

1 Effect of Different Seaweed Extracts and Compost on 2 Vegetative Growth, Yield and Nutraceutical Quality 3 of Cucumber fruit (*Cucumis Sativus*)

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22 **Abstract:** The objective of this research was to evaluate the production and phytochemical quality of
23 cucumber fruits (*Cucumis sativus*), in response to the foliar application of different seaweed extracts. This
24 study was carried out under shadow mesh conditions in the autumn - winter agricultural cycle at Instituto
25 Tecnológico de Torreón, Torreón, Coahuila. The experimental design was completely random, using six
26 treatments with six repetitions each. The treatments evaluated were: *Macrocystis pyrifera*, *Bryothamnion*
27 *triquetrum*, *Ascophyllum nodosum*, *Grammatophora* sp., *Macrocystis intergrifolia*, and a control treatment
28 with inorganic fertilization. The substrate used was a mixture of sand and vermicompost. The yield,
29 commercial quality and phytochemical compounds of the fruit were evaluated. Results showed that yield
30 using Steiner solution (6.75 kg m⁻²) was higher than that obtained with *Bryothamnion triquetrum* algae (6.07
31 kg m⁻²). Regarding the phenolic content, the extracts surpassed the control treatment, with *Macrocystis*
32 *pyrifera* and *Macrocystis integrifolia* being statistically equal, with values of 47.37 and 43.73 mg equiv. of Ac.
33 Gallic 100 g fresh weight, respectively. The antioxidant capacity by ABTS+ and DPPH+ methods was higher
34 using the treatment with algae *Macrocystis pyrifera* with 149.4 and 454.1 μM equiv Trolox / 100 g fresh base,
35 respectively. This treatment also presented the highest value of vitamin C with 5.07 mg / 100 g fresh base,
36 being 27% greater than the control treatment. Algae extracts increased the quality of the fruits by obtaining
37 the highest antioxidant capacity, making their use a viable option to minimize the application of conventional
38 fertilizers, thereby attenuating the effects on the environment and improving the health of the population.

39 Keywords: *Cucumis sativus*; nutraceutical; antioxidant; ABTS+, DPPH+

40

41 1. Introduction

42 The production of protected vegetables is one of the main export activities in Mexico because it
43 occupies fourth place worldwide and first place in the American continent [1]. The main crops
44 produced are tomatoes (70%), pepper (16%) and cucumber (10%); the production of these vegetables
45 is mainly located in the northwest (72%) and center of the country (19%) [2]. Cucumber is in great
46 demand in the United States of America (USA) and has had an unprecedented growth in recent years.
47 Its importation grew from 394 107 tons in 2002 to 459 242 in 2012 [3]. Of this import volume, 80%
48 comes from Mexico, which is the main exporting country to the USA [4]. Under protected agriculture
49 conditions, the production of cucumber is 2 to 9 times greater than in the open field, which depends
50 on technological levels, nutrition management and the control of climatological conditions which are
51 key factors for achieving high yields in this type of systems [5].

52 The nutrition of this crop is an important factor in maximizing its production. In several regions,
53 there is an indiscriminate use of agrochemicals and the excessive cultural practices have brought
54 serious consequences to the environment. In addition, the high costs of fertilizers in Mexico have put
55 them out of the reach of producers, resulting in low yields, low incomes and low quality produce [6].
56 Along with this, in the large market of imported vegetables, there is a growing tendency among the
57 consumer towards the acceptance of "innovative, different", suitable products that can contribute to
58 an ever-healthier diet with less environmental impact on natural resources [7]. The implementation
59 of biological control and organic nutrition with biofertilizers could reduce the dependence on
60 agrochemicals and thus make the production less hazardous to the environment and the consumer.
61 Additionally, the application of these products can reduce the use of chemical fertilizers, increase
62 performance and stress resistance by water and temperature tensions and positively influence their
63 growth and physiology [8].

64 Today, seaweed extracts (SWE) provide an alternative, as they not only contain nitrogen,
65 phosphorus and potassium, but also present molecules with a very wide range of structures and can
66 be composed by phytohormones or metabolically active plant extracts such as amino acids and
67 organic acids [9]. Researchers [10,11] point out that seaweed extracts contain a wide variety of plant
68 growth-promoting substances such as auxins, cytokinins, betaines, gibberellins, and organic
69 substances, including amino acids, macronutrients, and trace elements that improve crop yield and
70 quality.

71 SWE are natural bioactive materials soluble in water; they are also organic fertilizers that
72 promote the germination of seeds and that increase the development and yield of crops [12]. They
73 have been used as nutritional supplements, bio-stimulants or biofertilizers in agriculture and
74 horticulture [13]. In recent years, the use of marine algae extracts as biofertilizers has allowed the
75 partial substitution of conventional mineral fertilizers [14]. These can be used as liquid extracts for
76 foliar and soil applications, or in granular form as soil improvers and fertilizer [15]. SWE have also
77 been used to increase germination, fruit weight and yield in tomato [16]; to improve chlorophyll and
78 carbohydrates contents, fruit diameter, yield, and vitamin C in cucumber plants, [9]; and to increase
79 leaf area, number of leaves, fresh stems and biomass in beans [17]. In *Stevia* plants (*Stevia rebaudiana*
80 Bertoni), marine algae extracts have improved some morphological, anatomical and chemical
81 characteristics [18]. Investigation was carried out on the effects of three seaweed extracts viz.,
82 *Asparagopsis* spp, *Gelidium pectinatum* and green alga *Enteromorpha intestinalis*, commercial seaweed
83 extract Algreen and compost on cucumber [9]. Results showed that the use of *E. intestinalis*, *G.*
84 *pectinatum* or commercial seaweed extracts with compost is considered a suitable application to
85 improve vegetative growth and yield of cucumber plants. The effect of three commercial extracts of
86 the brown seaweed, *Ascophyllum nodosum*, on phytochemical content and yield in cabbage plants
87 under field conditions was evaluated [19]. They found that total phenolic content was higher in all
88 seaweed treated plants, with the highest increase recorded with AlgaeGree (3.5 l ha⁻¹). Similar
89 increases were recorded in total flavonoid content. In another study to investigate the effect of the
90 Seaweed Liquid Fertilizer (SLF), *Rosenvigea intricata* --with or without chemical fertilizer -- on seed
91 germination, growth, yield and pigment content [20], it was found that 20% of SLF -- with or without
92 chemical fertilizer -- shows a higher growth, yield, chlorophyll pigment and soil profile compared to
93 other concentrations. A significant improvement in the size of olives and the quality of olive oil was
94 recorded in trees sprayed with an *A. nodosum* extract fortified with added nitrogen and boron [21].
95 Anbuechzhian et al., [22], also mention that the biofertilizers derived from marine algae form great
96 biofertilizers and improve the soil quality and yields considerably. Marine algal species *Ascophyllum*,
97 *Ecklonia* and *Fucus* are commonly utilized as fertilizers containing amounts of nitrogen and potassium
98 comparable to animal manure and organic fertilizers, but with a low phosphorus content.
99 Application rates, frequency and timing of the treatments vary with species, season, geographical
100 location, and local environmental variables. Important ancillary benefits of seaweed products for
101 crop production include the amelioration of damage caused by insects and bacterial or fungal
102 diseases [23].

103 Due to the above and based on the importance of the growing protected horticulture in the
104 Comarca Lagunera, the objective of this research is to improve the yield and phytochemical quality
105 of cucumber via leaf application of different seaweed extracts.

106 2. Materials and Methods

107 This study was carried out in shade mesh at the Technological Institute of Torreon (ITT), located
108 at km 7.5 of the old Torreon - San Pedro road, Municipality of Torreon, Coahuila. The mesh is an
109 Agro Shade model of 250 m² with a maximum wind resistance of 120 km/h, and greater than 35 m²
110 load capacity. The mesh was 16X10 strands of stabilized monofilament of flat fabric at 50% shade.
111 The structure was manufactured with materials in accordance with the Mexican standard for the
112 design and construction of greenhouses NMX-E-255-CNCP-2008 [24].

113 The treatments evaluated were five biofertilizers formulated with seaweed extracts: T1 =
114 *Macrocystis pyrifera* (Macro), T2 = *Bryothamnion triquetrum* (Bryo), T3 = *Ascophyllum nodosum* (Asco), T4
115 = *Grammatophora* sp. (Gramma), T5 = *Macrocystis integrifolia* (Macro integri). They were compared with
116 a control treatment (T6) using Steiner Solution (SS).

117 2.1 Obtaining the seaweed extracts

118 Algae *Macrocystis pyrifera* and *Grammatophora* sp., were collected in the Pacific Ocean off the coast
119 of La Paz, Baja California Sur (29° 09'N, 110° 19'W), Mexico. The algae were washed with distilled
120 water and dried for 7-10 days at room temperature. For the preparation of the extracts, the dried algae
121 were ground in mortar and sieved. Subsequently, the aqueous extract was obtained by performing
122 an extraction with distilled water 1:5 (w/v) for 5 h with constant agitation at room temperature. It
123 was then centrifuged at 800 x g in a HealForce centrifuge (Beckman GS-GKR, USA) at 40 °C for 20
124 minutes. Subsequently, each sample was crushed in a mill and used to determine the chemical
125 composition by undergoing acid digestion.

126 Organic matter, N, F, K, Mg, Bo, Zinc and Si were determined by atomic absorption
127 spectrophotometry [25]. Then 100 g of each sample was added to 1 L of distilled water with constant
128 stirring for 15 min followed by autoclaving at 121 °C for 1h at 1.21 kg cm⁻² [26]. The hot extracts were
129 filtered through a Whatman No. 40 filter paper and stored. In the case of the *Bryothamnion triquetrum*,
130 *Ascophyllum nodosum* and *Macrocystis integrifolia* algae, commercial algae existed on the market under
131 the name of Fulvimax^{AT}, Seaplant^{AT}, y Gaia^{AT}, respectively.

132 The aqueous solution was applied at 10% concentration, via Foliar spray, at an average of 150
133 ml per plant according to the technical recommendation for horticultural crops. The applications
134 were started eight days after the transplant and, subsequently, every 2 weeks up until 80 days after
135 the transplant (dds).

136 The chemical characteristics of the algae are presented in Table 1. The experimental design was
137 completely randomized with six treatments and six repetitions resulting in 36 experimental units. An
138 analysis of variance (ANOVA) and the Tukey separation test (P≤0.05) were performed using the
139 statistical program SAS ver. 6.03 [27].

140

141 **Table 1.** Chemical composition of seaweed extracts evaluated in cucumber production.

| Compound | <i>Macrocystis</i> | <i>Bryothamnion</i> | <i>Ascophyllum</i> | <i>Grammatophor</i> | <i>Macrocystis</i> |
|------------------|--------------------|---------------------|--------------------|---------------------|---------------------|
| | <i>pyrifera</i> | <i>triquetrum</i> | <i>nodosum</i> | <i>sp.</i> | <i>integrifolia</i> |
| | % P/V | | | | |
| Organic material | 3.49000 | 4.15000 | n.a | 3.50000 | 2.94400 |
| Nitrogen | 0.14700 | 1.45000 | 3.00000 | 1.14500 | 0.02630 |

| | | | | | |
|------------|---------|---------|---------|---------|---------|
| Phosphorus | 0.00800 | 1.36000 | 0.10000 | 0.02000 | 0.00540 |
| Potassium | 0.07700 | 1.48000 | 5.30000 | 2.00000 | 1.31000 |
| Magnesium | 0.01360 | n.a | 0.15000 | 0.02000 | 0.00236 |
| Bore | 0.00061 | n.a | 0.0003 | n.a | 0.00093 |
| Zinc | 0.00013 | 0.00075 | n.a | 0.00025 | 0.00126 |
| Silicium | n.a* | 0.00027 | n.a | n.a | n.a |

142 Source: Technical specifications of the product. *n.a = not available.

143 2.2. Substrate use

144 The substrate was organic using the proportions 80% sand + 20% vermicompost. This substrate
145 has been evaluated in several studies showing the best results for this vegetable [28]. The chemical
146 and physical characteristics of the substrate are presented in Table 2.

147 **Table 2.** Chemical and physical composition of the organic substrate (sand: vermicompost) used in
148 the evaluation of algae extracts in the shade-grown cucumber culture.

| Nutrient | Sand + Vermicompost (80:20 v/v) | | |
|------------|---------------------------------|--------------------------|--------|
| | mg kg ⁻¹ | | |
| Nitrate | 13.72 | Texture (sand-silt-clay) | 97-1-2 |
| Phosphorus | 15.33 | CEC (meq/100g) | 9.00 |
| Carbonates | 3.30 | pH | 7.91 |
| Potassium | 198.5 | OM (%) | 7.04 |
| Iron | 1.76 | CE (dS m ⁻¹) | 3.21 |
| Copper | 0.49 | R | 2.95 |
| Zinc | 0.53 | ESP | 2.99 |
| Manganese | 2.91 | | |

149 Source: Analyzes carried out in the Laboratory of ITT and the Coop. Agrop. of Comarca Lagunera. CEC=
150 Cation exchange capacity, pH= Hydrogen potential, OM = Organic material, R=Rate; ESP=
151 Exchangeable sodium percentage.

152

153 A hydroponic substrate formulated with disinfected and sterilized river sand, along with
154 perlite in a ratio of 80:20 (v/v) was used for the control treatment. The algae and the control
155 treatments were established in polyethylene plastic pots, caliber 600 of 18 kg capacity.

156 2.3. Agronomic management of the crop

157 The experiment was carried out during the autumn-winter agricultural cycle. On September 1st,
158 cucumber (*Cucumis sativus* L.) American type parthenocarpic, hybrid, Hisham-EZ 1110 was planted

159 in 200-well polyethylene trays using the Premier Promix P6X peat moss substrate, using one seed per
160 cavity. The handling of the trays consisted in maintaining the relative humidity (80%) and adequate
161 temperature (25-27 °C) for its development. The transplant to the pots took place 15 days after
162 planting when the seedlings had a height between 5 - 7 cm and three to five true leaves. The pots
163 were accommodated in such a way that the population density was of 4 plants per square meter in
164 double rows separated by 30 cm, with a distance of 40 cm between pots and 1.5 m between rows.

165 The irrigation of the pots was on average 0.750 - 1.0 L of water per pot per day; this condition
166 varied depending on the evaporation and environmental conditions within the shade mesh. The
167 control treatment (SS) was irrigated with nutrient solution [29] every day applying an average of
168 0.750 L per pot. The solution (SS) was diluted at a rate of 30, 50, 75 and 100% concentration according
169 to the phenological stage of the culture. Organic products were used in the control of pests and
170 diseases, which included organic repellents. A dose of 40 mL of neem extract and garlic extract in 20
171 L of water was applied once a day in the morning at 8 day intervals for the control of whitefly (*Bemisia*
172 *tabaci*).

173 The agronomic management was in accordance with the usual practices of the producer of the
174 region. The stems were trained vertically, sustained with polypropylene agricultural raffia yarn. All
175 buds were eliminated below 40 cm of the main stem, and later on, all lateral buds were eliminated by
176 weekly pruning, leaving one fruit per axilla, until the plant reached 2.0 m in height.

177 2.4. Evaluated Variables

178 For the analyzes nine mature cucumber samples were taken in each treatment. The length of the
179 fruit was determined using a metal tape, with a precision of 1 mm. A Mitutoyo digital vernier, model
180 CD-6 " CS, with a precision of 0.01 mm was used for the diameter of the fruit in mm. The firmness of
181 the fruits was measured using an Exttech penetrometer (FHT200) with a maximum capacity of 20 kg
182 / 44.10 lb or 196.10 Newton, and the average of two measurements per fruit was reported in Newton
183 (N). The total soluble solids were measured with a 0 - 32% manual refractometer (Sper scientific
184 300001) and the results were reported in ° Brix. A model AUY Shimadzu digital balance was used to
185 weigh the fruit in grams, with a precision of 0.1 mg. The cucumber harvest was approximately 60
186 days after the transplant (daT); the fruits with commercial maturity were harvested, those with
187 smooth and straight cylindrical surfaces, a dark, uniform green color (with no yellowing),
188 considering the A category of 3.5 - 5.0 cm in diameter and 14.0 - 16.5 cm in length, according to the
189 Mexico Supreme Quality mark, as the norm for commercial production [30]. The yield was obtained
190 from all harvested fruits that fulfilled this consideration. The result was expressed in grams per plant
191 (g plant⁻¹) and kilograms per square meter (km⁻²).

192 193 2.5. Samples preparation for phytochemical quality analysis

194 The harvested fruits were washed with drinking water for 2 minutes to remove residues and
195 lyophilized for 5 days. Subsequently, the dry material was pulverized manually (using mortar and
196 pestle) and stored in plastic tubes at -18 °C until extracts were obtained.

197 2.5.1. Obtaining extracts for phytochemicals analysis

198 250 mg of dry sample were mixed in 5 ml of HPLC-grade methanol in plastic tubes with screw
199 caps, which were placed on a rotary shaker (ATR Inc., EU) for 24 hours at 20 rpm at room temperature
200 (25 °C). Then, the tubes were centrifuged at 3000 rpm for 5 minutes and the supernatant was removed
201 for analysis.

202 2.5.2. Total phenolic content

203 The total phenolic content was measured using a modification of the Folin-Ciocalteu method
204 [31]. The content was calculated by means of a standard curve using gallic acid (Sigma, St. Louis,
205 Missouri, USA) as the standard, and the results were reported in mg of equivalent gallic acid per g
206 of fresh base sample (mg equiv AG / g BF). The analyzes were performed in triplicate.

207 2.5.3. Equivalent antioxidant capacity in Trolox (ABTS+ method)

208 The equivalent antioxidant capacity in Trolox was evaluated by the in vitro ABTS+ (2,2'-azino-
209 bis (3-thylbenzthiozoline-6-sulphonic acid)) method published by Esparza et al. [31]. The results were
210 reported as equivalent antioxidant capacity in mM equivalent in Trolox per g fresh base (mM equiv
211 Trolox / g BF).

212 2.5.4. Equivalent antioxidant capacity in Trolox (DPPH+ method)

213 Also, the antioxidant capacity was evaluated through the DPPH+ (1,1-diphenyl-2-
214 picrylhydrazil) in vitro method using a modification of the method published by Esparza et al. [31].
215 The results were reported as equivalent antioxidant capacity in μM equivalent in Trolox per g fresh
216 base (μM equiv Trolox / 100 g BF).

217 2.6 Vitamin C content

218 The vitamin C content was determined using a modification of the chromatograph method cited
219 by Esparza et al. [31]. The extract obtained was filtered through a membrane filter of $0.45\ \mu\text{m}$ before
220 being injected into high-resolution chromatography equipment (Hewlett Packard, 1200 Series, Palo
221 Alto, Calif., USA) using the Chem Station software for CL (Agilent Technologies, Palo Alto, Calif.,
222 USA). The ascorbic acid (vitamin C) was eluted on a C18 Supelco column ($150\ \text{mm} \times 5.0 \times 0.5\ \text{cm}$) at
223 a flow rate of $0.5\ \text{ml} / \text{min}$, using a 50:50 mixture of phosphoric acid 0.5% and methanol
224 (chromatograph grade) as the mobile phase. The effluent was monitored at $254\ \text{nm}$ in a diode array
225 detector. The results were obtained using a standard curve for ascorbic acid (Sigma) and reported as
226 mg of ascorbic acid per g fresh sample. The analyzes were performed in duplicate.

227 **3. Results**

228 3.1. Fruit quality

229 The length and diameter of the fruit, as well as the firmness and concentration of soluble solids,
230 were better in the chemical treatment (SS), as seen in Table 3. The best treatment of algae extracts for
231 affecting the size of the fruit, was the *Bryothamnion triquetrum* algae, being statistically equal to the
232 control treatment. Regarding diameter, the *Macrocystis integrifolia* algae was statistically equal to
233 the control and, with respect to SST, the *Macrocystis pyrifera* algae showed the lowest value (see
234 Table 3). In the case of firmness, there were no differences between treatments. El-Sharony et al. [33],
235 reported that the use of algae extracts is very effective for improving fruit set, fruit retention, yield
236 and enhanced fruit quality.

237 **Table 3.** Diameter, length, total soluble solids and firmness of cucumber fruits in shaded house
238 conditions, with foliar applications of algae extracts.

| Treatment | Fruit quality | | Weight | Total soluble solids |
|---------------------------------------|-----------------------|----------|-----------------------|-----------------------|
| | Length | Diameter | | |
| | cm | | g fruit^{-1} | $^{\circ}\text{Brix}$ |
| T1(<i>Macrocystis pyrifera</i>) | 13.360 d [†] | 3.380 d | 154.330 e | 3.440 b |
| T2(<i>Bryothamnion triquetrum</i>) | 16.430 a | 4.520 c | 264.660 b | 4.810 a |
| T3(<i>Ascophyllum nodosum</i>) | 14.730 c | 4.640 b | 223.660 c | 4.590 a |
| T4(<i>Grammatophora sp.</i>) | 15.930 ab | 4.600 bc | 210.330 d | 4.880 a |
| T5(<i>Macrocystis integrifolia</i>) | 15.330 bc | 4.800 a | 208.330 d | 4.690 a |

| | | | | |
|----------------------|----------|---------|-----------|---------|
| T6(Steiner Solution) | 16.730 a | 4.820 a | 286.000 a | 4.760 a |
| MSD | 0.982 | 0.086 | 10.711 | 0.428 |
| SE | 0.293 | 0.0257 | 3.203 | 0.128 |
| CV | 3.790 | 1.280 | 4.450 | 4.900 |

239 † Values with equal letters within each column are statistically similar (Tukey, $P \leq 0.05$). MSD=Minimum
240 Significant Difference; SE=Standard Error; CV=Coef. Var.

241 In regard to the size of the fruit, DeGannes et al. [34] have indicated that the length of the
242 American cucumber fluctuates between 20 and 25 cm, and not under 15 cm [30]. For the diameter of
243 the fruit, Farrag et al. [35] found an average diameter of 3.17 cm with seaweed extract; likewise,
244 Wittwer and Honma [36] found that the diameter of cucumber fluctuates from 5.0 to 5.7 cm, and
245 should not exceed 6.0 cm. From the results, it can be observed that the foliar applications of the algae
246 extracts showed similar values to those obtained in the control treatment. This result can be attributed
247 to the benefits of algae when used in horticultural crops. In this regard, Bajpai [37] indicated that
248 algae present active organic compounds as growth regulators. Metting et al. [38] have indicated that
249 the physiological responses derived from the application of marine algae include greater nutrient
250 mobilization, the development of a vigorous root system, an increase in chlorophyll content, leaf area
251 and delay in fruit senescence.

252 Regarding the content of total soluble solids (SST), the treatments *Bryothamnion triquetrum*
253 (T2), *Ascophyllum nodosum* (T3), *Grammatophora* sp. (T4) and *Macrocystis integrifolia* (T1) were
254 statistically equal to the treatment with Steiner solution (SS), with an average value of 4.76 ° Brix, and
255 different to the treatment with *Macrocystis pyrifera* algae which decreased the SST by 29% (Table 3).
256 Colapietra and Alexander [39] observed an increment of up to 17 ° Brix in a table grape crop (cv Italia)
257 with foliar application of seaweed extracts *Ascophyllum nodosum*.

258 Similarly, Khan et al. [40] reported increments of 28% of total sugars in Perville cv Vine by
259 applying *Ascophyllum nodosum*. On the other hand, Kumari et al. [41] mentioned a higher
260 concentration of total soluble sugars in the tomato crop with soil and foliar applications of seaweed
261 extract *Sagarssum johntonii*.

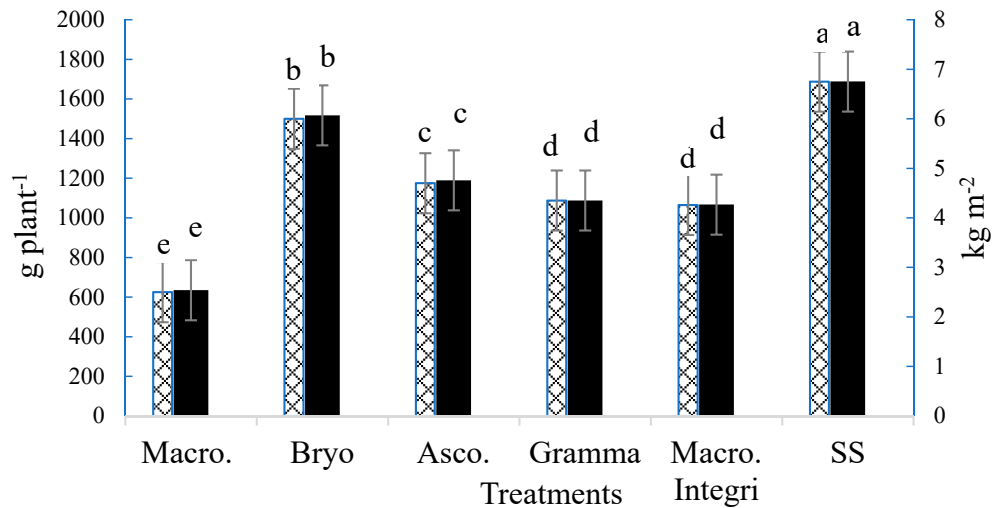
262 3.2. Weight of the fruit (g fruit⁻¹)

263 The weight of the American cucumber fruit should fluctuate between 300 to 400 grams. In this
264 study, the different algae extracts caused significant differences in weight ($P \leq 0.05$), with the control
265 treatment (SS) having the greatest weight. Regarding treatments with algae extracts, the
266 *Bryothamnion triquetrum* treatment was the best, with a 7% lower weight than that of the control,
267 but 41% higher than that of the treatment with the *Macrocystis pyrifera*, which obtained the lowest
268 weight. The differences in the weight of the American cucumber found in this study may be due to
269 different hybrids, different climate conditions and crop management. The specifications PC-021-2005
270 for the use of the official Mexico mark [30] establishes two types of cucumber in relation to its
271 attributes. The fruits obtained in this study fall into the classification B (diameter 5.1-6.5 cm and
272 length 14-16.5 cm) and, therefore, could compete in the national and International market.

273 3.3. Yield (g plant⁻¹)

274 The statistical analysis for this variable shows significant differences ($P \leq 0.05$) in the yield (g
275 plant⁻¹), which indicates that the foliar application of the marine algae extracts influenced the yield
276 (Figure 1). The control treatment with Steiner solution reported the highest yield, being 9% higher
277 than the best yield obtained with the algae extracts. The treatment with the *Bryothamnion triquetrum*
278 alga showed a 42% higher yield compared to the lower yield obtained with the alga *Macrocystis*
279 *pyrifera*. The higher yield obtained with *Bryothamnion triquetrum* may be due to its greater effect
280 on nutrient absorption. In this regard, Youssef et al. [42] have indicated that the higher production of

281 biomass and yield in cucumber plants could be attributed to the fact that the plants inoculated with
 282 biofertilizers have a greater capacity to maintain a high rate of net photosynthesis and a better
 283 nutritional status (high concentration of P, K, Mg, Fe, Zn and Mn) compared to non-inoculated plants.
 284 Likewise, Metting et al. [38] have mentioned that the physiological responses derived from the
 285 application of seaweed include greater mobilization of nutrients, the development of a vigorous root
 286 system, an increase in the chlorophyll content, leaf area and delay in the senescence of the fruit.



287

288 **Figure 1.** Effect of foliar application of seaweed extracts on the yield production (g plant⁻¹ and kg m⁻²) of cucumber plants established in shade mesh. Means followed by the same letter are not
 289 significantly different between treatments (Tukey; $P \leq 0.05$).
 290

291 It was observed that the control treatment (SS) presented the highest yield per square meter (kg
 292 m⁻²). The treatments with algae extracts showed values between 2.54 and 6.08 kg m⁻² (see Figure 1).
 293 Regarding the performance in protected systems in Mexico, by SIAP [43] reports, the average yield
 294 of cucumber is 98 t·ha⁻¹ (9.8 kg m⁻²). For example, the average yield for this shade mesh culture is
 295 17.6 kg m⁻² in the state of Baja California (Mexico) [44]. In the case of the Comarca Lagunera (Mexico),
 296 with medium-tech production technology in shade mesh and fertirrigation, for six plants, the yield
 297 fluctuates between 14 -17 kg m⁻² [45].

298 Comparing the benefits of the yields obtained in this experiment to those using chemical
 299 fertilization and algae, it can be mentioned that the latter are greater because they are friendly to the
 300 environment. The trend in the future is to produce organic cucumbers since they have a high demand
 301 by consumers. The payment of overpricing of these products in the foreign market could be higher
 302 in relation to the price of conventional products and, consequently, it would improve the cost-benefit
 303 ratio of their production.

304 3.4. Phytochemical quality

305 The phytochemical quality of the fruits (antioxidants, phenols and vitamin C) was influenced by
 306 the algae extracts ($p \leq 0.05$). Higher values were obtained with the *Macrocystis pyrifera* algae (Table
 307 4). Regarding the antioxidant capacity, in both determination methods (ABTS+ and DPPH+),
 308 *Macrocystis pyrifera* showed the highest value, followed by *Bryothamnion triquetrum*. Both values
 309 were statistically different from each other and different from the control treatment, exceeding it by
 310 30% and 4%, respectively. With the DPPH+ method, the treatment with chemical fertilization
 311 presented the lowest value. These results can be explained considering the nutrient deficiency that
 312 the substrate presents, particularly that of nitrogen for the plant. In this regard, Lorio [46] and Herms
 313 and Mattson [47] mentioned that plants produce higher amounts of sugars and secondary metabolites
 314 when they are subjected to a deficit of readily available nitrogen.

315 **Table 4.** Antioxidant capacity of cucumber produced in shade mesh with the application of different
 316 nutrient solutions derived from algae, using two methods (ABTS⁺ and DPPH⁺).

| Treatments | CAOX* | CAOX** |
|---------------------------------|-----------------------------|-----------------------------|
| | (Method ABTS ⁺) | (Method DPPH ⁺) |
| <i>Macrocystis pyrifera</i> | 149.4 a | 454.1 a |
| <i>Bryothamnion triquetrum</i> | 110.4 b | 373.9 bc |
| <i>Ascophyllum nodosum</i> | 90.6 c | 295.9 d |
| <i>Grammatophora sp.</i> | 77.0 d | 278.4 d |
| <i>Macrocystis integrifolia</i> | 120.4 b | 403.5 b |
| Steiner Solution (SS) | 105.9 b | 349.7 c |

317 * CAOX: Antioxidant capacity (ABTS⁺ method), results in μM equiv Trolox / 100 g fresh base.

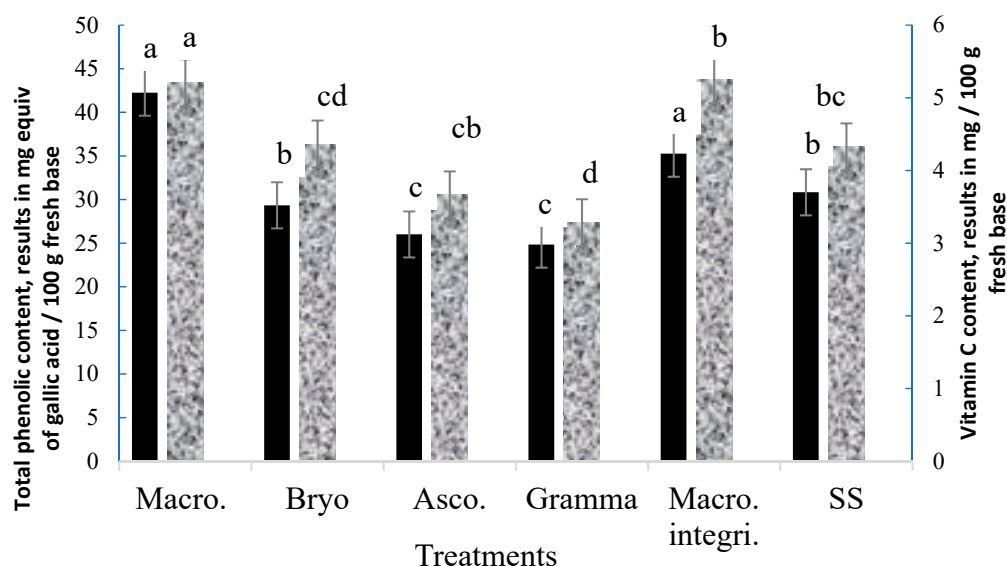
318 ** CAOX: Antioxidant capacity (DPPH⁺ method), results in μM equiv Trolox / 100 g fresh base.

319 Differences between means obtained by Tukey test ($P \leq 0.05$). Values in columns with different lowercase letters
 320 are statistically different ($P \leq 0.05$).

321 However, the results obtained contrast with those found by Santiago-López et al. [48] in which
 322 organic sources of fertilization were used for cucumber, obtaining antioxidant capacity values of
 323 779.9 to 1391.1 μM equiv Trolox / 100 g BF. On the other hand, Díaz-Méndez et al. [28] used different
 324 proportions of vermicompost and obtained antioxidant capacity values between 749.9 and 1015.2.
 325 Some studies have shown that high concentrations of NaCl in substrates increase the activity of
 326 antioxidant enzymes [49,50]. This result has been attributed to the synthesis of phenols by plants as
 327 a defense mechanism to counteract the negative effects of oxidative stress [51]. Therefore, phenolic
 328 compounds are the main drivers of antioxidant activity [52]. In Table 4, the phenolic content for the
 329 *Macrocystis pyrifera* algae had the highest value (47.37), which was expected since it generated the
 330 highest antioxidant capacity.

331 Cucumber provides a large amount of C vitamin (mg / 100 g fresh base) and, in this experiment,
 332 the treatments with the highest content were the algae *Macrocystis pyrifera* (5.07), *Macrocystis*
 333 *integrifolia*. (4.23) and the control treatment (3.70) (See Figure 2). The alga *Grammathosfera sp.*
 334 presented the lowest value -- 40% lower compared to *Macrocystis pyrifera*. Nagy and Pintér [53] have
 335 pointed out that the positive effect of seaweed extract can be explained by the high content of high-
 336 quality proteins, amino acids such as lysine and tryptophan, essential minerals, trace elements, B-
 337 complex vitamins and bio-constituents, especially cytokinins.

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Figure 2. Total phenolic and vitamin C content of cucumber produced in shade mesh with the application of different nutrient solutions derived from algae. Differences between means obtained by Tukey test ($P \leq 0.05$). Values with different lowercase letters are statistically different ($P \leq 0.05$).

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4. Conclusions

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Higher yields and commercial quality were obtained in cucumber fruits using chemical fertilization, mainly due to the higher concentration of N and the ionic balance of the nutritive solution. Of the treatments with algae extracts, *Bryothamnion triquetrum* was superior in total soluble solids, fruit firmness, fruit size and yield. However, the highest antioxidant capacity, phenols and vitamin C were obtained with the *Macrocystis pyrifera* extract. The algae extracts provided some phyto-nutrients for the nutrition of the cucumber plants and increased the quality of the fruits by obtaining greater antioxidant capacity. Therefore, its use is a viable option to minimize the application of conventional fertilizers, thereby attenuating the effects on the environment and improving the health of the population.

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