Melatonin induced changes in photosynthetic efficiency as probed by OJIP associated with improved chromium stress tolerance in canola (Brassica napus L.)

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ABSTRACT

Chromium toxicity is considered as a major problem for agricultural soil that reduced crop productivity by affecting photosynthetic tissues. Exogenous application of melatonin can alleviate the adverse effects of chromium toxicity on plant growth. However, little is known about its effect on thylakoidal protein complexes responsible for conversion of solar energy to biochemical energy. Chlorophyll fluorescence transients considered one of the best non-invasive and rapid method for the evaluation of photosynthetic (Photosystem II) efficiency of plants and plant health under environmental stress conditions. In the present study, three-week old plants of two canola cultivars AC-Excel and DGL were applied to melatonin (0, 1, 5, 10 μM) when grown under chromium stress (0, 50 and 100 μM) for further two weeks. Chromium stress reduced the growth (fresh and dry weights of shoots and roots) of both canola cultivars and exogenous application of 5 and 10 μM melatonin improved the growth of canola at 50 or 100 μM chromium stress. This improvement was greater in cv DGL than in AC-Excel. Increasing chromium decreased the photosynthetic pigments (chlorophyll a and chlorophyll b). However, 5 and 10 μM melatonin application improved chlorophyll a at 50 μM chromium stress. This improvement was greater in cv DGL than in AC-Excel. Increasing chromium decreased the photosynthetic pigments (chlorophyll a and chlorophyll b). However, 5 and 10 μM melatonin application improved chlorophyll a at 50 μM chromium stress. Structural stability and efficiency of photosystem II (PSII) measured as performance index (PIABS) and ratios of fluorescence (Fv/Fm, Fv/Fo) Fv decreased due to chromium stress. JIP-test parameters showed that chromium stress increased the absorption and trapping fluxes with decrease in electron transport fluxes which caused the damage to reaction centers (RC), detachment of oxygen evolving complex (OEC) from RC or inefficiency of electron transfer from OEC to RC. Such adverse effects were greater in cv AC-Excel. However exogenous application of melatonin improved PIABS, electron transport per reaction center (ET/RC), reduced variable fluorescence at J step (VJ) reflecting melatonin protected PSII from chromium stress induced damage by protecting OEC. Thus, OJIP fluorescence transients are quite helpful for understanding the intersystem electron transport beyond photosystem II in canola cultivars due to melatonin application under chromium stress. Findings: Exogenous application of melatonin alleviated toxic effects of chromium on plant growth of canola by modulating photosynthesis, enhanced photosystem II efficiency and regulation of electron transport flux to protect photo-inhibition of PSII from oxidative damage.

1. Introduction

Chromium is one of the heavy metals that are naturally occurring trace elements of soil and over accumulation of these trace elements in agricultural soil become major threat to food safety in several parts of the world (Ifon et al., 2019; Farooq et al., 2018). Excessive use of fertilizers and pesticides, leather tanning and mining causing high level of chromium contents in agricultural soils (Kotecha et al., 2019; Ulhassan et al., 2019).

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It has been investigated that chromium affects the chloroplast mainly in the form of poor lamellar, fewer grana and enhanced thylakoid lumen (Ali et al., 2013). Plants facing environmental abiotic stress conditions stimulates the formation of reactive oxygen species (ROS) in chloroplast such as photorespiration, cyclic electron flows through either PSI or PSII consequences down-regulation of PSII quantum yield regulated by xanthophyll cycle and proton gradient across thylakoid membrane (Gill et al., 2016). Photosystem II (PSII) is considered to be very sensitive and has great inhibitory effect under stress conditions (Souri et al., 2019), by inhibiting the interaction between electron donor and acceptor end photosystem II (PSII) and induces some changes in I–P normalized bands which are linked with electric trans-thylakoids potential produced by protons pump regulated by cyclic electron transport in PSI during electron transfer from PQ or PQH2 to PSI electron acceptor end i.e Fd and NADP.

At PSII chromium ions replaces the co-factor Ca2+ known to be very important for water-splitting, hence alters the structure and function of oxygen evolving complex. In addition to oxygen evolving complex chromium ions interacts with many essential electron acceptor proteins i.e Qb found in electron transport of PSI (Oves et al., 2016). Chromium also effects the efficiency of photosystem I (PSI) by interacting with monomeric and multimeric sub-units, therefore disturbs the overall pathway of electron in electron transport chain that ultimately lowers the energy conversion efficiency of PSI (Gill et al., 2015).

Photosystem II requires a certain amount of energy for the reaction center and continuity of electron transfer Qb to PSI. The energy requirements of PSII for electron flow in electron transport chain has been analyzed by observing the apparent activation energy change by fast chlorophyll rise. Apparent change in activation energy of thylakoid membrane can be observed by using O-J-I-P normalized bands which is linked with trans-thylakoids potential produced by protons pump regulated by cyclic electron transport in PSI during electron transfer from PQ or PQH2 to PSI electron acceptor end i.e Fd and NADP.

2. Methodology

A pot experiment was performed in wirenet house of Bahauddin Zakariya University, Multan, Pakistan with normal environmental conditions (30°N and 71°28E). Seeds of canola cultivars (DGL, AC Excel) were obtained from Ayub Agriculture Research Institute (AARI) Faisalabad. River washed sand was used as a rooting medium. In experiment 120 plastic pots with diameter 28 cm with 8 kg sand were used, five to seven seeds were sown in each pot. After germination of the plants thinning was carried out leaving 4 equally distant placed plants in each pot. After twenty days of germination, plants were treated by various levels of Cr (0, 50, 100 μM) with Hoagland nutrient solution (full strength). After 10 days of chromium treatment foliar application of melatonin with different concentration (0, 1, 5, 10 μM) mentioned as MTO, MT1, MT2, MT3 were applied to the plants. Effective chromium and melatonin concentrations were selected in preliminary experiment, where we found that 5, 10 μM of melatonin showed significant tolerance effect to plants under chromium stress. Experiment was designed according to CRBD (completely randomized block design) with three Cr levels and four melatonin levels, two cultivars and five replicates for each treatment.

After twenty days of treatment, plant fast chlorophyll a kinetic analysis was measured with FluorPen FP-100 (Photon System International, Czech Republic). Plants were harvested and separated into shoots and roots. Plant fresh biomass (root and shoot) were measured and then oven-dried at 70 °C for 48 h, then their dry weights were recorded.

2.1. Chlorophyll contents

For the estimation of chlorophyll contents of canola plants 0.2 g leaf tissue was homogenized in 80% acetone in pestle and mortar. Leaf extract volume after filtration was kept 10 ml in tubes. Absorbance were measured at different wave lengths i.e., 700, 663, 652, 645, and 470 nm by using double beam spectrophotometer (U-2900/2910 Hitachi) and calculate the chlorophyll and carotenoid contents using formulae as mentioned Arnon (1949).

2.2. Analysis of O-J-I-P fast chlorophyll a transients

Young and fully matured leaves (usually third leaf from the top) of canola plants were dark adapted for 20 min by wrapping aluminum foil and then exposed to saturating light pulse of 3000 μmol m–2 s–1 light intensity with 0.8 s duration over 4 mm leaf area using hand-held device FluorPen FP-100 (Photon System International, Czech Republic). Various basic fluorescence parameters (F0, Fv, Fm, Fv/Fm), relative maximum variable fluorescence (V1/Vm), ratios of fluorescence (Fv/Fm, Fv/ Fo) and other JIP-test parameters were calculated following Strasser et al. (2000). Raw fluorescence curves were normalized by Fo and double normalized by Fo and Fm of control and then plotted on logarithmic scale. Moreover, difference in double normalized curves of treated and control plants were calculated and then plotted on logarithmic scale to assess specific changes in PSII activity and electron transport flux.

The data obtained were statistically analyzed using three-way ANOVA with the help of statistical computer package CoStat 6.3 (Cohort, California USA). The percent of control of each JIP-test parameter was calculated and presented on one scale as radar plot to assess changes in different attributes due to chromium toxicity and changes due to melatonin.

3. Results

Chromium toxicity causes reduction (P < 0.001) in biomass (fresh and dry weight g/plant) (Figure 1), leaf number and quantum yield (Figure 2) of both the canola cultivars. While melatonin application improved the plant growth (fresh and dry biomass of shoots and roots, leaf number) and quantum yield of PSI in both canola cultivars. This improving effect of melatonin was more in 5μM melatonin treated AC-Excel plants.

Overall chlorophyll contents were significantly affected by chromium stress (P < 0.001) including chlorophyll a, chlorophyll b, chlorophyll a/b
and total chlorophyll of canola plants (Figure 3). While exogenous application of melatonin improved chlorophyll contents mainly 5 and 10 μM concentration significantly increased chlorophyll contents in chromium treated and non-treated plants. Significant increase in plant biomass and chlorophyll contents in AC-Excel shows more resistance as compared to DGL in chromium stress.

OJIP chlorophyll fluorescence transients were evaluated from dark adapted leaves of canola and were plotted on logarithmic scale. In chlorophyll fluorescence transient's some changes were observed due chromium and melatonin treatment in canola. Ok phase describes the uncoupling of oxygen evolving complex. ΔV_{OK} was high at 5μM melatonin level then 1 and 10 μM treatment. OJ is the light dependent phase which describes the information about the connection between antenna complex size and reaction center of PSII. Under chromium stress ΔV_{OJ} increases but more pronounced on 5 μM melatonin level as compared to the 1 and 10 μM melatonin level. On 1 and 10 μM melatonin plants showed little reduction in ΔV_{OJ} in control conditions that shows the plants are able to maintain the antenna complex of PSII and reaction center. OI phase describes the kinetic properties of reduction/oxidation of the PQ (Plastoquinone) pool. IP phase describes the electron flow from PQ to the PSI.

Data of all the parameters were recorded by normalized with their reference control and each variable with its reference control was standardized by fixing the value up to 100 (Tables 1 and 2). Chromium stress causes reduction in variable fluorescence at J step (VJ) in both canola cultivars. Foliar application of melatonin did not affect the variable fluorescence VJ in both canola cultivars. Whereas chromium stress and melatonin application did not affect the relative variable fluorescence at I step (VI) in canola plants shown in (Figures 4 and 5).

Among chlorophyll fluorescence ratios Fv/Fm explained that maximum primary yield of photochemistry of PSII did not show any change in presence of chromium stress and melatonin treatment in both canola cultivars, while Fv/Fl and Fm/Fl that are considered as most sensitive parameters of plant photosynthetic capacity were noted to be reduced due to chromium toxicity in both canola cultivars meanwhile exogenously applied melatonin treatment increased both of these parameters values in both cultivars. However, improved values of Fv/Fl and Fm/Fl because of melatonin treatment was observed more in AC-Excel under chromium treated that of non-treated plants. While increased values of Mo (primary photochemistry values) in AC-Excel cultivar was observed to be improved in foliar application of melatonin that of chromium treated plants of canola shown in (Figure 6). Similarly, total Area (PQ pool), redox state of multiple PQ turnover (Sm) and Q_{A} redox turnover until Fm actually (N values) was observed to be decreased in DGL only and melatonin application did not significantly affect the biophysical parameters.

Melatonin was applied to recover chromium toxicity damage on photosynthetic machinery of canola plants. OJIP transient curves were studied to observe melatonin induced modulation and repairing of structural and functional properties of PSII. Drop in transient curves at various points O-J, J-I and I-P occurs, indicates the toxic effect of chromium causing reduction in electron acceptor pool of PSII. Transient
curves were intense in melatonin treated plants under chromium stress that inhibits the electrons flow on donor side of PSII.

Chromium stress causes deficiency of primary photochemistry; decrease in Area over fluorescence curve, ultimately reduction in quantum yield of PSII (because of high F0, less and smaller active reaction centers (Fv/F0) reduction in electron flow beyond Quinone A proteins Qa), that causes electron deficiency at PSI (photosystem I). That causes reduction in electron flow in electron transport chain and acceptor site of PSII (Pool size of QA). Poor electron flow from PSII to PSI and low diffusion of PQ across the thylakoid membrane suggests the reduction of re-oxidation of Quinone A from QA to QA. Higher F0/Fm values in chromium treatment indicates high reduction rate of QA and re-oxidation of QB as well as photosystem I. To prevent photo-oxidative damage excess excited energy dissipation occurs in the form of thermal energy to maintain thermal balance for energy absorption and consumption, increased energy dissipation and dissipated energy flux (D1o/RC) values were observed in chromium treated plants.

Decreased values of Fm/Fo indicates structural damage of PSII in melatonin treated plants against chromium stress, while high Sm values (electron pool from PSII to PSI) because of chromium results in altered QA electron transfer efficiency and PQ activity. Decrease in overall performance index PIABS plants exposed to chromium results inactivity of reaction centers (RCs) enabling less restoration of total heat, ultimately reduction in size of antenna complex RC (ABS/RC). More reduced reaction centers favors high number of active reaction centers (ABS/RC) in chromium treated plants to enhance the more reaction centers turnover number for reduction of PQ pool. High transmission of electrons of reaction centers (TRo/RC) values shows less electron transfer from active reaction centers donating electron transfer from reduced QA to Qb reduces (Qa to QA) re-oxidation because of high electron abundance. Chromium treated plants showed increased ratios of energy dissipation of un-trapped excitation energy from all reaction centers that of increased open reaction centers. High Fo/Fv values showed effect of chromium on nutrients uptake efficiency i.e manganese (Mn) major component of OEC its deficiency results in structural changes of oxygen evolving complex (OEC).

Chromium treated plants showed increased values of Fm/Fo and Fv/Fo and decreased Fo/Fv suggesting melatonin induced high Fv causing decline of Fo, that results in modulation of acceptor site of PSII. Increased values of Fv/Fo indicates significant improvement of electron transport chain (ETC), Melatonin treatment improves Fm values suggests high Mn ion as an extrinsic proteins of oxygen evolving complex (OEC). High manganese ion leads to low electrons transfer from water to PSII causing structural changes in D1 protein that induces structural and functional changes of PSII. Melatonin effectively improves the electron transfer rate from oxygen evolving complex to D1 protein related to core reaction centers of PSII including psb A. Chromium stress causes decrease in ABS/RC, TRo/RC, ETo/RC and D1o/RC enabling photosystem II to tackle energy flux and balance of absorbed and trapped energy transfer from active reaction centers of photosystem II. However, melatonin improves overall photosystem II efficiency in terms of increased PIABS values supported by increased ‘Area’ in chromium stress.

In order to confirm the observed differences of semi quantitative evaluation was performed using differences of suitably normalized
transients from chromium treated and non-treated plants expressed in terms of OP, L and K bands. The kinetics difference of Melatonin treated canola plants in chromium treated and non-treated resulted negative bands of O-J and O-I region because of photochemical quenching of OJIP transients indicating excited electron transfer more faster to reduce PQ in respective control plants.

Canola plants cultivar DGL as compared to AC-Excel amplitude of kinetics difference of melatonin treated plants was greater in chromium transients for further two weeks.

Figure 3. Chlorophyll fluorescence with and without normalization of two cultivars of canola (Brassica napus L.) whereas (A), (B), (C), (D) represents DGL with melatonin (0,1,5,10 μM) and chromium (0,50,100 μM) while (E), (F), (G), (H) represents AC-Excel with melatonin (0,1,5,10 μM) and chromium (0,50,100 μM) respectively for further two weeks.

| SOV         | Df | Vj     | Vi     | Mo    | Fv/Fo | Fv/Fm | Phi_Po | Psi_O |
|-------------|----|--------|--------|-------|-------|-------|--------|-------|
| Cr          | 2  | 0.002* | 0.005* | 0.058* | 0.479** | 4.087** | 4.087* |
| MT          | 3  | 0.008*** | 0.01ns | 0.003ns | 0.702* | 0.001* | 0.001ns |
| Cr × Mt     | 6  | 0.003* | 3.53ns | 0.017ns | 0.562* | 0.001* | 0.001* |
| cultivars   | 2  | 117*   | 138ns  | 188ns  | 0.24*  | 1.023*  | 1.05ns  |
| Error       | 36 | 9.986  | 0.001  | 0.016  | 0.487  | 8.860  | 8.860  |
| Total       | 47 |        |        |        |        |        |        |

*, **, *** = significant at 0.05, 0.01 and 0.001 levels, respectively, ns = Non-significant.
treated plants (Figures 4 and 5). To understand the possible changes of chlorophyll fluorescence at each step of kinetics difference, i.e., L band that is energetic connectivity among PSII subunits, K band that is OEC activity were calculated and represented as $\Delta V_{OK}$, $\Delta V_{OJ}$ respectively. More negative amplitude of L-bands of melatonin treated plants in chromium stress conditions indicates greater energetic connectivity of PSII subunits that of control plants. Whereas L and K ($\Delta V_{OJ}$) bands in cultivar DGL 5μM concentration was higher than AC-Excel plants in Figure 4. Chlorophyll fluorescence with double normalization of two cultivars of canola (Brassica napus L.) whereas (A), (B), (C), (D) represents DGL with melatonin (0, 1, 5, 10 μM) and chromium (0, 50, 100 μM) while (E), (F), (G), (H) represents AC-Excel with melatonin (0, 1, 5, 10 μM) and chromium (0, 50, 100 μM) respectively for further two weeks.

Table 2. Analysis of variance (ANOVA) of OJIP transient parameters.

| SOV     | Df | Phi_Eo | Phi_Do | PI_ABS | ABS_RC | TRo_RC | ETo_RC | DIO_RC |
|---------|----|--------|--------|--------|--------|--------|--------|--------|
| Cr      | 2  | 8.135* | 4.087ns| 3.457ns| 0.713**| 0.450***| 0.197***| 0.030**|
| MT      | 3  | 0.007***| 0.001* | 0.895ns| 0.117***| 0.086*  | 0.085***| 0.010* |
| Cr × Mt | 6  | 0.002* | 0.001* | 1.201ns| 0.027** | 0.017** | 0.005** | 0.005**|
| cultivars| 2  | 1.002* | 1.23ns | 0.98   | 0.21*  | 0.95** | 0.08*  | 0.02*  |
| Error   | 36 | 0.001  | 8.860  | 1.320  | 0.064  | 0.040  | 0.007  | 0.006  |
| Total   | 47 |        |        |        |        |        |        |        |

*, **, *** = significant at 0.05, 0.01 and 0.001 levels, respectively, ns = Non-significant.
chromium treated and non-treated plants. Both the cultivars differ in amplitude of negative K band because of melatonin treatment can be observed in control conditions, where AC-Excel cultivar showed lower amplitude of L-band. Whereas, changes in I–P band can be measured by two normalized methods which are linked with electric trans-thylakoids potential produced by protons pump regulated by cyclic electron transport in PSI during electron transfer from PQ or PQH2 to PSI electron acceptor end i.e Fd and NADP.

Foliar application of melatonin increased the amplitude of I–P curve in both canola cultivars in control conditions. Suggesting melatonin might acts vital role by enhancing electron pool (carrier of photosystem I site), reduced from upcoming electron from PQ pool in canola plants. Whereas I–P band measured as VIP = [(Ft−FI)/(Fm−FI)] indicates chromium stress reduction rate of constant value of AC-Excel melatonin enhances the rate of constant value in AC-Excel cultivar that of DGL. Chromium stress induced biophysical changes derived from chlorophyll fluorescence curve of canola cultivars explained as in radar plot as shown in (Figure 6).

The derived fluxes of specific energy including ABS/RC (absorbance flux per reaction center), ETo/RC (trapped energy flux per reaction center), ETo/RC (electron transport flux per reaction center) and Dlo/RC dissipation energy flux per reaction center all of these fully reduced in chromium stressed plants of both cultivars and melatonin treatment only improved all these OJIP parameters in cultivar AC-Excel cultivar then DGL (Figure 6) this rapid electron transfer (reduction) rate becomes faster due to chromium toxicity, because of inactive reaction centers that of control plants. In fact the relationship between P_{ABS} and relative electron transfer yield are considered as plants capacity to transfer light energy into NADH (chemical energy) that is utilized subsequently into metabolisms i.e photosynthesis (Gómez et al., 2019). In this regard our results suggested that exogenously applied melatonin under chromium stress have higher capacity to convert light energy to chemical energy which can be used to further CO₂ to carbohydrates. Conversion of light energy into chemical energy was observed to be higher in cultivar AC-Excel then DGL (Figure 6).

4. Discussion

Despite the significant efforts and advancements in abiotic stress research, soil metal toxicity remains one of the most detrimental factors in agriculture (Farooq et al., 2016). In the present study we examined the chlorophyll fluorescence measurements followed by OJIP test corresponded well with melatonin induced enhanced PSII activity under chromium stress. This relationship, at least in chromium stress may be in direct damage of PSII reaction centers or other elements of photosynthetic electron transport chain. In the present study chromium reduced the number of active reaction center, while further steps for photosynthetic electron transport chain were less affected. While, P_{total} of both stress and control conditions were clearly discriminated for canola plants but in control conditions in a way similar to quantum yield of reduction of PSI electron transport chain, that means in both control and stress conditions net assimilation rate is to some extent connected with overall

![Radar plot of OJIP transient parameters (A), (B), (C), (D) represents DGL cultivar treated with (0, 1, 5, 10 μM Melatonin) and chromium (0, 50,100 μM) for two weeks.](image-url)

Figure 5.
photochemical efficiency, where in control conditions it is more closely related to activity of thylakoid electron transport chain site acceptors. Our results confirmed that chromium induced changes in OJIP parameters are very sensitive to any change in PSII activity, where melatonin efficiently can improve PSII activity in chromium stress (Farouk and Al-Amri, 2019). These parameters reflect any possible change in quantum yield (Fv/Fm) of PSII that is key indicator of plant photosynthetic ability either in control or stressful conditions (Li et al., 2019). Melatonin application in chromium stress prevents pigment degradation that helps in improving the overall photosynthetic process. Plants exposed to metals in root region causes inhibition of growth by producing reactive oxygen species that ultimately leads to plants death (Mizushima et al., 2019).

All the chlorophyll fluorescence transient data of canola plants explained the reduced of primary photochemistry fluorescence and photo electrochemical quenching at O-J and J-I step under chromium stress in both canola cultivars where melatonin application increases the photosynthetic activity by compensating reduction rate of PSI and electron acceptor site of step I-P site in both Canola cultivars especially in AC-Excel than DGL.

In addition Fo normalized transient and relative variable fluorescence transients of F0 and Fm verified our results (Figure 6), for detailed analysis whole difference of kinetics at each step from OJ-JI-IP was performed (Figures 3, 4, and 5). Low fluorescence values in I-band against chromium stress conditions showed the loss of energetic connectivity to some extent due to chromium stress. Both canola cultivars showed maximum ability of resistance at K-band showed for donor and acceptor sides of PSIIm imbalance at 1000μs in chromium stress. While, an increase in K-band peaks from 1000-2000 μs showed reduced oxygen evolving complex (OEC) performance because of electron flow imbalance from (OEC) to reaction center at acceptor site of PSII in chromium stress. However, at 1000–2000 μs decreased fluorescence curve at K-band under melatonin treatment against chromium stress (Figures 4 and 5) suggested that both the canola cultivars showed maximum resistance to electron imbalance at donor and acceptor sides of PSII.

While O-I step (describes redox properties of PQ pool) decreased/ negative fluorescence (ΔVOI) curve described the involvement of melatonin in maintaining the maximum PQ reduction rate in both cultivars against chromium stress. Meanwhile, fluorescence curve at I-P step that indicates the electron transfer rate from PQH2 to electron accepter end of PSI, melatonin treated plants showed positive increased in fluorescence transient values at I-P step in chromium stress, while decrease in chromium treated plants suggested that exogenous melatonin application enhances PQ redox rate ultimately lowering the chromium stress and increased PQ pool size in both the canola cultivars. Decreased fluorescence transient curve at I-P phase eventually happens because of sharp decline of leaf water status in chromium stress that might reaches to maximum tolerance level of pants. Chlorophyll fluorescence transients and their different ratios at each step of OJIP considered as key indicators for PSII efficiency evaluation.

Chromium stress significantly reduces Fo (minimum fluorescence level) that eventually increases energy excitation and transfer rate from
antenna complex to reaction center ultimately leads to low Fo. However, melatonin treated plants also have reduced Fo values, resulting increase in electron transfer efficiency from antenna complex to PSII reaction center. While Fm (maximal fluorescence) values were also reduced in chromium stress that explains the reduction in electron transfer to PSII acceptor site, indicating induced changes in Q<sub>A</sub> reduction rate.

Foliar application of melatonin enables the plants to maintain balance of Plastoquinone redox state by transferring electron to PSI. Melatonin treated plants showed reduction in ABS/RC values suggested, increased size of antenna complex of active reaction centers. However, PI most sensitive parameter of OJIP indicates the conformational changes and confirms the vitality canola plants of PSII. While exogenously applied melatonin in chromium stress increased the PI values and possible link between ETo/RC and log Pi was that suggests the utilization of PAR which reduces the CO<sub>2</sub> into sugars in natural environmental conditions. However, melatonin induced improvement in photosynthetic efficiency of AC-Excel cultivar that of DGL is because of its genetic potential but its effect on the exact site of photosynthetic apparatus is still unclear. In a semi-quantitative observation of melatonin treatment with and without chromium stress on different parts of photosynthetic apparatus of canola cultivars.

5. Conclusion

In current era tremendous increase in environmental toxicity because of excessive use of fertilizer and anthropogenic activities toxic metal contents have increased in agricultural soil, that especially affects agricultural crops. To overcome these kind of challenges exogenous application of hormones as melatonin used in our study with minute concentration effectively can increase the plant growth, development and tolerance against certain environmental stress. There is need to focus on exogenous application of growth enhancing agents that enables plants especially agricultural crops to increase their yield and tolerance against toxic elements. In present study chlorophyll fluorescence measurements revealed that chromium stress reduced energy trapping efficiency by damaging the oxygen evolving complex, however exogenous application of melatonin protected the oxygen evolving complex of PSII and helped out in maintaining PSII activity. Given that melatonin showed positive effect on plants and it is expected that in future melatonin could have potential role in developing photosynthetically efficient stress tolerant transgenic crops.

Declarations

Author contribution statement

Ahsan Ayyaz: Conceived and designed the experiments; Wrote the paper.
Misbah Amir, Yamma Noor, Aneela Kanwal: Performed the experiments.
Sarah Umer, Saima Gul, Muhammad Javed: Analyzed and interpreted the data.
Muhammad Iqbal, Hussan Bano, Zafar Ullah Zafar, Ayesha Khalid: Contributed reagents, materials, analysis tools or data.
Habib R Athar: Contributed reagents, materials, analysis tools or data; Wrote the paper.
Muhammad Ahsan Farooq: Performed the experiments; Wrote the paper.

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Competing interest statement

The authors declare no conflict of interest.

Additional information

No additional information is available for this paper.

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