

Original Research Article

Impact of biofertilizer combinations on quality, yield, and economic returns of strawberry in central region of Punjab

ABSTRACT

~~Strawberries~~ Strawberry are is a globally cherished fruit, celebrated for their sweet flavor, vibrant color, and rich nutritional profile. The use of biofertilizers in strawberry cultivation is gaining importance in India due to their eco-friendly and sustainable nature. Biofertilizers enhance soil fertility by promoting beneficial microbial activity, which improves nutrient availability and uptake by strawberry plants. This leads to healthier crops with better yields and quality, while also reducing using artificial by reduced application of inorganic fertilisers, thus benefiting It ultimately benefits both the environment and farmers' long-term productivity. The current inquiry was conducted at experimental farm, Mata Gujri College, Shri Fatehgarh Sahib, Kharora (Punjab), India during 2023-24. In this research, there are eight bio fertilizer treatments combination and three replications were laid out in RBD. Among them The experimental results revealed that T₈ (RDF + Azotobacter @ 2g/plant + PSB @ 2g/plant + Azospirillum @ 2g/plant) significantly increased the plant height (16.82 cm), plant spread (32.78 cm), berry-fruit length (45.88 mm), berry-breadth (38.96 mm), berry weight (12.76 g), TSS (9.42 °Brix), anthocyanin content (26.14 mg/100g) and yield attributes i.e., yield plant⁻¹ (238.88 g) and yield hectare⁻¹ (13.14 t/ha). It also ~~concluded~~ found that the plants treated with T₈ (RDF + Azotobacter @ 2g/plant + PSB @ 2g/plant + Azospirillum @ 2g/plant) gives us had maximal-maximum leaf nutritional content, namely, - nitrogen (3.09 %), phosphorus (1.37-%), potassium (3.39-%), calcium (2.37-%) and magnesium (0.39-%) and also gives-gave highest net return i.e., Rs 16,61,790 Rs/ha and-with highest B:C ratio i.e.-of 3.77, - as economics point of view Hence, it can be concluded that the treatment T₈ found-to-be-more is most suitable for improving the quality, yield and economic return of strawberry in central region of Punjab.

Keywords: leaf nutrient, fruit quality, net returns, nutrient, microbial activity

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Strawberry (*Fragaria × ananassa* Duch.) is a perennial herbaceous plant ~~in of the~~ Rosaceae family ~~that which~~ is octaploid by nature, possessing chromosomal number $2n=8x=56$ (Hossain *et al.*, 2019). *F. chiloensis* Duch. and *F. virginiana* Duch., two native American species, are hybridised to create ~~it the cultivated form of strawberry.~~ ~~and b~~ Botanically it is an aggregate fruit which is highly perishable in nature. ~~There are about sixteen wild strawberry species worldwide, but only four have been identified in India i.e., F. Vesca, F. Chiloensis, F. Daltoniana and F. Nilgerrensis.~~ Strawberry ~~It~~ is one of India's most prominent tropical fruits that may reach heights over 12,000 feet above mean sea level in both arid and wet climate regions. Fruits, ~~also is~~ a great source of endogenous metabolites, phenols, carotenoids, vitamins, and flavonoids, among other natural antioxidants. (Jideani *et al.*, 2021). In strawberries, indiscriminate and prolonged use of inorganic chemical fertilisers and pesticides raises production costs while contaminating the environment, water, and soil by volatilisation, runoff, leaching, and denitrification. It negatively affects the fertility, vegetation, and fauna of the soil. Beneficial microorganisms found in bio-fertilizers includes *Azospirillum*, *Azotobactor*, *Pseudomonas*, *Bacillus*, *Mycorrhizae fungi*, *Aspergillus*, and *Penicillium*. ~~ean~~ ~~These microorganisms play vital role to~~ boost nutrient availability, absorption, and biological activity in the rhizosphere to affect plant growth and health. According to (Reddy *et al.*, (2020), fixing nitrogen from the atmosphere, solubilising phosphorus, and mobilising iron, zinc, and phosphorus to different degrees through the synthesis of secondary metabolites, phytohormones, and antibiotics that help host plants withstand abiotic stress conditions and disease. Bio-fertilizers are living microorganisms (useful bacteria and fungi) which, by nitrogen-fixing, significantly contribute to enhancing soil fertility, solubilise insoluble soil phosphates, potassium removable and improving plant production (Kumar *et al.*, 2019a; Nosheen *et al.*, 2021). They contribute significantly to crop productivity and are environmentally beneficial. According to Mishra and Barolia (2020), bio-fertilizers or microbial inoculants, are carrier-based preparations contain advantageous microbes that increase biological activity in the rhizosphere with the goal of enhancing soil fertility and facilitating plant growth. In other words, biofertilizers depend on renewable energy sources, which are more eco-friendly than ~~commercial inorganic~~ fertilizers (Kaur and Purewal, 2019).

MATERIALS AND METHODS

The current ~~inquiry investigation~~ was executed at experimental farm, Mata Gujri College, Shri Fatehgarh Sahib, Kharora (Punjab), India during 2023-24. This study examined

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the influence of several bio-fertilizers viz., T₁ (Control), T₂ (RDF + *Azotobacter* @ 2g/plant), T₃ (RDF + *PSB* @ 2g/plant), T₄ (RDF + *Azospirillum* @ 2g/plant), T₅ (RDF + *Azotobacter* @ 2g/plant + *PSB* @ 2g/plant), T₆ (RDF + *Azotobacter* @ 2g/plant + *Azospirillum* @ 2g/plant), T₇ (RDF + *PSB* @ 2g/plant + *Azospirillum* @ 2g/plant) and T₈ (RDF + *Azotobacter* @ 2g/plant + *PSB* @ 2g/plant + *Azospirillum* @ 2g/plant). The experiment was divided into 24 sub-plots with dimensions of 15 m × 1 m. Every plot had three replications and eight treatments set up in an under completely Randomized Block Design (CRBD). The significance of variation among the treatments was determined using the ANOVA and critical difference (C-D) tests was tested at the 5% probability level of significance.

VEGETATIVE GROWTH ATTRIBUTES OF PLANT

Plant height: A metre scale was used to measure the plant's elevation from the primary leaf apex to the crown and the findings stated in terms of centimetres (cm). The final number was determined based on the average plant height of five plants.

Plant spread: Using a metre scale, the plant spread was measured in centimetres (cm) from East-east to West-west and North-north to South-south direction. The average of five plants in each direction was then computed to represent the mean value.

Leaf area: A leaf area meter was used to measure five leaves from each plant. To calculate the overall leaf area of a single plant and the outcome should be expressed in cm and calculated by multiplying the average area of each leaf by the number of leaves.

FLOWERING CHARACTERS OF PLANT

Days taken to first flower: The interval (40%) between the opening date of first flower opening date and the planting date has been noted in order to calculate the number of days required to generate the first flower.

Number of flowers per plant: The overall quantity of flowers was noted at 10-day intervals, and the average quantity of flowers per plant was calculated to ascertain the quantity of flowers per plant.

FRUIT QUALITY PARAMETER OF PLANT ATTRIBUTES

Berry-Fruit length: A digital vernier calliper was used to measure the length of the fruit, and the mean of the observations was computed and expressed in millimetres.

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Berry-Fruit breadth: The fruit diameter was measured using Digital Vernier Callipers on the same fruits used to measure length, and the mean was given in millimetres.

Berry-Fruit weight: After weighing the ten fruits on the above list, the total weight of the fruits was calculated and expressed in grammes.

Total soluble solids: The strained juice of the fruits was analysed for TSS using a 'Erma-hand refractometer' (0 to 32 °Brix). The refractometer was calibrated with distilled water prior to use and to measure the TSS, a few juice-drops of juice were put on the prism. A temperature correction was performed when it was above or below 20°C (AOAC, 2002). To accurately estimate TSS, an average of three readings per treatment were recorded and expressed as °Brix.

Acidity (Titrable acidity): The level of acidity was evaluated by diluting a known weight of dry material/fresh fruit (which was weighted and pulverised thoroughly with a pestle and mortar) and titrating it against standard 0.1 NaOH, using phenolphthalein as an indicator. Pale pink appearance of the solution indicates the final titrate value. The acidity is determined using the following formula and represented in percentages.

$$\text{Acidity (\%)} = \frac{1 \times \text{Eq. Wt of acid} \times \text{Normality of NaOH} \times \text{Titre}}{10 \times \text{Weight of sample}} \times 100$$

Anthocyanin content: Harborne's (1973) approach was used to determine the quality of anthocyanin pigments in berries present in the fruit. One gram of berry-fruit pulp was immersed in a known volume of methanol containing 1 percent hydrochloric acid. The goods solution was were permitted to remain overnight— at Subsub-zero temperatures in a deep freezer. were used, and Next day, using a Spectronic-20 colorimeter, the absorbance of the resultant red solution was determined at 530 nm. The intensity of color was measured and represented in absorbance units per gram of fresh fruit.

Leaf analysis

Collection and preparation of leaf samples

A composite sample of hundred leaves for each orchard/cropping system were taken during the month of January from the plant with a total of 20 leaves from each treatment. From the plants that were marked, leaves, including petioles from the mid-terminal shoots of the growth for this season, were gathered. The leaf samples were carried directly to the

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laboratory and properly cleaned with tap water before being treated with 0.1N HCl and distilled water to eliminate dust particles, as described by Chapman (1964). For surface drying, the washed leaf samples were placed on sheets of filter paper. After that, they were put in paper bags and left for 48 hours at $65 \pm 5^\circ\text{C}$ to dry in a hot air oven. The dehydrated samples were finely pulverized and homogeneous. The ground samples were subsequently stored in butter paper bags and kept in cold and shady ~~spot-place~~ to estimate nutritional components. ~~Sampling and the preparation for leaf chemical analysis was carried out using standard procedure.~~

Digestion of leaf samples

Leaf samples (1_g) ~~were-was~~ processed in intense H_2SO_4 with a mixture of K_2SO_4 (400 parts), CuSO_4 (20_parts), HgSO_4 (3_parts), and Se (1_part) for total nitrogen measurement. The samples were kept boiling until they turned a pale blue tint. Following cooling, the samples were diluted with 100 millilitres of distilled water. The appearance of a slight crimson tint marked the conclusion. To compute P, K, Ca, and Mg, 0.5_g of leaf sample was digested in a di-acid mixture of HNO_3 and HClO_4 in a 4:1 ratio, adopting all appropriate precautions as recommended by Piper (1966).

Nitrogen: The total N was calculated using Micro-Kjeldhal's method (Jackson, 1967), and the outcomes were represented as percent nitrogen on a dry weight basis.

Phosphorus: The Vanado-molybdate Phosphoric Yellow Colour Method (Jackson, 1967) was used to measure total phosphorus. ~~Pipette~~ 10_ml of digested aliquot was pipetted into a 25_ml volumetric flask, then ~~add~~ 5_ml of vanado molybdate reagent was added in to it. After adding ~~purified-distilled~~ water to dilute the solution to 25 ml, it was left to develop colour for 30 minutes. Following colour development, the concentration of phosphorus in the solution was measured using ~~a~~ 470 nm wavelength in Nukes UV-VIS spectrophotometer, with a blank run in parallel to adjust zero absorption. The leaf P was represented in percent on a dry weight basis.

Potassium, Calcium, Magnesium: Total potassium content was assessed utilising the Agilent 5110 ICP-OES as suggested by Jackson (1973) and the results were expressed in per cent (%). Using an Analyst 400 Atomic Absorption Spectrometer, total calcium and magnesium were measured. ~~The outcomes were displayed as dry weight percentages.~~

Results and discussion

Vegetative growth attributes

The data collected for the various growth attributes are displayed in Table.1, which stated that the highest plant height (16.82 cm), plant spread (32.78 cm), number of leaves plant⁻¹ (20.00), leaf area (121.53 cm²) were ~~increased significantly and~~ recorded ~~highest within~~ T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant). However, the application of T₁ (control) resulted the ~~smallest minimum~~ plant height (10.66 cm), plant spread (22.90 cm), number of leaves plant⁻¹ (11.87), and leaf area (101.02 cm²). According to Beer *et al.* (2017), The capacity of a biofertilizer to produce antibacterial and antifungal compounds is linked to its growth-promoting properties, in addition to its N-fixing capabilities, growth regulators, and siderophores, resulting in increased vegetative development of strawberries. The use of biofertilizers such as PSB and *Azospirillum* improves biological N₂ fixation and phosphorus availability, both are essential for vigorous vegetative growth (Kumar *et al.*, 2019b; Kumar *et al.*, 2015). These findings are consistent with those of Gupta and Tripathi (2012), Lata *et al.* (2013), and Negi *et al.* (2021), Kumar *et al.* (2021) and Kumar *et al.* (2023).

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Physico-chemical attributes of ~~berry~~the fruit

~~According to the present analysis, there was a~~ considerable rise in fruit length, ~~berry~~ width, and ~~berry~~ weight ~~was observed under this experiment~~. The highest ~~berry~~ fruit length (45.88 mm), ~~berry~~ breadth (38.96 mm), ~~berry~~ weight (12.76 g) ~~along with the quality attributes viz.;~~ total soluble solid (9.42 °Brix), total sugar (7.58 %), reducing sugar (4.38 %) and anthocyanin content of ~~the fruit~~berries (26.14 mg/100g) were recorded in treatment T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant). However, ~~lowest berry minimum fruit~~ length (34.31 mm), ~~berry~~ breadth (31.43 mm), ~~berry~~ weight (8.38 g), ~~Total total~~ soluble solid (6.90 °Brix), total sugar (5.68 %), reducing sugar (3.73 %) and anthocyanin content of ~~berries the fruit~~ (16.39 mg/100g) ~~was were resulted estimated~~ in treatment T₁ (control). The appropriate supply of macro and micronutrients may have contributed to the improved fruit physical ~~as well as biochemical~~ quality., ~~as well as~~ Further, hormones that promote growth ~~that are~~ ~~also~~ generated by different biofertilizers used in varied combinations with RDF treatments. This may be related to more fruit filling from a more balanced food intake, which may have enhanced the plant's metabolic processes and increased the synthesis of proteins and carbohydrates (Kumar *et al.*, 2024; Nazir *et al.*, 2015). The release of growth-promoting compounds, the increased availability of P and the

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improved ability of microbial inoculants to fix atmospheric N that quicken physiological processes like carbohydrate synthesis could all be responsible for greater ascorbic acid content [in the harvested fruits](#) (Beer *et al.*, 2017). Rise in anthocyanin pigment following the administration of nitrogen and *Azotobacter* is consistent with the results of [\(Rana *et al.*, 2003\)](#) [Kumar *et al.* \(2019a\)](#), who noticed strawberry's increased red pigment [ev. Chandler](#) following the simultaneous treatment of nitrogen and bio-fertilizers. The given results are consistent comparing said by [Kumar *et al.* \(2023\)](#), Singh *et al.* (2018) and Jain *et al.* (2017).

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Flowering characteristics

The outcome demonstrated from Table 2 showed the plants need a minimum of 58.99 days to generate their first blooming which were treated with T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant). However, maximum number of days taken to [produce induce](#) first flowering (70.55days) was noted in T₁ (control). The highest number of flowers plant⁻¹ (24.13) was recorded in T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant) and lowest number of flowers plant⁻¹ (14.60) were recorded in T₁ (control). According to [\(Bashandy *et al.*, 2019\)](#), *Azotobacter* and *Azospirillum* reduced the number of days for the first flower to bloom compared to the control in both years. It's feasible that [using application of](#) bio-fertilizers makes nitrogen more available to plants, forcing them to transition from the vegetative to reproductive phase earlier. Strawberry plants [infected treated](#) with biofertilizers took the fewest days to produce their first blossom, according to research by Tripathi *et al.* (2016). Similar outcomes have [also](#) been documented by Singh *et al.* (2015), Kushwah *et al.* (2018), [Kumar *et al.* \(2019a\)](#) and Jaiswal *et al.* (2023).

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Yield characteristics

Table 3 represented the combination of biofertilizers [showed had](#) a notable variance in the quantity of fruits, yield plant⁻¹, and yield hectare⁻¹, where highest number of fruits plant⁻¹ (18.73), yield plant⁻¹ (238.88 g) and yield hectare⁻¹ (13.14 t/ha) was found in the plants that received [application treatment of T₈](#) (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant (T₈). However lowest number of fruits plant⁻¹ (10.87), yield plant⁻¹ (91.04 g) and yield hectare⁻¹ (5.01 t/ha) was noted in [control the plants treated with T₁](#) ([control](#)). Increased production could also be attributed to improved nutrient absorption and utilisation due to the combined use of inorganic and organic fertilisers, as well as bio-fertilizers [\(Kumar *et al.*, 2024\)](#). The biofertilizer's positive effect on harvesting duration and

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fruit yield could be attributed due to advantageous effects on vegetative ~~development-growth~~ and flowering, which probably gave the inoculated plants access to more photosynthates for a longer ~~amount-period~~ of time, boosting fruit yield and lengthening the harvesting season- (Gupta and Tripathi, (2012). Similar outcomes were ~~presented-reported~~ by Jain *et al.* (2017), Singh *et al.* (2015), and Reddy and Goyal (2020).

Leaf nutrient content

The results demonstrated that the maximal leaf nutritional content; namely, nitrogen (3.09%), phosphorus (1.37%), potassium (3.39%), calcium (2.37%) and magnesium (0.39%) ~~was-were~~ recorded in T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant) and minimum nitrogen (2.21%), phosphorus (0.94%), potassium (2.15%), calcium (1.70%) and magnesium (0.11%) ~~was-were~~ recorded in T₁ (control) (Table 4). The intake of any nutrient from the soil has a significant impact on how much of it accumulates in the leaves. Weinbaum *et al.* (1978) discovered the presence of leaves was necessary for nitrate uptake in prune trees, and they addressed this phenomenon in terms of the supply of leaf carbohydrates. In a similar vein, adding FYM and vermicompost that had been enhanced with *Pseudomonas* and *Azotobacter* increased the phosphorus levels in strawberry leaves- (Negi *et al.*, 2021). Yadav (2010) found that increased nitrogen application led to an increase in nitrogen and phosphorus content in peaches. Furthermore, *Azotobacter* and PSB boosted leaf phosphorus content, indicating that biofertilizers may have produced a specific microbial community in the root rhizosphere zone to improve phosphorus absorption. Singh *et al.* (2012) and Verma and Rao (2013) found that organic manures and biofertilizers had a similar favourable effect on the NPK content of strawberry leaves. These findings unequivocally demonstrate that bacterial strains would have facilitated the correct uptake of soil nutrients and that the nutritional condition of leaves was positively correlated with increased availability of soil nutrients.

Soil health attributes

In light of the current study's findings, the application of bio-fertilizers in combination with RDF had a substantial effect on soil electrical conductivity, organic carbon, and soil pH. Lowest soil pH (7.13) and soil electrical conductivity (0.36 dS/m) were measured after applying T₈ ~~treatment~~ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant). However, maximum soil pH (7.70), soil electrical conductivity (0.65 dS/m) was resulted in treatment T₁ (control). On the other side highest organic carbon (0.64%), available

nitrogen (349.93_kg/ha), phosphorus (59.17_kg/ha) and potassium content (150.77 kg/ha) ~~is~~ was recorded in T₈ (RDF + *Azotobacter*_@_2g/plant + *PSB*_@_2g/plant + *Azospirillum*_@_2g/plant) and minimum organic carbon (0.41%), available nitrogen (271.57_kg/ha), phosphorus (36.70_kg/ha) and potassium content (127.97_kg/ha) ~~is-was~~ observed in T₁ (control) (Table 5). Bio-fertilizers may have boosted biological nitrogen fixation and phosphate solubilisation, ~~resulting-resulted~~ in an increase in available N and P content. Singh *et al.* (2012) also indicated that bio-fertilizers improved plant growth and had a direct impact on N₂-fixation and phosphorous mobilisation in strawberry plants. Addition of phosphate-solubilizing microorganisms to inorganic fertilisers also had a considerable impact on the accumulation of accessible phosphorous in the soil. These results align with ~~Singh et al. (2010) and~~ Sau *et al.* (2017) ~~and~~ Kumar *et al.* (2019a).

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Economic attributes

Economics of strawberry cultivation influenced by different treatments of bio-fertilizers is presented in table 6. The cost of cultivation of strawberry cv. Winter Dawn computed, in open field circumstances, treatment-wise, for a single season ~~on afor oe~~ hectare ~~of land area~~. The result indicated that net return was higher in treatment T₈ (RDF + *Azotobacter*_@_2g/plant + *PSB*_@_2g/plant + *Azospirillum*_@_2g/plant). Data pertaining to economics of different treatments ~~shows-showed~~ that maximum gross income (21,02,150 ₹/ha), net return per hectare (16,61,790 ₹/ha) and B:C ratio (3.77) for strawberry production was ~~observed-calculated for strawberry production~~ with the application of RDF + *Azotobacter*_@_2g/plant + *PSB*_@_2g/plant + *Azospirillum*_@_2g/plant. Whereas, the minimum net return per hectare (3,98,969 ₹/ha), gross income (8,01,128 ₹/ha) and B:C ratio (0.99) ~~were observed~~ was estimated with treatment T₁ (control) (Table 6). The increased yield of high-quality fruits may account for the rise in benefit-to-cost ratio (Kumar *et al.* 2018). These results concur with Hazarika *et al.* (2016), Pardeep and Saravanam (2017) and Jaiswal *et al.* (2023) ~~which support the result obtained in the present investigation~~.

Conclusion

~~From overall study, we obtained that bio fertilizers are more essential for cultivation of strawberry. The overall results, gave~~ From the entire experiment, it can be concluded that treatment T₈ (RDF + *Azotobacter*_@_2g/plant + *PSB*_@_2g/plant + *Azospirillum*_@_2g/plant) is the best treatment for strawberry cultivation with respect to in all the attributes. The treatment significantly increased the vegetative growth (plant height, plant spread, number of

leaves, leaf area), flowering attributes (days taken for first flowering and no. of flowers per plant) in strawberry plant. In case of physico-chemical attributes, soil attributes and yield attributes, the maximum result was also ~~found~~ obtained with T₈ (RDF + *Azotobacter* @ 2g/plant + PSB @ 2g/plant + *Azospirillum* @ 2g/plant) as compared to all other treatment combination and also gave us maximum net return (₹ 16,61,790) and highest B:C ratio.

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Table 1: Effect of biofertilizer on vegetative growth attributes of strawberry cv. Winter Dawn

Treatment	Plant height (cm)	Plant spread (cm)	Number of leaves per plant	Leaf area (cm ²)
T ₁	10.66 ^e	22.90 ^b	11.87 ^f	101.02 ^b
T ₂	13.59 ^c	27.93 ^e	14.73 ^d	113.45 ^d
T ₃	12.22 ^d	27.22 ^f	13.60 ^e	110.37 ^f
T ₄	11.69 ^d	26.57 ^g	13.13 ^e	106.76 ^g
T ₅	15.01 ^b	31.48 ^b	18.73 ^b	119.10 ^b
T ₆	14.89 ^b	29.98 ^c	17.07 ^c	116.53 ^c
T ₇	13.9 ^c	29.18 ^d	15.27 ^d	112.02 ^e
T ₈	16.82 ^a	32.78 ^a	20.00 ^a	121.53 ^a
C.D.(0.05)	0.70	0.62	0.65	0.62
SE(m)	0.23	0.20	0.21	0.20
SE(d)	0.32	0.29	0.30	0.28

Such as: T₁: (Control), T₂: (RDF + Azotobacter @ 2g/plant), T₃: (RDF + PSB @ 2g/plant), T₄: (RDF + Azospirillum @ 2g/plant), T₅: (RDF + Azotobacter @ 2g/plant + PSB @ 2g/plant), T₆: (RDF + Azotobacter @ 2g/plant + Azospirillum @ 2g/plant), T₇: (RDF + PSB @ 2g/plant + Azospirillum @ 2g/plant), T₈: (RDF + Azotobacter @ 2g/plant + PSB @ 2g/plant + Azospirillum @ 2g/plant)

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Table 2: Effect of biofertilizer on flowering and yield characteristics of strawberry cv. Winter Dawn

Treatment	Days taken to first flowering	Numbers of flower per plant	Number of fruits per plant	Yield per plant (g)	Yield per hectare (t/ha)
T ₁	70.55 ^a	14.60 ^g	10.87 ^g	91.04 ^g	5.01 ^g
T ₂	63.33 ^d	19.53 ^d	14.87 ^e	165.41 ^d	9.09 ^d
T ₃	67.77 ^{bc}	17.80 ^e	14.40 ^e	153.41 ^e	8.44 ^e
T ₄	68.55 ^b	16.73 ^f	13.53 ^f	132.86 ^f	7.31 ^f
T ₅	60.77 ^c	23.00 ^b	17.73 ^b	198.57 ^b	10.92 ^b
T ₆	62.55 ^d	21.60 ^c	16.80 ^c	187.57 ^c	10.32 ^c
T ₇	66.66 ^c	20.73 ^c	15.80 ^d	165.10 ^d	9.08 ^d
T ₈	58.99 ^e	24.13 ^a	18.73 ^a	238.88 ^a	13.14 ^a
C.D.(0.05)	1.75	0.92	0.74	7.76	0.43

SE(m)	0.57	0.30	0.24	2.54	0.14
SE(d)	0.85	0.43	0.34	3.59	0.19

Such as: T₁(Control), T₂(RDF + *Azotobacter*@2g/plant), T₃(RDF+PSB@2g/plant), T₄(RDF+*Azospirillum*@2g/plant), T₅(RDF + *Azotobacter*@2g/plant + PSB@2g/plant), T₆(RDF + *Azotobacter*@2g/plant + *Azospirillum*@2g/plant), T₇(RDF +PSB@2g/plant + *Azospirillum*@2g/plant), T₈(RDF + *Azotobacter*@2g/plant + PSB@2g/plant + *Azospirillum*@2g/plant)

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Table 3: Effect of biofertilizer on physico-chemical attributes of strawberry cv. Winter Dawn

Treatment	Berry length (mm)	Berry breadth (mm)	Berry weight (g)	TSS (°Brix)	Total sugar (%)	Reducing sugar (%)	Anthocyanin content (mg/100g)
T ₁	34.31 ^e	31.43 ^e	8.38 ^e	6.90 ^c	5.68 ^d	3.73 ^d	16.39 ^d
T ₂	40.42 ^c	35.77 ^b	11.13 ^b	8.13 ^b	6.12 ^d	4.04 ^b	20.29 ^c
T ₃	39.71 ^c	34.11 ^c	10.65 ^c	8.17 ^b	6.76 ^{bc}	4.08 ^b	17.65 ^d
T ₄	35.18 ^c	33.03 ^{cd}	9.82 ^d	8.37 ^b	6.63 ^c	3.82 ^{cd}	17.92 ^d
T ₅	42.27 ^b	36.74 ^b	11.20 ^b	8.95 ^{ab}	7.18 ^b	4.29 ^a	22.57 ^b
T ₆	39.11 ^{cd}	36.08 ^b	11.16 ^b	8.53 ^b	7.01 ^{bc}	4.05 ^b	23.93 ^{ab}
T ₇	37.52 ^d	32.79 ^d	10.45 ^c	8.15 ^b	6.98 ^{bc}	3.96 ^{bc}	18.74 ^{cd}
T ₈	45.88 ^a	38.96 ^a	12.76 ^a	9.42 ^a	7.58 ^a	4.38 ^a	26.14 ^a
C.D.(0.05)	1.74	1.21	0.36	0.81	0.40	0.15	2.28
SE(m)	0.57	0.39	0.12	0.26	0.13	0.05	0.74
SE(d)	0.80	0.56	0.16	0.37	0.19	0.07	1.05

Such as: T₁(Control), T₂(RDF + *Azotobacter*@2g/plant), T₃(RDF+PSB@2g/plant), T₄(RDF+*Azospirillum*@2g/plant), T₅(RDF + *Azotobacter*@2g/plant + PSB@2g/plant), T₆(RDF + *Azotobacter*@2g/plant + *Azospirillum*@2g/plant), T₇(RDF +PSB@2g/plant + *Azospirillum*@2g/plant), T₈(RDF + *Azotobacter*@2g/plant + PSB@2g/plant + *Azospirillum*@2g/plant)

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Table 4: Effect of biofertilizer on leaf nutrient content of strawberry cv. Winter Dawn

Treatment	Nitrogen (%)	Phosphorus (%)	Potassium (%)	Calcium (%)	Magnesium (%)
T ₁	2.21 ^h	0.94 ^f	2.15 ^h	1.70 ^g	0.11 ^f
T ₂	2.63 ^e	1.18 ^{cd}	2.76 ^e	2.02 ^d	0.20 ^{de}
T ₃	2.53 ^f	1.14 ^{de}	2.54 ^f	1.92 ^e	0.14 ^f
T ₄	2.47 ^g	1.09 ^e	2.30 ^g	1.86 ^f	0.17 ^{ef}
T ₅	2.93 ^b	1.28 ^b	3.21 ^b	2.21 ^b	0.31 ^b
T ₆	2.82 ^c	1.21 ^c	3.11 ^c	2.13 ^c	0.27 ^{bc}

T₇	2.71 ^d	1.23 ^{bc}	2.99 ^d	2.03 ^d	0.24 ^{cd}
T₈	3.09 ^a	1.37 ^a	3.39 ^a	2.37 ^a	0.39 ^a
C.D.(0.05)	0.04	0.04	0.04	0.05	0.02
SE(m)	0.014	0.013	0.015	0.015	0.007
SE(d)	0.02	0.02	0.02	0.02	0.01

Such as: T₁(Control), T₂(RDF + *Azotobacter*@2g/plant), T₃(RDF+PSB@2g/plant), T₄(RDF+*Azospirillum*@2g/plant), T₅(RDF + *Azotobacter*@2g/plant + PSB@2g/plant), T₆(RDF + *Azotobacter*@2g/plant + *Azospirillum*@2g/plant), T₇(RDF +PSB@2g/plant + *Azospirillum*@2g/plant), T₈(RDF + *Azotobacter*@2g/plant + PSB@2g/plant + *Azospirillum*@2g/plant)

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Table 5: Effect of biofertilizer on soil health attributes of strawberry cv. Winter Dawn

Treatment	Soil pH	Soil EC (dS/m)	Organic carbon (%)	Available nitrogen (kg/ha)	Available phosphorus (kg/ha)	Available potassium (kg/ha)
T₁	7.70 ^a	0.65 ^a	0.41 ^d	271.57 ^g	36.70 ^g	127.97 ^b
T₂	7.40 ^{bc}	0.51 ^{bc}	0.53 ^{bc}	333.40 ^c	44.83 ^f	137.80 ^d
T₃	7.43 ^b	0.55 ^b	0.50 ^{bc}	323.37 ^e	51.07 ^d	133.83 ^f
T₄	7.43 ^b	0.54 ^{bc}	0.47 ^c	296.83 ^f	46.23 ^e	132.53 ^g
T₅	7.20 ^{cd}	0.42 ^d	0.56 ^b	344.53 ^b	57.43 ^b	147.10 ^b
T₆	7.30 ^{bcd}	0.49 ^c	0.54 ^b	342.57 ^b	47.53 ^c	143.50 ^c
T₇	7.47 ^b	0.55 ^b	0.49 ^{bc}	326.83 ^d	53.50 ^c	135.87 ^c
T₈	7.13 ^d	0.36 ^d	0.64 ^a	349.93 ^a	59.17 ^a	150.77 ^a
C.D.(0.05)	0.20	0.06	0.02	2.08	1.31	1.29
SE(m)	0.06	0.02	0.01	0.68	0.43	0.42
SE(d)	0.09	0.03	0.01	0.96	0.61	0.59

Such as: T₁(Control), T₂(RDF + *Azotobacter*@2g/plant), T₃(RDF+PSB@2g/plant), T₄(RDF+*Azospirillum*@2g/plant), T₅(RDF + *Azotobacter*@2g/plant + PSB@2g/plant), T₆(RDF + *Azotobacter*@2g/plant + *Azospirillum*@2g/plant), T₇(RDF +PSB@2g/plant + *Azospirillum*@2g/plant), T₈(RDF + *Azotobacter*@2g/plant + PSB@2g/plant + *Azospirillum*@2g/plant)

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Table 6: Effect of biofertilizer on economic attributes of strawberry cv. Winter Dawn

Treatment	Gross income	Net Return	B:C
T₁	8,01,128.5	3,98,969	0.99
T₂	14,55,643	10,18,583	2.33

T₃	13,50,020	9,12,960	2.09
T₄	11,69,127	7,30,967	1.67
T₅	17,47,398	13,09,238	2.99
T₆	16,50,587	12,11,327	2.76
T₇	14,52,886	10,13,626	2.31
T₈	21,02,150	16,61,790	3.77

Such as: T₁(Control), T₂(RDF + *Azotobacter*@2g/plant), T₃(RDF+PSB@2g/plant), T₄(RDF+*Azospirillum*@2g/plant), T₅(RDF + *Azotobacter*@2g/plant + PSB@2g/plant), T₆(RDF + *Azotobacter*@2g/plant + *Azospirillum*@2g/plant), T₇(RDF +PSB@2g/plant + *Azospirillum*@2g/plant), T₈(RDF + *Azotobacter*@2g/plant + PSB@2g/plant + *Azospirillum*@2g/plant)

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