INTRODUCTION

Sage has the oldest history among medicinal plants. This plant belongs to the family Lamiaceae. The leaves of sage have been applied in pharmaceutical, perfumery, and food industries for centuries (Altindal et al., 2016). Secondary metabolites (SMs) indirectly affect the primary processes of metabolism and directly influence the plant adaptation to environmental stresses (Mohammadi et al., 2018; Verma & Shukla, 2015). SMs significantly ameliorate plant growth and development in several environmental factors, thereby acting
as main primitive metabolites (Zandalinas et al., 2017). Water stress is the main ecological issue which leads to altered essential oil (EO) yield of Cuminum cyminum (Bettaieb et al., 2011). Also, water deficit stress (WDS) affects several physiological processes of plant including photosynthesis, respiration, transpiration, cell turgidity, stomatal conductance, and light absorption, finally resulting in reduced crop production (Hazrati et al., 2017). Photosynthetic processes in crops are largely affected by stresses, while photosynthesis is the most basic physiological process (Ashraf & Harris, 2013). There is evidence that WDS has a considerable effect on the performance of photosynthesis (Hazrati et al., 2016). The Chlorophyll (Chl) content of leaves is the important component of pigments that can straight affect the photosynthetic potential and so, initial production (Gitelson et al., 2003). Environmental stresses affect pigments and can prevent photosynthesis. Photosynthetic rate and Chl concentration change in plants under environmental stress (Ashraf & Harris, 2013). Many reports have indicated variations in the leaf Chl content under WDS. Indeed, pigments’ concentration and structure are determinatives of light absorption efficiency (Anjum et al., 2011; Porcar-Castell et al., 2014).

According to some reports, Chl concentration and carotenoid vary according to environmental conditions (Horton & Ruban, 2005). As reviewed by Müller et al. (2001), the light energy imbibed with Chl molecules can be managed in three ways: photosynthesis, heat, or vary according to environmental conditions (Horton & Ruban, 2005). According to many studies, photosynthetic efficiency of photosystem II (Fv/Fm) is the most basic physiological process (Murchie & Lawson, 2013). Environmental stresses affect pigments and can prevent photosynthesis. Photosynthetic rate and Chl concentration change in plants under environmental stress (Ashraf & Harris, 2013). Many reports have indicated variations in the leaf Chl content under WDS. Indeed, pigments’ concentration and structure are determinatives of light absorption efficiency (Anjum et al., 2011; Porcar-Castell et al., 2014).

According to some reports, Chl concentration and carotenoid vary according to environmental conditions (Horton & Ruban, 2005). As reviewed by Müller et al. (2001), the light energy imbibed with Chl molecules can be managed in three ways: photosynthesis, heat, or fluorescence. The most important chlorophyll fluorescence parameters include non-photochemical quenching (NPQ), Fv/Fm, photochemical quenching (qP), and FPSII which are applied in plant stress physiology studies (Murchie & Lawson, 2013). According to many reports, photosynthetic efficiency of photosystem II (Fv/Fm) is the most widely applied chlorophyll fluorescence measuring parameter (Hazrati et al., 2016). As reviewed by Paknejad et al. (2007), PSII parameter indicates the quantum yield of noncyclic PSII photochemistry in plants that are stressed, and Fv/Fm is a fine indicator of light inhibition in stressed plants. Flow of electrons at PSII is indicated by this parameter (Moseki & Dintwe, 2011). NPQ parameter is the non-photochemical quenching and is relevant to radiant energy loss as heat (Maxwell & Johnson, 2000). NPQ and qP would increase and decrease, respectively, under WDS (Ashraf & Harris, 2013). Also, for better understanding xanthophyll cycle activity, NPQ can be checked (Ralph & Gademann, 2005). Many physiological processes rely on cellular K, such as maintaining cellular turgor, regulating membrane permeability, regulating ion balance, enhancing photosynthesis, and influencing protein synthesis (Maathuis & Amtmann, 1999). The physical similarities between Na and K may cause Na to compete with K for entry into the symplast, resulting in a K deficiency. A high ratio of K to Na at binding places in the cytoplasm could inhibit enzyme functions and metabolic processes that rely on K (Maathuis & Amtmann, 1999). Other effective environmental factors include the seasons of year that influence the synthesis of chemical substances in some medicinal plants. Changes across different seasons may lead to corresponding shift in and/or accumulation of some EO compounds (Koptur, 1985). Souza et al. (2018) observed the volatile oil yield of seasonality in Spiranthera odoratissima. Zutic et al. (2014) indicated the significant effect of cutting time on the EO yield of sage. Several studies have been conducted on effective application of elicitors for the production of target SMs in plants (Emami Bistgani et al., 2017).

Polyamines (PAs) such as PUT, spermine, spermidine, and cadaverine as well as a class of phyt hormorne-like aliphatic amine composites regulate many physiological processes such as development and growth, leaf senescence, plus abiotic and biotic plant stress reactions (Mohammadi et al., 2018). PAs contribute to the regulation of physical and chemical properties of membranes and nucleic acids (Aloisi et al., 2016). PAs can protect plants under abiotic stresses (Bouchereau et al., 1999). PAs can induce biosynthesis of secondary metabolites, increase permeability plus integrity of the plasma membrane, and inhibit the chlorosis under environmental stresses (Alcázar et al., 2010, 2011; Gupta et al., 2013; Kusano et al., 2007; Mohammadi et al., 2018). Some reports have indicated that PAs positively affected photosynthetic pigments in Morus alba, Cucumber sativus, and Phaseolus vulgaris (He et al., 2002; Nassar et al., 2003).

However, the mechanisms of prohibition of photosynthesis by water deficit and PUT remain poorly defined. Hence, the current study was designed to check the role of exogenous (exo) PUT and WDS on leaf gas exchange, greenness index, chlorophyll fluorescence parameters, pigments, Na, K, EO content of sage at two cutting times.

2 | MATERIALS AND METHODS

2.1 | Position description and experimental design

1200 of sage seedlings was cultivated in Tarbiat Modares University’s Agriculture Faculty located in Tehran, Iran (1200 m upper sea level and 35° 70’ N, 51° 40’ E), in 2 years (2017-2018). Thirty-year mean annual temperature and rainfall are 22°C and 232.6 mm, respectively (see meteorological data in Supplementary Information 1). The experiments were as split–split plot adjustment in a randomized complete block design with three repetitions. The main plot was the water-deficit stress (WDS), the subplot was PUT, and the sub-subplot was the cutting time. The WDS was as follows: 20%, 40%, 60%, and 80% ASWD. WDS treatments were carried out based on the maximum allowable depletion (MAD) from the percentage of available soil water (%ASWD). Irrigation started after soil water reached a threshold level (Bahreininejad et al., 2013; Govahi et al., 2015). Time Domain Reflectometry probe (TDR) (Model TRIME-FM, Germany) was used to measure soil water level at a root zone of sage (depth of 50 cm). Four concentrations of PUT (distilled water (0), 0.75, 1.5, and 2.25 mM) were used. All aboveground parts in each plant were exo sprayed 50 cm upper of plant. Foliar application of PUT was carried out twice each year, 1 week before using WDS in each cutting in 2 years. Furthermore, the foliage was handpicked in spring (cutting 1) and summer (cutting 2). No PUT and WDS were used in the first month of the growth cycle, because plants should have been formed of similar masses of foliage, before applying treatments. The cutting 1 and 2 carried out before flowering. The foliage of plants was cut
8–10 cm over the soil surface. Collections were carried out in 2 years (2017–2018).

2.2 | Essential oil extraction

Dried samples (100 gr) were extracted via the hydrodistillation method in a Clevenger device with double-distilled water (1000 ml). The collected surplus of aqueous EO was dried over anhydrous sodium sulfate. Then, the weight of pure EO was determined, and its percentage was computed.

2.3 | Leaf gas exchange and leaf greenness quantification

An Li-6400XT Portable Photosynthesis System (Li-Cor Inc.) was utilized to determine the net photosynthetic rate (Pn) and stomatal conductance (gs) of plants per plot after each cutting time. The chamber was corrected to 25°C (temperature), ambient CO2 concentration (Ca) was 380 μM/mol, and photosynthetic photon flux density was 1250 μM/m²/s. For estimating the leaf greenness, GREENNESS INDEX-502 was used on the same leaves mentioned previously (Konica Minolta).

2.4 | Chlorophyll fluorescence parameters

The maximum performance of PSII (Fv/Fm) and nonphotochemical quenching (NPQ), photochemical quenching (qP) was determined applying a modulated fluorescence chlorophyll meter (MINI-PAM, Walz). Leaves were dark-adapted via leaf clips (15 min). After dark adaptation, the highest photochemical performance (Fv/Fm), actual photochemical performance (ΦPSII) = (Fm'-Fs)/Fm', qP = (Fm'-Ft)/(Fm'-Fo'), and NPQ =1- (Fm'-Fo')/(Fm-Fo) were computed as well, and the light severity applied to calculate the yield of ΦPSII and NPQ was 1.200 μmol/m²/s (Roháček, 2002).

2.5 | Photosynthetic pigments

To estimate the chlorophyll and carotenoid, the procedure of Lichtenthaler (1987) was used. Briefly, 0.5 g of fresh leaves sage was powdered applying mortar and pestle containing 10 ml of acetone (80% V/V). Then, the extract was centrifuged for 10 min (12,000 rpm). The light absorption was read at 645, 663, and 470 nm by a UV-vis spectrophotometer. Photosynthetic contents were expressed as mg/g/fw.

2.6 | Endogenous putrescine analysis determination

Endogenous (endo) PUT extraction and thereafter HPLC measurement were performed following the procedure of Lütz et al. (2005). First, samples were injected into injector loop (20 μl) in RP-C18 Column (15 cm × 4 mm i.d.), particle size (5 μm), at 30°C using Methanol: Water linear gradient from ratio of 50:50 to 80:20 (v/v) for 30 min. The last ratio was retained at 1 ml/min. PUT was detected by measuring the fluorescence intensity of samples (254 nm) and then comparing their peak times with those of standard PUT.

2.7 | Estimation of Na⁺, K⁺, and leaf area index (LAI)

For estimating sodium and potassium, the procedure of Ahanger et al. (2015) was used. The Na⁺ and K⁺ contents were determined by flame photometry (Jenway-Flame Photometer Models PFP7). To estimate the area index, the leaf area meter (Delta-T Devices Ltd.) was employed after using treatments in the end of each cutting time.

2.8 | Statistical analysis

The procedure as type3 in MIXED method of SAS v. 9.4 (SAS Institute) was applied to analyze data. The significant effects of WDS, PUT, cutting time, and their two- and three-way interactions were considered fixed effects, while years, replicates ×years and years ×WDS × PUT ×cutting time were considered chance effects. The PDIFF choice of least square means adjusted for the Tukey-Kramer was applied for mean comparisons. The interactions among experimental factors were separated by slicing procedure. The significance of linear and quadratic regression models (p < .05) was examined with polynomial orthogonal contrasts. Pearson’s correlation coefficients were specified applying the CORR method.

3 | RESULTS

3.1 | Dry weight, LAI, and endogenous putrescine

Based on the results analysis of variance, effects of water deficit, PUT, and cutting time were significant for the dry weight, LAI, and endo PUT of sage (Supplementary Information 2). Also, dry weight and LAI were higher in cutting 2 than in cutting 1. Endo PUT was higher in cutting 1 than in cutting 2 (Table 1). The maximum dry weight (234.76 g/m²) and LAI (0.83) were obtained in 20% ASWD. Indeed, there was a descending trend in dry weight and LAI by incrementing severity of water-deficit stress (WDS) (Figure 1a,c). The response of dry weight and LAI to PUT can be expressed with a quadratic model, indicating that dry weight and LAI would reach their maximum (219.04 g/m² and 0.72) under 0.98 and 1.14 mM of PUT, respectively (Figure 1b,d). There was an incrementing trend in endo PUT with increasing concentration of PUT. The maximum endo PUT (195.55 nmol/g FW) was obtained under 2.25 mM of PUT (Supplementary Information 3).
3.2 | Essential oil content and yield

Based on the analysis of variance, the main effects of WDS, PUT, and cutting time were on EO content and EO yield (Supplementary Information 2). The EO content and yield were higher in cutting 1 and cutting 2, respectively (Table 1). The response of EO content and EO yield to WDS can be expressed by a quadratic model, showing that the EO content and EO yield would reach their maximum (1.45% and 260.47 g/m²) under 75.5% and 34.86% ASWD, respectively (Figure 2a,c). The response of EO content and EO yield to foliar application of PUT can be expressed with a significant quadratic model (p ≤ .01), suggesting that EO content and EO yield would reach their maximum (1.39% and 279.62 g/m²) under 1.34 and 1.16 mM of PUT (Figure 2b,d).

3.3 | Photosynthetic gas exchange parameters, chlorophyll fluorescence, and greenness index

Based on the results analysis of variance, cutting time affected Pn, Gs, Fv/Fm, and NPQ significantly (Supplementary Information 2). Pn and Gs were higher in cutting 1 than in cutting 2. In contrast, Fv/Fm and NPQ were higher in cutting 2 than in cutting 1 (Table 1). The main effect of WDS was on Pn, Gs, and spad. Also, the main effect of PUT was in Fv/Fm and spad. None of the treatments was significantly different on qP (Supplementary Information 2). The maximum Pn (16.2 μmol/m² s⁻¹) and Gs (0.057 mmol/m² s⁻¹) were obtained in 20% ASWD. Indeed, there was a descending trend in Pn and Gs by increasing severity of WDS (Figure 3a,b).

The response of Fv/Fm to WDS can be expressed with a linear model, indicating that Fv/Fm would reach its most by the increasing severity of WDS. A maximum Fv/Fm (0.66) was observed for 80% ASWD (Figure 3c). The response of NPQ and spad to WDS can be expressed by quadratic model, showing that NPQ and spad would

![Figure 1](image-url)
attain their most (0.63 and 72.01) in 38.33% and 84.13% ASWD, respectively (Figure 3d and Supplementary Information 4). The response of Fv/Fm, ΦpsII, and spad to PUT can be expressed with a quadratic model, indicating that Fv/Fm, ΦpsII, and spad would reach their maximum (0.72, 0.24, and 56.1) under 1.27, 1.17, and 1.18 mM of PUT, respectively (Figure 3e,f, and Supplementary Information 5).
3.4 Chlorophyll a, b, total, and carotenoid

The result showed that the main influence of WDS and PUT was on Chl a, Chl b, Chl t, and carotenoid. Also, the significant effect of cutting time was on Chl a, Chl t, and carotenoid (Supplementary Information 2). Chl a, Chl t, and carotenoid were higher in cutting 1 than in cutting 2 (Table 1).

The response of Chl a and Chl t to WDS can be expressed by a quadratic model, showing that Chl a and Chl t would reach their maximum (0.08 and 0.17 mg g\(^{-1}\) FW) in 60% and 70% ASWD, respectively (Figure 4a,e). The response of Chl b to WDS can be expressed with a linear model, indicating that Chl b would attain its most with an increasing severity of WDS. The maximum Chl b (0.11 mg g\(^{-1}\) FW) was observed for 80% ASWD (Figure 4c). The highest carotenoid (0.63 and 0.74 mg g\(^{-1}\) FW) was obtained in 20% and 40% ASWD, respectively (Figure 4g). The response of Chl a, Chl b, Chl t, and carotenoid to PUT can be expressed by a quadratic model, showing that Chl a, Chl b, Chl t, and carotenoid would reach their maximum (0.08, 0.10, 0.18, and 0.78 mg g\(^{-1}\) FW) under 1.25, 1.17, 1.27, and 1.31 mM of PUT, respectively (Figure 4b,d,f,h).

3.5 Sodium, potassium, potassium/sodium ratio

Based on the results, that main effect of cutting time and PUT was significant for Na, K, and K/Na. Also, the main effect of WDS on Na and K (Supplementary Information 2). K and K/Na were higher in cutting 1 than in cutting 2. In contrast, Na was maximum in cutting 2 than in cutting 1 (Table 1).

The response of Na to WDS and PUT can be expressed by a quadratic model, indicating that Na would attain its most (16.48 mg/g DM) under 63.96% ASWD and minimum (13.40 mg/g DM) under 1.20 mM of PUT, respectively (Figure 5a,b). The response of K and K/Na to WDS can be expressed with a quadratic model, suggesting that K and K/Na would reach their maximum (90.58 mg/g DM and 13.40 mg/g DM) under 63.96% ASWD and 50.40% ASWD, respectively (Figure 5c). Also, the response of K and K/Na to PUT can be expressed by a quadratic model, showing that K and K/Na would reach their maximum (90.96 mg/g DM and 7.36) by applying 1.21 and 1.19 mM PUT, respectively (Figure 5d,f,h).

4 DISCUSSION

Slight water-deficit stress (WDS) (34.86% ASWD) would be required for achieving the maximum EO content of sage. In the present study, the maximum EO yield of sage was produced under potential yield conditions in 75.5% ASWD. The lowest dry weight and LAI of sage were produced under 80% ASWD. Analogous to our results, the dry weight diminished with increasing WDS where the EO yield of sage was the highest under 60% ASWD (Govahi et al., 2015). Also, analogous to our results, Nowak et al. (2010) announced that the main part of monoterpene increased, and biomass decreased in sage under WDS. Our results indicated that there was a high correlation between dry weight and LAI (0.72, p < .01) and EO yield (0.94, p < .01) (Supplementary Information 6). Also, dry weight, LAI, and EO yield were higher in cutting 2 than in cutting 1, in which these variations could be imputed to inter- and intraseasonal weather changes resulting from years and cutting times tested. Over both years, rainfall was higher by almost 0.64 mm in cutting 1 than in cutting 2, while temperature was lower by almost 13.47°C (Supplementary Information 1). The shift in EO yield suggests a possible corresponding shift and/or accumulation of EO in response to seasonal conditions. Indeed, the EO could increase, decrease, or disappear during cutting 1 and cutting 2. Thermoregulation is among the main causes, as the EO hydrophobic compounds could increase during the hot periods to protect the plant from desiccation (Kamatou et al., 2008). The genus salvia is one of the important known genera in the Labiatae family due to monoterpene. Indeed, terpenes biosynthesis is composed of two separate ways, including methylerthritol 4-phosphate (MEP) and mevalonate (MVA), occurring in plastids and cytoplasm in plants. As reviewed by Müller Verma and Shukla (2015), the MEP pathway is involved in the synthesis of carotenoids, isoprene, mono- and diterpenes, plant hormones, phytol, the side chain of Chl, tocopherol, phylloquinone, plastoquinones, etc. The polyamines in plants are found in cytoplasm and organelles such as mitochondria, chloroplasts, and vacuoles (Kusano et al., 2008). Based on the results, the highest dry weight, LAI, EO yield, and EO content were obtained under potential yield conditions with application of 0.98, 1.14, 1.16, and 1.34 mM of PUT, respectively. There was a high correlation between endo PUT and EO content (0.51, p < .01) and Gs (0.42, p < .05) (Supplementary Information 6). As an explanation, perhaps PUT enters the leaves by penetrating the cuticle or via the stomata, before entering the plant cell, where they can be practical in metabolism and are mainly carried to other portions via plasmodesmata. Hence, polyamines and monoterpene were probably produced in the pathway of methylerthritol 4-phosphate (MEP) and EO content affected by PUT.

In this study, the lowest LAI, Pn, and Gs were observed under 80% ASWD. Also, there was a high correlation among Pn and LAI (0.48, p < .01) and Gs (0.81, p < .01) (Supplementary Information 6). Probably, the minimum LAI resulted due to minimum of Pn under 80% ASWD. In addition, 80% ASWD-induced reduction of Pn was accompanied by a reduction of Gs, suggesting that the effect of WDS on Pn can be due to stomatal agents (Yang & Lu, 2005). Based on our results, the levels of PUT were not accompanied by a considerable corresponding shift in Pn and Gs, indicating that stomatal guidance was not the only agent for PUT-induced shifts in photosynthesis. Skowron et al. (2021) also reached similar results. The decreased photosynthesis can be due to nont stomatal limitations including the stagnation in PSII activity and electron transport (das Neves et al., 2008; Xia et al., 2004). Results showed that there was a negative correlation among NPQ and Gs (−0.37, p < .05), Fv/Fm and Pn (−0.38, p < .05), and Fv/Fm and Gs (−0.71, p < .01) (Supplementary Information 6). Indeed, there was a negative correlation among photosynthetic gas exchange parameter and chlorophyll fluorescence parameter.
Electron transfer in the photosynthesis process is necessary for assimilation. The catalytic protein PSII is the first component of the electron transport chain (Liao et al., 2013). Based on our results, the highest Fv/Fm was obtained under 80% ASWD. Also, Fv/Fm was higher in cutting 2 than in cutting 1. Indeed, elevation results indicated the protection of light in sage as well as higher Fv/Fm was higher in cutting 2 than in cutting 1. As a result, environmental stress makes the electron transfer chain saturated and leads to protons accumulating, whereby NPQ would show increase (Porcar-Castell et al., 2014). By increasing this value, the chloroplasts are more capable of reversing the effects of WDS, as these organelles can decompose the excess excitation energy (Li et al., 2014). There was a negative correlation among NPQ and Chl a (−0.59, p < .01), Chl b (−0.38, p < .05), Chl t (−0.50, p < .01), and carotenoid (−0.55, p < .01) (Supplementary Information 6). In addition, NPQ and Na were higher in cutting 2 than in cutting 1. As a result, environmental stress makes the electron transfer chain saturated and leads to protons accumulating, whereby NPQ would show increase (Porcar-Castell et al., 2014). By increasing this value, the chloroplasts are more capable of reversing the effects of WDS, as these organelles can decompose the excess excitation energy (Li et al., 2014). There was a negative correlation among NPQ and Chl a (−0.59, p < .01), Chl b (−0.38, p < .05), Chl t (−0.50, p < .01), and carotenoid (−0.55, p < .01) (Supplementary Information 6). Hence, it could be concluded that excess excitation energy is mostly used to produce Chl and carotenoids compared to the production of NPQ in WDS conditions in sage. There was a negative correlation among NPQ and EO content (−0.56, p < .01) (Supplementary Information 6). Also, NPQ and EO were higher in cutting 2 and cutting 1, respectively. Indeed, the synthesis of secondary metabolites and the biosynthesis of highly decreased compounds like isoprene are mainly involved in the waste of surplus photosynthetic energy (Wilhelm et al., 2011). Fv/Fm, ΦPSII, and spad exhibited a shift in pattern analogous to that of PUT, showing their most important effects on incrementing Fv/Fm, ΦPSII, and spad at
concentrations of 1.27, 1.17, and 1.18, respectively. Indeed, Fv/Fm and ΦpsII improved after the PUT application; however, NPQ was not affected. These findings propose that, probably, the PUT improved the photosynthetic system by optimizing energy broadcast and by enhancing structure and function.

The maximum concentration of endo PUT was obtained by 2.25 mM of PUT. Based on our results, there was a high correlation between endo PUT and ΦpsII (0.55, p < .01) and Gs (0.42, p < .05) (Supplementary Information 6). This correlation could be ascribed to the role of PUT in stimulating plant growth by photosynthesis. There was a negative correlation between endo PUT and NPQ plus Fv/Fm (−0.64, −0.72, p < .01) (Supplementary Information 6). Similar results were obtained by Razzaq et al. (2014). In the present study, there was a high correlation among endo PUT and Chl a (0.58, p < .01) Chl t (0.43, p < .01), and carotenoid (0.60, p < .01) (Supplementary Information 6). Also, PUT, Chl a, Chl t, and carotenoid were higher in cutting 1 than in cutting 2. Indeed, stomatal and photosynthetic activity improved with increasing K levels with PUT. According to our results, Iqbal and Ashraf (2005) reported that PUT could adjust ion homeostasis, sorption, and translocation of poisonous ions. Indeed, the regulating effect of PUT on the ion balance is due to the aggregated endo PUT rather than rivalry among cationic PUT and Na at the absorption status (Ndayiragije & Lutts, 2006).

According to regression results, the maximum and minimum Na concentration was observed in 63.96% ASWD and 1.20 mM of PUT, respectively. Also, K and K/Na would reach their maximum by applying 1.21 and 1.19 mM of PUT, respectively. Instead, there were positive and negative correlations among endo PUT and Na (−0.43, p < .01), K (0.58, p < .01), and K/Na (0.59, p < .01) (Supplementary Information 6). There was a positive correlation among K/Na and Pn (0.59, p < .01) and Gs (0.51, p < .01) (Supplementary Information 6). Also, K, K/Na, Pn, and Gs were higher in cutting 1 than in cutting 2. Indeed, stomatal and photosynthetic activity improved with increasing K levels with PUT. According to our results, Iqbal and Ashraf (2005) reported that PUT could adjust ion homeostasis, sorption, and translocation of poisonous ions. Indeed, the regulating effect of PUT on the ion balance is due to the aggregated endo PUT rather than rivalry among cationic PUT and Na at the absorption status (Ndayiragije & Lutts, 2006).

5 | CONCLUSION

The study indicated that the maximum EO yield of sage was produced under potential yield conditions in 75.5% ASWD.
Greenness index, Chl, and carotenoid increased by the intensity of WDS. The lowest LAI, Pn, and Gs were observed under 80% ASWD. But, the highest Fv/Fm was obtained under 80% ASWD. The reduced photosynthesis can be due to stomatal factors and nonstomatal limitations including the stagnation in PSII activity and electron transport. Regression results showed that the NPQ increased under 38.33% ASWD. There was a negative correlation between NPQ and K and K/Na. Indeed, environmental stresses make the electron transfer chain saturated and increment proton accumulation, whereby NPQ would increase. There was a negative correlation between NPQ and Chl and carotenoid. So, it could be concluded that excess excitation energy is mostly used to produce Chl and carotenoids compared to the production of NPQ under WDS conditions in sage. There was a negative correlation between NPQ and EO content. Also, NPQ and EO were higher in cutting 2 and cutting 1, respectively. Nonetheless, the dissipation of surplus photosynthetic energy is mainly accomplished by the biosynthesis of highly reduced compounds such as isoprene and secondary metabolites. PUT showed that Fv/Fm and ΦPSII improvement, however, did not affect NPQ. These findings propose that, probably, PUT optimized energy broadcast and improved the structure and function of the photosynthetic system, and thereby it could be promoting primary electron transfer in PSII. PUT incremented the accumulation of Chl a, Chl b, Chl t, and carotenoid, indicating that PUT improved the transport of the photosynthetic matter. The maximum concentration of endo PUT was obtained with 2.25 mM of PUT. There were positive correlations between K/N and endo PUT, Pn, and Gs. Indeed, stomatal and photosynthetic activity improved with increasing K levels with the application of PUT.

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DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author by reasonable request.

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