The effect of supplementary light on the photosynthetic apparatus of strawberry plants under salinity and alkalinity stress

Mohammad Reza Malekzadeh Shamsabad1, Majid Esmaeilizadeh1, Hamid Reza Roosta2, Mohammad Reza Dehghani3, Piotr Dąbrowski4 & Hazem M. Kalaji5,6

Considering the destructive effect of stresses on the photosynthetic apparatus of plants and the important role of light in photosynthesis, we investigated the effect of complementary light on the photosynthetic apparatus under salinity and alkalinity stress conditions. Light-emitting diodes (LEDs) in monochromatic blue (460 nm), monochromatic red (660 nm), dichromatic blue/red (1:3), white/yellow (400–700 nm) at 200 μmol m⁻² S⁻¹, and without LED treatment were used. The stress treatments were in three stages: Control (no stress), Alkalinity (40 mM NaHCO₃), and Salinity (80 mM NaCl). Our results showed that salinity and alkaline stress reduced CO₂ assimilation by 62.64% and 40.81%, respectively, compared to the control treatment. The blue light spectrum had the highest increase in water use efficiency (54%) compared to the treatment without supplementary light. Under salinity and alkalinity stress, L, K, and H bands increased and G bands decreased compared to the control treatment, with blue/red light causing the highest increase in L and K bands under both stress conditions. In salinity and alkalinity stress, white/yellow and blue/red spectra caused the highest increase in H bands. Complementary light spectra increased the G band compared to the treatment without complementary light. There was a significant decrease in power indices and quantum power parameters due to salt and alkalinity stress. The use of light spectra, especially blue, red, and blue/red light, increased these parameters compared with treatment without supplementary light. Different light spectra have different effects on the photosynthetic apparatus of plants. It can be concluded that using red, blue spectra and their combination can increase the resistance of plants to stress conditions and be adopted as a strategy in planting plants under stress conditions.

Light quality detection and measurement are essential for plant growth and development. Under natural growth conditions, due to environmental factors such as clouds, rain, and other climatic factors, natural light may not be enough for optimal plant growth. Under these conditions, artificial light sources in the controlled cultivation of plants promote high yield and production of quality products. Light-emitting diodes (LEDs) can also be used as light sources for planting systems and photobiological research. These LEDs have a low mass, small size, long lifetime, and narrow spectral output. The use of light-emitting diodes as light sources in agriculture has been increasing in recent years.

Photosynthetic pigments absorb light energy and convert it into chemical energy, a process in the photosynthetic apparatus. Blue and red spectra play an important role in photosynthesis. Blue and red lights have been shown to induce stomatal opening. An important tool for researching the effects of various stresses on photosynthesis in plants is chlorophyll a fluorescence (Chl) kinetics. The photochemical efficiency of plant photosynthesis is represented by fluorescence kinetics. This tool provides important information about the...
functional and structural properties of the components involved in photosynthetic electron transfer. The use of combined tools (Photosynthetic Efficiency Analysis of plants and Measurement of Plant Gas Exchange) permitted the exploration and description of responses to stress. The JIP test has been successfully used to study the photosynthetic apparatus of various plant species under abiotic stresses.

With increasing population and degradation of natural environments, soil salinization has become a serious global problem. Approximately 7% of the world’s land (more than 900 million hectares) is at risk of salinity and alkalinity. Salinity and alkaline stresses as two abiotic stresses have severely limited the development of global agriculture. Therefore, the study of plant resistance mechanisms in salinity and alkaline stress conditions is of great practical importance for the development of strategies for tolerance to salinity and alkalinity stresses. It is possible to increase plant tolerance to stress conditions for sustainable agricultural development by using methods. Salinity stress affects the transfer of electrons from the reaction centers (RCs) to the plastoquinone pool. Salinity stress disrupts electron transfer, and adversely affects photosynthetic efficiency. These destructive effects reduce the efficiency of photosynthesis. Salt accumulation in mesophytic cells prevents carbon uptake, thus increasing the internal CO₂ concentration and ultimately decreasing the stomatal conductance.

One of the criteria for studying stress tolerance in plants is the attributes of plant gas exchange. Plant gas exchange properties are among the most important physiological characteristics that significant changes in these properties occur due to salinity stress. Growth of different strawberry cultivars with alkaline stress decreased due to reduction of chlorophyll content and inhibition of photosynthesis and electron transfer was significantly inhibited, which led to a decrease in photosynthetic yield index. In an experiment, salinity stress reduced chlorophyll content. It reduced vegetative and reproductive growth in the Paros strawberry cultivar, and inhibition of chlorophyll biosynthesis, acceleration of its degradation, and oxidative damage caused by salinity were considered as the main reasons for chlorophyll content reduction.

Much research has been done on the growth response of plants to light quality, and its effect on plant photosynthesis. For example, the work of Hyo et al. investigated the effect of LED light on strawberries in the greenhouse and growth chamber. They showed that different spectrums of LED lights in various conditions have different effects on the growth and development of strawberry plants. They suggested that growing strawberries in a greenhouse with LED lighting is more efficient than planting in a growth chamber. Much research has been done on the effects of blue and red spectra on plants, but it is not easy to reach a consensus on the effect of an optical spectrum due to reduction of chlorophyll content and inhibition of photosynthesis and electron transfer was significantly inhibited, which led to a decrease in photosynthetic yield index. In an experiment, salinity stress reduced chlorophyll content. It reduced vegetative and reproductive growth in the Paros strawberry cultivar, and inhibition of chlorophyll biosynthesis, acceleration of its degradation, and oxidative damage caused by salinity were considered as the main reasons for chlorophyll content reduction.

Table 1. The concentration of nutrients used in the nutrient solution of this experiment.

| Macronutrients | Concentration (mg L⁻¹) | Micronutrients | Concentration (mg L⁻¹) |
|----------------|------------------------|----------------|------------------------|
| N              | 128                    | Fe             | 5                      |
| P              | 58                     | Mn             | 2                      |
| K              | 211                    | Zn             | 0.25                   |
| Ca             | 104                    | B              | 0.7                    |
| Mg             | 40                     | Cu             | 0.07                   |
| S              | 54                     | Mo             | 0.05                   |

This work aimed to understand better the mechanism of photosynthetic apparatus response to salinity and alkaline stresses under different light spectra. We tried to answer the question of which spectrum or spectra of complementary light can improve the performance of the photosynthetic apparatus under salinity and alkaline stress conditions. We tried to show how complementary light spectra affect the photosynthetic apparatus of the Paros strawberry cultivar under salinity and alkalinity stress conditions. We hypothesize that the use of complementary light can have a positive effect on the photosynthesis apparatus of strawberry plants. The manipulation of the light spectra used by plants has a significant positive effect on the efficiency of the photosynthesis apparatus in stressed plants. Therefore, selecting the appropriate light spectrum can reduce the destructive effects of salinity and alkalinity stress. We used LEDs as complementary lighting in the greenhouse and under stress conditions. Expanding the use of LEDs as complementary lighting in greenhouses and stress conditions is important for commercial horticulture and plant research.

**Materials and methods**

**Plant material and growth conditions.** In 2020, the experiment was conducted in the research greenhouse at Vali-e-Asr University of Rafsanjan- Iran (Latitude: 30°21'17.6004" N, Longitude: 56°0'9.738" E, Elevation: 1545.924). Bare root plants of strawberry (Fragaria × ananassa Duch, cv. Paros) were obtained from a nursery in Karaj, Iran. Plants were planted in a pot of 4 l containing cocopeat and perlite (70:30 ratio). Each treatment included three pots, and each pot was an experimental unit in which three plants were planted. Pots with a width of 30 cm were used to plant three plants. Plants were cultivated in the greenhouse with a temperature of 25/15 ± 2 °C (day/night), a photoperiod of 11.13 h (light/dark), and relative humidity of 50 ± 10%. During this period, plants were fertigated with the Morgan nutrient solution (Morgan 2003) (Electrical Conductivity (EC): 1.4 dS m⁻¹ and pH 6.5) (Table 1). Plants were treated with five light levels and two stress levels, including control (without...
out stress), alkalinity (40 mM NaHCO₃), and salinity (80 mM NaCl). The alkalinity and salinity treatments were applied twenty days after planting. 100 ml of NaHCO₃ and NaCl were added to each pot to maintain uniform stress for the plants. Stresses continued until the end of the experiment and the completion of data collection.

**Light environment.** Plants were grown under LED systems (length: 100 mm, width: 5 mm, and height: 5 mm) with 24W LED tubes (Parto Roshd Novin Company, Iran grow light, Iran) of different spectral ranges: monochromatic blue (B) (with peak 460 nm), monochromatic red (R) (with peak 660 nm), dichromatic blue/red (1:3), white/yellow (400–700 nm) (Fig. 1) and without LED treatment.

Photon flux density (PPFD) of LED lights was at 200 μmol m⁻² s⁻¹ per 11-h light period. Directly above each plant, LED lighting systems were mounted 30 cm apart.

**Evaluation of chlorophyll fluorescence.** We used a portable photosynthetic efficiency analyzer (PEA, Hansatech Inc. Co., UK) sixty days after planting to measure and calculate chlorophyll fluorescence parameters (PEA, Hansatech Inc. Co., UK). For this purpose, fully mature leaves of each pot were adapted to a dark time for 15 min by fixing special tags on each leaf before measurement. Then the sensor cup was mounted on the leaf for calculation. The chlorophyll a fluorescence transient was caused by a saturating photon flux density of 3,500 μmol(photon) m⁻² s⁻¹ given by three light-emitting diodes (peak 650 nm) to produce fluorescence curves ranging from Fₚ to Fₘ (Fₚ fluorescence at time t after the onset of actinic illumination; Fₚ = Fₚₐ₃ ₜₐₘᵢₐ₉ minimum fluorescence intensity; Fₚ = Fₚ₃ₙ₉ₙₚ₉ₜₚ₉ fluorescence intensity at the I-step; Fₚ = Fₚ₃ₙ₉ₚ₉ₜₚ₉ₚ₉ maximum fluorescence intensity at peak P of OJIP) for all treatments. The PSII parameters obtained from the OJIP transient were evaluated based on the Strasser et al.¹⁴ methods. Parameters for chlorophyll fluorescence are listed in an Suppl. Appendix Table.

**Leaf gas exchange.** Plant gas exchange parameters include net CO₂ assimilation rate (A, μmol CO₂ m⁻² s⁻¹), intrinsic water-use efficiency (WUEi, μmol CO₂ mol H₂O⁻¹), stomatal conductance (Gs, mol H₂O m⁻² s⁻¹), transpiration (E, mmol H₂O m⁻² s⁻¹), Sub-stomatal CO₂ concentration (Ci, μmol CO₂ mol⁻¹) and, instantaneous carboxylation efficiency (A/Ci) were measured using a portable photosynthesis system (ADC BioScientific Ltd, Hoddesdon, UK) after 60 days after planting. Around 9:00 AM and 12:00 AM, measurements were performed on completely expanded leaves.
| Light sources | Stress | A (μmol CO₂ m⁻² s⁻¹) | E (mmol H₂O m⁻² s⁻¹) | Gₛ (mol H₂O mol⁻¹) | Cᵢ (μmol CO₂ mol⁻¹) | WUEᵢ (μmol CO₂ mol⁻¹ H₂O⁻¹) | A/Cᵢ |
|--------------|--------|----------------------|----------------------|------------------|---------------------|---------------------|------|
| Blue         | Control | 20 ± 0.6             | 9.91 ± 0.29          | 0.280 ± 0.00     | 167 ± 7.8          | 5.2 ± 2.0           | 0.12 ± 0.005    |
|              | Salinity| 5.99 ± 0.29          | 8.5 ± 0.63           | 0.114 ± 0.003    | 255 ± 8            | 2.5 ± 3.36         | 0.023 ± 0.004   |
|              | Alkalinity| 12.5 ± 0.37         | 4.97 ± 0.49          | 0.190 ± 0.005    | 207 ± 2.9          | 3.1 ± 0.84         | 0.06 ± 0.002    |
| Red          | Control | 18 ± 0.33            | 13.2 ± 0.13          | 0.280 ± 0.005    | 93 ± 4.2           | 1.9 ± 0.3           | 0.192 ± 0.008   |
|              | Salinity| 8.32 ± 0.15          | 5.9 ± 0.42           | 0.160 ± 0.005    | 218 ± 12           | 1.2 ± 0.1          | 0.038 ± 0.002   |
|              | Alkalinity| 9.96 ± 0.13          | 8.15 ± 0.39          | 0.166 ± 0.008    | 164 ± 5.4          | 1.8 ± 0.3           | 0.06 ± 0.001    |
| Blue/Red     | Control | 27.6 ± 0.16          | 8.92 ± 0.25          | 0.363 ± 0.008    | 142 ± 3            | 2.3 ± 0.5           | 0.194 ± 0.003   |
|              | Salinity| 12.7 ± 0.64          | 7.17 ± 0.26          | 0.183 ± 0.003    | 222 ± 3.5          | 1.2 ± 0.1          | 0.057 ± 0.003   |
|              | Alkalinity| 14.3 ± 0.2           | 7.86 ± 0.52          | 0.180 ± 0.005    | 166 ± 3.9          | 1.8 ± 0.1          | 0.086 ± 0.002   |
| white/yellow | Control | 16.9 ± 0.34          | 10.1 ± 0.45          | 0.226 ± 0.008    | 209 ± 0.57         | 2.0 ± 0.1          | 0.08 ± 0.001    |
|              | Salinity| 5.89 ± 0.11          | 5.59 ± 0.12          | 0.116 ± 0.003    | 255 ± 1.2          | 0.9 ± 0.3          | 0.023 ± 0.004   |
|              | Alkalinity| 8.47 ± 0.51          | 7.24 ± 0.36          | 0.143 ± 0.003    | 215 ± 2.3          | 1.6 ± 0.4          | 0.039 ± 0.002   |
| Ambient      | Control | 14.4 ± 0.24          | 7.42 ± 0.31          | 0.240 ± 0.005    | 222 ± 1.4          | 3.3 ± 0.3          | 0.064 ± 0.001   |
|              | Salinity| 4.94 ± 0.21          | 3.88 ± 0.03          | 0.100 ± 0.001    | 278 ± 1.4          | 1.8 ± 0.3          | 0.017 ± 0.007   |
|              | Alkalinity| 6.2 ± 0.19           | 4.64 ± 0.12          | 0.120 ± 0.001    | 272 ± 10           | 1.9 ± 0.4          | 0.022 ± 0.005   |

Table 2. Effect of light spectra and stress on leaf gas exchange parameters of strawberry cv. Paros. Values are means ± SE of three replicates. Bars with different letters show significant differences at P ≤ 0.05 (LSD). Significance according to ANOVA, ns, *, **, *** nonsignificant and significant P ≤ 0.05, 0.01, 0.001, respectively. Control (non-stress), salinity (80 mM NaCl) and alkalinity (40 mM NaHCO₃). Photon flux density for the complement spectrum (PPFD) 200 ± 10 mmol m⁻² s⁻¹. SAS software version 9.4 was used for data analysis (https://www.sas.com/en_us/home.html).

Experimental design and data analysis. The experiment was factorial, consisting of a randomized, complete design with two factors in three replicates and three single plants per pot. All data were analyzed using SAS software version 9.4 (SAS Institute, Cary, NC, USA. https://www.sas.com/en_us/home.html). When variance analysis (ANOVA) indicated significant treatment effects, significant mean variations (P < 0.05) were calculated using the LSD Multiple Range Test. Biophysical parameters were determined with “PEA Plus” software version 1.12 (http://www.hansatech-instruments.com). Pearson’s correlation coefficient was applied to determine the relationships among the parameters studied. Principal component analysis was performed using MATLAB software version R2015b (https://www.mathworks.com). A correlation plot was drawn with Origin Pro software version 2021 (https://www.originlab.com/2021). The biplots were made using Excel software version 2016 (https://www.microsoft.com).

Statement of compliance. The authors confirm that all the experimental research and field studies on strawberry plants, including the collection of plant material, complied with relevant institutional, national, and international guidelines and legislation. Also, obtained licenses for the preparation of Bare root plants of strawberry.

Results

Leaf gas exchange analyses. The analysis of ANOVA showed that plant gas exchange parameters were significantly affected by different light spectra, stress, and their interaction effects. Net CO₂ assimilation (A) of plants was influenced considerably by salinity and alkalinity stress and different light spectra. Net CO₂ assimilation decreased under salinity and alkalinity stress compared to the control. Under salinity stress, blue/red and red light had the greatest effect on increasing CO₂ assimilation under stress conditions and had the lowest percentage of reduction compared to the control treatment. Blue/red light has the largest level of CO₂ assimilation under alkalinity stress. There was the least decrease in CO₂ assimilation in blue light relative to the control treatment, and the highest percentage reduction was found without LED treatment (Table 2).

A comparison of the mean interactions of different light spectra and stress on plant gas exchange parameters of strawberry plants showed that transpiration rate (E) and stomatal conductance (Gₛ) decreased under salinity and alkalinity stress. The lowest percentage reduction in transpiration rate (E) under salinity stress was observed in blue and blue/red light with 16% and 19% reduction compared to the control treatment, respectively. Under alkaline stress, the lowest percentage of transpiration (E) was observed in blue/red light with a decrease of 11% compared to the control treatment. The red light had the greatest effect on the Gₛ parameter under salinity conditions and caused the least decrease (42%) compared to the control treatment. Under alkalinity stress, blue light had the greatest effect on the E parameter.
Internal CO₂ (Ci) concentrations under stress conditions were significantly affected by different light spectra. Under stress conditions, this parameter increased, and the application of light treatment decreased Ci. Blue/red light increased the instantaneous carboxylation efficiency (A/Ci) parameter compared to other light spectra. Salinity and alkalinity stress reduced water use efficiency (WUEi) compared to the control treatment. Under all stress conditions, blue/red light significantly increased WUEi. The lowest WUEi was observed in the without supplementary light treatment.

Prompt chlorophyll a fluorescence. According to the results, stress and light spectrum had a significant effect on the fluorescent transients. Salinity and alkalinity stress reduced the fluorescent transients compared to non-stress conditions. Different light spectra had a positive effect on the fluorescence curve. Under salinity stress, the blue/red spectrum had a significant effect on increasing the fluorescence curve, especially at points I and P compared to other light spectra. Whereas under alkaline stress conditions, red and white/yellow light had the most significant effect on increasing the fluorescent transients. Blue and blue/red light also caused a significant increase in the fluorescent transients compared to the treatment without supplementary light treatment.

Chlorophyll fluorescent transients and calculated curves. To help visualize the influence of the stress and light spectra on the transient dynamics, the curves were plotted as relative variable fluorescence, $V_t = (F_t - F_o)/(F_M - F_o)^{14}$. The changes in the OJIP fluorescence were calculated by subtracting the values of the fluorescence (O–P) recorded in plants under stress from those recorded for control plants. We observed significant changes in the prompt fluorescence of plants under stress and different light spectra at the J (Vj) and I (Vi) steps. The characteristics of OJIP fluorescence transients recorded in strawberry plants under different light spectra differed from those recorded without LED treatment. The results showed that the relative variable fluorescence curve in plants under salinity and alkalinity stress was higher than in control plants. Also, salinity stress was higher than alkalinity stress (Fig. 3).

To do a complete analysis of the different light spectra and stress-induced shifts in the OJIP curve, the differential curves were plotted separately for the main band that occurred during the transient O–P: The L, K, H, and G bands. The Curves for these bands were plotted by subtracting the normalized fluorescence values (between O and K, O and J, J and I, and I and P, respectively) observed in control plants from those recorded in plants under different light spectrum and stress conditions.
In ambient light, significant changes in the K band were observed under the influence of alkaline and salt stress caused smaller but also visible changes (Fig. 4). Alkaline stress also caused changes during the L band, which was not observed in the plants treated with salt. In non-stressed plants under the influence of different light conditions, no significant differences were observed in both bands. Under the influence of salt stress, both bands had the highest course in blue/red light and the lowest in red and blue light. Under alkaline stress, both bands had a higher course under blue/red light, while under the other light conditions, the curves had the same course as in ambient.

Both stresses caused a change in the H and G bands (Fig. 5). However, the H band increased, but the G band decreased compared to the control (non-stressed plants). In the ambient treatment, in non-stressed plants, red and blue/red spectra, and salt-treated plants, white/yellow spectra significantly increased the H band. Under alkaline stress, the blue/red spectrum increased the H band more than other light spectra. In the control treatment, the blue and red spectra increased the G band compared to the treatment without supplementary light. Under salinity stress, blue/red and blue light could significantly increase the G band, and under alkaline stress, white/yellow light had the greatest effect on this band (Fig. 5).

**JIP-test parameters, calculated from chlorophyll fluorescence transients.** OJIP transients have been translated into biophysical parameters: the basic parameters derived from the extracted data, normalized data, specific energy fluxes (per active PSII reaction center), performance indexes, quantum yield for primary photochemistry, and slopes and integrals (Strasser et al. 2010). The values of the measured parameters have been normalized to those of the control plants. On radar plots, the deviation of the activity pattern of plants under stress and different light spectra from control plants were shown (Fig. 6).
Figure 4. Differential curves of $\Delta W_K$ and $\Delta W_L$ of chlorophyll a fluorescence of strawberry cv. Paros. (A,B) In ambient light (control, salt stress, alkalinity stress); (C,D) non-stressed in different light spectra (ambient, blue, red, blue/red, white/yellow); (E,F) under salt stress in different light spectra (ambient, blue, red, blue/red, white/yellow); (G,H) under alkalinity stress in different light spectra (ambient, blue, red, blue/red, white/yellow). Biophysical parameters were determined with "PEA Plus" software version 1.12 (http://www.hansa-tech-instruments.com). The graphs were drawn using Microsoft Excel version 2016 (https://www.microsoft.com).
Figure 5. Differential curves of ΔWₚₐ and ΔWₑ of chlorophyll a fluorescence of strawberry cv. Paros. (A,B) In ambient light (control, salt stress, alkalinity stress); (C,D) non-stressed in different light spectra (ambient, blue, red, blue/red, white/yellow); (E,F) under salt stress in different light spectra (ambient, blue, red, blue/red, white/yellow); (G,H) under alkalinity stress in different light spectra (ambient, blue, red, blue/red, white/yellow).

Biophysical parameters were determined with "PEA Plus" software version 1.12 (http://www.hansa-tech-instruments.com). The graphs were drawn using Microsoft Excel version 2016 (https://www.microsoft.com).
Figure 6. JIP-test parameters normalized on radar plots. (A) The effect of ambient light conditions on non-stressed plants; (B) the effect of salt stress at different light conditions; (C) the effect of alkalinity stress at different light conditions; (D) the effects of salinity and alkaline stress at ambient light conditions. Biophysical parameters were determined with "PEA Plus" software version 1.12 (http://www.hansa-tech-instruments.com). The graphs were drawn using Microsoft Excel version 2016 (https://www.microsoft.com).

Table 3. ANOVA results of different light spectra and stress on JIP-test parameters in strawberry cv. Paros. *, **, ***Significance at the 0.05, 0.01, and 0.001 probability levels, respectively, ns, not significant. SAS software version 9.4 was used for data analysis (https://www.sas.com/en_us/home.html).
According to ANOVA results (Table 3), stress, light, and their interaction effects changed the JIP-test parameters significantly. Salinity and alkalinity stress induced an increase in Fv, and decreased Fv'/f, and as a result, a decrease in Fv and area. In salt-treated plants, blue light caused a significant rise in Fv'/Fv and area parameters. Red light had the most significant effect on reducing the Fv parameter. Under alkaline stress, red light had the most significant effect on increasing Fv'/Fv and area parameters. The performance indexes (PlABS and PlTotal) were significantly affected by treatments. Both stress treatments caused a significant decrease in these parameters and showed that performance index parameters are susceptible to stress conditions. Under salinity stress, red and white/yellow light had the greatest effect on PlABS and PlTotal parameters, respectively. Under alkaline stress, using red and blue/red light spectra significantly increased these parameters compared to without supplementary light treatment. Salinity and alkalinity stress significantly reduced Quantum yield parameters (φoPo, φoRO, ΨEo, δR). The blue, red, and their combination spectra reduced the effects of stress and increased the quantum performance parameters compared to the without supplementary light treatment. Both stress treatments affected reaction centers, and ABS/RC, D1/RC, and TRo/RC parameters increased. ETo/RC and especially REo/RC decreased under stress conditions, and salinity stress had a more significant effect on these parameters. Under salinity stress, blue light spectra had a significant effect on the REo/RC parameter. Under alkaline stress, blue/red light spectra had the greatest effect on ETo/RC and REo/RC parameters. The Vf, dVf/dt, and dVf/dt parameters increased in the plants treated with NaCl and NaHCO3. Different light spectra reduced the effects of stress on these parameters compared to treatment without complementary light (Fig. 6).

**Correlation analysis.** Based on the statistical analysis, it was observed that there is a significant correlation between some chlorophyll and fluorescence and the CO2 assimilation parameter (A). Figure 7 shows the correlations between the CO2 assimilation parameter (A) and the fluorescence chlorophyll parameters (ABS/RC, TRo/RC, Eto/RC, REo/RC, PlABS, and PlTotal). The CO2 assimilation parameter (A) correlated positively with Eto/RC, REo/RC, PlABS, and PlTotal.

**Principal component analysis.** First, the data were standardized to zero mean and unit variance. PCA was performed to summarize the variations of 15 parameters during five-light spectra treatments at the control treatment and the two stress levels, separately. In the control treatment, the performed PCA explained 97/54% of total variations of five light spectra treatments (Fig. 8A). This value was 82.08 and 82.79% for salinity and alkalinity treatments, respectively (Fig. 6B,C). Most of the variations were explained by the first component. Thus, the effect of five light spectra treatments on the variations of the 15 parameters was linear. Based on two principal components, the 15 evaluated parameters at the control treatment were divided into three groups, and the salinity and alkalinity stress treatments into five and four separate groups, respectively (Fig. 8). Regardless of the direction of the effect, the parameters p11, p12, and p15, respectively, had the least contribution to the first principal component of the variations caused by five light spectra treatments at the control treatment. These parameters were p15, p1, p11, p3, and p2 at salinity stress treatment, and alkalinity stress treatment were p15, p12, p1, p2, p3, and p11. The other parameters have the most contribution to the first principal component in control and two stress treatments. Among these parameters, at the control treatment, parameter p13, and two stress treatments, parameter p10 had the most contribution in the first principal component. Thus, in studying the effect of light spectra treatments on various parameters, it is necessary to pay special attention to these two parameters.

**Discussion**

The use of LED as a light source is important in terms of improving plant productivity because not only light intensity but also its spectral composition affects plant life processes. Therefore, in this work, we compared four different light spectra in terms of photosynthesis performance and chlorophyll fluorescence parameters of strawberry cv. Paros under salinity and alkalinity stress conditions.

Inhibition of photosynthesis in saline and alkaline stresses on different plant species was investigated27,28. In this experiment, we investigate the effect of complementary light spectra on increasing the tolerance of strawberry plants to salinity and alkalinity stress. The oxygen-evolving complex (OEC) was described by Schreiber et al.29 as one of the most sensitive components of the electron transport chain of the photosynthetic apparatus. Usually, its decreased efficiency is caused by an electron transport disorder. Our results showed that under salinity and alkalinity stress, the absorption and trapping of electrons decreased. Based on these results, the amount of ABS/RC and TRO/RC increased in both stresses, which is due to the loss of active reaction centers. Under salinity conditions, due to the dissociation of LHCl and PSII, the efficiency of trapping electrons in the PSII reaction center decreases30. Based on this knowledge, there is known that photosynthesis is restrained by both stresses. A study of alkalinity and salinity stress showed an additional impact of alkalinity stress due to increased pH. The high pH condition around the roots will precipitate metal ions, P, and Mg precipitation is caused by alkalinity stress, leading to inhibition of chlorophyll synthesis31. Photosystem II (PSII) has also been shown to play a major role in the reaction of photosynthesis to stress conditions32,33, which was confirmed by these studies. Our results showed that the maximum quantum yield of primary PSII photochemistry (φPo) decreased under the influence of both stresses. In tomato seedlings, PSII electron transfer is blocked by salinity-alkalinity stress34. PSII activity showed that the maximum quantum yield of primary PSII photochemistry (φPo) decreased under the influence of both stresses. The use of LED as a light source is important in terms of improving plant productivity because not only light intensity but also its spectral composition affects plant life processes. Therefore, in this work, we compared four different light spectra in terms of photosynthesis performance and chlorophyll fluorescence parameters of strawberry cv. Paros under salinity and alkalinity stress conditions.
Figure 7. Correlation plot between the plant gas exchange (A, WUEi) and prompt fluorescence parameters. (A) In control conditions (non-stressed plants); (B) under salt stress; (C) under alkalinity stress. The size and color intensity of circles are proportional to Pearson's correlation coefficient at p < 0.01. Red circles indicate positive correlations, while blue are negative correlations. In the correlogram scale from −1 to +1, Pearson's correlation coefficient for variables is on the vertical and horizontal axis. Asterisk indicates values that are statistically different at p < 0.05. The correlation plot was drawn with Origin Pro software version 2021 (https://www.originlab.com/2021).
Blue light is mainly used in commercial research and horticulture because it is one of the most important spectra in the process of photosynthesis. At the same time, the lack of one of them (red or blue light) reduces the efficiency of photosynthesis. The stress of salinity and alkalinity affects the amplitude of both the F₀ and Fₐ parameters. Fₐ increases and Fₐ decreases, indicating no electron transfer from P₆₈₀ to Q⁻ and energy dissipation of PSII excited states. The results showed that both salinity and alkalinity stress increased the energy dissipation of PSII, but salinity stress had a greater effect on energy dissipation. The rise in the initial level of fluorescence (F₀) observed could be attributed to the increase in the slope of the initial resulting from the K-step. Our results (Fig. 4) showed that alkalinity stress caused a significant rise in Fₐ compared to salinity stress with an increasing K band. A significant delay of Fₐ can indicate inhibition of electron flow by PSII. It can result from non-photochemical quenching, D₁ protein degradation, or PSII RC inactivation. In salt-stressed wheat leaves, PIABS decreased due to ionic and osmotic stress, and the decline of PIABS was associated with a decrease in (FM – FJ)/FV. D1 protein degradation, or PSII RC inactivation in salt-stressed wheat leaves, PIABS decreased due to ionic and osmotic stress, and the decline of PIABS was associated with a decrease in (FM – FJ)/FV. Inactivation or inhibition of electron transfer on the donor or acceptor side of PSII by the inactivation of ferredoxin NADP⁺-reductase was shown by a lower δRo level. Alkalinity stress in strawberry plants has been reported to decrease PI and Sm, similar results have been reported under mixed salinity-alkaline stress conditions by Deng et al. The total number of photons absorbed from all reaction centers is expressed by chlorophyll molecules divided by the total number of active reaction centers by ABS/RC. It is affected by the ratio of active/inactive RCs, and the ABS/RC ratio also increased as the number of inactive centers increased, which decreases the transport of electrons in active RC (ETₒ/RC) and reduces the final acceptor in PSI (REₒ/RC). Plants were grown in blue and blue/red light, electron transport flux per reaction center, and the probability that the trapped exciton would transfer the electron in the electron transport system beyond QA⁻ was increased following the stress condition (Fig. 8). This indicates that plants grown in blue and blue/red light were more capable of carrying electrons from absorbed photons through the electron transport chain and beyond QA⁻. This indicated that the energy level in reaction centers was positively regulated by plants grown in blue and blue/red light following stress exposure.

Figure 8. Principle component biplot of JIP test parameters based on variations of 5 levels of light spectra under (A) control (non-stress); (B) salinity stress; (C) alkalinity stress. P1: ABS/RC; P2: Dio/RC; P3: TRo/RC; P4: Eto/RC; P5: REo/RC; P6: ϕP₂/Θ; P7: VEₒ/RC; P8: ϕEₒ/RC; P9: δRo; P10: ϕRo; P11: PIABS; P12: PIₙₚₚ; P13: Vₒ; P14: Vᵢ; P15: Sm. Principal component analysis was performed using MATLAB software version R2015b (https://www.mathworks.com).

According to studies, changes in the spectrum of light have a significant impact on the physiological processes of the strawberry plant. These conclusions were confirmed by Hogewoning et al. or Macedo et al. We also found that under salinity and alkalinity stress, blue and blue/red light had a positive effect on CO₂ assimilation. Salinity and alkalinity stress reduced water use efficiency (WUE) compared to the control. The blue/red light had a significant effect on WUE, and increased it in all stress conditions. For chlorophyll synthesis, blue light is essential, but red light is also involved in this process. Plants absorb blue and red light, mostly (about 90%). Blue light leads to an increase in stomata dilation and, consequently, to an increase in transpiration, and red light inhibits stomata dilation, which leads to a reduction in transpiration. The response of stomatal guard cells to a decrease in intercellular CO₂ concentration and the direct reaction of guard cell chloroplasts to red light leads to red light-induced stomatal opening. Blue light is mainly used in commercial research and horticulture because it is one of the most important spectra in the process of photosynthesis. At the same time, the lack of one of them (red or blue light) reduces the efficiency of photosynthesis.

Photosynthesis is impaired by salinity by stomatal closure, photochemical reaction destruction, and carbon assimilation. The stomatal closure is the first defense of the plant against salinity–alkalinity stress. Due to the increased resistance of diffusion in the pores and through mesopholic cells, the supply of CO₂ in stressful environments decreases. The reduction of CO₂ disrupts the electron transfer chain. In the isolated epidermis, it is
accepted that red light causes stomatal opening. This reaction depends on the chloroplasts of the guard cells.

Both C_{i} and guard cell chloroplasts play a role in the synergistic effect of blue and red light on stomatal opening. The blue-light reaction can be indirectly induced by Mesophyll photosynthesis. For blue-light signaling in guard cells, Ca^{2+} was indicated to be responsible. Both blue-light-dependent H^{+} pumping and stomatal opening were inhibited by calmodulin antagonists.

A good correlation was observed between the increased capacity to assimilate CO_{2} and the increased value of P_{I_{abs}}, P_{I_{tot}}, E_{to}/RC, and R_{Eo}/RC. The finding provided evidence of the correlation of OJIP fluorescence variations with changes in general photosynthesis capacity during stress in field or laboratory conditions. The coordinated control of the entire photosynthesis system takes place in such a way as to maintain an internal balance between the efficiency of the photosynthetic light phase reaction and the efficiency of the reactions leading to CO_{2} assimilation.

**Conclusions**

From our studies, it can be concluded that the photosynthesis apparatus of strawberry plants are sensitive to both types of stressors. The influences of the adverse conditions could be mitigated by proper choice of lighting (quality). Under salt and alkalinity stress, blue/red and blue light had a significant effect on the assimilation of CO_{2} and photosynthetic electron transfer chains. Using blue and blue/red light as complementary light in the greenhouse can increase the tolerance of plants to salinity and alkalinity stresses. Chlorophyll fluorescence measurements were a reliable tool for monitoring and early detection of the changes caused by the stressor. Previous studies have mainly focused on the effect of light intensity on plant growth and development. However, there is limited information on the effects of different light spectra on plant photosynthetic efficiency and plant gas exchange under stress conditions. It is suggested that this research be done other plants and cultivars and in different spectra and intensities of light under stress conditions. This research will be helpful for improving the growth and development of plants in greenhouses and under stress conditions. In greenhouse crops and under abiotic stresses, it is possible to enhance the growth and development of plants by applying complementary light spectra.

**Data availability**

All data generated or analyzed during this study are included in this published article (and its supplementary information files).

Received: 22 March 2022; Accepted: 25 July 2022
Published online: 02 August 2022

**References**

1. Carvalho, R. E., Campos, M. L. & Azevedo, R. A. The role of phytochrome in stress tolerance. *J. Integ. Plant Biol.* 53, 920–992 (2011).
2. Manivannan, A., Soundararajan, P., Halimah, N., Ko, C. H. & Jeong, B. R. Blue LED light enhances growth, phytochemical contents, and antioxidant enzyme activities of Rehmannia glutinosa cultured in vitro. *Hortic. Environ. Biotechnol.* 56, 105–113 (2015).
3. Johkan, M., Shoji, K., Goto, F., Hashida, S. & Yoshitaka, T. Blue light-emitting diode light irradiation of seedlings improves seedling quality and growth after transplanting in red leaf lettuce. *HortScience* 45, 1809–1814 (2010).
4. Fan, X. et al. Effects of different light quality on growth, chlorophyll concentration and chlorophyll biosynthesis precursors of non-heading Chinese cabbage (*Brassica campestris* L.). *Acta Physiol. Plant.* 35, 2721–2726 (2013).
5. Assmann, S. M. Signal transduction in guard cells. *Annu. Rev. Cell Biol.* 9, 345–375 (1993).
6. Allakhverdiev, S. I. & Murata, N. Environmental stress inhibits the synthesis of novel proteins involved in the photodamage-repair cycle of photosystem II in *Synechocystis* sp. PCC 6803. *Biochim. Biophys. Acta-Biologener.* 1657, 23–32 (2004).
7. Kalaji, H. M. et al. Frequently asked questions about in vivo chlorophyll fluorescence: Practical issues. *Photosynth. Res.* 122, 121–158 (2014).
8. Dąbrowski, P. et al. Exploration of chlorophyll a fluorescence and plant gas exchange parameters as indicators of drought tolerance in perennial ryegrass. *Sensors* 19, 2736 (2019).
9. Kalaji, H. M. et al. Chlorophyll fluorescence as a tool to monitor physiological status of plants under abiotic stress conditions. *Acta Physiol. Plant.* 38, 102 (2016).
10. Rapacz, M., Sasal, M., Kalaji, H. M. & Kościelniak, J. Is the OJIP test a reliable indicator of winter hardiness and freezing tolerance of common wheat and triticale under variable winter environments? *PLoS ONE* 10, e0134820 (2015).
11. Shabalala, S. Learning from halophytes: physiological basis and strategies to improve abiotic stress tolerance in crops. *Ann. Bot.* 112, 1209–1221 (2013).
12. Li, J. et al. Soil salinization research in China: Advances and prospects. *J. Geogr. Sci.* 24, 943–960 (2014).
13. Fang, S., Hou, X. & Liang, X. Response mechanisms of plants under saline–alkali stress. *Front. Plant Sci.* 12, 545 (2021).
14. Strasser, R. J., Tsimilli-Michael, M., Qi, S. & Goltsev, V. Simultaneous in vivo chlorophyll fluorescence and prompt delayed fluorescence and 820-nm reflection changes during drying and after rehydration of the resurrection plant *Haberlea rhodopensis*. *Biochim. Biophys. Acta-Biologener.* 1797, 1313–1326 (2010).
15. Mehta, P., Jayoo, A., Mathur, S. & Bharti, S. Chlorophyll a fluorescence study revealing effects of high salt stress on photosystem II in wheat leaves. *Plant Physiol.* 118, 16–20 (2010).
16. Sayed, O. H. Chlorophyll fluorescence—A practical guide. *J. Exp. Bot.* 51, 659–668 (2000).
17. Ashraf, M. Some important physiological selection criteria for salt tolerance in plants. *Flora-Morphol. Distrib. Funct. Ecol. Plants* 199, 361–376 (2004).
18. Raza, S. H., Athar, H. R., Ashraf, M. & Hameed, A. Glycinebetaine-induced modulation of antioxidant enzymes activities and ion accumulation in two wheat cultivars differing in salt tolerance. *Environ. Exp. Bot.* 60, 368–376 (2007).
19. Malekzadeh-Shamsabad, M. R., Roosta, H. R. & Esmailizadeh, M. Responses of seven strawberry cultivars to alkalinity stress under soilless culture system. *J. Plant Nutr.* 44, 168–180 (2021).
20. Yaghubi, K., Ghaderi, N., Vafaee, Y. & Javadi, T. Potassium silicate alleviates deleterious effects of salinity on two strawberry cultivars grown under soilless pot culture. *Sci. Hortic.* (Amsterdam) 213, 87–95 (2016).
Van Heerden, P. D. R., Swanepoel, J. W. & Kruger, G. H. J. Modulation of photosynthesis by drought in two desert scrub species

Scientific Reports | (2022) 12:13257 | https://doi.org/10.1038/s41598-022-17377-8

The authors declare no competing interests.
