition in biological systems through high-throughput elemental profiling (Michaletti et al., 2018). Inorganic components of cellular and organismal systems are also represented by this term (Michaletti et al., 2018). It has several uses in forward and reverse genetics, mutant screening, ion absorption, compartmentalization, transport, and exclusion processes, and therefore aids in the understanding of drought and other abiotic stress mechanisms in plants (Shelden and Roessner, 2013). Integration of ionomics with other omics, such as genomics or metabolomics, might help researchers find possible genes and networks that increase crop resilience to physiological and environmental stress (Colmsee et al., 2012; Satismruti et al., 2013; Singh et al., 2013; Wu et al., 2013; Huang and Salt, 2016; Guo et al., 2017).

Ionomics investigations in wheat under drought stress is yet to be published (Budak *et al.*, 2015). Ionomics is a relatively new functional genomics approach with a small number of research accessible, but spatial and technologically advanced ion profiling will be critical in the future to understand drought tolerance signalling networks (Budak *et al.*, 2015; Shah *et al.*, 2018).

2.6, Phenomics approach

Phenomics is described as the collecting of highdimensional phenotypic data on an organism-wide scale to characterise phenotypes (Houle et al., 2010). Plant phenomics is a fast-growing subject that includes anything from high-throughput field phenotyping to cellular imaging (Kaur et al., 2021). Field-based high-throughput phenotyping (HTP) has received greater attention in the previous decade, especially to forecast agronomic and physiological features (Crain et al., 2018). The new phenomics era has given scientists the tools they need to decode the data encoded in plant genomes (Finkel, 2009; Yang et al., 2020). The capacity to collect comprehensive sets of field data and the fast speed of plant phenotyping based on phenomics has enhanced the selection phenomena of possible elite advanced lines that perform well under stress situations (Montes et al., 2007).

The new phenomics era has given scientists the tools they need to decode the data encoded in plant genomes (Finkel, 2009; Yang et al., 2020). Handheld infrared thermometers were used to assess canopy temperature in a high-throughput manner to avoid plot-to-plot fluctuation by Deery et al., (2016) along with thermal imaging sensors in wheat for water and heat stress. The ability of several sensors to predict drought tolerance capacities of genotypes using water stress indices was tested by Becker et al., (2017) utilizing passive and active hyperspectral reflectance sensors in wheat under drought stress. The usage of Unmanned Aerial Systems (UAS)-based HTP as a technique for collecting high-spatiotemporal data is quickly expanding (Haghighattalab et al., 2016). Using an unmanned aerial vehicle (UAV) imagery system in 248 elite durum wheat

Condorelli et al.. (2018)measured leaf chlorophyll content, leaf rolling, dry biomass, Normalized Difference Vegetation Index (NDVI), and QTLs for NDVI under water deficit conditions. Bhandari et al., (2021) concluded that the combination of UAS-based remote sensing, agronomy, wheat breeding, and data analytics might lead to the development of digital tools that could be used to select genotypes for drought resistance and increase wheat breeding programmes' genetic gain.

3. Conclusion

Global climate change has resulted in the emergence of complex stress combinations and their effects on crop growth and production in modern agriculture. Climate change is a multifaceted issue with long-term consequences in the form of many abiotic pressures. Among them, water scarcity is receiving attention as abiotic stress because of its negative impact on plant growth and development, as well as a considerable loss in plant yield and biomass, resulting in worldwide food insecurity. Drought is one of the primary issues for attaining the potential vield since most of the worldwide wheat production area is in arid and semi-arid regions. It limits plant growth and development, which delays fruiting and grain filling, resulting in smaller and fewer wheat grains. Drought reduces wheat yields by altering the balance between the formation of ROS in plant cells, resulting in ROS overproduction and oxidative stress. Droughtinduced ROS damage a wide range of macromolecules, including proteins and nucleic acids, causing plant death. It has an overall effect plant production by interfering on with physiological, metabolic. and biochemical processes. Certain breeding approaches, molecular and genomics views, with a specific emphasis on the omics technology modification, i.e., metabolomics. proteomics. genomics. phenomics transcriptomics. glycomics, and approaches, are of considerable use in improving drought stress tolerance in plants. However, a combination of advanced technologies and technical improvements is required for a better

understanding of drought tolerance mechanisms and the development of drought tolerant wheat plants.

To fully comprehend complex abiotic stress tolerance features, more research is needed to combine data from omics investigations. As a result. investigations in genomics, transcriptomics, proteomics, metabolomics, and, particularly, phenomics, should focus more on the response to abiotic stressors.

Compliance with Ethical Standards

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