ADAPTIVE RESPONSES OF TWO DISTINCT WHEAT VARIETIES TO DROUGHT STRESS THROUGH MICROBIAL-AUGMENTED VERMICOMPOST

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Abstract

Wheat, a vital cereal crop, is predominantly grown in arid and semi-arid regions, making it susceptible to drought stress. The adverse effects of drought on wheat can potentially be mitigated through the application of cellulolytic microbes-enriched vermicompost. To explore this, two field studies were conducted at the Student Research Farm, Agronomy Department, University of Agriculture Faisalabad, during 2020-21 and 2021-22. The research examined the impact of cellulolytic microbesenriched vermicompost on the agronomic, physiological, and enzymatic antioxidant traits of wheat under varying soil moisture levels. The treatments included: (a) Three soil moisture levels: well-watered (D0, 70% field capacity), moderate drought (D1, 45% field capacity), and severe drought (D2, 30% field capacity). (b) Two wheat varieties: Galaxy-13 (drought-sensitive) and Faisalabad-08 (drought-tolerant). (c) Four microbial enriched vermicompost levels: VT0 (control, no vermicompost), VT1 (6 t/ha wheat straw vermicompost), VT2 (6 t/ha rice straw vermicompost), and VT3 (4 t/ha cow dung vermicompost). The results revealed that the highest crop growth and yield were achieved with 4 t/ha cow dung vermicompost (VT3), followed by rice straw vermicompost (VT2) and wheat straw vermicompost (VT1). The lowest performance was observed in the control (VT0) across both wheat cultivars. Faisalabad-08 consistently outperformed Galaxy-13 in growth and yield under both moderate and severe drought conditions. The findings further highlighted that drought stress significantly reduced agronomic, physiological, biochemical, and growth traits of wheat. However, the application of vermicompost enhanced these characteristics and improved yield, even under water-limited conditions.

Key words: Drought levels, Field experiment, Microbes, Straw, Vermicompost, Wheat cultivars.

Introduction

After corn and rice, wheat (Triticium aestivum L.) is amongst the most commonly produced staple crop in a variety of climates. Globally, China ranked first with a reported 134.30 million tons of wheat yield, followed by India, Russia, the United States, and France, with Pakistan ranked seventh among other developing nations (FAO, 2017). It covers an area of approximately 219.52 million hectares worldwide and touches production of almost 733.91 million tons (WAP, 2018). Wheat contributes almost 1.70% to the GDP of Pakistan and its value addition is 8.7% in agriculture sector of country and wheat production was projected approximately 27 million tons on the total cultivated area of 9.20 million hectares (Govt. Pakistan, 2019-20).

The biochemical, physiological, morphological, and yield attributes of plants are negatively influenced by many constraint factors, such as harsh climatic conditions, erratic weather, sowing time, salinity, heavy metals, pest attack and irrigation water availability. Drought is the major restriction factor of all these factors, which is usually correlated with plant yield reduction rather than other constraint factors (Anjum & Tanveer, 2016; Flexas et al., 2004).

Water deficit stress arises when the moisture in the plant is limited and thus the plant reabsorbs the water from the rhizosphere, thus reducing its natural functioning under the impact of water scarcity. Plants susceptible to extreme drought stress severely interrupt their several morpho-physiological functions, such as leaf area, photosynthetic content, plant biomass, plant growth

and development, yield and yield contributing parameters. In plants, water shortage greatly represses the mechanism of photosynthesis associated with glucose formation. Photochemistry and photosynthetic metabolism are impaired, and CO₂ accessibility is also decreased under restricted water regimes (Lawlor & Cornic, 2002; Akram, 2011; Anjum et al., 2017). In the field trial, the absorption of CO_2 by the plant facing water stress conditions normally decreases (Chaves et al., 2002; Hussain et al., 2018; Mitchell et al., 1998). Plants exposed to drought stress conditions considerably stimulate free radical mechanisms that impairs the plant's photosynthesis system, protein biosynthesis, and other metabolites. Plants have an innate mechanism to reduce drought stress injury by restricting their growth, such as shoot and leaves (Chaves, 1991). According to reports, different genotypes of wheat respond differently to water constraint (Tang et al., 2002; Khakwani et al., 2011).

The production of crop is exaggerated by water deficit stress by modifying many metabolic processes, including reduced rate of carbon absorption, enhanced oxidative damage, and decreased gaseous exchange of leaf (Hussain et al., 2018; Anjum et al., 2017). It also retards various enzymatic processes, ions uptake, and leaf development due to which crop productivity severely hampers (Sharma & Garg, 2018; Todaka et al., 2017). Water deficit stress is a serious issue as it takes hold of several physiological processes of crop growth and production. To eradicate such problem, however, it is highly recommended to incorporate organic amendments such as vermicompost and cultivation of drought-tolerant cultivars (Ji et al., 2010).

We have a lot of farm waste in the form of crop residues, tree residues and FYM, if this waste is managed and used wisely, it helps to enhance quality of soil and levels of nutrient, which helps us to fulfil the increasing population's demand of food. Farmers usually use agriculture wastes as fuel and heat due to a shortage of energy and coal, wasting an abundant supply of nutrient organic matter. Organic matter in the form of agricultural wastes is good source of food for insects and microorganism which can create a serious problem for sustainable agriculture. Vermicomposting techniques can effectively be used to manage agriculture wastes like wheat straw, cow dung, rice straw and paper waste *etc* (Jyotsana *et al.*, 2010). The process of vermicompost preparation is commonly described as the aerobic

wheat straw, cow dung, rice straw and paper waste etc (Jyotsana et al., 2010). The process of vermicompost preparation is commonly described as the aerobic degradation and modification of solid organic residues by manipulation of the biological activity of earthworm and other mesophilic microflora (Garg & Gupta, 2009). The vermicompost produced by earthworm activity enriches predominantly with certain immobilized microflora, growth-regulating hormones, vitamins, macro/ micronutrients, and chitinase, lipase, amylase, and protease are examples of degrading enzymes. With the introduction of another antagonistic microflora, these enzymes have already been secreted by earthworm can degrade the organic substrate (Barik et al., 2011).

Many studies have focused on Bacillus subtilis, which is found in the gut of the Eisenia fetida earthworm. All of these antagonistic bacteria secrete enzymes that degrade cellulolytic matter, such as endo-beta-1, 4-glucanase, cellulase, protease and amylase (Amita Paul et al., 2017; Farooq et al., 2009). Several researchers in cereal crops, wheat and maize have assessed the beneficial effects of vermicompost alone and combined with other organic fertilizers under non-drought, moderate drought, and severe drought stress situations. Previous research has revealed that the use of vermicompost can mitigate water deficit stress due to its high ventilation, porosity, strong water preservation, and drainage capability (Hosseinzadeh et al., 2016). Vermicompost's microorganisms improve the roots' ability to absorb water. Vermicompost also enhances soil water preservation and increases fertilizer solution (Hosseinzadeh et al., 2016). Vermicompost contains soluble sugars, sorbitol, betaine, amino acids, and further organic acids, in addition ions of phosphorus, nitrogen, calcium, zinc, boron, magnesium, sulphur, and iron (Hosseinzadeh et al., 2016; Aslam et al., 2022; Aslam et al., 2023). Vermicompost is like peat with a fine structure, strong aeration, porosity, microbial activity, drainage, and a high-water holding capability, as well as a high nutrient content that is ideal for improving plant and soil health (Pathma and Natarajan, 2012). The availability of enzymes and hormones is needed to improve plant health and eliminate pathogens. During the vermicomposting process these enzymes and hormones release from earth worm's gut (Gajalakshmi & Abbasi, 2004).

In our earlier research (Ahmad *et al.*, 2022; Ahmad *et al.*, 2022a; Ahmad *et al.*, 2024), we explored the effects of plant-based (wheat straw, rice straw) and animal-based (cow dung) vermicompost, enriched with cellulose-degrading bacteria, on the physiological and biochemical

traits of wheat seedlings under water deficit conditions in pot experiments. The primary objective of these experiments was to determine the optimal vermicompost application rate that maximized plant performance, which was subsequently tested in the field in this study. However, the impact of animal and plant based vermicompost, supplemented with cellulolytic microbes, on wheat growth and productivity under drought stress remains unclear. To fill this knowledge gap, this study sought to evaluate whether cellulolytic microbe-enriched vermicompost could alleviate the adverse effects of drought on wheat productivity under field conditions. We hypothesized that this treatment would enhance drought resistance and improve the physiological, biochemical, and yield-related attributes of wheat. Considering the aforementioned details the two-year trial was directed in field having following particular objectives. i). To assess the role of cellulolytic microbes enriched vermi-fertilizer (prepared from wheat straw, cow dung and rice straw) application on wheat for drought tolerance. ii). To estimate the outcome of vermicompost application on agronomic, physiological, and enzymatic antioxidant traits of wheat grown under drought stress conditions.

Materials and Method

The experiment was conducted at Agronomic Research Area, University of Agriculture, Faisalabad. An optimized amount of wheat straw, rice straw and cow dung vermicompost were applied under field conditions at the time of sowing using randomized complete block design with split-split plot arrangement and were replicated thrice. Seed bed well prepared according to the demand of crop. Wheat crop was sown@ 120 kg ha⁻¹ seed rate. Recommended fertilizers were applied @ 110: 100 kg ha⁻¹ NP according to soil analysis. All other management practices were kept uniform except the treatments under study. The net plot size was $4.5m \times 2.25m$. For vermicomposting the epigeic species of earth worms Eisenia fetida were used. The active strains of cellulose degrading microbes i.e., C-03, C-18 and C-21 were used for wheat straw, cow dung and rice straw vermicompost respectively. After completing all the practices treatments were applied in their respective plots. The factor wise experimental plan is as under: Factor A: Drought levels (main plot); $D_0 = 70\%$ of field capacity (no drought), $D_1 =$ 45% of field capacity (mild drought), $D_2=30\%$ of field capacity (severe drought). Factor B: Wheat cultivars (sub plot); V_1 = Faisalabad-08 (drought tolerant), V_2 = Galaxy-13 (drought sensitive). Factor C: Cellulolytic microbes and earth worms based vermicompost (sub-sub-plot); $VT_0 =$ Control, $VT_1 = 6$ t/ha wheat straw vermicompost enriched with cellulose degrading microbes, $VT_2 = 6$ t/ha rice straw vermicompost enriched with cellulose degrading microbes, $VT_3 = 4$ t/ha cow dung vermicompost enriched with cellulose degrading microbes.

Meteorological data of experimental site: The weather in respect of temperature (°C), relative humidity (%) and rainfall (mm) of the experimental site for the year 2020-21 and 2021-22 is given in the (Fig. 1).

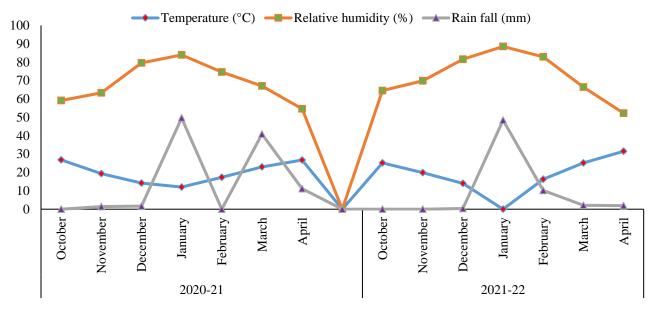


Fig. 1. Meteorological data of the experimental site for the growing seasons 2020-21 and 2021-22.

Data to be recorded: Data on various agronomic, physiological and enzymatic antioxidants traits of the plants grown in the field were recorded using standard procedures that are described as under.

Agronomic parameters

(i). Plant height (cm): Ten plants were picked from each sub-plot to calculate the plant height. Using a meter rod from the soil surface to the top of the plant, height was measured and then the average plant height was determined.

(ii). Number of total tillers m⁻²: The number of total tillers m⁻² was reported using a quadrate of 1m⁻². In each sub plot, drop it randomly. Tillers were counted and the average was calculated.

(iii). Spike length (cm): Ten plants were picked randomly from each sub-plot and the spike length was determined with the help of foot rod from the beginning of the spike to the top of it and average was calculated.

(iv). Number of spikelets per spike: Ten plants were picked from each subplot, and the number of spikelets from each spike were counted. The average number of spikelets per spike were determined.

(v). Number of grains per spike: Ten plants were picked randomly from each sub-plot. Their spikes were isolated and manually threshed. The number of grains were measured and then averaged from each spike.

(vi). 1000 grain weight (g): Grains were separated during the threshing and 1000 grains were counted. They were weighted on the electronic balance. Weight was measured in grams.

(vii). Biological yield (t ha⁻¹): A sample of plants from 1 m² of area was collected from each sub-plot. Weight on the

electronic balance and the conversion of biological yield into (t ha⁻¹) was calculated.

(viii). Grain yield (t ha⁻¹): From each sub plot, an area of one m² was selected and harvested. Threshed manually and took the weight of the grain by electronic balance, and the yield of the grain was converted to t ha⁻¹.

(ix). Straw weight (t ha⁻¹): Straw yield per plot of each treatment was calculated and expressed in t ha⁻¹.

(x). Harvest index (%): By dividing grain yield over biological yield, the harvest index was determined.

$$HI (\%) = \frac{Grain yield}{Biological yield} \times 100$$

Physiological parameters

(i). Leaf water potential (Ψ_w) [-MPa]: To measure the leaf water potential, at booting stage a fully emerged flag leaf was incised. With water potential apparatus (Chas W. Cook & Sons. Birmingham B 42, ITT England) leaf water potential was determined by adopting the procedure defined by Scholander *et al.*, (1964). With the cut surface popping out of the opening, a single leaf (flag leaf) was sealed in the pressure chamber. Pressure filled gas was exerted to the leaf till the xylem components visible on the cut section. To avoid any evaporative losses sampling was carried out between 6.00 A.M. to 8.00 A.M. The leaves were placed as early as possible in the pressure chamber and separate measurements were taken on three leaves from the control and stress treatments.

(ii). Leaf osmotic potential (Ψ s) [-MPa]: The leaf used in water potential measurement was frozen for seven days in the freezer below -20 °C. Then, following a frozen leaf material thawing, inserting by syringe. The extracted sap was used directly to assess the osmotic potential using an osmometer (Wescor 5500).

(iii). Turgor pressure (Ψ p) [MPa]: The difference of water potential and osmotic potential is measured as turgor pressure.

$$(\Psi p) = (\Psi w) - (\Psi s)$$

(iv). Canopy Temperature (°C): It indicates the direct measurement of energy emitted by wheat plants. IRIS, infrared temperature sensors was used to measure it. It gives data on plant metabolic activity *i.e.* plant water status and water use (Pettigrew, 2004; Singh *et al.*, 2018).

(v). Photosynthetic rate (An) $[\mu mol m^{-2} s^{-1}]$: IRGA, infrared gas analyzer was attached to plant leaves to measure photosynthetic rates (Singh *et al.*, 2018; Rosolem *et al.*, 2019). In each treatment, five measurements were collected from five different plants and then, their mean was taken.

(vi). Transpiration rate (*E*) [mmol m⁻² s⁻¹]: Transpiration rate (E) was measured on leaf attached to the plant by using IRGA, infrared gas analyzer (Singh *et al.*, 2018; Rosolem *et al.*, 2019). Five measurements were recorded from five different plants in each treatment and then their average was taken.

(vii). Stomatal Conductance (gs) [μ mol m⁻² s⁻¹]: The stomatal conductance was measured using an open system LCA-4ADC portable infrared gas analyzer (Analytical Development Company, Hoddeson, England). The measurement was carried out with the following adjustments: leaf surface area 6.25 cm², ambient CO₂ concentration (Cref) 371 μ mol mol⁻¹, temperature of leaf chamber (Tch) varying from 25-28°C, leaf chamber volume gas flow rate (v) 296 mL min⁻¹, leaf chamber molar gas flow rate (U) 400 μ mol s⁻¹, ambient pressure (P) 97.95 kPa, PAR (Qleaf) at leaf surface maximum up to 770 μ mol m⁻² s⁻¹. Measurement of stomatal conductance (gs) was made on the youngest fully emerged leaf (normally flag leaf from top) of each plant.

(viii). Sub-Stomatal CO₂ concentration (C_i) [µmol CO₂ mol air⁻¹]: The fully expanded leaves, using a CIRAS-2 (PP system®) portable gas exchange system (CO₂ and H₂O) connected to a gas exchange chamber (Parkinson Leaf Cuvette). The system measured CO₂ concentration (Parkinson *et al.*, 1980; Parkinson & Porometry, 1983).

Antioxidant enzymes activities

Enzyme extraction: Using a 50 mM cooled phosphate buffer (pH 7.8) for the extraction of antioxidant enzymes, fresh flag leaves (0.5 g) were ground in the pestle and mortar. Following filtration via cheese cloth, the homogenate was centrifuged at 15000 x g for 20 min at 4° C, and the supernatant was used for enzyme assays.

(i). Superoxide dismutase (SOD) [U mg⁻¹ protein]: SOD activity was assayed according to the Giannopolitis & Ries (1977) process; by finding photo chemical reduced of nitroblue tetrazolium (NBT) at 560 nm. SOD activity was calculated by adding 3 mL of total reaction 50 μ M NBT solution (NBT dissolved in ethanol), 13 μ M riboflavin solution, 13 mM methionine solution, 75 nM EDTA

solution, 50 Mm phosphate buffer (pH 7.8). The reaction solutions were put in a chamber under 30 W fluorescent lamps illumination. The reaction initiated when the fluorescent lamps were switched on and stopped 5 minutes later when they were switched off. The blue formazane formed by photoreduction of NBT was measured at 560 nm as an increase in absorbance. The non-leaf extract of reaction mixture was taken as control and placed under the light. The blank solution with the same exact reaction mixture (including enzyme extract) was, however, placed in the dark. A UV-visible spectrophotometer was used to read the absorption of the irradiated solution at 560 nm (IRMECO U2020). One unit of SOD was specified as the amount of enzyme needed to inhibit the rate of reduction of NBT at 560 nm by 50% compared to tubes that do not have the plant extract.

(ii). Peroxidase (POD) [U mg⁻¹ protein]: The Peroxidase behavior was determined by using the oxidation of guaiacol and characterized as 0.01 absorbance shift min⁻¹ mg⁻¹ protein. To form the reaction mixture in 100, 400, 500 μ L of enzyme extract, guaiacol (20mM), H₂O₂ respectively and 2 ml Phosphate (50 mM) were combined. The reaction mixture's absorbance at 470 nm was measured every 20 sec for up to 5 minutes. Activity of POD is express in U mg⁻¹ protein (Chance & Maehly, 1955).

(iii). Catalase (CAT) [U mg⁻¹ protein]: A UV-visible spectrophotometer was used to observe the CAT behavior detected by decomposition and change in absorption due to Hydrogen peroxide every 30 s for 5 min at 240 nm. The reaction mixture for CAT was 900 μ L H₂O₂ (5.9 mM) and 2 mL phosphate buffer, respectively (50 mM). The reaction was initiated by adding an extract of 100 μ L of enzyme to the reaction mixture. The Catalase activity was expressed as U mg⁻¹ protein (Chance & Maehly, 1955).

Statistical analysis

The recorded data regarding physiological, enzymatic antioxidants, yield and yield related agronomic traits of the experiment was statistically evaluated by applying the method of Fisher's analysis of variance (ANOVA). Tukey's HSD test was used ($p \le 0.05$) to compare significant treatments means using Statistic version 8.1 (Analytical Software ©, 1985-2005) according to Steel *et al.*, (1997) and graphical presentation was done by Excel 2016 and Sigma Plot 10.0.

Results

Agronomic parameters: Different moisture levels and cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost application and two contrasting wheat genotypes showed significant effect on plant height, number of total tillers, spike length and spikelets per spike during both years of study. The data regarding the abovementioned attributes affected by moderate drought (45% FC), severe drought (30% FC) and VT (Table 1, Fig. 2). Less plant height, number of total tillers, spike length and spikelets per spike were calculated in drought prone field as compared to well-watered field due to adverse effect of

drought. The highest value was recorded at optimized cellulolytic microbes enriched vermicompost levels cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) in moderate and severe drought fields in both cultivars. At well-watered conditions the plant height, number of total tillers, spike length and spikelets per spike were observed statistically at par with each other in both wheat cultivars. In moderate and severe drought stressed field, Faisalabad-08 produced more agronomic attributes than Galaxy-13. Similar response was observed in both wheat cultivars under different moisture levels and vermicompost application during both years 2020-21 and 2021-22 experiments).

The data referred to the number of grains per spike, 1000-grains weight, biological yield and grain yield are presented in (Table 2, Fig. 3). In drought stress significantly affected the above stated agronomic attributes in wheat plants grown at moderate drought stress (45% FC) and severe drought stress (30% FC) as compared to well-watered conditions (70% FC) in both the wheat cultivars. Cellulolytic microbes enriched vermicompost application significantly increased the number of grains per spike, 1000-grains weight, biological yield and grain yield of both cultivars. Maximum value was recorded at optimized cellulolytic microbes enriched vermicompost levels cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) and it was significantly higher from control at both under drought and well-watered conditions. Comparing the cultivars, Faisalabad-08 performed better than Galaxy-13 under moderate and severe drought conditions while under well-watered conditions both cultivars had statistically at par results during both year studies.

The data related the straw weight and harvest index affected by drought stress and vermicompost is shown in (Table 3, Fig. 4). Less straw weight and harvest index obtained in moderate and severe drought affected field as compared to well-watered conditions due to adverse effect of drought stress during 2020-21 and 2021-22 field trials. The highest value was recorded at optimized cellulolytic microbes enriched vermicompost levels, cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) in drought induced and well-watered fields in both cultivars. When we check at well-watered conditions plot, straw weight and harvest index were observed statistically at par with each other in both wheat cultivars during both year studies. Under drought prone field, Faisalabad-08 formed more straw weight and harvest index than Galaxy-13.

Physiological parameters: Table 4, Fig. 5 indicated that leaf water potential, osmotic potential and turgor potential were less at moderate and severe drought plots in comparison to well-watered plot in both wheat cultivars during 2020-21 and 2021-22. The minimum results were calculated under severe drought conditions. Vermicompost application ameliorated the unpleasant effect of drought and caused a significant increase in leaf water potential, osmotic potential and turgor potential. The maximum value was recorded where cellulolytic microbes enriched vermicompost levels, cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) were used under moderate drought, severe drought and well-

watered conditions. Among the cultivars, Faisalabad-08 had higher water potential, turgor potential and osmotic potential than Galaxy-13 observed under drought stress conditions, while, at well-watered conditions both cultivars were showed statistically at par during both year trials.

Canopy temperature in both wheat cultivars in the presence or absence of vermicompost (Table 4, Fig. 5) was also affected more at severe drought conditions. The canopy temperature significantly decreased with the vermicompost application under moderate and severe drought stress. The minimum temperature of canopy was achieved in response to cellulolytic microbes enriched cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) and maximum value was recorded in control (without vermicompost application) in Faisalabad-08 as well as in Galaxy-13. Comparing the cultivars, at moderate and severe drought stress, Faisalabad-08 showed lower canopy temperature than Galaxy-13 and proved to be drought tolerant variety.

data regarding physiological The attributes (photosynthetic rate, transpiration rate, stomatal conductance and sub-stomatal CO₂ concentration) was influenced by drought stress and vermicompost amendment are shown in (Table 5, Fig. 6). Drought stress significantly ($p \le 0.05$) reduced the rate of photosynthesis, transpiration, stomatal conductance and also sub-stomatal CO2 concentration in the moderate and severe drought tempted field when we compared with well-watered field. Vermi-fertilization augmented physiological attributes both in moderate drought, severe drought and well-watered field conditions. The maximum value was recorded where cellulolytic microbes enriched cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹) was used and minimum value was recorded where no vermi-fertilization was done. Comparing the cultivars, Faisalabad-08 (drought tolerant) had higher photosynthetic rate, transpiration rate, stomatal conductance and sub-stomatal CO2 concentration than Galaxy-13 (drought sensitive).

Enzymatic antioxidants: Interactive effect of drought and vermicompost on catalase activity, superoxide dismutase and the peroxidase was recorded statistically significant during both years 2020-21 and 2021-22. The data about enzymatic antioxidants were influenced by moderate drought, severe drought and vermicompost are shown in (Table 6, Fig. 7). According to data more catalase, superoxide dismutase and also peroxidase activity was observed in drought stressed field as we compared to wellwatered conditions. As drought stress enhanced the enzymatic antioxidants started to increase, but according to data record their increase was less to the requirement of the crop. The highest value was recorded when cellulolytic microbes enriched cow dung VT (4 t ha⁻¹) followed by rice straw VT (6 t ha⁻¹) and also wheat straw VT (6 t ha⁻¹) was used, and minimum value was recorded where no vermifertilization was applied in drought involved field in both cultivars. Under well-watered conditions the antioxidants enzymes were not influenced by vermicompost levels. In moderate and severe drought field, Faisalabad-08 generated more catalase, superoxide dismutase and peroxidase than Galaxy-13 whereas in well-watered field both cultivars produced similar antioxidant enzymes.

			2020-21	<u>1-21</u>			2021-22		
SOV	DF	Plant height	al	Spike length	Spikelet's per	Plant height	8	Spil	Spikelet's per
		(cm)	(m ⁻²)	(cm)	spike	(cm)	(m ⁻²)	(cm)	spike
Drought stress (DS)	0	1554.04**	55311.80**	102.12^{**}	305.54**	1050.29^{**}	24293.80**	81.70**	382.59**
Vermicompost (VT)	c	156.24**	3252.00**	20.64^{**}	131.94^{**}	213.75**	2344.20**	22.30^{**}	69.97**
Wheat (W)	1	120.12^{**}	806.70**	8.68**	20.05^{**}	100.35^{**}	654.00**	6.87^{**}	36.12**
$DS \times VT$	9	$0.26^{\rm ns}$	428.80**	0.48^{**}	0.26^{**}	0.70^{ns}	200.80^{**}	0.03 ^{ns}	0.67ns
$DS \times W$	2	44.04**	327.90**	2.85**	9.84**	15.85^{**}	232.60**	3.04**	4.62^{**}
$VT \times W$	С	0.01 ns	1.60 ^{ns}	0.07^{ns}	0.01 ^{ns}	0.09 ^{ns}	$0.70^{\rm ns}$	0.01 ^{ns}	0.19^{ns}
$DS \times VT \times W$	9	$0.04^{\rm ns}$	3.90^{ns}	$0.04^{\rm ns}$	$0.08^{\rm ns}$	$0.14^{\rm ns}$	2.10^{ns}	$0.04^{\rm ns}$	0.03^{ns}
Error	46	1.78	52.70	0.14	0.43	1.37	16.30	0.23	0.43
			2020-21				2021-22	2	
SOV	DF	Grains per	1000-gr	Biol	U	Grains per spike	1000-gr	Biological yield	Grain yield
		spike	(g)	$(t ha^{-1})$	$(t ha^{-1})$	•	(g)	$(t ha^{-1})$	(t ha ⁻¹)
Drought stress (DS)	0	762.09**	1013.54**	71.45**	20.01^{**}	1438.43**	927.04**	63.72**	15.48^{**}
Vermicompost (VT)	с	143.82**	133.87**	4.61^{**}	1.43**	148.38**	121.38**	3.54**	1.22^{**}
Wheat (W)	1	33.34**	23.35**	2.42**	0.58**	100.35^{**}	70.01**	2.16^{**}	0.59**
$DS \times VT$	9	0.02^{ns}	0.23^{ns}	$0.14^{\rm ns}$	0.05**	1.25^{ns}	0.30^{ns}	0.15^{ns}	0.08^{**}
$DS \times W$	0	27.26**	18.35**	1.21^{**}	0.18^{**}	22.93**	28.76**	0.68*	0.15**
$VT \times W$	c	0.08 ^{ns}	0.01 ^{ns}	0.02^{ns}	0.00^{ns}	0.53^{ns}	0.05^{ns}	0.00^{ns}	0.00^{ns}
$DS \times VT \times W$	9	0.22^{ns}	0.07^{ns}	0.01 ^{ns}	0.01 ^{ns}	$0.34^{\rm ns}$	0.13 ^{ns}	$0.00^{ m ns}$	0.00^{ns}
Error	46	0.72	0.89	0.10	0.00	1.05	1.15	0.13	0.00
Significant at 0.01 le	vel of sig	gnificance; *Signi	**Significant at 0.01 level of significance; *Significant at 0.05 level of significance, ns, non-significant	cance, ns, non-sigr	nificant				
ble 3. Mean sum	of squa	rres regarding t	Table 3. Mean sum of squares regarding the effect of soil applied c	ellulolytic micro	obes enriched w	heat straw, rice	cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost on the agronomic	ermicompost on	the agronom
	•)		of wheat cultivars under different drought levels.	s under differei	nt drought levels	,)
				7070 71			CC 1000		

ACC 3		2020-21	-21	2021-22	-22
200	JU -	Straw weight (t ha ⁻¹)	Harvest index (%)	Straw weight (t ha ⁻¹)	Harvest index (%)
Drought stress (DS)	2	16.07^{**}	231.03**	16.45^{**}	146.91**
Vermicompost (VT)	ŝ	0.90**	21.92**	0.60^{**}	17.67^{**}
Wheat (W)	1	0.62**	6.33**	0.49*	8.86**
$DS \times VT$	9	$0.05^{ m ns}$	0.82^{ns}	0.02^{ns}	0.81 ^{ns}
$DS \times W$	2	0.50**	1.01 ^{ns}	0.19 ^{ns}	2.49 ^{ns}
$VT \times W$	ŝ	0.02^{ns}	$0.17^{ m ns}$	0.00^{ns}	0.08^{ns}
$DS \times VT \times W$	9	0.00^{ns}	$0.44^{\rm ns}$	0.00^{ns}	0.00 ^{ns}
Error	46	0.07	0.44	0.10	0.54

6

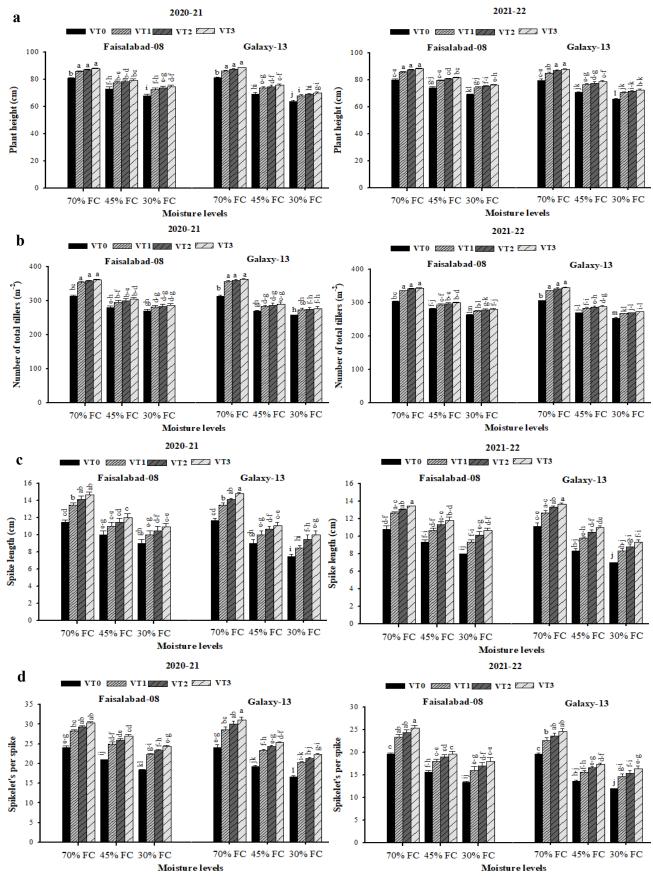


Fig. 2. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on the plant height (a), number of total tillers (b), spike length (c) and spikelets per spike (d) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT_0 = Control, $VT_1 = 6$ t/ha wheat straw vermicompost enriched with cellulose degrading microbes, $VT_2 = 6$ t/ha rice straw vermicompost enriched with cellulose degrading microbes.

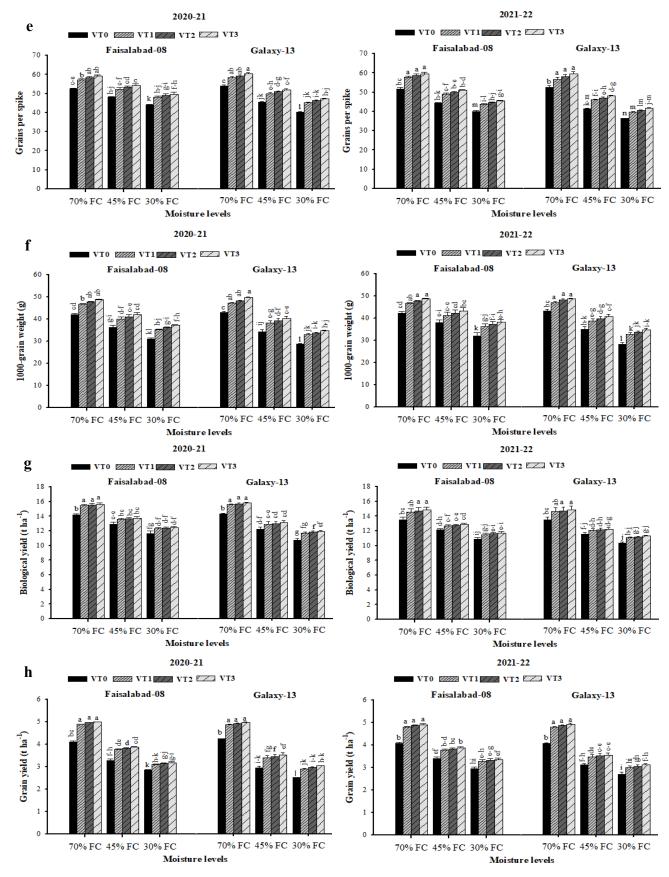


Fig. 3. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on the grains per spike (e), 1000-grain weight (f), biological yield (g) and grain yield (h) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT0 = Control, VT1 = 6 t/ha wheat straw vermicompost enriched with cellulose degrading microbes, VT2 = 6 t/ha rice straw vermicompost enriched with cellulose degrading microbes.

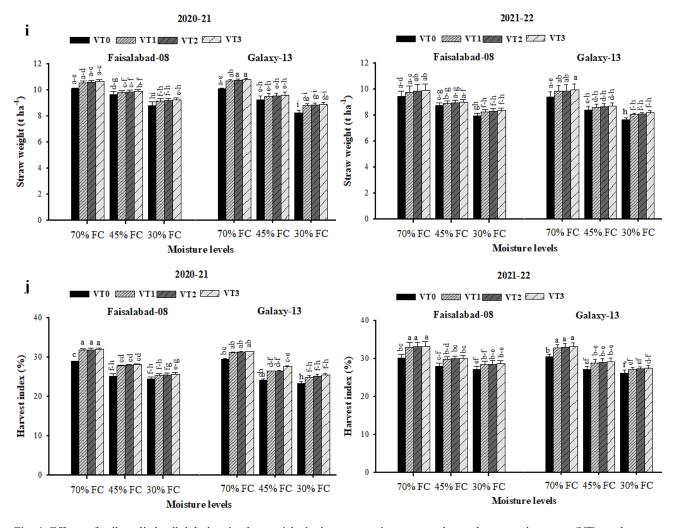


Fig. 4. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on the straw weight (i) and harvest index (j) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT0 = Control, VT1 = 6 t/ha wheat straw vermicompost enriched with cellulose degrading microbes, VT2 = 6 t/ha rice straw vermicompost enriched with cellulose degrading microbes, VT3 = 4 t/ha cow dung vermicompost enriched with cellulose degrading microbes.

Discussion

Reduced agricultural yields in semiarid and arid regions are mostly attributable to drought stress (Taheri-Asghari et al., 2009). Wheat is a prevalent crop in this region, although it has been badly affected by the drought. In fields treated to mild water deficit (45% FC) and severe water shortage (30% FC), yields of both cultivars and their components were found to be decreased, as shown by the present field experiment. Dryness and low tissue water potential may reduce crop development and yield by interfering with a number of biochemical and physiological processes, including turgor pressure, photosynthesis, enzymatic activities, and transpiration (Nayyar & Gupta, 2006; Patade et al., 2011; Ansari et al., 2017). Multiple researchers had demonstrated that vermicompost has aid plant growth regardless of abiotic (environmental) and biotic (plant disease and insect) obstacles (Jat et al., 2006; Hosseinzadeh et al., 2016). There is evidence that vermicompost may reduce drought stress by promoting plant growth in arid areas (Hafez et al., 2021; Aslam et al., 2023). Fertilizing fields with cellulolytic microbe-enriched

cow dung VT (4 t ha⁻¹), followed by rice straw VT (6 t ha⁻ ¹) and wheat straw VT (6 t ha⁻¹) significantly increased yield of grain and biological yield in both drought-stressed and adequately irrigated fields, demonstrating the vermicompost's beneficial effects. A multitude of characteristics, including height of plant, per unit area tillers, spikelet length, per spike spikelet, and weight of grain, affected yield. When these parameters are increased, the biological output may be substantially increased. Similarly to how the total quantity of grain harvested is the outcome of the cumulative behavior of the yield components, it is also a measure of the efficiency and effectiveness of a technological package. When vermicompost treatment was optimized in both droughtstricken regions and farms with sufficient irrigation, yield and contributing factors rose dramatically. Aslam et al., (2019) found that increasing the rate of VT feeding in the field increased the wheat plants' dry matter production, providing increased weight to the data provided here. The addition of vermicompost increased wheat output in two ways: the percentage of ripe grains and the quantity of grains collected per unit of land.

$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$				7	2020-21			2021-22	2		
** $0.31**$ $0.07**$ $0.00**$ $0.00**$ ** $0.01**$ $0.00**$ $0.00**$ $0.00**$ ** $0.01**$ $0.00**$ $0.00**$ $0.00**$ ** $0.01**$ $0.00**$ $0.00**$ $0.00**$ ** $0.01**$ $0.00**$ $0.00**$ $0.00**$ ** $0.00**$ $0.00**$ $0.00**$ $0.00**$ * $0.00**$ $0.00**$ $0.00**$ $0.00**$ * 2.73 2.73 2.03 0.00 * 2.73 2.73 2.03 0.00^{**} e 2.73 2.73 2.03 0.00^{**} e $3.70*$ $1.38*^{*1}$ $1.34*^{*}$ $1.34*^{*}$ $0.11*$ $0.20**$ $0.11*$ $0.20**$ 0.00^{**} $0.01*$ $0.01*$ $0.01*$ 0.00^{**} 0.00^{**} $0.01*$ $0.01*$ $0.01*$ 0.00^{**} 0.00^{**} $0.01*$ $0.01*$ $0.01*$ 0.00^{**} 0.00^{**} 0.00^{**} <th>SOV</th> <th>DF</th> <th>Leaf water pote (Yw) [-MPa</th> <th>antial Leaf osmotic po (</th> <th>itential Turgor potential</th> <th></th> <th>Leaf water potential (Yw) [-MPa]</th> <th>Leaf osmotic potentia (Ys) [-MPa]</th> <th>I Turgor potentis (\Pm p) [MPa]</th> <th></th>	SOV	DF	Leaf water pote (Yw) [-MPa	antial Leaf osmotic po (itential Turgor potential		Leaf water potential (Yw) [-MPa]	Leaf osmotic potentia (Ys) [-MPa]	I Turgor potentis (\Pm p) [MPa]		
** $0.01**$ $0.00**$ $0.00**$ $0.00**$ * $0.01**$ $0.00**$ $0.00**$ $0.00**$ * $0.00**$ $0.00**$ $0.00**$ $0.00**$ * $0.00**$ $0.00**$ $0.00**$ $0.00**$ * $0.00**$ $0.00**$ $0.00**$ $0.00**$ * $5.55*$ $7.87*$ $0.00*$ $0.00*$ * 2.73 2.03 0.00 $0.00*$ * 2.73 2.03 0.00 $0.00*$ theat straw, rice straw and cow dung vermicompost on the physiologic stratement devels. 2.73 0.00 theat straw, rice straw and cow dung vermicompost on the physiologic stratement devels. $2.34*$ 0.00 the different drought levels. $2.14*$ $1.349*$ $1.349*$ * $0.11*$ $0.01*$ $0.00*$ $0.00*$ * 0.01 $0.01*$ $0.00*$ $0.00*$ * $0.11*$ $0.11*$ $0.00*$ $0.00*$ * $0.01*$ $0.00*$ $0.00*$ $0.00*$	Drought stress (DS)	2	0.33**	-		49.29**	0.31**	0.07**	0.08**	36.84**	
* 0.01 ** 0.00 ** 0.00 ** * 0.00 ** 2.54 ** 0.00 ** * 0.00 ** 0.00 ** 0.00 ** * 5.55 ** 5.55 ** 0.00 ** * 5.55 ** 7.87 ** 0.00 ** * 5.55 ** 7.87 ** 0.00 ** * 0.00 ** 0.00 ** 0.00 ** * 0.00 ** 0.00 ** 0.00 ** * 0.00 ** 0.00 ** 0.00 ** * 0.11 ** 0.00 ** 0.00 ** * 0.11 ** 0.00 ** 0.00 ** * 0.11 ** 0.00 ** 0.00 ** 0.00 ** 0.01 ** 0.01 ** 0.00 ** 0.00 ** 0.01 ** 0.01 ** 0.00 ** 0.00 ** 0.01 ** 0.01 ** 0.00 ** 0.00 ** 0.00 ** 0.00 ** 0.00 ** 0.00 ** 0.01 ** 0.01 ** 0.00 ** 0.00 ** 0.01 ** 0.01 ** 0.00 ** * 0.11 ** 0.01 ** 0.00 **	Vermicompost (VT)		0.01^{**}	0.00^{**}	0.00**	14.64^{**}	0.01^{**}	0.00^{**}	0.00^{**}	16.44^{**}	
* 0.37m 2.54m 0.00** * 0.00** 3.70m 0.00m * 0.00** 0.00m 0.00m * 5.55m 7.87m 0.00m * 5.55m 7.87m 0.00m * 5.55m 7.87m 0.00m * 5.55m 7.87m 0.00m * 2.73 2.03 0.00m * 2.73 2.03 0.00m * 2.73 2.03 0.00m * 2.14m 1.34m 1.34m * 7.98** 0.14m 1.34m * 7.98** 0.00m 0.00m * 7.98** 0.00m 0.00m * 7.98** 0.00m 0.00m * 7.98** 0.00m 0.00m * 9.00m 0.00m 0.00m * 9.00m 0.00m 0.00m * 9.01** 0.00m 0.00m * 9.01** 0.00m 0.00m * 0.01**	Wheat (W)	-	0.02**	0.00**	0.00**	4.01**	0.01**	0.00**	0.00**	6.72**	
* 0.00^{**} 0.00^{**} 0.00^{**} s 5.55^{ns} 7.87^{ns} 0.00^{ns} 2.73 2.73 0.00^{ns} 0.00^{ns} end drought levels. 2.73 2.03 0.00^{ns} end drought levels. 2.73 2.03 0.00^{ns} end drought levels. 2.73 2.03 0.00^{ns} (40) (unol Co, m ² s ¹) (monol H ₂ O m ² s ¹) (monol m ² s ²) 1.34^{ns} (10) (monol Co, m ² s ¹) (monol m ² s ²) 1.34^{ns} 1.34^{ns} 1.34^{ns} (10) (monol Co, m ² s ¹) (monol H ₂ O m ² s ²) $(0.00^{ns}$ 0.00^{ns} 0.00^{ns} (11) (monol Co, m ² s ¹) (10) (monol m ² s ²) (10) m ² s ²) 1.34^{ns} 1.34^{ns} (11) (monol Co, m ² s ¹) (10) (monol H ₂ O m ² s ²) (10) m ² s ²) (10) (10) (11) (monol Co, m ² s ²) (10) m ² s ²) (10) (10) (10) (11) (monol Co, m ² s ²) (10) m ² s ²) (10) (10) (10) (11) (monol Ro) (10) m ² s ²) $(10$	$DS \times VT$	9	0.00 ^{ns}	4.16**	3 70 ^{ns}	0.19 ^{ns}	0.37 ^{ns}	2.54ns	0.00 ^{ns}	0.06 ^{ns}	
m 1.30 3.70 0.00 s 5.55 7.87m 0.00 feat straw, rice straw and cow dung vermicompost on the physiologic ent drought levels. 2.73 0.00 feat straw, rice straw and cow dung vermicompost on the physiologic ent drought levels. 2021-22 0.00 r^1_{1} Photosynthetic rate 3.0 Transpiration rate $r(E)$ Stomatal conductance (s) 1.82** r^1_{1} $7.98**$ $2.021-22$ Stomatal conductance (s) 1.82** r^1_{1} $7.98**$ 0.00^m 0.00^m 0.00^m r^1_{1} $7.98**$ 0.00^m 0.00^m 0.00^m r^1_{1} 0.01^m 0.01^m 0.00^m 0.00^m 0.00^m r^1_{1} 0.01^m 0.01^m 0.00^m 0.00^m 0.00^m r^1_{1} r^2_{1} r^2_{1} r^2_{1} r^2_{1} r^2_{1}	DS × W	о с	0.00**	5 01**	8 66**	4 76**	0.00**	0.00**	0.00**	0.00	
x 5.55 a 7.87 b 0.00 b theat straw, rice straw and cow dung vermicompost on the physiologic ent drought levels. 2.73 b 0.00 b ent drought levels. 2.02 b 2.02 b 0.00 b ent drought levels. 2.02 b 800 matal 0.00 b (CO ₂ Photosynthetic rate Transpiration rate (E) 800 matal 0.00 b (CO ₂ Photosynthetic rate Transpiration rate (E) 0.00 matal 9.134 b 0.00 b (CO ₂ Photosynthetic rate Transpiration rate (E) 800 matal 9.134 b 0.00 b σ_1 σ_1 σ_1 σ_2 σ_1 σ_2 <	$\mathbf{VT} < \mathbf{W}$	1 (1	0.00 n	1 Q 5 IIS	5 55 US	0.05ns	1 2 Qus	2 70ns	0 00 us	0.0505	
2.73 2.03 0.00 theat straw, rice straw and cow dung vermicompost on the physiologic ent drought levels. 2.03 0.00 ent drought levels. 2.03 0.00 $(C0_3)$ Photosynthetic rate Transpiration rate (E) stomatal $(C0_3)$ Photosynthetic rate Transpiration rate (E) stomatal (10) (µmol Co, m ⁻² s ¹) (µmol H ₂ O m ⁻² s ¹) (µmol m ⁻² s ¹) (10) (10) (µmol Co, m ⁻² s ¹) (10) (µmol M ⁻² s ¹) (10) (µmol M ⁻² s ¹) (10) (µmol M ⁻² s ¹) (11) (11) (µmol Co, m ⁻² s ¹) (11) (µmol M ⁻² s ¹) $(0.00^m M^{-2} s^{1})$ (11) (µmol M ⁻² s ¹) (11) $(0.01^m M^{-2} s^{-1})$ $(0.01^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ (10) (11) $(0.01^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ (11) $(0.01^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ (11) $(0.01^m M^{-2} s^{-1})$ $(0.01^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ $(0.00^m M^{-2} s^{-1})$ <th cold<="" td=""><td>$M \times VT \times W$</td><td>2</td><td>0.00</td><td>2 2 2 Ans</td><td>1 11ns</td><td>0.00 suco</td><td>5 55ns</td><td>0/.C</td><td>0.00</td><td>0.02 0.04ns</td></th>	<td>$M \times VT \times W$</td> <td>2</td> <td>0.00</td> <td>2 2 2 Ans</td> <td>1 11ns</td> <td>0.00 suco</td> <td>5 55ns</td> <td>0/.C</td> <td>0.00</td> <td>0.02 0.04ns</td>	$M \times VT \times W$	2	0.00	2 2 2 Ans	1 11ns	0.00 suco	5 55ns	0/.C	0.00	0.02 0.04ns
Alter straw, rice straw and cow dung vermicompost on the physiologic ent drought levels. 2021-22 CO5 Photosynthetic rate Transpiration rate (E) conductance (gs) (CO5 0.11* 0.029** 0.029** 0.029** (A10) (µmol Co5, m ⁻³ s ⁻¹) (µmol M ⁻³ s ⁻¹) (µmol M ⁻³ s ⁻¹) (µmol M ⁻³ s ⁻¹) (A10) (µmol Co5, m ⁻³ s ⁻¹) 0.01** 0.029** 0.029** 0.029** (A11) (µmol Co5, m ⁻³ s ⁻¹) 0.01** 0.00** 0.00** 0.00** 0.01** 0.01** 0.00** 0.00** 0.00** 0.00** 0.01* 0.01** 0.00 0.00** 0.00** 0.00** 0.00** 0.01** 0.01** 0.01** 0.00** 0.00** 0.00** 0.00** 1.10** 0.01** 0.01** 0.00** 0.00** 0.00** 0.00** 0.00** 0.00** <td< td=""><td>Error</td><td>46</td><td>0.00</td><td>2.20</td><td>1.13</td><td>0.25</td><td>2.73</td><td>2.03</td><td>0.00</td><td>0.24</td></td<>	Error	46	0.00	2.20	1.13	0.25	2.73	2.03	0.00	0.24	
centration and cow dung vermicompost on the physiologic inder different drought levels. 2021-22 Sub-stomatal CO ₂ Photosynthetic rate 2021-22 Sub-stomatal CO ₂ Photosynthetic rate Transpiration rate (E) conductance (gs) CO ₂ m0 and in (G) [µmol (An) (µmol CO ₂ m ² s ¹) (mmol H ₂ O m ² s ¹) conductance (gs) 20087.20** 7.60** 7.93** 0.50** 0.34*** 0.34*** 20087.20** 0.11* 0.11** 0.34*** 0.34*** 0.00** 210.90** 0.11* 0.01** 0.00** 0.00** 0.00** 0.00** 210.90** 0.11* 0.01** 0.00** 0.00** 0.00** 0.00** 1.70* 0.01** 0.01** 0.00** 0.00** 0.00** 0.00** 8.30 0.01** 0.01** 0.01** 0.00** 0.00*** 0.00*** 1.70** 0.01** 0.01** 0.01** 0.00*** 0.00*** 0.00**** 1.70** 0.01** 0.01** 0.01*** 2021-22 202	**Significant at 0.01	leve	l of significance;	*Significant at 0.05 le	vel of significance, ns, noi	n-significant					
Inder different drought levels. 2021-22 Sub-stomatal CO ₂ Photosynthetic rate CO ₂ mol air ⁻¹ Transpiration rate (E) Stomatal (mmol m ⁻² s ⁻¹) 20087.20** 49.71** 19.50** 13.49** 20087.20** 49.71** 19.50** 13.49** 20087.20** 9.71** 19.50** 13.49** 20087.20** 7.88** 0.11** 0.28** 20087.20** 0.11** 0.28** 0.34** 20087.20** 0.11** 0.22** 0.34** 470.20** 0.11** 0.22** 0.00* 97.60** 0.00* 0.00** 0.00** 0.00** 1.70* 0.01** 0.00** 0.00** 0.00** 1.70* 0.01** 0.00** 0.00** 0.00** 8.30 0.01** 0.00** 0.00** 0.00** 8.30 0.01** 0.01** 0.00** 0.00** 8.30 0.01** 0.03** 5021-22 0.00 177** 56.91** 0.92*** <t< td=""><td>TADIC D. MEAL SUIL</td><td>S 10 1</td><td>quares regarum</td><td>g une entect of son app</td><td>men cenniolytic micrope</td><td>s enfricheu wheat su</td><td>aw, rice suraw and co</td><td>w uung vermicompos</td><td>t on the physiolog</td><td>ucal parameters of</td></t<>	TADIC D. MEAL SUIL	S 10 1	quares regarum	g une entect of son app	men cenniolytic micrope	s enfricheu wheat su	aw, rice suraw and co	w uung vermicompos	t on the physiolog	ucal parameters of	
Sub-stomatal CO ₂ Photosynthetic rate CO ₂ mol air ⁻¹ Transpiration rate (E) Stomatal (muol m ⁻² s ¹) centration (C) [µmol $(4n)$ (µmol Co ₂ m ⁻³ s ¹) (mmol H ₂ O m ⁻³ s ¹) stomatal (mmol m ⁻² s ¹) 20087.20** 49.71** 19.50** 13.49** 20087.20** 49.71** 19.50** 13.49** 20087.20** 7.60** 0.11** 0.06** 20087.20** 0.11** 0.20*** 0.34** 20087.20** 0.11** 0.00** 0.00** 210.90** 0.11** 0.20** 0.34** 210.90** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** 1.70** 0.01** 0.00** 0.00** <					°	under different drou	ght levels.				
Sub-stomatal CO, contact (C) Photosynthetic rate (An) (µmol Co ₂ m ⁻³ s ¹) Transpiration rate (E) (mmol H ₂ O m ⁻³ s ¹) Stomatal (mmol rate (E) (µmol m ⁻³ s ¹) CO2 mol air ¹ (An) (µmol Co ₂ m ⁻³ s ¹) (mmol H ₂ O m ⁻³ s ⁻¹) (mmol rate (E) (µmol m ⁻³ s ⁻¹) Stomatal (µmol m ⁻³ s ⁻¹) CO2 mol air ¹ $A971^{+*}$ 19.50^{+*} 13.49^{+*} 13.49^{+*} 7005 0.11^{18} 0.11^{18} 0.02^{+*} 0.36^{+*} 7.60^{+*} 0.11^{18} 0.01^{18} 0.00^{-8} 0.00^{-8} 7.60^{+*} 0.11^{18} 0.00^{-8} 0.00^{-8} 0.00^{-8} 210.90^{+*} 0.00^{-8} 0.01^{18} 0.00^{-8} 0.00^{-8} 0.30^{18} 0.01^{18} 0.00^{-8} 0.00^{-8} 0.00^{-8} 1.70^{18} 0.01^{18} 0.00^{-8} 0.00^{-8} 0.00^{-8} 1.70^{18} 0.01^{18} 0.00^{-8} 0.00^{-8} 0.00^{-8} 1.70^{18} 0.01^{18} 0.01^{18} 0.00^{-8} 0.00^{-8} 1.00^{18} 0.01^{18} <t< td=""><td></td><td></td><td></td><td>-</td><td></td><td>100 - 1000 - 1000 - 10 - 1000-1000000</td><td></td><td>2021-22</td><td></td><td></td></t<>				-		100 - 1000 - 1000 - 10 - 1000-1000000		2021-22			
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	SOV	DF	Photosynthetic rate (An) (µmol Co ₂ m ⁻² s ⁻¹)	Transpiration rate (E) (mmol H ₂ O m ⁻² s ⁻¹)	Stomatal conductance (gs) (μmol m ⁻² s ⁻¹)	Sub-stomatal CO ₂ ncentration (C _i) [µmol CO ₂ mol air ⁻¹]		Transpiration rate (E) (mmol H ₂ O m ⁻² s ⁻¹)	Stomatal conductance (gs) (µmol m ⁻² s ⁻¹)	Sub-stomatal CO ₂ concentration (C _i) [µmol CO ₂ mol air ⁻¹]	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Drought stress (DS)	2	86.46**	25.80**	17.64	20087.20**	49.71**	19.50**	13.49**	185442.40**	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Vermicompost (VT)	С	11.21^{**}	4.39**	1.92	1481.60^{**}	7.98**	2.14**	1.82^{**}	1413.10^{**}	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Wheat (W)	-	3.73**	1.47**	1.17	470.20^{**}	3.42**	0.61^{**}	0.86^{**}	1241.70^{**}	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	$DS \times VT$	9	0.06^{ns}	0.04*	0.00	97.60**	0.11^{ns}	0.11^{*}	0.02**	57.20**	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	$DS \times W$	0	1.87^{**}	0.76^{**}	0.57	210.90^{**}	1.68^{**}	0.29^{**}	0.34^{**}	281.80^{**}	
$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	$VT \times W$	С	0.03^{ns}	0.00^{ns}	0.00	0.30^{ns}	0.00^{ns}	0.00^{ns}	0.00^{ns}	7.10^{ns}	
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	$DS \times VT \times W$	9	0.02^{ns}	0.00^{ns}	0.00	1.70 ^{ns}	0.01^{ns}	0.01 ^{ns}	0.00^{ns}	2.70^{ns}	
$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$	Error	46	0.10	0.01	0.00		0.05	0.03	0.00	4.20	
$ \begin{array}{ c c c c c c c c c c c c c c c c c c $	**Significant at 0.01 lev Table 6. Mean sur	wel of m of	significance; *Sign squares regar	nificant at 0.05 level of sig ding the effect of so	mificance, ns, non-significan il applied cellulolytic 1 iovidente of wheet cul	t microbes enriched tivers under diffe	wheat straw, rice s	traw and cow dung	vermicompost	on the en	
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$					2020-21				1-22		
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	SOV		DF	Catalase activity	Superoxide dismutas					Peroxidase	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$				(U mg ⁻¹ protein)	(U mg ⁻¹ protein)	(U mg ⁻¹ prote	10000			U mg ⁻¹ protein)	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Drought stress	; (DS		63.75**	58966.80**	2605.85**	8		•	2433.75**	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Vermicompost	TV)	_	1.59^{**}	454.50**	56.91**	0.92**	326.6(**(54.00**	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Wheat (W	(1	1.62^{**}	415.70^{**}	32.00**	1.00^{**}	648.0(**0	42.78**	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	DS×VT		9	0.37^{**}	94.10^{**}	15.64^{**}	0.17^{**}	66.20	**	10.85^{**}	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	DS×W		2	0.45**	101.00^{**}	13.54**	0.41^{**}	216.10	**0	8.53**	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$VT \times W$		S	0.01^{ns}	1.10^{ns}	0.11^{ns}	0.00^{ns}	0.30	us	0.24^{ns}	
46 0.01 32.20 0.62 0.01 3.00	DS×VT×W	N	9	0.00^{ns}	0.80^{ns}	0.15^{ns}	0.00^{ns}	0.80	us	0.09^{ns}	
	Error		46	0.01	32.20	0.62	0.01	3.0(0	0.47	

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 Error
 46
 0.01
 32.20
 0

 **Significant at 0.01 level of significance; *Significant at 0.05 level of significance, ns, non-significant

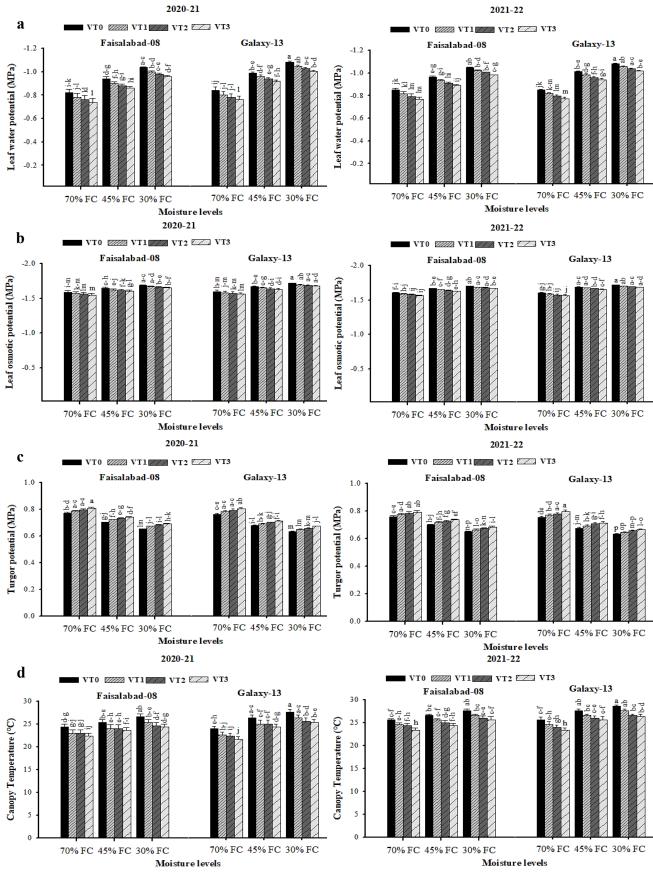


Fig. 5. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on the leaf water potential (a), osmotic potential (b), turgor potential (c) and canopy temperature (d) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT0 = Control, VT1 = 6 t/ha wheat straw vermicompost enriched with cellulose degrading microbes, VT2 = 6 t/ha rice straw vermicompost enriched with cellulose degrading microbes.

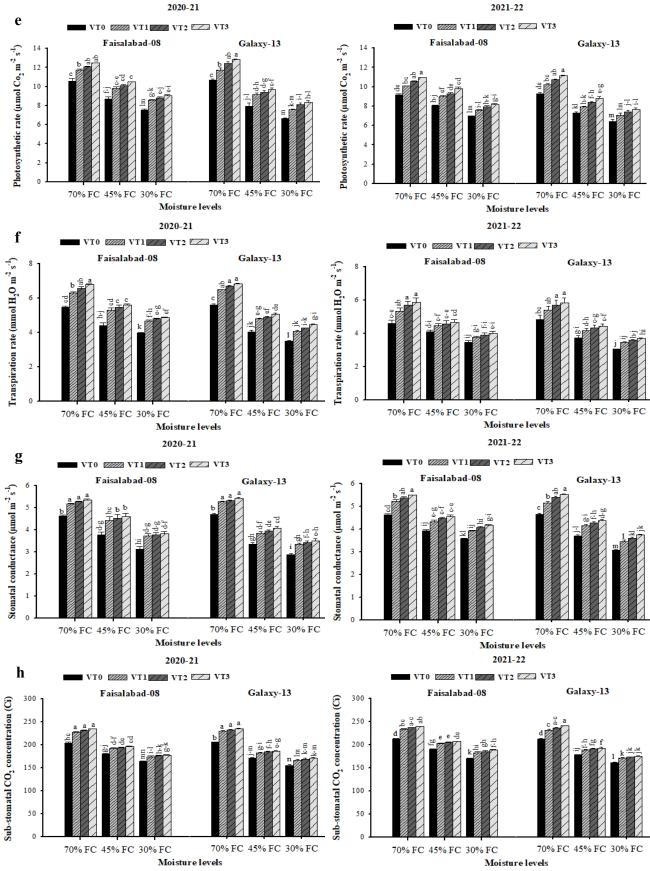


Fig. 6. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on photosynthetic rate (e), transpiration rate (f), stomatal conductance (g) and sub stomatal CO₂ concentration (h) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT0 = Control, VT1 = 6 t/ha wheat straw vermicompost enriched with cellulose degrading microbes, VT2 = 6 t/ha rice straw vermicompost enriched with cellulose degrading microbes.

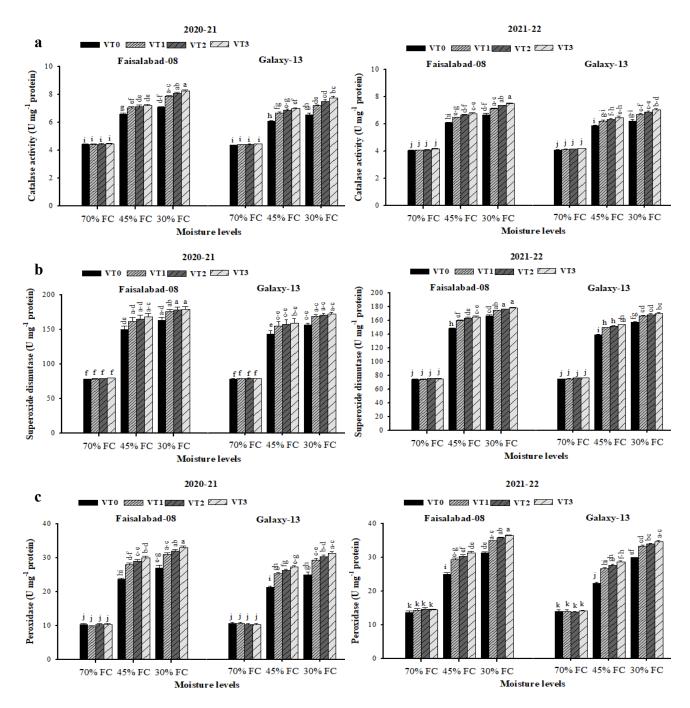


Fig. 7. Effects of soil applied cellulolytic microbes enriched wheat straw, rice straw and cow dung vermicompost (VT) on the catalase activity (a), superoxide dismutase (b) and peroxidase (c) of two wheat cultivars Faisalabad-08 and Galaxy-13 under different drought levels, well-watered condition (70% field capacity-FC), moderate drought (45% FC) and severe drought (30% FC). Bars with different small letters show significant differences at 5% probability level according to Tukey's HSD test. VT0 = Control, VT1 = 6 t/ha wheat straw vermicompost enriched with cellulose degrading microbes, VT2 = 6 t/ha rice straw vermicompost enriched with cellulose degrading microbes.

Since vermicompost treatment in wheat boosted relative growth rate, Hafez *et al.*, (2021) hypothesized that the constructive impacts of vermi-fertilization on yield of wheat may be attributable to the plants' higher hydration and photosynthetic rates. Observations indicate that increases in photosynthetic or net assimilation rates increased water efficiency. It was revealed that Faisalabad-08 (a droughtresistant cultivar) and Galaxy-13 (a drought-sensitive cultivar) had the highest grain yield, harvest index, height of plant, biological yield, tillers number, grains number, and spikelets spike⁻¹ in well-watered fields, moderate drought conditions, and severe drought conditions, respectively by addition of vermicompost. Vermicompost facilitates the uptake and transfer of essential minerals for plant development and energy generation (Sinclair & Vadez, 2002; Tara, 2003; Kmeťová & Kováčik, 2014).

Both wheat cultivars exhibited substantial increases in photosynthetic rate, osmotic potential, water potential of leaf, transpiration rate, leaf turgor potential, sub-stomatal CO₂ concentration and stomatal conductance after treatment of vermicompost in 2020-21 and 2021-22. Galaxy-13 has shown less dramatic physiological responses to mild and

severe drought stress than Faisalabad-08. In reaction to water stress, plants shut their stomata, lowering transpiration, cellular CO₂ concentrations, and net photosynthesis, according to a considerable body of research (Flexas & Medrano 2008; Mssacci et al., 2008). The reduction in photosynthesis seen under situations of water stress has been connected to the inhibition of a variety of metabolic processes, including ATP production and Rubisco activity (Mssacci et al., 2008). In dry environments, two kinds of components impede photosynthesis. The first group consists of causes of stomatal closure, which lowers CO₂ absorption from leaves and subsequent transport to chloroplasts, hence decreasing photosynthesis (Pagter et al., 2005). Non-stomatal limiting variables include the suppression of synthesis ribulose-1, 5-bisphosphate, decline in transfer photosynthetic electron to PSII, and the decrease in chlorophyll content (Pagter et al., 2005). The need for NADPC to accept electrons decreases under situations of water scarcity, absence of oxidation, and use of NADPH molecules from the light reaction in photosynthesis. Through the ETC pathway, molecules of oxygen form free radicals such as superoxide, hydroxyl and hydrogen peroxide (Sairam & Saxena, 2000). ROS cause oxidation of liposomal lipids, modification of structural proteins, oxidation and inactivation of sulfhydryl groups (-SH), bleaching of pigments like chlorophyll and carotenoids, and strikes on photosystems (Sairam & Saxena, 2000). These factors may explain why photosynthesis is slowed in arid settings. Iron is added to soil through vermicompost (Davatgar et al., 2009). Because negatively charged groups like phenolic acid and carboxylic acid are present, vermicompost's humic compounds have a high potential for absorbing metals (Matos & Arruda, 2003). The ironcontaining prosthetic group facilitates the activation of antioxidant enzymes, such as catalase, superoxide dismutase and peroxidase, which may be important for removing ROS in plants, according to Flexas & Medrano (2008). Vermicompost's high porosity, high ventilation capacity, efficient drainage, and water storage all lead to decreased stomatal closure and an increase in photosynthesis-required CO₂ (Arancon *et al.*, 2004; Ahmad *et al.*, 2025).

Inadequate water supply may result in oxidative damage. Vermicompost treatments and their combinations exhibited significant ($p \le 0.05$) changes in the levels of antioxidant enzymes, including peroxidase (POD), catalase (CAT), and superoxide dismutase (SOD). Vermicompost increased the POD, SOD, and CAT proportions of leaves under water deficiency stress comparing the control (70% FC). The leading SOD enzyme activity was found in plants having treatment of cow dung VT (4 t ha⁻¹) containing cellulolytic bacteria, followed by rice straw VT (6 t ha⁻¹) and wheat straw VT (6 t ha⁻¹). Utilizing vermicompost may improve a plant's ability to absorb cations like potassium and calcium. Andersen et al. (1992) revealed that K protects plants by reducing transpiration, while Ca acts as an enzyme activator. Boosted activities of SOD, POD and CAT membrane integrity and lowered enhanced lipid peroxidation by reducing ROS levels. As a result of drought stress, rice plants treated with vermicompost showed increased levels of the three antioxidant enzymes, namely catalase (CAT), peroxide oxidase (POD), and superoxide dismutase (SOD) (Garcia et al., 2014; Wang et al., 2017).

A mycorrhizal relationship with plant roots, an increase in soil-dwelling bacteria as nitrogen stabilizers, fungal spores, and actinomycetes, and a host of other helpful microbes are further advantages of vermicompost (Kale et al., 1992). For instance, mycorrhizal fungi may store carbohydrates from a plant's roots and use them later in their metabolism. Sugars may be provided as appropriate osmolytes to manage osmotic pressure in the roots and increase water stress resistance (Huerta et al., 2010). Around a plant, the incorporating of natural material in the form of vermicompost increases microbial activity and the soil's capacity to create CO2 (Marinari et al., 2000). According to the findings of this field investigation, vermicompost may help wheat withstand to drought. Vermicompost treated with cellulolytic bacteria significantly improved enzymatic antioxidants, physiological features, yield contributing factors, and yield when applied to both wheat cultivars under wellwatered and dry field situations.

Conclusion

Topsoil used cellulolytic microbes augmented vermicompost made from wheat straw, rice straw, and cow dung improved growth, stabilized nutrients, increased the actions of SOD, POD, and CAT contents, and lessened the problems caused by moisture deficits at yield and yieldrelated traits of wheat. Ultimately, during a drought, these changes increased wheat yield. The most widely accepted, efficient, cost-effective, and environment friendly treatments are cellulolytic microbe-enriched cow dung VT (4 t ha⁻¹), rice straw VT (6 t ha⁻¹), and wheat straw VT (6 t ha⁻¹). This is because it has shown stimulatory effects in yield enhancement of both wheat cultivars (Faisalabad-08 and Galaxy-13) under drought. It is advised that farmers incorporate vermicompost into their modern crop production techniques, particularly in drought-prone areas, as it can serve as a valuable nutrient source for all field crops in both normal and water-scarce conditions. Vermicompost is made from wheat straw, rice straw, and cow dung by earth worms and cellulolytic microbes. Due to its high nutrient content, vermicompost increases the availability of both macro- and micronutrients and serves as a backup source of nutrients for biofortification.

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