Salinity stress has hindered the growth and yield of crops globally. The demands for inducing salt stress tolerance by natural and biological sources with potent antioxidants and growth-promoting metabolites have been the main focus of the recent era. Therefore, the current research was conducted to extract salt stress tolerance-ameliorating metabolites and growth-promoting hormones from the marine brown macroalgae Sargassum wightii Greville ex J. Agardh with maximum antioxidant potential used as a liquid fertilizer for okra (Abelmoschus esculentus L.). In the current study, the biochemical analysis showed that Sargassum aqueous extract (SAE) was rich in growth-promoting metabolites, antioxidants, and hormones. Meanwhile, overaccumulation of glycine betaine attracted the focus of the current research dealing with salt stress tolerance amelioration in A. esculentus. The plants supplemented with SAE (2% and 4%) and 0.04% ascorbic acid (AsA) alone and in combinations were subjected to sodium salt stress (NaCl; 75 mM). Results revealed that SAE efficiently promoted the vegetative and reproductive growth of plants by elevating the growth-promoting metabolites and hormones in comparison to control plants. Ionic contents (Na+, K+, Ca2+, and Mg2+) and ratios (K+/Na+, Mg2+/Na+, and Ca2+/Na+) were modulated in SAE-treated plants. SAE also increased the level of carbohydrates, proteins, lipids, carotenoids, and proline and decreased the level of hydrogen peroxide and abscisic acid in salt-treated plants compared with the control groups. Enzymatic activities of catalase, ascorbate peroxidase, and guaiacol peroxidase were also enhanced by SAE treatment upon salt stress. The SAE-mediated stress tolerance amelioration and the positive growth response of A. esculentus were further accelerated by AsA (0.04%) supplementation used in combination with SAE (2% and 4%).
The current study revealed a novel report of the antioxidant and metabolite-rich algal extract (S. wightii) formulation along with AsA that induced salt stress tolerance and promoted the overall growth performance of A. esculentus by rebalancing the ionic and metabolic status.

Keywords: marine macroalgae, seaweed, Abelmoschus esculentus, ion homeostasis, antioxidants, salt stress tolerance, bioactive metabolites, biostimulant

1 INTRODUCTION

Abiotic stresses deteriorate approximately 50% of crop yields. Unexpectedly, drastic fluctuations in the global climate impose numerous environmental stresses (Rane et al., 2021), and due to rapid urbanization, water stress is also associated with salt, drought, and temperature stress and water shortages that prime to various morphological and physiological deformities that cause a reduction in plant growth (Chele et al., 2021). The soil which has greater than 4 dS m⁻¹ electrical conductivity is considered saline soil (Alqahtani et al., 2018). Due to salinity, approximately 4 × 10^4 ha of land in over 100 countries is becoming unfit for agricultural purposes (FAO and ITPS, 2015). Salt stress adversely impacts plant physiology by affecting metabolic, osmotic, and ionic status, due to extreme uptake of Na⁺ and Cl⁻ leading toionic toxicity and deficiency of nutrients that boost macromolecules’ degradation by reactive oxygen species (ROS) production. Being sessile, it is inevitable for plants to avoid salinity stress imposed by saline soil. Salt-tolerant plants show increased activity of antioxidant enzymes such as catalase, superoxide dismutase, peroxidase, and ascorbate peroxidase (Umar and Siddiqui, 2018), which are called halophytes. Salt-tolerant plants have adapted several other mechanisms to tolerate salt stress such as i) the ability to biosynthesize and accumulate compatible osmolytes that control turgor to prevent ultrastructural damage, maintain ion homeostasis, and regulate water uptake to enhance water use efficiency; ii) adaptation for flexible growth plasticity (morphological and developmental pattern change); iii) enhancement in photosynthetic activity; iv) detoxification of ROS through antioxidant enzymes and non-enzyme molecules; and v) induction of phytohormones and growth-promoting metabolic products, including glycine betaine and proline. However, these adaptive strategies are not sufficient to induce tolerance to rapidly increasing salinity in the soil. Moreover, unfortunately, not all plant species are halophytes; therefore, salt stress has been found to negatively affect the biomass and yield of susceptible plants including Spartina alterniflora, Brassica oleracea, Jerusalem artichoke, Gossypium hirsutum, Helianthus tuberosus, Catapodium rigid, Pistacia vera, Orzya sativa, Lycopersicon esculentum, Fragaria ananassa, Triticum aestivum, and Abelmoschus esculentus (Roy and Chowdhury, 2020). Abelmoschus esculentus is the most popular mallow crop and a common food crop in Asia. Its finger-like fruits are a rich source of vitamins, minerals, and dietary fiber and are low in calories. The mucilage’s properties have enormous medicinal value. The rationale behind the current study was the fact that a 90% loss (6.5 dS m⁻¹) in A. esculentus yield has been reported under high salt levels (Mushtaq et al., 2020). Global food demand requires the production of salt-stress-tolerant, nutritionally rich crops including A. esculentus. Therefore, researchers have been exploiting different approaches to ameliorate salt stress tolerance in plants. Scientists have been searching for stress-resistant genes to reconstruct the stress-tolerant response in plants by transgenic approaches. Recently, transcriptome analysis, performed for exploring the molecular mechanism of melatonin-induced salt tolerance in A. esculentus, has revealed the induction of transcription factor genes including MYB, WRKY, and NAC. The genes controlling nitrogen, sulfur, alanine, aspartate, and glutamate metabolism were also significantly induced (Zhan et al., 2021). Although genetic engineering is a powerful strategy for generating ideal plants of desired traits for food production, it is not applicable in agriculture due to its controversial status concerning the risk to human health as well as the agricultural environment. Thus, rather than genetic alteration, understanding the molecular processes for induction of salt stress tolerance-associated pathways provides for enhanced stress resistance efficiency in plants. For example, exogenous application of various priming mediators (trehalose, sodium nitroprusside, manganese sulfate, sodium selenate, sodium silicate, calcium chloride, glycine betaine, proline, nano-silica, α-tocopherol, vitamins, and nanoparticles) has been used for salt stress tolerance amelioration in crops. Recently, glycine betaine has also been found to induce salinity tolerance in plants by modulating the physiological, antioxidant, and ionic status (Dustgeer et al., 2021). The foliar spray of α-tocopherol has been used to mitigate salt stress tolerance in A. esculentus by modulating the antioxidant potential (Naqve et al., 2021). The potential mitigation effect of ZnO nanoparticles has also been found to mitigate the effect of salt stress in A. esculentus (Alabdallah and Alzahrani, 2020). Menadione sodium bisulfite is reported by Akbar et al. (2021) to regulate physiological and biochemical responses for salinity stress alleviation on wheat (T. aestivum L.).

Growth regulators such as polyamine (diamine putrescine, tetramine spermine, triamine spermidine) and gibberellic acid have also received attention from seed priming chemicals due to their strong influence on germination and growth of the plant (Sofy et al., 2020). Growth promotion has also been noticed under salinity stress by exogenously applied gibberellic acid in maize through modulating the morphophysiological, biochemical, and molecular attributes (Shahzad et al., 2021).
Exogenous melatonin is reported to mitigate salinity-induced damage in olive seedlings by modulating ion homeostasis, antioxidant defense, and phytohormone balance (Zahedi et al., 2021). Exogenous application of melatonin also proved to be effective for the alleviation of severe salt stress (300 mM NaCl) of okra (Zhan et al., 2021).

However, the use and synthesis of these chemicals are time-consuming, expensive, and mostly not environment-friendly. Furthermore, the efficiency of these chemical fertilizers, priming mediators, and growth regulators differs with different environmental stresses and plant species. Scientists, therefore, are suggesting an alternative promising approach to use biological sources such as seaweeds for inducing salt stress tolerance in various plants. Seaweeds are a group of macroscopic marine algae that are saltwater tolerant and rich in growth regulators and micro- and macronutrients, essential for plant growth promotion (Bulgari et al., 2019).

The application of seaweed extracts in plants has been reported to increase chlorophyll content, photosynthetic activity, transpiration rate, stomatal conductance (Al-Ghamdi and Elansaey, 2018), the electron transfer rates of photosystems I and II (Digruber et al., 2018), plant height, leaf numbers, root width, and length with an overall increase in biomass (Ali et al., 2019).

Previously, foliar application of ascorbic acid significantly increased chlorophyll a, chlorophyll b, carotenoids, and the total photosynthetic pigments at a dose of 400 mg L⁻¹, in common beans under water stress conditions (Gaafar et al., 2020). Therefore, the current research was also planned to use an optimal concentration (0.04%) of ascorbic acid (AsA) solution for foliar spray on A. esculentus under salt stress of 75 mM. Recently, AsA as an antioxidant was also used as a priming agent and mediator for salt stress tolerance amelioration, and foliar application showed powerful potential to reduce salt toxicity in A. esculentus (Saheed, 2020). Thus, AsA (0.04%) was used as a positive control. Meanwhile, Sargassum aqueous extract (SAE, 4%) was used in combination with AsA (0.04%) to evaluate the enhancement in growth response and amelioration of salt stress tolerance response of A. esculentus. It is also well known that salt stress affects A. esculentus plant growth and development as well as its physiological attributes like photosynthesis rate and stomatal conductance and overall growth of plants in a dose-dependent manner, with 25 and 50 mM NaCl being less toxic and 75 mM NaCl being toxic (Shahid et al., 2011). Therefore, under the current study, 75 mM NaCl was supplemented to A. esculentus plants for salt stress induction.

The mechanisms behind positive influences driven by marine macroalgae (seaweed extracts) upon salt stress tolerance have not been studied for A. esculentus so far. Therefore, the current work was designed to investigate the potential of marine algal extract as a bioimulant for growth promotion as well as stress tolerance in A. esculentus. The present study was aimed to explore whether S Sargassum wightii aqueous extract may induce salt stress tolerance in A. esculentus and what could possibly be the essential metabolites, ions, and hormones in SAE to influence the, i) the essential metabolic reshuffling in A. esculentus plants itself upon salt stress, ii) the modulation of growth-controlling hormones, and iii) the antioxidant potential. The findings of the current research allowed us to explore the in-depth physiological, biochemical, metabolic, and ionic reshuffling for salt stress tolerance amelioration in A. esculentus.

2 METHODS

2.1 Biological Material and Chemicals

The use of biological material in the current research has fully complied with institutional, national, and international guidelines and legislation. Certified A. esculentus variety (Sabz Pari) was obtained from Early Wood Seed (Pvt) Limited, Multan, Pakistan. Seaweed (marine macroalgae) was collected from the Karachi coast (24°48′N latitude and 66°59′E longitude), Pakistan, in May 2019. The sample was washed first with seawater followed by fresh water thoroughly to remove the epiphytes and other contaminants in the laboratory, Department of Botany, University of Karachi, Pakistan. The sample was air-dried in shade (to avoid thermal degradation of the metabolites) for 7 days. Dried samples were carefully transported to the Plant Physiology and Molecular Biology Laboratory, Department of Botany, Abdul Wali Khan University, Mardan, Pakistan. Marine algal species (seaweed) were identified as S. wightii, according to Chapman and Gellenbeck (1989), and taxonomically classified according to Papenfuss (1955). The identification was based on morphological (external and internal characteristics) as well as ecological (distribution and habitat) features. The chemicals and reagents were bought from Sigma-Aldrich (Taufkirchen, Germany), Merck (Darmstadt, Germany), and Fluka (Buchs, Switzerland).

2.2 Preparation of Sargassum Aqueous Extract

SAE was prepared following the method of Erulan et al. (2009). Fresh sample of collected seaweed was dried and ground to form a fine powder. The fine powder and distilled water were mixed at a ratio of 1 kg:1 L, boiled for an hour, collected in conical flasks using Whatman No. 1 filter paper, and covered with aluminum foil to store at 29°C ± 1°C under dark conditions for 24 h. The filtrate was considered as 100% concentration.

2.3 Compositional Analysis of Sargassum Aqueous Extract

The filtrate (10 ml) was collected, oven-dried at 45°C, and recorded as the total yield. The reconstituted extract (2 mg/ml) was stored at −20°C for composition analysis. Total antioxidant activity was quantified according to Prieto et al. (1999). Total phenolic content was measured following the method of Singleton and Rossi (1965). The method for DPPH radical scavenging activity of SAE was adapted from Wen and Chen (1995). The ferric reducing antioxidant power (FRAP) assay was performed for measuring the reducing power of the SAE following the method of Benzie and Strain (1996). Hormonal quantification from SAE was done according to Benitez Garcia et al. (2020), and glycine betaine quantification was estimated in the Sargassum
aqueous extract as mentioned by Grieve and Grattan (1983). Elemental analysis of the SAE was performed by a procedure adapted from Nazarudin et al. (2021). A 10% seaweed extract was used for quantifying the concentrations of macro- (K\(^{+}\), Na\(^{+}\), Mg\(^{2+}\), and Ca\(^{2+}\)) and micronutrients (Fe\(^{2+}\), Cu\(^{2+}\), Mn\(^{2+}\), and Zn\(^{2+}\)) using the ICE 3000 atomic absorption spectrophotometer (Thermo Scientific, USA).

### 2.4 Abelmoschus esculentus Growth Conditions and Experimental Setup

The experiments were carried out in the Botanical Garden, Plant Physiology and Molecular Biology Laboratory, Department of Botany, Abdul Wali Khan University, Mardan, Pakistan. Seeds were sterilized for 1 min with 0.1% HgCl\(_2\) and then washed with distilled water. Ten seeds per pot were sown; however, the uniformly growing plants were selected and thinned out to five, after 15 days of germination. Plants were irrigated with an equal amount of tap water every day. Each earthen pot with 12 × 12 × 30 cm (length × width × height) size was filled with sandy loam soil. A completely randomized design (CRD) was assigned for all 84 pots which were divided into 12 various groups representing the various treatments such as supplementation of SAE, NaCl salt, and AsA.

The 3-week-old plants were subjected to salt (NaCl) treatments by irrigation with 75 mM salt water with 10.5 dS m\(^{-1}\) electrical conductivity, twice a week regularly. SAE (2% and 4%) + AsA (0.04%) were supplied by foliar spray (4 ml/treatment) on each plant at the 3rd, 6th, and 10th week after germination. The foliar application was applied on the abaxial and adaxial surfaces at sunset to ensure SAE and AsA uptake by the leaves. At the end of the grand period of growth (plants), samples were taken for biochemical analysis (Figure 2).

The experimental setup with treatments is mentioned below.

1. Control (water)
2. SAE (2%)
3. SAE (4%)
4. Control (AsA 0.04%)
5. SAE (2%) + AsA (0.04%)
6. SAE (4%) + AsA (0.04%)
7. NaCl (75 mM)
8. NaCl (75 mM) + SAE (2%)
9. NaCl (75 mM) + SAE (4%)
10. NaCl (75 mM) + AsA (0.04%)
11. NaCl (75 mM) + SAE (2%) + AsA (0.04%)
12. NaCl (75 mM) + SAE (4%) + AsA (0.04%)

### 2.5 Phenotypic and Physiological Analysis

#### 2.5.1 Measurement of Growth Parameters
At the termination of the experiment, the yield was recorded by measuring vegetative (shoot length, root length, no. of intact leaves, fresh and dry weight of intact leaves) and reproductive attributes (number of pods per plant, pod fresh weight, dry weight of fruit, pod length, number of seeds per pod and weight of seeds per pod, number of seeds per plant, weight of seeds per plant, and weight of 100 seeds).

#### 2.5.2 Relative Water Content and Leaf Water Loss
The relative water content (RWC) of the fresh leaf (leaf number 12 counted from the bottom of each plant) was calculated by using freshly weighed leaves subjected to a rehydration process in deionized water for 2 h and turgor weight was measured. The dry weight of the leaf samples was measured after incubating the samples in a preheated oven. The formula given below was used to calculate RWC:

Relative Water Content; RWC(%) = (Fresh weight − Dry weight) / (Total weight − Dry weight) × 100

Leaf water loss was determined by measuring the initial weight (W\(_1\)) of the fresh-leaf samples (3.0 g) from leaf number 12 of each plant (counted from the base). The sample was kept for 2 h at 30°C and then weighed again (W\(_2\)). Leaf water loss (LWL) was calculated using the following formula:

Leaf Water Loss; LWL(%) = W\(_1\) − W\(_2\)/W\(_1\) × 100

#### 2.5.3 Extraction and Estimation of Chlorophyll and Carotenoids
Total chlorophyll and carotenoids in fresh-leaf samples (3.0 g) from leaf number 12 of each plant (counted from the base) were estimated and calculated through a process described by Maclachlan and Zalik (1963).

#### 2.5.4 Analyses of Metabolites
Metabolic quantification was performed by using fresh-leaf samples (3.0 g) from leaf number 12 of each plant (counted from the base). For total sugars, quantification was done by adding phenol and H\(_2\)SO\(_4\) to the sample. The optical density was recorded at 485 nm. Total protein in the leaves was determined according to Lowry et al. (1951). Lipid content was determined as measured by Van Handel (1985). Proline quantification was done according to Bates et al. (1973). Total-phenolics were estimated as mentioned by Aziz et al. (2021b). Tannin was estimated and calculated through a process described by Makkar et al. (1993). Lycopene and β-carotene were determined following the method of Nagata and Yamashita (1992). Total terpenoid was measured following the method of Fan and He (2006). Indoleacetic acid (IAA) content was measured following the method mentioned by Rauf et al. (2021). SA content was analyzed following the method of Warrier et al. (2013). Endogenous AsA was determined by Jagota and Dani (1982). GA\(_3\) and abscisic acid (ABA) content was analyzed by the procedure of Ergun et al. (2002).

#### 2.5.5 Antioxidant Enzyme Activities
Total antioxidant capacity was determined by using fresh samples (3.0 g) from leaf number 12 of each plant (counted from the base), crushed in methanol. Three milliliters of solution with H\(_2\)SO\(_4\) (0.6 mM), sodium phosphate buffer (28 mM), and ammonium molybdate (4 mM) was added to the sample extract. After incubation, absorbance was measured at 695 nm against the blank. Catalase was determined following the method of Luck.
Guaiaicol peroxidase was determined by the procedure of Haida and Hakiman (2019). Ascorbate peroxidase was determined following the method of Nakano and Asada (1981).

2.5.6 Determination of H$_2$O$_2$
H$_2$O$_2$ determination was found following the method of Mishra and Agrawal (2014). Fresh-leaf samples (3.0 g) from leaf number 12 of each plant (counted from the base) were crushed in trichloroacetic acid (0.1%). After centrifugation, potassium phosphate buffer (10 mM, pH = 7) and potassium iodide (1 M) were added to the samples and taken at an absorbance of 390 nm. ROS detection was performed through the method described by Yokawa et al. (2016).

2.5.7 Ionic Analysis
The concentration of various ions (Ca, Mg, K, Na) were quantified in dried ground plant samples. Approximately, 0.5 g of ground dried plant materials were finely powdered and transferred to digesting vials. For the digestion process, 6.5 ml of acid solution (HNO$_3$, H$_2$SO$_4$, HClO$_4$) with a ratio of 5:1:0.5, was added to each sample. After soaking for 12 h the samples were burned at 300°C for 3 h till the appearance of white fumes. Distilled water was added to the sample for dilution and then filter with Whatman filter paper No. 1. The ionic contents were then determined using an atomic absorption spectrophotometer (ICE 3000; Thermo Scientific, USA) (Aziz et al., 2021a).

2.6 Statistical Analysis
The experiments were carried out in three biological replicates. Two-way ANOVA was used to analyze the data with a significance level of $p < 0.005$. To compare the means of all values, SPSS 20 (SPSS Inc., Chicago, IL, USA) was used for performing DMRT (Duncan multiple range test) at $p < 0.005$.

3 RESULTS

3.1 Characterization of Sargassum wightii
Sargassum wightii composition analysis showed a high level of total sugars (4.3 mg/g FW), proteins (0.145 mg/g FW), lipids (0.15 mg/g FW), lycopene (0.16 mg/g FW), and β-carotene (0.5 mg/g FW). High total phenolics (7 μg/g FW) and flavonoids (8 μg/g FW), proline (0.003 μg/g FW), and tannins (144 μg/g FW) were also produced. A higher content of cytokinin (0.191 mg/100 ml), GA$_3$ (12.2 mg/100 ml), auxin (0.11 mg/100 ml), and glycine betaine (0.17 μg/g DW) was noticed. Glycine betaine is an important osmolyte in maintaining osmotic balance and is reported to counter salinity stress by maintaining a high K$^+$/Na$^+$ ratio and antioxidant defense via limiting Na$^+$ uptake. Sargassum wightii also showed high total antioxidant capacity (57 mg AAE/g), DPPH (76%), and FRAP (54 mg AAE/g). Results for the measurement of the mineral composition of S. wightii revealed the following: sodium (Na$_7$) (875 mg/g DW), potassium (K$^+$) (101.05 mg/g DW), calcium (Ca$_{10}$) (100 mg/g DW), magnesium (Mg$^{2+}$) (22 mg/g DW), Fe$^{3+}$ (16 mg/g DW), Cu$^{2+}$ (0.5 mg/g DW), Mn$^{2+}$ (0.7 mg/g DW), and Zn$^{2+}$ (0.3 mg/g DW) (Figure 1).

3.2 Effect of Sargassum wightii Aqueous Extract on the Vegetative Growth of Abelmoschus esculentus Under Salt Stress
Plant height, root length, and leaf number are the main components in plant architecture and play a main role associated with biomass production. The results of the present study displayed that vegetative growth parameters (shoot length, root length, number of intact leaves, fresh and dry weight of intact leaves) were significantly ($p < 0.005$) reduced in A. esculentus plants after the application of salt stress compared with control plants. Simultaneously, SAE (2% and 4%) supplementation significantly ($p < 0.005$) increased the growth parameters of A. esculentus plants, compared with control plants as well as those grown under salt stress (Figure 2).

3.3 Effect of Sargassum wightii Aqueous Extract on the Reproductive Attributes of Abelmoschus esculentus Under Salt Stress
Current data revealed that salt supplementation exhibited a significant ($p < 0.005$) decline in different reproductive parameters such as total intact pods/plant, seed number/pod, seed weight/pod, the weight of 100 seeds, total seeds/plant, seed weight/plant, and pod length of A. esculentus plants, compared with control plants. On the contrary, results revealed that SAE (2% and 4%) application improved the reproductive attributes in both control and saline stress conditions. Moreover, it was found that exogenous application of SAE (2% and 4%) proved significantly ($p < 0.005$) effective in improving different reproductive attributes of A. esculentus plants, compared with control plants as well as those grown under salt stress (Figure 3).

3.4 Effect of Sargassum wightii Aqueous Extract on the Physiological Attributes of Abelmoschus esculentus Under Salt Stress
Current results revealed that salt-stressed A. esculentus plants showed a significant ($p < 0.005$) reduction in the level of chlorophyll a, chlorophyll b, and total chlorophyll compared with control plants. On the other hand, foliar application of SAE (2% and 4%) exhibited improvement in chlorophyll a, chlorophyll b, and total chlorophyll contents of A. esculentus plants, compared with control plants as well as those grown under salt stress. Current data showed that salt stress caused a significant ($p < 0.005$) increase in total carotenoid levels in A. esculentus leaves compared with control plants. Treated with SAE (2% and 4%) and AsA further increased total carotenoid levels and were noticed in both non-saline and saline conditions. Salt stress also caused a significant ($p < 0.005$) reduction in the relative water content and acceleration in leaf water loss of A. esculentus plants compared with control plants. On the other hand, SAE (2% and 4%) promoted a significant ($p < 0.005$) accumulation in the relative water content and reduction in leaf water loss of A. esculentus plants, compared with control plants as well as those grown under salt stress (Figure 4).
3.5 Effect of *Sargassum wightii* Aqueous Extract on the Metabolic Attributes of *Abelmoschus esculentus* Under Salt Stress

The obtained results exhibited that salinity stress significantly \( p < 0.005 \) decreased the total carbohydrates, proteins, lipids, and proline content compared with control plants, while SAE (2% and 4%) significantly \( p < 0.005 \) induced the total carbohydrates, proteins, lipids, and proline content of *A. esculentus* both under normal and stressed conditions (Figure 5). The obtained data from the current study revealed that 75 mM salt stress caused a significant \( p < 0.005 \) increase in total phenols, flavonoids, tannin, lycopene, β-carotene, and terpenoid contents of *A. esculentus* plants under both normal and stressed conditions significantly \( p < 0.005 \) (Figure 6).

3.6 Effect of *Sargassum wightii* Aqueous Extract on the Hormonal Attributes of *Abelmoschus esculentus* Under Salt Stress

The results of the current study indicated that 75 mM salt stress caused a significant \( p < 0.005 \) decrease in IAA, GA, AsA, and salicylic acid levels, while a significant increase was noticed in the ABA level of *A. esculentus*, compared with the control plants. A reverse pattern of hormonal accumulation was noticed upon further increased total phenols, flavonoids, tannin, lycopene, β-carotene, and terpenoid contents of *A. esculentus* plants under both normal and stressed conditions. 

**FIGURE 1** | Chemical composition of *Sargassum wightii*. (A) Primary metabolites; (B) secondary metabolites; (C) hormonal content; (D) antioxidant capacity; (E) ionic analysis. Quantitative data represent means ± SD (n = 7). FW, fresh weight; DW, dry weight.
SAE (2% and 4%) application compared with control plants (Figure 7).

### 3.7 Effect of *Sargassum wightii* Aqueous Extract on the Antioxidant Capacity and ROS Status of *Abelmoschus esculentus* Under Salt Stress

Soil salinity causes the generation of oxidative stress through an increase in ROS. Different antioxidants especially enzymes play a vital role in protecting the cell by scavenging ROS. In the present study, data indicated that salt stress promoted a significant ($p < 0.005$) increase in total antioxidants such as catalase, guaiacol peroxidase, and ascorbate peroxidase compared with the control. On the other hand, in SAE (2% and 4%) supplementation, a further significant increase in total antioxidants was noticed in *A. esculentus* plants, compared with controls as well as those grown under salt stress (Figure 8). The current investigation highlighted that salt stress caused a significant ($p < 0.005$) increase in hydrogen peroxide levels in *A. esculentus* plants compared with control. Elevated DAB level further confirmed an overaccumulation of H$_2$O$_2$ in *A. esculentus* leaf tissue compared with control upon salt supplementation. The current results indicated the positive role of SAE (2% and 4%) in the reduction of hydrogen peroxide level and DAB intensity in *A. esculentus* under both normal and stressed conditions (Figure 9).

### 3.8 Effect of *Sargassum wightii* Aqueous Extract on the Mineral Status of *Abelmoschus esculentus* Under Salt Stress

To cope with salinity-induced water stress, the accumulation of inorganic solutes is the most suitable strategy to maintain its turgor which ultimately balances the cell water potential by enhancing water uptake. During the present investigation, the result showed that salt stress causes a significant ($p < 0.005$) overaccumulation in Na$^+$ concentration and reduction of K$^+$, Ca$^{2+}$, and Mg$^{2+}$ in *A. esculentus* plants compared with control. The foliar application of SAE (2% and 4%) led to a significant ($p < 0.005$) increase in K$^+$, Ca$^{2+}$, and Mg$^{2+}$ ion concentrations with a reduction in Na$^+$ ions in *A. esculentus* plants, compared with control plants as well as those grown under salt stress. Thus, increased K$^+$/Na$^+$, Ca$^{2+}$/Na$^+$, and Mg$^{2+}$/Na$^+$ ratios were noticed upon foliar application of SAE (2% and 4%) compared with control plants and those treated with salt stress (Figure 10).

### 4 DISCUSSION

Salt stress, among the major forms of abiotic stresses in plants, inhibits plant growth due to excessive absorption of Na$^+$ and Cl$^-$,
causing ion imbalances and physiological disorders. It causes osmotic stress, which lowers plant development by reducing water uptake, lowering turgor pressure, increasing stomatal closure, and reducing photosynthetic activity. Excessive Na\(^+\) generates certain ionic toxicities; therefore, controlling Na\(^+\) exclusion as well as conveyance is critical. The lesser the absorption of other nutrients like K\(^+\) and Ca\(^{2+}\), the higher the levels of Na\(^+\) and Cl, resulting in nutritional imbalance. Plants may produce ROS as a result of salt stress. Increased ROS causes more lipid peroxidation, protein breakdown, and DNA mutations (Mittal et al., 2012; Riaz et al., 2020).

Plants cannot avoid salt stress since they are sessile; thus, they must constantly adapt and evolve a range of ways to mitigate the negative consequences of salt stress. The production of non-enzymatic antioxidants (ascorbic acid) to detoxify ROS, the formation of suitable osmolytes such as proline or glycine betaine, and the accumulation of ABA are only a few of these strategies. The activity of various antioxidant enzymes, including catalase, guaiacol peroxidase, glutathione reductase, and ascorbate peroxidase, is also increased in salt-tolerant cultivars exposed to salt stress. Similar to several salt-sensitive crops, A. esculentus is also salt sensitive as salt supplementation of NaCl 75 mM has been known to negatively affect the growth and development of A. esculentus as well as its physiological attributes like photosynthesis rate and stomatal conductance and its overall growth (Shahid et al., 2011).

Seaweed extracts are used by researchers as low-cost alternatives to synthetic antioxidants with greater efficacy, eco-friendliness, and non-toxic nature that enhance phytohormone production and the moisture-holding capacity of plants as well as the overall growth of cereals, grasses, vegetables, fruits, and spices (Van Oosten et al., 2017; Sohn et al., 2021), and they can also be exploited as soil fertilization as well as plant stress tolerance amelioration. The application of a betaine-rich extract of Ascophyllum nodosum in sweet pepper and tomato showed an increased chlorophyll content.

**FIGURE 3** | Effect of SAE on different reproductive growth parameters. (A) Total pods/plant; (B) total seed number/pod; (C) total seed number/plant; (D) total seed weight/pod; (E) weight of 100 seeds; (F) total seed weight/plant; (G) average pod length; (H) phenotypes of pods. Error bars of quantitative data represent ± SEM (**\(p < 0.005\)).
extracts with higher betaine compounds showed a reduction in photosynthetic activity and promoted chlorophyll degradation. Similarly, an accelerated photosynthetic and transpiration rate and stomatal conductance were observed in *Asparagus* plants treated with *A. nodosum* (Al-Ghamdi and Elansaey, 2018). Treatment of willow plants with an extract of *Ecklonia maxima* accelerated the electron transfer rates of photosystems I and II (Digruber et al., 2018). Tomato plants supplemented with seaweed extracts showed better plant

**FIGURE 4** | Effect of SAE on pigment composition and water content of *Abelmoschus esculentus* grown under salt stress. (A) Chlorophyll a; (B) chlorophyll b; (C) total chlorophyll; (D) carotenoids; (E) relative water content; (F) leaf water loss. Error bars of quantitative data represent ± SEM (**p < 0.005). LWL, leaf water loss; FW, fresh weight.
height and increased leaf numbers, root width, and length with an overall increase in biomass (Ali et al., 2019).

Seaweed extracts have also been proven to promote the performance and quality of horticultural crops by increasing antioxidant capacity such as S. wightii, Kappaphycus alvarezii, A. nodosum, Sargassum johnstonii, and Sargassum latifolium along with Ulva lactuca resulting in a better yield of Vigna sinensis and Phaseolus radiata, L. esculentum, Paspalum vaginatum, and T. aestivum, respectively (Elansaey et al., 2017).

Sargassum spp. are the most common types of brown marine algae naturally growing in the saline environment and are halotolerant. The chemical composition of seaweed extracts relies on the algal species, growth environment, method of preparation, and environmental conditions, as well as its general composition including polysaccharides, minerals, amino acids, and growth hormones affecting the performance and development of plants. Considering their natural adaptation to survive and grow in a saline environment, the marine macroalgae (S. wightii) evaluated in the present study were found to have a strong antioxidant capacity along with a high potential for the production of growth-promoting metabolites and hormones. A pilot-scale pot experiment was conducted under the current research to evaluate and compare the potential of this antioxidant-rich SAE on A. esculentus growth under salt stress. The present observations revealed that foliar spray of SAE significantly improved both vegetative and reproductive growth attributes of A. esculentus, while a clear decline in optimum yield was noticed due to the drastic effects of salt stress. The positive impact of SAE resulting in optimum yield is consistent with those reported for the application of seaweed on other crops (Hernández-Herrera et al., 2018). Previously, foliar application of ascorbic acid was significantly also known to increase the photosynthetic activity and growth in common beans under water stress conditions (Gaafar et al., 2020). AsA as an antioxidant was also used as a priming agent and mediator for salt stress tolerance amelioration, and its foliar

![FIGURE 5](image-url)
application showed a powerful potential to reduce salt toxicity in *A. esculentus* (Saheed, 2020). In the current research, AsA (a strong antioxidant) was also used alone (as positive control) and in combination with SAE for foliar spray on *A. esculentus* under salt stress. The evaluation revealed that AsA as an antioxidant further supplemented and amplified the antioxidant potential of SAE on *A. esculentus* under salt stress, as the foliar application of AsA and SAE showed a powerful potential to reduce salt toxicity in *A. esculentus*. This complementary enhancement effect was also prominently visible in the growth response as well as stress tolerance response of *A. esculentus*.

Previously, it was also reported that seaweed extract supplementation by foliar application to the leaves increased the nutrient content of plants, along with the accumulation of metabolites and growth-promoting hormones (Uthirapandi et al., 2018). In a study by Kalaivany et al. (2019) on *Vigna..."
they observed a high pod number with *Sargassum* treatment and considered it as inorganic fertilizer which provides sufficient nutrients to the plant for its proper growth.

Moreover, chemical composition analysis of seaweeds also revealed that seaweed extracts are a rich source of antioxidants that counteract environmental stresses and play an important role in the prevention of toxic free radicals for improved growth and productivity of crops by increasing nutrient availability and uptake (Roy and Chowdhury, 2020). *Sargassum* extracts have been used as a biostimulant to induce abiotic stress tolerance in *Solanum lycopersicum*, *Capsicum annuum*, *F. × ananassa*, *T. aestivum*, and *O. sativa* (reviewed by Ali et al., 2021). The positive effect of algal extracts on plant growth and stress tolerance is supported by their high content of growth regulators and antioxidants.

**FIGURE 7** | Endogenous hormone levels. (A) IAA; (B) GA₃; (C) ABA; (D) AsA; (E) salicylic acid. Error bars of quantitative data represent ± SEM (**p < 0.005). FW, fresh weight.
extracts on photosynthetic pigments has been reported by other authors as well (Uthirapandi et al., 2018). The current research also revealed a high level of total sugars, total proteins, total lipids, lycopene, and β-carotene contents in SAE composition analysis, which is attributed to the positive morphological and physiological effects on A. esculentus plant to such growth regulators. The positive effect of SAE on photosynthetic pigments and promoting photosynthesis machinery may also justify enhanced vegetative growth attributes such as shoot length, root length, fresh biomass, and dry biomass as shown in the results. The sugar accumulation in the SAE could be the reason for the induction of signaling components for biosynthesis pathways of phytohormones for A. esculentus growth promotion under salt stress.

SAE supplementation is known to induce photosynthetic potential by improving the chlorophyll composition of leaves as Ramya et al. (2010) also found that the application of a liquid extract of S. wightii on Cyamopsis tetragonoloba under stress conditions increased the level of photosynthetic pigments due to the high absorption of magnesium ion that is a major constituent for chlorophyll biosynthesis. Generally, seaweeds contain high levels of iron, magnesium, and nitrogen which help increase chlorophyll synthesis. On the other hand, these ions are important in reducing lipase and lipoxygenase enzyme activities, thus protecting membrane breakdown.

The present study revealed that SAE application amplified the carotenoid production of A. esculentus plants under salt stress, consistent with the previous reports where Raphanus sativus treated with salt showed an increase in total carotenoid levels (Kasim et al., 2016). Increased carotenoids are known to protect the chloroplast from photo-oxidative damage induced by ROS production under salinity stress (Latef et al., 2017).

Water status in plants can be best determined through the relative water content of the leaves. In the present research, SAE-treated plants caused an improvement in plant water status and increased relative water content. Previously, a reduction in relative water content was also observed in plants such as Thymus vulgaris and Thymus daenensis Celak. (Bistgani et al., 2019) under salinity stress. In plants under salt stress, negative

**FIGURE 8** | Different antioxidant enzyme activities. (A) Total antioxidants; (B) catalase; (C) ascorbate peroxidase; (D) guaiacol peroxidase. Error bars of quantitative data represent ± SEM (**p < 0.005). FW, fresh weight.
effects were created on osmotic potential, water potential, and relative water content in the leaves. Elansaey et al. (2017) observed the same results in experimental plants under salt stress due to a reduction in osmotic potential. The favorable impact of AsA on cucumber plants under normal and stressful circumstances was described by Naz et al. (2016), as it promotes water absorption and turgor capacity by maintaining moisture in plant tissues.

High salt concentrations change the water status of plants with a reduction of cell division/elongation with retarded growth due to limited carbon metabolism, leading to cell death by ROS production and deterioration of different biochemical processes. Bioactive compounds of seaweed extract impart a positive impact on plant overall growth by optimizing metabolite production (stress alleviating amino acids), membrane permeability, and transport of osmolytes/ions for enhancing abiotic stress tolerance through modifying the water capacity and turgor pressure in plants. Glycine betaine and proline (osmolytes) perform a vital role in osmoregulation (Annunziata et al., 2019), and they can interchange with molecules to modify the structures, maintain the veracity of membranes upon stresses, conserve the activity of macromolecules, scavenge the ROS, stabilize the photosynthetic activity, and initiate the gene expression responsible for controlling oxidative stress responses in G. hirsutum L. (Hamani et al., 2020), Zea mays L. (Chen et al., 2020), and rice (Hafez et al., 2019).

Consistent with previous reports, the present study also showed that SAE exhibited a high level of glycine betaine and proline which might be one of the reasons for salt stress tolerance amelioration in A. esculentus. Glycine betaine is also reported to counter salinity stress by maintaining a high K⁺/Na⁺ ratio and antioxidant defense via limiting Na⁺ uptake in the common bean (Phaseolus vulgaris L.) (Sofy et al., 2017). Our data also revealed that an increase in K⁺/Na⁺ ratio, antioxidant potential, and photosynthetic pigments might be attributed to the effect of glycine betaine supplied to A. esculentus plants via SAE application.

Proline is another important amino acid that is produced in plants under different stresses and helps in plant development. Proline accumulation under salinity stress regulates C and N
source, osmosis, plasma membrane regulation, production of proteins, and destruction of free OH radicals. Current data revealed that salt stress drives an increase in proline level in *A. esculentus* as compared with control, which upon SAE supplementation in salt stress is further increased. According to Sofy et al. (2017), proline level enhancement in barley plants under salinity stress is a stress indicator signal but protects plants as it is involved in the alleviation of lipid peroxidation. The current study also supported the fact as SAE chemical composition has shown higher endogenous proline content too, which justifies the salt stress tolerance amelioration in *A. esculentus*.

It was previously known that stress-alleviating secondary metabolites and antioxidants act as the first line of defense for plants under different abiotic stresses. Cherry tomatoes with the application of AsA have also shown an increase in lycopene acting as a stress indicator as well as an antioxidant (Abdelgawad et al., 2019). According to different studies, it is evident that AsA is involved in the production of xanthophylls, antioxidants, carotenoids, lycopene, and chlorophyll in plants (Noctor et al., 2012). SAE application with antioxidant potential also showed pronounced influence in improving the lycopene content of *A. esculentus* plants. Akladious and Mohamed (2018) have also found an increase in lycopene content in pepper plants under salt stress. Researchers reported a rise in phenols, protein, flavonoids, tannins, terpene (particularly camphor and cineole), and antioxidant capacity after applying seaweed and AsA to plants, like chalcone isomerase (flavonoid biosynthesis enzyme) activity. Terpene is a class of secondary metabolites having antioxidant properties proven in different *in-vitro* assays (Aziz et al., 2018). Tannins have been proven to have a significant impact on plants in terms of inducing ROS-scavenging qualities under stress, and high tannin levels in plants induced by seaweed application improve the uptake of potassium ions, which has a positive influence on plant development and salt stress reduction (Baatour et al., 2018). SAE treatment had a significant impact on the terpene and tannin levels of *A. esculentus* plants, according to current findings. SAE treatment had a favorable effect on increasing phenolic and flavonoids in *A. esculentus* in this research, which is consistent with earlier results. For example, an increase in phenols under abiotic stress protected plants and played an important part in the regulation of stress by scavenging the ROS (Naikoo et al., 2019).

Hormones act as signaling elements as well. The hormonal contents present in seaweed extract positively influence the phytohormonal activity of plants that positively drives plant growth (Vinoth et al., 2019; Polat et al., 2021). Hormonal analysis of SAE under the current research showed a high level of auxin, cytokinin, and GA content, while *A. esculentus* plants treated with SAE also have revealed an increase in endogenous growth-promoting hormones with better overall growth response under salt stress in comparison to control plants. Thus, the growth-promoting impact of seaweed on growth parameters might be attributed to the presence of growth-promoting hormones present in SAE. The cytokinin-rich seaweed extracts are reported to induce photochemical efficiency in *Agrostis*...
paulstris (Zhang and Ervin, 2004) and heat stress tolerance in
Agrostis stolonifera (Zhang et al., 2010). Consistent with current
observations, previous reports have indicated that seaweed
(Sargassum) extract helped increased leaf number (Kaladharan
et al., 2019) and biomass in plants, after the application of seaweed
extracts with a higher content of auxin, phosphorus, and other
growth (Hasanuzzaman et al., 2020), minimized the
conditions. Exogenous AsA application has promoted plant
translocator to regulate oxidation
enzyme cofactor, antioxidant restorer, ROS scavenger, and signal
hormone biosynthesis. It also plays a vital role as a redox buffer,
senescence, cell division, defense, and the yield of plant and
hormone biosynthesis. It also plays a vital role as a redox buffer,
AsA is an important growth regulator which acts positively
under abiotic stress and modulates photosynthesis, growth, senescence, cell division, defense, and the yield of plant and
hormone biosynthesis. It also plays a vital role as a redox buffer,
enzyme cofactor, antioxidant restorer, ROS scavenger, and signal
translocator to regulate oxidation–reduction, especially in stress
conditions. Exogenous AsA application has promoted plant
growth (Hasanuzzaman et al., 2020), minimized the
production of H2O2 upon stresses (El Sebai et al., 2016), and
increased chlorophyll contents and photosynthetic activity in
tomatoes under salt stress (Chem et al., 2021).

An increase in plant height, leaf number, and fresh and dry
biomass has been recorded after seaweed application due to the
presence of a high amount of nutrients such as phosphorus, in
radish (Mahmoud et al., 2019) and V. unguiculata (Kalaivany et
al., 2019). Chemical analysis of SAE under the current research also
showed a high ionic level of macro- and micronutrients (sodium,
potassium, calcium, magnesium, iron, copper, zinc). The
growth enhancement effects of SAE could also be attributed to
these macro- and microelements that made up the chemical
composition of SAE as found in this study and in agreement with
previous investigations as well. It is also known that seaweed
extracts contain minerals that plants can readily assimilate to
stimulate growth and productivity, such as zinc, copper, boron,
and IAA, improving cell division, cell elongation, and root growth
in plants. Moreover, consistent with previous reports, SAE
supplementation also modulated the ionic status (K+/Na+, Mg2+/
Na+, and Ca2+/Na+ ratios) in A. esculentus grown under salt stress
that led to a significant reduction of salt-induced toxicity.

5 CONCLUSION

K+/Na+, Mg2+/Na+, and Ca2+/Na+ ratios in A. esculentus grown
under salt stress were increased by foliar application of antioxidant-rich algal extract (S. wightii), noticeably reducing the
salt toxicity and providing growth promoting effects. Such
growth-promoting and salt stress-alleviating effects induced by
SAE are considered a potentially effective strategy for its use as an
agricultural biocatalyst and biostimulant. Thus, exploitation of
the antioxidant and metabolite-rich algal extract of S. wightii
induced salt stress tolerance in A. esculentus by rebalancing the
ionic and metabolic status of plants under stress. SAE is found
in this research as an eco-friendly, sustainable, feasible, and
growth-promoting biological tool for ameliorating salt
stress tolerance in A. esculentus. However, future studies are
needed to identify the signaling elements driving the salt stress
responses in A. esculentus to fully understand the molecular
mechanisms activated under salt stress. It is recommended that
complete genome, transcriptome, and proteome analyses and
characterization of salt stress response genes in A. esculentus be
performed to elucidate the molecular basis controlling salt responses.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be
made available by the authors, without undue reservation.

AUTHOR CONTRIBUTIONS

HG, MR, MH, and MA designed the experiments. ZK, MR, and
HG performed the main experiments. AR, AT collected and
provided the seaweed sample. ZK, HG, MR, MA, participated in
the writing the main manuscript. HG, MR, and MA analyzed the
data. ZP, AU-D, SK and ZS assisted in the critical intellectual
review of the manuscript. MH and I-JL provided financial and
scientific support for the research project. All authors
contributed to the article and approved the submitted version.

FUNDING

The authors declare that current research was supported by the
National Research Foundation of Korea (NRF) grant funded by
the Korean government (MSIT) (No. 2022R1A2C1008993).
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