INTRODUCTION

The *Raphanus sativus* L. is a root vegetable widely grown for its nutritional, culinary and medicinal uses that belongs to Brassicaceae. It grows in temperate climates at altitudes between 190 and 1240 m. Its roots are thick and of various sizes, forms, and colors. Red Radish belongs to the variety radicular. The edible fleshy axis is derived from the hypocotyl and upper radicle tissues. Radish is widely used in salad preparations and contains a considerable amount of antioxidants, vitamin C, and health-promoting compounds such as glucosinolates and phenolic compounds. Various parts of the radish plant, including roots, seeds, and leaves, have been used for medicinal purposes. Its health-promoting properties have been attributed to polyphenolic compounds. *R. sativus* is popularly used to treat liver and respiratory illnesses (Paredes, 1984), reduce cancer development (Ku *et al.*, 2008) and contains a range of digestive enzymes (Cho *et al.*, 2009). A comprehensive review of the plants’ active constituents and its therapeutic properties could be found in the publication of (Gutierrez & Perez, 2004). Radish has also been used in naturopathic medicine as a laxative, stimulant, and digestive aid, as well as in the treatment of stomach disorders (Kapoor, 2000). Red radish cultivars are a potential source of natural colorants due to the presence of anthocyanins, which have high stability. Anthocyanins have well-known health benefits, including the ability to scavenge free radicals, inhibit cancer and diabetes, prevent neuronal and...
cardiovascular diseases, and suppress inflammation (Hwang et al., 2012). Great attention is being given to post-harvest quality of vegetables due to rising consumer awareness about diet-health connections. Quality parameters of stored vegetables are determined by both pre- and postharvest factors (Kader, 2003). So, it is assumed that manipulation of growth and metabolism of plants during preharvest stage may have significant impact on postharvest quality of their stored products.

Humic acid (HA) are the most significant constituents of organic matter in soils and have a relevant role in the cycling of many elements in the environment and in soil ecological functions (Senesi et al., 1996). Foliar sprays of HA promote growth, increase yield and quality in a number of plant species (Karakurt et al., 2009). Moreover, humic acid influence respiration process, the amount of sugars, amino acids and nitrate metabolism (Boehme et al., 2005). Nardi et al. (2002) characterized the effects of HA on physiology of higher plants. According to them, HA positively influencing the uptake of some nutrients, especially nitrate, may influence both respiration and photosynthesis, may display a hormone-like activity, and exhibit stimulatory effects on plant cell growth and development.

Seaweed products exhibit growth-stimulating activities, and the use of seaweed formulations as bio modulators in crop production is well established. Seaweed components such as macro- and microelement nutrients, amino acids, vitamins, cytokinins, auxins, and abscisic acid (ABA)-like growth substances affect cellular metabolism in treated plants leading to enhanced growth and crop yield (Ordog et al., 2004). In addition, it has been reported that seaweed extract enhanced the plant’s antioxidant defense system of wheat plants subjected to drought stress through elevating both the enzymatic and non-enzymatic components of the plant’s antioxidant defense system (Kasim et al., 2015).

Essential oils are well known for their antioxidant and antimicrobial properties that prevent food degradation and alteration (Justesen & Knuthsen, 2001). They possess antimicrobial properties and prevent growth of bacteria, fungus and virus during storage and protect cells from senescence, thereby increase shelf life of vegetables. Thyme oil contains more than 40% phenolic constituents (thymol, carvacrol and caffic acid) which have strong antiseptic effects. Essential oils chemical composition of Thymus vulgaris and Mentha piperita were identified by GC-MS where the most components of thyme were thymol (44.7%), cymin (18.6%) and tripenene (16.5%) whereas menthol and menthane are the main components of peppermint oil (Porte & Godoy, 2008). Application of 4% thyme and basil oils had a positive effect on broccoli seed germination, which was ascribed to their antioxidant and disinfecting effects (Nguefack et al., 2005).

The objectives of the current investigation was to assess the effect of bio stimulators seaweed extract (SWE), humic acid (HA), thyme oil as well as peppermint oil on certain quality-related biochemical constituents during both pre- and post-harvest stages.

MATERIALS AND METHODS

Two field experiments were conducted during the two consecutive growing seasons of 2015 and 2016 at EL-Bramoof Farm, Mansoura city, Dakahlia Governorate. Seeds of Raphanus sativus L. (var. Cherry red) were secured from Vegetables Department, Horticulture Research Institute, ARC, Egypt and planted on October 22nd during both growing seasons. Soil samples at 30 cm depth were taken for estimating the experimental soils physical and chemical characteristics (Table 1) according to (Black et al., 1965).

**Table 1.** Physical and chemical characteristics of the experimental soil (average of the two growing seasons).

| Texture | Organic matter% | Total carbon% | Ec (dsm-1) | pH | N% | P% | K% |
|---------|----------------|---------------|------------|-----|-----|-----|-----|
| Clay    | 1.8            | 1.82          | 0.75       | 8.65 | 0.19 | 0.015 | 0.14 |

Experimental design

The experimental design was randomized complete block design with three replicates. Each replicate contained three rows, six meters in length and 3 m in width, with the total experimental area of 162 m². Within the row, the distance between plants was 20 cm.

Sampling procedures

Plants were sprayed twice, 20 and 30 days after sowing (DAS) onto foliage with solutions of Seaweed Ecklonia maxima extract (SWE) at the rate of 4 ml L⁻¹, Humic acid (HA) at the rate of 30 ml L⁻¹, Peppermint oil (P.O) at the rate of 50 ml L⁻¹, and Thyme oil (T.O) at the rate of 50 ml L⁻¹. Control plants were sprayed with tap water. Tween 20 was added to sprayed solutions at the rate of 0.05 % as a wetting agent. Seaweed extract and humic acid were obtained from Shouara Company, Egypt. Peppermint and thyme oils were obtained by hydro-distillation for 2-3 h in the Post-harvest Lab., Hort. Res. Station, Agric. Res. Cent., Egypt, using modified Clevenger apparatus according to (Guenther, 1965). Samples were collected 40 DAS to determine the contents of anthocyanin, total phenols, total flavonoids, ascorbic acid, total carbohydrates, total free amino acids as well as the activity of Peroxidase and Polyphenol oxidase.

Anthocyanin determination was determined according to (Chiriboga & Francis, 1973). Total phenols were measured calorimetrically at 650 nm using Folin-ciocalteu reagent according to (Bray & Thorpe, 1954). Total flavonoids content was estimated calorimetrically at 510 nm as described by Heimler(2006). Ascorbic acid was determined by the 2,6-Dichlorophenolindophenol dye procedure of (Freed, 1966). Total carbohydrates were determined using phenol sulphuric acid method according to (Dubois et al., 1956). Total free amino acids were determined according to the procedure of (Salnikow et al., 1996).
Peroxidase extraction and assay were performed according to the procedure of (Maxwell & Bateman, 1967). Polyphenol oxidase (PPO) extraction and assay were performed according to (Galeazzi et al., 1981).

Nutrients concentration

Sulfur estimated by using atomic absorption spectroscopy (AAS) was subjected by Kirkbright & Wilson, (1974). Potassium was determined (mg/g) by using flame photometer (Model 400M.X) according to (Chapman & Pratt, 1961).

Storage parameters

Plants were uprooted 60 DAS and samples from roots (ten roots from each replicate) were packed in polyethylene bags and stored at controlled storage conditions (5˚C, relative humidity 95%) for three months. Afterwards, postharvest quality and storage parameters represented by weight loss percentage, post-harvest decay percentage and dry matter percentage were estimated.

Weight loss percentage (WLP %) was estimated according to the following equation:

\[ \text{WLP} \% = \left( \frac{\text{Initial} - \text{Final Weight}}{\text{Initial Weight}} \right) \times 100 \]

Post-harvest decay percentage (PDP %) was estimated according to the formula:

\[ \text{Number of Decayed Fruits/ Number of Total Fruits} \times 100 \]

Dry matter percentage (DMP %) was estimated as described by Lee (1981). After drying in the oven at 60˚C until weight stability, moisture percentage was calculated according to the equation.

Moisture content percentage

Fresh weight of root before storage – Dry weight of root after storage / Dry weight of root after storage × 100; then dry matter (%) was estimated as (100 - % moisture percentage).

Statistical analysis

Data were subjected to analysis of variance using GENSTATE software (version 11.1.0.1575). To compare differences between means, least significant differences (LSD) at 5% were calculated according to (Gomez et al., 1984). Correlation matrix between biochemical analysis after storage in refrigerator at (5˚C/Rh 95%) and storage parameters at the 0.05 and 0.01 level (2-tailed) on PASWStatistics18 software. The sign of the correlation coefficient determines whether the correlation is positive or negative. The magnitude of the correlation coefficient determines the strength of the correlation subjected by Evans, (1996) suggests absolute r-value 0.00-0.19 (very weak), 0.20-0.39 (weak), 0.40-0.59 (moderate), 0.60-0.79 (strong) and 0.80-1.0 (very strong).

RESULTS AND DISCUSSIONS

All applied bio modulators increased total anthocyanin’s concentration in the roots either pre- or post-harvest compared to control (Table 2). The highest anthocyanins concentrations were obtained in response to treatment with thymus oil followed by humic acid in pre-harvest stage. In roots that stored in the specified storage experimental conditions, total anthocyanin concentration was decreased, either in biomodulators-treated or untreated plants compared with before storage. However, in post-harvest stage anthocyanins concentrations were highest in response to the treatment with peppermint oil followed by seaweed extract i.e. peppermint oil and seaweed extract caused the least decline in anthocyanins concentration in stored roots, in order. Data presented in table (2) show that ascorbic acid concentration in pre-stored roots was increased in response to thyme oil and seaweed extract, though the increase was insignificant in case of seaweed extract treatment. On the other hand, humic acid and peppermint oil did not affect ascorbic acid concentration in the roots before storage. In stored roots, ascorbic acid concentration was decreased either in bio modulators-treated or untreated plants. However, the magnitude of decline was lower in roots of bio modulators-treated plants. So, after storage, ascorbic acid concentration in roots was significantly higher in all biomodulators treatments compared with control. The lowest loss percentage was recorded in response to peppermint oil (29.7 %) followed by humic acid (31.7 %).

Table 2. Effect of applied bio modulators on total anthocyanin and ascorbic acid content (mg g⁻¹ F.W) in roots of *Raphanus sativus* plants before and after storage.

| Parameters | Treatments | Anthocyanin (mgg⁻¹.F.W) | Ascorbic acid (mgg⁻¹.F.W) |
|------------|------------|-------------------------|---------------------------|
|            | Before     | After                   | Total loss%               | Before       | After       | Total loss% |
| Seaweed extract | 8.34       | 3.71                    | 55.52                     | 24.13        | 13.97       | 42.11        |
| Humic acid  | 8.51       | 2.65                    | 68.86                     | 20.50        | 14.00       | 31.71        |
| Peppermint oil | 8.22       | 4.54                    | 44.77                     | 20.52        | 14.42       | 29.71        |
| Thyme oil   | 8.55       | 2.60                    | 69.59                     | 37.10        | 22.57       | 39.16        |
| Control     | 7.16       | 1.44                    | 79.89                     | 20.41        | 9.23        | 78.54        |
| LSD at 0.05 | 0.60a      | 0.29d                   | -                         | 6.32a        | 1.50b       | -            |

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Table 3. Effect of applied bio modulators on total phenols and flavonoids (mg g⁻¹ D.W) roots of Raphanus sativus plants before and after storage.

| Parameters     | Treatments             | Total phenols (mg g⁻¹ F.W) | Total flavonoids (mg g⁻¹ F.W) |
|----------------|------------------------|-----------------------------|--------------------------------|
| Before         | After                  | Total loss%                 | Before                        | After                  | Total loss%                 |
| Seaweed extract| 14.49                  | 12.08                       | 16.63                         | 0.33                   | 0.29                        | 12.12                        |
| Humic acid     | 13.41                  | 10.78                       | 19.61                         | 0.40                   | 0.35                        | 12.50                        |
| Peppermint oil | 14.85                  | 12.45                       | 16.16                         | 0.39                   | 0.34                        | 12.82                        |
| Thyme oil      | 13.00                  | 10.42                       | 19.85                         | 0.47                   | 0.44                        | 6.38                         |
| Control        | 12.50                  | 9.51                        | 23.92                         | 0.31                   | 0.26                        | 16.13                        |
| LSD at 0.05    | 2.30⁻                   | 1.02⁻                       | -                             | 0.01⁻                  | 0.03⁻                       | -                             |

Table 4. Effect of applied biomodulators on total carbohydrates (mg g⁻¹ D.W) and free amino acids (mg/100g.D.W) in dried roots of Raphanus sativus plants before and after storage.

| Parameters     | Treatments             | Total carbohydrates (mg g⁻¹ D.W) | Total free amino acid (mg/100g.D.W) |
|----------------|------------------------|---------------------------------|-------------------------------------|
| Before         | After                  | Total loss%                     | Before                        | After                  | Total loss%                     |
| Seaweed extract| 127.50                 | 104.80                         | 17.80                         | 12.09                  | 5.64                        | 53.35                         |
| Humic acid     | 127.03                 | 90.83                         | 28.50                         | 12.49                  | 4.99                        | 60.05                         |
| Peppermint oil | 126.60                 | 101.92                        | 19.49                         | 13.56                  | 4.34                        | 67.99                         |
| Thyme oil      | 112.12                 | 84.33                         | 27.46                         | 12.66                  | 5.20                        | 58.93                         |
| Control        | 110.90                 | 79.00                         | 28.76                         | 12.03                  | 3.30                        | 72.57                         |
| LSD at 0.05    | 22.64⁻                  | 25.74⁻                        | -                             | 3.73⁻                  | 0.49⁻                       | -                             |

Table 5. Effect of applied biomodulators on potassium concentration (mg g⁻¹ D.W.) and sulphur percentage in dried roots of Raphanus sativus plants before and after storage.

| Parameters     | Treatments             | Potassium (mg g⁻¹ D.W.) | Sulfur (%) |
|----------------|------------------------|-------------------------|------------|
| Before         | After                  | Total loss%             | Before      | After                  | Total loss%             |
| Seaweed extract| 12.48                  | 11.00                   | 11.86      | 1.43                   | 0.93                    | 34.97                      |
| Humic acid     | 13.00                  | 10.00                   | 23.08      | 1.20                   | 0.90                    | 25.00                      |
| Peppermint oil | 13.64                  | 10.09                   | 26.03      | 1.50                   | 1.15                    | 23.33                      |
| Thyme oil      | 13.26                  | 10.07                   | 24.06      | 1.29                   | 1.01                    | 21.71                      |
| Control        | 12.37                  | 9.00                    | 27.24      | 1.10                   | 0.50                    | 54.55                      |
| LSD at 0.05    | 0.15⁻                  | 0.71⁻                   | -          | 0.48⁻                  | 0.35⁻                   | -                           |

All applied bio modulators increased total flavonoids in roots during both pre- and post-harvest stages as well as total phenols only during post-harvest stage (Table 3). The only bio modulator that increased total phenols in the roots before storage was Peppermint oil. Both total flavonoids and phenols were decreased after storage, but the decrease was lower in case of total flavonoids. The lowest decrease in total phenols was recorded in response to Peppermint oil whereas the least decrease in total flavonoids was recorded in response to thyme oil treatment.

Generally, total carbohydrates concentration in roots before storage was increased in response to bio modulators treatments, though the increase was insignificant (Table 4). In addition, roots of bio modulators-treated plants after storage contained higher concentrations of total carbohydrates compared with control roots, though the differences did not reach the significance level. In all treatments, total carbohydrates concentrations were decreased after storage, with the least decrease in response to seaweed extract followed by peppermint oil treatment.

Total free amino acids concentrations in roots treated with all bio modulators before storage were not significantly different than that in control roots (Table 4). On the other hand, roots of bio modulators-treated plants after storage contained significantly higher concentrations of total free amino acids compared with control. Storage caused a decrease in total free amino acids concentrations in roots of bio modulators-treated as well as control plants. In stored roots of seaweed extract-treated plants, the decrease in concentration of total free amino acids was the least, followed by thyme oil.
Roots of biomodulators-treated plants either before or after storage contained higher concentrations of both potassium (K) and sulphur (S) compared with control roots (Table 5). However, the recorded increase in S concentration before storage was insignificant in response to all biomodulators treatments. In roots of biomodulators-treated as well as control plants, concentrations of both potassium K and S were decreased after storage. The decrease was of the least magnitude in response to seaweed extract treatment in case of K and in response to peppermint oil in case of S.

In roots of biomodulators-treated plants, the activity of peroxidase (POD) was increased either before or after storage compared with that in roots of control plants. A similar trend in the activity of polyphenol oxidase (PPO) was recorded in roots after storage. On the other hand, before storage, PPO activity in roots was increased in response to seaweed extract and peppermint oil treatments whereas decreased in response to humic acid and thyme oil treatments. Storage affected the activity of both POD and PPO, but in different manner. Where POD activity was decreased after storage, PPO activity was increased (Fig. 1). The least decrease in POD activity was recorded in response to thyme oil treatment, whereas the highest increase in PPO activity was achieved in response to humic acid treatment.

**Figure 1.** Quality parameters of stored roots of *Raphanus sativus* plants after storage for three months in refrigerator at 5˚C and 95% relative humidity as affected by biomodulators treatments.

**Effect of biomodulators on post-harvest quality parameters of stored roots**

After two months-storage, stored roots had visible decay symptoms manifested by rotting, water stains and unusual smell due to growth of bacteria and fungus on roots. However, quality parameters of stored roots were differed between biomodulators treatments and control. All applied biomodulators decreased weight loss and post-harvest decay percentages hence, roots of biomodulators-treated plants contained higher dry matter compared with the roots of untreated plants. Weight loss as well as post-harvest decay percentage was minimum in response to peppermint treatment, followed by thyme oil treatment. Nevertheless, dry matter percentage in stored roots was highest due to seaweed extract followed by humic acid treatments (Figure 2).

**Pearson’s correlation matrix between roots biochemical analyses after storage and their post-harvest quality parameters**

In table (6) showed the output of Pearson’s correlation matrix between biochemical analysis after storage in refrigerator at (5˚C/Rh 95%) for three months and storage parameters of radish plants in first season 2015. Dry matter and post-harvest decay showed very strong significant negative as follow (-0.90 and -0.99) respectively relationship with sulfur concentration at the 0.05 and 0.01 level. Total anthocyanin revealed very strong significant positive as follow (0.98 and 0.92) relationship with total phenols and carbohydrates at the 0.05 and 0.01 level. Ascorbic acid, total phenols, flavonoids and anthocyanin recorded very strong significant positive as follow (0.95, 0.97, 0.89 and 0.91) respectively relationship with total flavonoids, carbohydrates, potassium concentration and peroxidase enzyme at the 0.05 and 0.01 level.
Figures 2. Effect of applied bio modulators on the activity (U g⁻¹ F.W.) of Peroxidase (POD) and Polyphenoloxidase (PPO) in roots of *Raphanus sativus* plants before and after storage.

Table 6. Pearson's correlation matrix between roots biochemical analyses after storage and their post-harvest quality parameters.

| Biochemical Parameters | DM% | PDP% | WLP% | T.A  | ASA  | T.P  | T.F  | T.C  | T.A  | K    | S    | POD  | PPO  |
|------------------------|-----|------|------|------|------|------|------|------|------|------|------|------|------|
| DM%                    | 0.88| -0.30| -0.61| -0.83| -0.57| -0.73| -0.52| -0.57| -0.35| -0.48| -0.90| -0.74| -0.72|
| PDP%                   |     |      |      |      |      |      |      |      |      |      |      |      |      |
| WLP%                   |     |      |      |      |      |      |      |      |      |      |      |      |      |
| T.A                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| ASA                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| T.P                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| T.F                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| T.C                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| T.A                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| K                      |     |      |      |      |      |      |      |      |      |      |      |      |      |
| S                      |     |      |      |      |      |      |      |      |      |      |      |      |      |
| POD                    |     |      |      |      |      |      |      |      |      |      |      |      |      |
| PPO                    |     |      |      |      |      |      |      |      |      |      |      |      |      |

Correlation is significant at the 0.05 level (2-tailed). DM: dry matter; PDP: post-harvest decay; WLP, weight loss; TA, total anthocyanin; TP, total phenols; ASA, ascorbic acid; TF, total flavonoids; TC, total carbohydrates; TA, total amino acids; K, potassium; S, sulphur; POD, peroxidase; PPO, polyphenoloxidase.

Senescence is the most important internal factor that causes damage to vegetables whereas low temperature is the best way to delay post-harvest deterioration of fruits and vegetables. In this context, (Bayouni, 2008) concluded that the higher post-harvest decay percentage in late harvesting stage of fruits is due to higher rate of respiration, more skin permeability for water loss and high susceptibility to decay. Dry matter acts an index to evaluate the fruit quality due to endure prolonged storage (Jackson & Harker, 1997). Generally, low temperature can maintain organic matter and vitamins levels in short and long term storage because most enzyme activities decrease and the expression of many genes inhibited during cold storage. Low temperature markedly delays senescence of broccoli during storage (Javanmardi & Kubota, 2006). In addition, (Shen et al., 2013) reported that refrigeration can reduce deterioration and extend shelf life of fresh fruits by delaying the metabolic processes. Dry matter production represents a balance between photosynthesis and respiration. Respiration rate, which has long been used to measure metabolic process in stored produce (Kittosc & Law, 1968; Scholz, 1962), is governed by water availability, temperature, O₂ concentration, microbial contamination, mechanical damage, among other factors. Porter et al., (2003) reported that respiration rate of chinese cabbage was higher at 20°C than at lower temperatures.

Synthetic chemicals are commonly used in the control diseases of stored plant products however; these chemicals may cause toxic residues in treated plants (Isman, 2000). Alternatively, the use of natural products like herbal extracts and essential oils can reduce harm to public health and preserve the environment from pollution. These eco-friendly plant-derived products generally possess a broad spectrum of activity against several pathogens and pests (Bakkali et al., 2008) Essential oils are natural complex...
compounds with a strong smell and produced as secondary metabolites in aromatic plants. In nature, essential oils act as antibacterial, antiviral, antifungal and insecticidal. It contains compositions of terpenes, sesquiterpenes, aldehydes, ketones and phenolic compounds (Burt, 2004). The main chemical components of thyme oil are thymol, carvacrol, α-pinene, β-pinene, borneal, linalool, β-simine and camphene (Dew, 1984). The composition of volatile oil of M. piperita is menthol, monoterpen and methofuran (Dew, 1984) These constituents act as inhibitor for pathogens growth during cold storage and reduce senescence in treated plants.

In the current investigation, thyme as well as peppermint oil decreased weight loss and post-harvest decay percentages hence, preserved roots dry matter during cold storage (Figure 2). These findings are in harmony with those of previous studies (Aminifard & Mohammadi, 2013; Alkhani et al., 2009; Amin, 2016; Bakry et al., 2013; Geransayeh et al., 2015). Antioxidants play an important role in scavenging reactive oxygen species (ROS) that appear during storage-related senescence of vegetables (Hounsome et al., 2009). Antioxidant capacity is directly correlated with phenolic compounds (Koh et al., 2009; Tavarini et al., 2008). Therefore, plants with enhanced levels of antioxidants can resist oxidative damage (Navarro et al., 2006). The results of the present investigation revealed that generally, the applied biomodulators diminished the decline in concentrations of anthocyanins, ascorbic acid, phenols and flavonoids as well as in the activity of the antioxidant enzyme, peroxidase. This implies that stored roots of biomodulators-treated plants contained higher levels of antioxidants compared with roots of untreated plants. Similar conclusions were made based on the results of previous studies (Del Nobie et al., 2007; Kramchote et al., 2012; Wang et al., 2008; Znidaric & Pozrl, 2006). How elevated levels from these constituents preserve roots quality during storage could be discussed as follows:

Anthocyanin belongs to flavonoids group and located in a cell vacuole (Friend & Rhodes, 1981). It plays important role as an antioxidant. The anthocyanin pigment inhibits active oxygen radicals formed by exposure to stress conditions that occur during storage periods due to pathological injuries i.e., enzymatic browning, molds and water stais on fruits. The results of some in vitro experiments indicated that anthocyanin pigments scavenges superoxide, active oxygen radicals (Yamasaki et al., 1996) and hydrogen peroxide (Leng & Qi, 2003). On the other hand, anthocyanin pigment is affected by the amount of sugar due to a lower water activity (Hounsome et al., 2009). Andreu et al. (1988) reported that anthocyanins concentration was decreased at the end of the cold storage period, which implies that they are consumed in scavenging storage-related ROS.

Ascorbic acid is a major component of the plant defense system (Hodges & Forney, 2000) that protect biological processes of the plant from ROS generated during biotic and abiotic stresses (Foyer & Noctor, 2011). In addition, it has a pivotal role in eliminating H2O2 through the glutathione-ascorbate cycle that operates in the cytosol, mitochondria, plastids and peroxisomes where it involves ascorbate, glutathione, NADPH and the enzymes linking these metabolites during long term storage. Since glutathione, ascorbate and NADPH are present in high concentrations in plant cells it is assumed that the glutathione-ascorbate cycle plays a key role for H2O2 detoxification. Peroxidases also contribute to H2O2 removal in plants. The destruction of ascorbic acid is one of the most serious problems facing plants under stress conditions, whether in the soil or when storing in the short or long term (Jeney et al., 2008). The oxidized form of ascorbic acid is more prone to decomposition during storage in low temperature. Ascorbic acid decreases during storage due to its consumption in the ROS-detoxification process especially at temperature above 0 °C (Aijobila et al., 2009) through oxidation in the presence of ascorbate oxidase enzyme, and changes backto its active form (Lee & Kader, 2000). The antioxidant role of ascorbic acid in mitigating stress-related metabolic abnormalities was evident from the results of other studies (Al-Karaki, 2000; Aminifard & Mohammadi, 2013; Raafat et al., 2012; Serrano et al., 2005; Wang et al., 2008).

Post-harvest stresses e.g. mechanical injuries and others during storage led to enzymes activity that responsible for phenolic compounds deterioration (Yang et al., 2011). Phenolic compounds and enzymes do not interact with each other because of their different locations within cells consequently; shelf life of vegetables and fruits is increased. However, when membranes are damaged, destruction and oxidation of phenolic compounds are initiated, causing breakdown of defense line in the cell against free radicals (Toor & Savage, 2006; Yang et al., 2011). Similar conclusions were made by (Amin, 2016; Wang et al., 2008). Flavonoids are the most important group of phenolic compounds in plants (Hounsome et al., 2009) that assist in scavenging free radicals and inhibiting oxidative stress, thereby decreasing cell membrane deterioration during long storage (Koh et al., 2009) which low temperature could decrease the membrane lipid degradation and POD activity as a result, it can slow down flavonoids oxidation. These results are in agreement with those obtained by (DuPont et al., 2000) on lettuce, (Yuan et al., 2010) on broccoli plants and (Geransayeh et al., 2015) on strawberry fruits.

Carbohydrates are among the most prevalent organic compounds and represent the most important sources of energy in the plant. This energy from the sun was obtained in photosynthesis process (anabolism) and it is released through respiration (catabolism). Finally, the carbohydrates serve as the starting compounds for the biogenesis of many other cell constituents. Total carbohydrates represent the production of both sugars and starch within the cell. This study showed increased photosynthesis and increased plant growth rate with adding HA and SWE treatments in the open field led to an increase in the rate of carbohydrate content. These results are in harmony with those recorded by Bakry et al. (2013) on two wheat cultivars (Tehranifar
B. (1977). The effects of applied in the field experiment led to a significant increase in storage of fruits due to increased carbohydrates anabolism process within plant cells. (Blunden & Wildgoose, 1977) mentioned that seaweed extract applied on potato plants recorded significantly increased in the yield of tubers for both the varieties King Edward and Pentland Dell due to plants hormones such as cytokinins represent the most important compound in SWE that affect on all phytochemicals before storage on the other hand, oxidative damage, enzymatic browning enzymatic browning is responsible for the degradation of phenolic compounds especially with long storage time at specified temperature where this phenomenon and increased the rate of respiration cause decreasing phytochemicals which increased within storage intervals. Accordingly, the rate of broccoli plants respiration effect on carbohydrate levels when storage in cold temperature at 5°C. These results are in agreement with those reported by Vigna radiate plants that also supported by Majidi et al. (2014) reported that decrease in fruit metabolic activities results in a decrease in fruit water loss and carbohydrates depletion rate and for this reason senescence process delayed in fruits in short-term storage at specific temperature.

CONCLUSION

It can be concluded that biochemical constituents of red radish plants recorded significantly increase with adding humic acid, seaweed extract and essential oils before storage compared with control plants. On the other hand, the same biomodulators in this study decrease the values of biochemical analysis after storage at 5°C for three months compared with untreated roots as a result of oxidative damage. All applied biomodulators can be decreased the weight loss and postharvest decay percentage compared with control plants. Dry matter percentage that reflected on the marketing rate recorded significantly increase compared with untreated plants.

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