INTRODUCTION

Agriculture is exposed to frequent periods of drought and limitation of water resources what is expected to exert an adverse impact on plant growth and crop productivity (Shao et al., 2008). The Mediterranean regions are experiencing periods of intense drought, leading to the extension of arid zones (Gao & Giorgi, 2008). In view of gradually depleting irrigation water resources throughout the world, it is highly imperative to investigate the effects of water deficit in plants consuming large amounts of water like tomato (Solanum lycopersicum). In fact, water plays a crucial role in determining the yield of processing tomato but it is likely that water

ORIGINAL RESEARCH

Effects of water deficit on leaves and fruit quality during the development period in tomato plant

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Abstract

In nature, plants are often exposed to a multitude of environmental constraints that severely limit crop productivity. Water deficit is one of the factors that most affects agricultural production. The aim of this work is to evaluate the effect of water deficit on morphology, development, nutritional behavior, as well as chlorophyll fluorescence and certain important metabolic parameters (soluble sugars, organic acids, starch, carotenoid, and vitamin C) of the cultivated tomato (Solanum lycopersicum cv Plovdiv). In this study, the water supply was reduced by 60% compared to control conditions. The conditions of water deficit showed that the size of the different organs (leaves, fruits) was reduced. A reduction in the number, width, and length of the leaves, respectively, 9%, 36%, and 37%, then the leaf surface was also observed. Reduction of fluorescence (Fo, Fm, and Fv) and total index performance were among the other symptoms of plants with water deficiency. For fruit, we observed a significant decrease in diameter, fresh weight, and moisture content during the cell division period, the cell expansion period, and the fruit ripening period. In contrast, the composition of the Plovdiv fruit changed only during cell division and expansion phase. On the other hand, the water deficit induces an increase in the total carotenoid and vitamin C content of the fruits. Besides, water deficit induced a reduction of fruit size, moisture content, and production dry matter during different phases of development. Decrease levels of soluble sugars and organic acid but increase in vitamin C and carotenoid content.

KEYWORDS
fruit, growth, leaves, metabolic parameters, tomato, water deficit

1 INTRODUCTION

Agriculture is exposed to frequent periods of drought and limitation of water resources what is expected to exert an adverse impact on plant growth and crop productivity (Shao et al., 2008). The Mediterranean regions are experiencing periods of intense drought.
2 | MATERIALS AND METHODS

2.1 | Plant material & experimental conditions

The study was performed on S. lycopersicum. L type genotype: Plovdiv XXIVa, is a cultivated tomato plants, Plovdiv seeds were provided by the Genetic Resource center of INRA, Avignon (France). This Genotype showed important allelic variability (SNP differences on chromosomes 3, 4, 5, 7, 8, 9, 11, and 12) (Causse et al., 2013).

The trial was conducted during winter 2014 in a glasshouse located near Avignon, France. 40 plants were grown in pots (Plant/ pots) filled with compost (substrate 460, Klasmann, Champety, France) distributed in two rows (control and stressed plants) at a density of 1.3 plant m⁻² and 20 plants were actually used for the experiment. Plants were supplied daily with a nutrient solution (Liquoplant Rose, Plantin, Courthézon, France) diluted between 0.4 % and 0.8 % according to the plant development stage, which corresponds to an average electroconductivity of 1.8 mS cm⁻¹ for the whole period. Flowers were pollinated three times a week using an electrical bee. Day–night temperature control was set at 25–15 °C. Over the whole trial period, the air temperature and relative humidity remained relatively stable (on average the day temperature ranged between 20.4 and 24.5 °C, the night temperature between 15.1 and 19.7 °C, and the air humidity between 56% and 72%). Control plants were irrigated, according to current practices, in order to maintain soil humidity and drainage around 70% (maximum water retention capacity of the substrate) and 15%, respectively. Soil humidity was measured every two days in all pots using water content sensors (WCM-control, Grodan, Roermond he Netherlands). During the study, water supply was reduced by 60% compared to control conditions. Preliminary experiments demonstrated that this level or irrigation induced a moderate water stress based on several plant indicators (leaf conductance, stem and leaf water potentials, and the specific leaf area (SLA). Moreover, it was observed that four days were necessary to reach stable soil humidity after the beginning of water restriction in the conditions of our trial (Ripoll et al., 2016). Therefore, treatments were applied four days before the beginning of development phase. The substrate water content was measured in all pots every day and maintained around 25% for treated plants, and more than 60% for control plants.

2.2 | The measured parameters

2.2.1 | Morphological parameter

Plant leaf number, width, and leaf length were measured every 3 days out of 10 plants per treatment. Mature leaves nonsenescent, which were initiated during the WD treatments, were harvested on each plant and their specific leaf area was measured. Leaf area was measured with a Planimeter (Li- 3,100 C Area Meter, Li-Cor, Lincoln, NE, USA) and leaf dry weight was measured after seven days at 70 °C.
Medyouni et al. in a ventilated oven. All fruit measurements were made on fruits harvested at different stages of fruit development. Ten fruits per treatment were served to measure. Fruit size, fresh weight, and water content were measured immediately after harvest.

2.2.2 | Physiological parameter

Fluorescence Chlorophyll parameters (Table 1) were measured on dark-adapted leaves (30 min.) using a fluorimeter (HANDY-PEA, Hansatech, King’s Lynn, UK). Dark-adaptation allowed the PSII electron acceptor pool to be gradually re-oxidized to a point where all PSII reaction centers are capable of undertaking photochemistry. Measurements were carried out with an induction period of 1 s and leaves were illuminated to a light level of 3,000 μmol photons m⁻² s⁻¹. The measurements were carried out on nonsenescent mature leaves, at around 11 a.m. Fluorescence was measured weekly during the entire culture period. For the measurement of cations, plant material was dried at 80°C and digested with nitric acid [1% (v/v) nitric acid (HNO₃)] according to the method of Wolf (1982).

2.2.3 | Mineral analysis

For the measurement of cations, plant material was dried at 80°C and digested with nitric acid [1% (v/v) nitric acid (HNO₃)] according to the method of Wolf (1982). K⁺, Ca²⁺, and Mg²⁺ were analyzed by flame emission using a spectrophotometer (Eppendorf Geratebau Netherler).

2.2.4 | Biochemical parameter

Fruits and leaves were frozen in liquid nitrogen and kept at ~80°C prior to biochemical analysis soluble sugars, starch, organic acids, and carotenoids. Soluble sugars (glucose, fructose, and sucrose) and organic acids (i.e., citric acid, malic acid, and quinic acid) were extracted according to the method described by Gomez et al. (2002) and analyzed by HPLC (Waters 410, Part WAT070390, Milford, U. S. A.). Ascorbic acid content was measured according to the method described by Stevens et al. (2006), and the absorbance was read at 550 nm using a Multiscan Ascent MP reader (Labsystems, Thermo Fisher Scientific, Courtaboeuf, France). Carotenoids (i.e., lycopene, phytoene, beta-carotene, and lutein) were extracted according to the method described by Serino et al. (2009) and assayed by HPLC with a UV–vis detector (UV6000LP, Thermo Separation Products, Riviera Beach, U. S. A.).

2.3 | Statistical analysis

Statistical analyzes were performed using the SPSS for Windows software, version 21.0. Mean values and standard error (SE) were obtained from at least 10 measurements for physiological parameters (DW, WC, leaf area, number, width, and length of leaves) and 5 measurements for biochemical parameters. A P value under 5% was considered statistically significant. Duncan’s multiple range test was used to perform means’ comparisons.

3 | RESULTS

3.1 | Water deficit effect on leaf morphology

After three months, the results showed that water deficit treatment decreased number of leaves (Figure 1a). The number of stressed leaves was reduced by approximately 9% compared to control leaves. Our results showed that treatment of water deficit decreased the leaf length and leaf width as compared to control plants (Figure 1b, c). Our results showed that treatment of water deficit decreased the

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**TABLE 1** Chlorophyll fluorescence parameters

| Parameter | Description |
|-----------|-------------|
| Fo | The first reliable fluorescence value after the onset of actinic illumination |
| Fm | Maximum value under saturating illumination |
| Fv | Maximum variable Chlorophyll fluorescence |
| Fo/Fm | A parameter related to changes in heat dissipation in the photosystem II antenna |
| Fv/Fm | The maximum photochemical efficiency of light harvesting in PSII |
| Fv/ Fo | Quantum yield of primary PSII photochemistry, represents the contribution to the PI of the light reactions for primary photochemistry |
| VJ | The fluorescence at J step (2 ms), F2ms |
| VI | The fluorescence at I step |
| ABS/RC | Specific fluxes or specific activities |
| DI0/RC | Specific fluxes or specific activities |
| TRI0/RC | Specific fluxes or specific activities |
| ETI0/RC | Specific fluxes or specific activities |
| PIABS | Performance index |
| Pilt | Performance index |
length and width of leaves compared to control plants (Figure 1b, c), this reduction is respectively by of 21% and 20%.

Our results showed that water deficit decreased dry weight for tomato as compared to control plants (Figure 2a). The dry weight was reduced by approximately 36% compared to control.

The leaf area and the area density of tomato plants were reduced not significantly with water deficit. In fact, leaf area was reduced by approximately 28% compared to control leaf area. In the same way, area density of stressed plants was reduced about 37% than control plants (Figure 2 b, c).

3.2 | Water deficit effect on leaf physiology

3.2.1 | Fluorescence parameters

Illumination of a dark-adapted leaf induces characteristic changes in fluorescence intensity. $F_o$ and $F_M$ respectively represent the intensity of the minimum fluorescence (all the reaction centers are oxidized or open) and maximum (all the reaction centers are reduced or closed). The rapid increase in chlorophyll fluorescence yield between $F_o$ and $F_M$ during the first second of intense illumination was used to analyze electron transport in the PSII. Chlorophyll fluorescence was measured on the leaf at stages10. Analysis of variance showed that irrigation regime, time significantly affected all measured traits. The intensity of the minimum $F_o$ and maximum fluorescence $F_M$ estimated, in tomato seedlings subjected to water deficit decreases compared with control plants. The highest Fv was decreased to plants subjected to water deficit, The FV was reduced by approximately 6% compared to control. Also, the Fv/Fm ratio significantly decreased with increasing water deficit stress severity during all the sampling times (Table 2). After a period of water deficit, a significant increase was observed for $V_i$ and $V_j$ and no difference compared to the control was observed for the parameters of the activity of the flows $ABS/RC$, $D_{tr}/RC$, $TR_{tr}/RC$, $ET_{tr}/RC$. The water deficit induced a significant increase in $IP_{ABS}$, this increase was about 12% compared to the witness. Under a water deficit, Plovdiv tomatoes showed a significant decrease in $PI_3$ and it varied between 4.067 (control plants) and 2.409 (stressed plant).

3.2.2 | Mineral analysis

The water deficit decreased the mineral content in leaves of tomato plants. Content of $K^+$ and $Ca^{2+}$ were decreased by water deficit, which suggests a deficit in providing the plants with essential ions to the growth. The water deficit reduced a $K^+$ and $Ca^{2+}$ concentration in leaves respectively by, 16% and 5% (Figure 3). However, water deficit induced an increase in $Mg^{2+}$ content compared of control plants (Figure 0.3).

3.2.3 | Water deficit effect on leaves biochemical parameters

After three months of irrigation treatment, different metabolite contents: soluble sugars, organic acids, and ascorbic acid (AsA) were measured in leaves of tomato plants Plovdiv (Figure 4). The level of

![Figure 1](image-url)
acids in leaf decreased significantly with 19%, in detail the citric acid decreased with 19% and the malic acid decreased with 18% as compared of control plants. Also, the water deficit increased significantly the leaf sugar level sugar: glucose 17% and fructose 22% as compared of control plants. Furthermore, the water deficit was decreased significantly the content starch with 31% compared to control leaves. Total AsA content of leaves decreased below that of the controls, the water deficit treatment, significantly decreased the AsA by 15% compared to control.

### 3.3 Water deficit effect on fruit morphology

Fruit diameter, fresh weight, dry weight, and water content were affected by the WD treatments (Table 3). The significant increase in fresh weight, fruit diameter, dry weight, and water content was observed during the cell division phase respectively (22%, 47%, 50%, and 53% respectively). During the phase of cell expansion of the fruits. Only a significant increase in fresh weight and water content (21% and 54% respectively by). On the contrary, the treatments did

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**TABLE 2** Variation of $F_o, F_m, F_v$. Fluorescence Reports, Specific energy fluxes and Performance index of the leaves of tomato plants *Solanum lycopersicum* (cv. Plovdiv) cultivated in the presence of water (CT: control treatment) or under water restriction conditions (ST: stressed treatment), measurements were taken from the vegetative stage to the maturation stage, about 10 plants per treatment

| Fluorescence parameters | $F_o$ | $F_m$ | $F_v$ |
|-------------------------|-------|-------|-------|
| CT                      | 6,189.35 ± 162.74a | 34,950.14 ± 1,015.30a | 28,953.17 ± 1,015.46a |
| ST                      | 5,996.97 ± 117.72a | 33,335.66 ± 905.45b | 27,146.32 ± 1,009.32b |

| Fluorescence reports | $F_o/F_m$ | $F_v/F_m$ | $F_v/F_o$ | $V_i$ | $V_j$ |
|----------------------|-----------|-----------|-----------|-------|-------|
| CT                   | 0.175 ± 0.006a | 0.825 ± 0.006a | 4.865 ± 0.180a | 0.432 ± 0.019a | 0.808 ± 0.016a |
| ST                   | 0.191 ± 0.010b | 0.808 ± 0.010b | 4.492 ± 0.219b | 0.392 ± 0.019b | 0.754 ± 0.024b |

| Specific energy fluxes | $ABS/RC$ | $DI_v/RC$ | $TR_v/RC$ | $ET_v/RC$ |
|------------------------|----------|-----------|-----------|-----------|
| CT                     | 1.664 ± 0.058a | 0.296 ± 0.0231a | 1.368 ± 0.039a | 0.773 ± 0.033a |
| ST                     | 1.669 ± 0.104b | 0.338 ± 0.046a | 1.330 ± 0.061a | 0.807 ± 0.053a |

| Performance index | $PI_{ABS}$ | $PI_{I}$ |
|-------------------|------------|----------|
| CT                | 4.382 ± 0.395a | 4.067 ± 1.479a |
| ST                | 5.029 ± 0.501b | 2.409 ± 0.426b |

Note: Bar indicates standard error ($p = .05$). Same letter above the bars denotes that the difference between means were not significant.
not impact the diameter, height, and dry weight. In Plovdiv fruits, during the maturation phase diameter, height, fresh weight, and dry weight had not affected by water deficit. Whereas the water content decreased about 31% compared to control.

### 3.4 Water deficit effect on fruit physiology

Whatever the ages of fruits, the deficit induced a significantly decrease in K⁺ content as compared to control. In addition, the decrease of Ca²⁺ content is important in 21- and 28-day-old fruits, noted that this concentration is 0.04 meq for control fruit ages 21 days while it is only 0.02 m eq in fruit stresses (Figure 4), the reduction was about 46%. For fruits aged 28 days the concentration is 0.05 m eq for control fruit and 0.02 meq for stressed fruits, the reduction was about 51%. The water deficit treatment significantly decreased K⁺ content in fruit of tomatoes. The water deficit treatment significantly decreased K⁺ content in fruit of tomatoes. The water deficit treatment significantly decreased K⁺ content in fruit of tomatoes. The water deficit treatment significantly decreased K⁺ content in fruit of tomatoes. This decrease of K⁺ is important in 35- and 42-day-old fruits, for example, the reduction of aged fruits 35 days K⁺ content was about 1.66 m eq for control fruits and 1.41 meq for stressed fruits, the reduction was about 15% (Figure 5).
Effect of water deficit on the biochemical parameters of the fruits

Total sugars, total acid, AsA, and carotenoid were measured at cell expansion phase and maturation phase showed a general decrease in treated plants as compared to control plants. Contrasting responses were observed between the two ages of the fruits (Table 4).

In fact, for the green fruit, content of fructose (19%), and glucose (22%) were decreased significantly respectively 19%, 22% compared to control plants. Moreover, green fruit increased citric acid (10%), and thus total acid (25%) as compared to their in control plants. Concerning the ascorbic acid (AsA) showed higher content of vitamin C (20%) then those of control plant. Finally, the lycopene (24%), phytoene (11%) and thus total carotenoids (23%) were increased in green fruit after water restriction treatment compared to control plants.

The sugar content was reduced in the red fruits compared to the control plants. It is found that sucrose is the more decreased sugars. The reductions are significant about 54% compared to controls. The level of organic acid in red fruits decreased with 9%, the malic acid decreased significantly more is about 33% compared to citric acid. The water deficit increased the carotenoid content in red fruits of tomato plants. In contrast to green fruits, Beta-carotene and Lutein were present in red fruits, with significantly higher values compared to control fruit. This reduction is respectively 63% and 73%. Which suggesting the water deficit increasing the synthesis of carotenoids in mature fruits. Moreover, in the presence of water deficit, the Vitamin C was accumulated in red fruits less than green fruits. This significant increase is of the order of 38% compared to control red fruit.

DISCUSSION

Water deficit causes numerous disturbances of many functions of the cell and the whole plant. Sensing mechanisms, yet to be identified, initiate the responses to water deficit, which occur at the molecular, metabolic, cellular, physiological, and developmental level. In the present work, we will study the effect of water deficit on the tomato crop to understand the relationship between the water deficit determined by a restriction on the amount of water and the development of the plant and the quality of the fruit. Leaf is an essential organ because it has an important role in the regulation of respiration, in the synthesis of organic matter contributing to nutrition and therefore to the growth of the plant. The response of the plant to environmental conditions and more particularly to the water deficit, which is the most limiting stress of growth, has an impact on the leaf of tomato structure. It differs according to the nature and the duration of the stress (Anjum et al., 2011). The present study showed that after 3 months, water deprivation decreased number, widths, and length of leaves (Figure 1). With regard to the effect of WD on leaf area, the treatment resulted in a

### TABLE 3

| Phase of cell division | Cell expansion phase | Maturation phase |
|------------------------|----------------------|------------------|
| **Diameter (mm)**      |                      |                  |
| CT                     | 6.17 ± 0.91a         | 31.73 ± 4.21a    |
| ST                     | 4.76 ± 1.07b         | 34.82 ± 1.13b    |
| **Height (mm)**        |                      |                  |
| CT                     | 7.12 ± 1.12a         | 32.4 ± 3.38a     |
| ST                     | 5.78 ± 1.01b         | 35.34 ± 2.09b    |
| **Fresh weight (g)**   |                      |                  |
| CT                     | 0.17 ± 0.07a         | 19.99 ± 6.77a    |
| ST                     | 0.09 ± 0.04b         | 15.65 ± 4.90b    |
| **Dry weight (g)**     |                      |                  |
| CT                     | 0.02 ± 0.01a         | 1.52 ± 0.53a     |
| ST                     | 0.01 ± 0.005b        | 1.47 ± 0.54a     |
| **Water content**      |                      |                  |
| CT                     | 8.36 ± 0.54a         | 21.56 ± 6.63a    |
| ST                     | 4.43 ± 0.81b         | 9.83 ± 1.20b     |

Note: Bar indicates standard error (p = .05). Same letter above the bars denotes that the difference between means were not significant.

### FIGURE 5

Ca^{2+} and K⁺ contents in the fruit of tomato plants Solanum lycopersicum (cv. Plovdiv) cultivated in the presence of water (CT: control treatment) or under water restriction conditions (ST: stressed treatment). Bar indicates standard error (p = .05)
TABLE 4 Relative differences in metabolite contents (soluble sugars, organic acids, ascorbic acid (AsA) and carotenoids) in the fruit red and green of tomato plants Solanum lycopersicum (cv. Plovdiv) cultivated in the presence of water (CT: control treatment) or under water restriction conditions (ST: stressed treatment), Bar indicates standard error (p = .05)

|                      | Green fruits |                      | Red fruits |                      |
|----------------------|--------------|----------------------|------------|----------------------|
|                      | CT           | ST                   | CT         | ST                   |
| Glucose              | 13.12 ± 2.48a| 10.18 ± 1.21b       | 21.21 ± 1.3a| 20.89 ± 1.27a       |
| Fructose             | 11.16 ± 3.17a| 9.03 ± 0.9b         | 18.58 ± 1.1a| 17.17 ± 1.56a       |
| Sucrose              | 0.91 ± 0.17a | 1.20 ± 0.17b       | 1.12 ± 0.08a| 2.47 ± 0.40b        |
| Total sugars         | 25.02 ± 5.81a| 20.42 ± 1.80b      | 40.91 ± 2.37a| 40.32 ± 2.54a       |
| Citric acid          | 2.75 ± 0.53a | 3.04 ± 1.1a        | 3.38 ± 0.31a| 3.65 ± 0.12a        |
| Malic acid           | 1.88 ± 0.34a | 1.61 ± 0.53a      | 2.53 ± 0.29 a| 1.69 ± 0.35b        |
| Total acid           | 4.00 ± 3.11a | 4.65 ± 1.62a      | 5.90 ± 0.54a| 5.34 ± 0.23a        |
| Lycopene             | 3.16 ± 0.43a | 4.17 ± 1.21b      | 0.37 ± 0.05a| 0.32 ± 0.05a        |
| Beta-carotene        | –            | –                   | 22.65 ± 8.82a| 61.51 ± 8.82b       |
| Phytoene             | 2.00 ± 0.15a | 2.62 ± 0.29b      | 1.82 ± 0.24a| 2.82 ± 0.26b        |
| Lutein               | –            | –                   | 7.12 ± 3.22a| 27.90 ± 4.94b       |
| Total carotenoids    | 5.16 ± 0.84a | 6.78 ± 1.99b      | 31.12 ± 0.18a| 92.23 ± 0.58b       |
| Total AsA            | 18.09 ± 5.70a| 22.80 ± 3.79b     | 13.24 ± 1.40a| 21.36 ± 1.70b       |

Same letter above the bars denotes that the difference between means were not significant.
vitamin C content in the leaves of treated plants as control (Figure 4), this may be confused with the hypothesis that AsA is not limited by photosynthesis or sugar availability, but because of environmental condition (Gautier et al., 2008). Similar research showed that water deficit increased Vitamin C levels in the leaves, suggesting a positive interaction of ascorbic acid in the protection of chloroplasts and other cell compartments (Zhu, 2001). On leaves, vitamin C content is high in young leaves still growing and decreases in mature and pre- nescent leaves (Bulley et al., 2009; Chen et al., 2003; Li et al., 2010).

Fruit fresh and dry masses, diameter, height, water content, mineral nutrition, concentration in soluble sugars, organic acids, AsA, and carotenoids were measured during the development phases of the tomato fruit (Phase of cell division, cell expansion phase, and maturation phase). Several studies reported the negative effect of water deficit on cell division on many species such as grape berries (Vitis vinifera), (McCarthy et al., 2002; Ojeda et al., 2001) and olives (Olea europaea L), (Gucci et al., 2009). In the present study, the water deprivation for three months decreased the fruit diameter, height, and water content respectively as compared to control tomato plants. Similar work (Wang & Gartung, 2010) reported that water deficit decreased fruits seize, weight, and water content, indeed, that fruit decreased proportionally to the intensity of WD in peach trees. Also, in our results, we found that WD caused a reduction in the weight of fruit as well the dry matter of tomato fruit respectively by about 43% and 26% as compared to control tomato plants (Table 3). The application of the water deficit throughout the tomato fruit development phase (cell division phase, cell expansion phase, and the maturation phase) induces a decrease in the concentration of K⁺ and Ca²⁺. The effect of WD on the mineral composition of tomato fruits remains poorly documented in literature. Soluble sugars and organic acids (primarily malic and citric acids) are major osmotic compounds that accumulate fleshy fruits (Ripoll et al., 2014). Acids citric, malic acids which were found in fresh tomato fruit, promotes gastric secretion, acts as a blood purifier and works as intestinal antiseptic (Pruthi, 1993), and determine taste and represent more than half of the total dry matter in tomatoes. Generally, the predominant organic acid in ripe fruits varies between species (Etienne et al., 2013). The sugars that have accumulated in stressful conditions and can be simple sugars (glucose, fructose), alcohol sugars (glycerol, sorbitol, pinnitol) or complex sugars (trehalose raffinose and fructans) (Bohnert & Jensen, 1996.), In the present study, fruit composition in soluble sugars, organic acids, carotenoids and vitamin C were determined for green and red fruits. The water deficit was at the origin of the decrease in the accumulation of organic acids in green fruits (Table 4). Similar studies on tomatoes have shown a decrease in sugars under the conditions of WD applied during the cell division phase (Ripoll et al., 2016). Plant tissues accumulate organic acids (Hummel et al., 2010) to reduce their osmotic potential and prevent the reduction of cell turgor pressure. The accumulation of solutes or osmolytes helps maintain an osmotic balance at the cell level under dehydration conditions (Bray et al., 2000). Vitamin C is present in all plants and compartments as it plays important roles in plants. It is very essential for plant growth and development. In our study, the water deficit increased the vitamin C content of the green and red fruit of the treated plant as a control. Application of water stress during the ripening phase increased vitamin C levels (Table 4). The results suggest a positive interaction of ascorbic acid in cell compartments (Zhu, 2001). Tolerance to water deficit is correlated with AsA accumulation, which plays an important role in ROS (reactive oxygen species) detoxification (Wang et al., 2012). Indeed, vitamin C is a major antioxidant for the plant, capable of neutralizing the active forms of oxygen. These results were similar to those of Ripoll et al. (2016) who found that the application of water stress during the maturation phase increased the vitamin C levels. The results suggest a positive interaction of ascorbic acid in cell compartments (Zhu, 2001). In the plant, carotenoids, secondary pigments collecting light energy, antioxidants and precursors of hormones (abscisic acid and strigolactones), contribute to several major physiological functions and contribute to the adaptation of the plant to micro climatic variations (Gomez-Roldan et al., 2008). In this study, the application of water deficiency during the ripening phase affected the composition of Plovdiv fruits. The composition of fruits in organic acid and soluble sugars has been barely modified by the water deficit due to a decrease in malic acid and glucose and fructose, similar results have been reported by (Ripoll, et al. 2016). In the plant, carotenoids, secondary pigments that collect light energy, antioxidants, and hormone precursors (abscisic acid and strigolactones), contribute to several major physiological functions and contribute to the adaptation of the plant to micro-climatic variations (Gomez-Roldan et al., 2008). In this study, water deficit increased carotenoid content (Table 4) for green and red fruits. Similar results were observed in tomatoes (Ripoll et al. 2016). In addition, De Pascale et al. (2007) showed that carotenoids and ascorbic acid were involved in the detoxification of reactive oxygen species that accumulate in response to different constraints. Variations in carotenoids and AsA would result from stress-induced cellular redox changes (Fanciullino et al., 2014).

5 | CONCLUSION

In the present study, water deficit treatment induced independent response pathways in leaves and fruits. Reduction of dry matter production, reduction of leaf area to reduce the amount of water used. water deficit reduces the accumulation of mineral ions in the leaves. Significant decrease in the levels of organic acids and vitamin C in the leaves. At the fruit level, water stress induces a reduction in fruit size, moisture content and dry matter production during the three phases of fruit development. The nutritional quality of the fruit is not affected by the water deficit. In general, soluble sugars and organic acids are stable. The nutritional quality associated with carotenoids and AsA levels could be positively affected. The action of the water deficit on tomato varieties (Plovdiv) is negative. We have also noted a certain regulation of growth to reduce the amount of water used. In addition, moderate moisture defect affects the nutritional...
distribution of leaves and fruits. In addition, the water deficit improves the production of carotenoids and vitamin C in fruits.

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CONFLICT OF INTEREST
The authors declare that the research was conducted in the absence of any commercial or relationships that could lead to a conflict of interest.

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