

Article

A Chemical Investigation of the Antioxidant Capacity of Extracts from Red Macroalga *Gracilaria domingensis*

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Abstract: Extracts that were obtained with solvents of increasing polarity (hexane, dichloromethane, methanol, 80% methanol, and water) from the red macroalga *Gracilaria domingensis* were evaluated by reducing power with ferric reduction antioxidant power (FRAP) and Folin–Ciocalteu (FC) assays, lipid peroxidation inhibition by β -carotene-linoleic acid assay, and metal chelating ability based on the iron-ferrozine system. The highest antioxidant capacity was reported for the hexane (Hx) extract by the FRAP, metal chelating, and lipid peroxidation inhibition assays. An activity-guided fractionation of the Hx extract was carried out for the identification of its active constituents. The primary components were the most active antioxidant compounds. Despite the high antioxidant activities, the Hx extract was not active in the FC assay. In this assay, the activities were found in the methanol (M) and 80% methanol (80M) extracts. The FC assay is commonly used to measure the total phenolic compounds. However, no phenolic compounds were detected by GC-MS and HPLC analyses in the M and 80M extracts. Thus, non-phenolic components influenced the FC assay. The M and 80M extracts showed high content of mycosporine-like amino acids (MAAs). A fraction contained two MAAs (porphyra-334 and shinorine) ($156 \text{ mg GAE} \cdot \text{g}^{-1}$) showed a similar performance to the values that were found for well-known antioxidants (BHT = $156 \text{ mg GAE} \cdot \text{g}^{-1}$ and Trolox = $166 \text{ mg GAE} \cdot \text{g}^{-1}$) and 30 times higher than those of the original extracts ($\sim 5 \text{ mg GAE} \cdot \text{g}^{-1}$) in the FC assay. Thus, MAAs contribute to the antioxidant activities that were observed in the FC assay within the studied samples. Together, these results advance our understanding of the antioxidant properties of algal extracts.

Keywords: antioxidant; mycosporine-like amino acids; total phenolic content; Folin–Ciocalteu; *Gracilaria*



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1. Introduction

Reactive oxygen species (ROS) are produced in the organism naturally as a byproduct of metabolism. An imbalance between ROS production and antioxidant protection creates oxidative stress, which can lead to biomolecule damage. Antioxidants play a crucial role in biological systems since they prevent or delay the oxidation of these biomolecules. Evaluating the antioxidant capacity provides relevant information that may be used in different research fields, such as studies of the abundance and distribution of species in ecology [1] and understanding of the physiological state of species under stress [2]. Besides, in vitro antioxidant assays are low-cost and require simple laboratory procedures.

There are several in vitro assays to measure the antioxidant capacities. These in vitro assays can evaluate different mechanisms of antioxidant action [3]. Assays that are based on hydrogen atom transfer (HAT) measure the free-radical scavenging capacity of an antioxidant by hydrogen donation [4]. The inhibition of β -carotene bleaching that is caused by peroxidation of linoleic acid is a well-known HAT-based assay. In the assays that are based on electron-transfer (ET) reaction, the free radical is stabilized by donating an electron from an antioxidant [5]. The ability to reduce ferric iron (Fe^{3+}) to ferrous iron (Fe^{2+}) measured on the FRAP (ferric reduction antioxidant power) and the ability to reduce the Folin–Ciocalteu (FC) reagent that was measured in the FC assay are based on ET reactions.

In the literature, the FC assay is commonly used for the quantification of the total phenolic content (TPC). Despite the popularity for this use, the electron transfer reactions to the FC reagent are not performed exclusively by the phenolic compounds [6,7], so the scientific community has discussed the use of the FC assay for measuring concentrations of TPC [8]. Yet, a third kind of assay can measure the ability of the antioxidant to chelate transition metals. Metal chelating assays measure the ability to sequester transition metals that cause free radical generation. A chelator agent such as ferrozine is used in these assays.

Due to the diversity of mechanisms of antioxidant action and radical types, an antioxidant may react in an assay, but it may be not sensitive in others. Consequently, more than one assay should be used to evaluate the antioxidant capacity of a sample. This capacity is related to the interaction effects (synergistic, additive, and antagonistic) between compounds. Detterweiler et al. [9] proposed an index to evaluate how much the interaction between the extract components is responsible for biological activities. Interactions between antioxidant components are often in complex mixtures such as extracts [10]. Thus, the activity-guided fractionation can provide us with a better understanding of the antioxidant properties of crude extracts.

Antioxidant capacity has been studied in numerous genera of macroalgae. Part of these studies searches for natural antioxidants and extracts for health, food, and cosmetic applications [11,12] and other topics such as the evaluation of the effect of abiotic stresses, or variation among seasons. For example, the antioxidant capacity from *Gracilaria cornea* (Rhodophyta) increased under UV exposure [13], and the antioxidant capacity differed between dry and rainy seasons for *Sargassum vulgare* (Ochrophyta) and *Palisada flagellifera* (Rhodophyta), with the highest antioxidant activities in the rainy season for the first and in the dry season for the last [14].

Gracilaria (Gracilariaceae, Rhodophyta) is a genus of red macroalga comprising of about 197 species that are distributed nearly worldwide [15]. This genus includes species of economic value due to the agar, a phycocolloid that is widely used in the food and biotechnology industries [16]. The antioxidant potentials from *Gracilaria* species are reported mainly for crude extracts [17], with promising results for some species. Methanolic extracts from *Gracilaria corticata* and *Gracilaria edulis* have significant activities in the FRAP assay when compared to BHT (butylated hydroxytoluene) [18], and some crude extracts from *Gracilaria gracilis* have higher or similar reducing activities than BHT and gallic acid. [19].

Compounds with antioxidant properties that are already identified in the genus include bromophenols from *Gracilaria edulis* [20], carotenoids from *Gracilaria birdiae* and *Gracilaria tenuistipitata* [21,22], floridoside from *G. tenuistipitata* [23], citric acid from *Gracilaria vermiculophylla* [24], and mycosporine-like amino acids (MAAs) from several species [13,25–27]. Phenolic compounds are among the most potent antioxidants in nature, and they are found in abundance in land plants and brown algae. Benzoic acid, gallic acid, gentisic acid, p-hydroxybenzoic acid, protocatechuic acid, salicylic acid, syringic acid, and vanillic acid are phenolic compounds that have been identified in *Gracilaria* [17].

Gracilaria domingensis (Kützinger) Sonder ex Dickie is distributed in the Western Atlantic Ocean from Mexico to northeast Brazil. The antioxidant potential of *G. domingensis* was evaluated in two studies. One study evaluated only crude extracts [21], and another study focused on the algal nutritional potential using bleached samples [25]. Unlike previous studies, we investigate the components that are responsible for the antioxidant capacities of the crude extracts using activity-guided fractionation. This procedure allows locating the compounds that are active. To this end, we selected assays that evaluate the main mechanisms of antioxidant action: electron-transfer reaction (FRAP and Folin–Ciocalteu assays), hydrogen atom transfer reaction (β -carotene-linoleic acid assay), and the ability to chelate transition metals based on the measurement of the iron-ferrozine complex.

2. Material and Methods

2.1. Algal Samples

Gracilaria domingensis was collected in the intertidal zone at “Morro de Pernambuco” Beach in the city of Ilhéus in Bahia state (14°48′21.6″ S, 39°01′25.6″ W). The fresh sample (10 kg) was rinsed in tap water and triaged for removal of epibionts. DNA barcoding (*COI-5P*—mitochondrial cytochrome *c* oxidase subunit I gene) and morphological characters were used for taxonomical identification of the species [28,29]. The samples were initially air-dried at room temperature in a shaded environment (~one day), and then in an air circulation oven at 40 °C (~two days).

2.2. Preparation of the Crude Extracts and Fractionation of the Active Extracts

Figure 1 depicts the general scheme of the extraction and fractionation procedure. The dried samples were ground in a knife mill (30-mesh; Fortinox® STAR FT 80, Piracicaba, Brazil). The algal powder (1 kg) was macerated at 50 ± 5 °C for 8 h in organic solvents (1:10, *w/v*) in increasing order of polarity starting with hexane, followed by dichloromethane, methanol, and 80% methanol. The final residue was macerated in ultrapure water (1:20, *w/v*) at 50 ± 5 °C for 8 h. The dried methanolic and hydromethanolic extracts were desalted using methanol precipitation by mixing with one volume of methanol. After the homogenization, the sample was allowed to sit at room temperature for 15–30 min, and the supernatant was collected. The organic extracts were concentrated in a rotary evaporator (<45 °C), and the aqueous extract was freeze-dried. There were five crude extracts that were obtained: hexane (Hx = 4.2 g), dichloromethane (DCM = 2.9 g), methanol (M = 65 g), 80% methanol (80M = 29 g), and aqueous (Aq = 340 g).

The Hx and M extracts were partitioned and fractionated. The Hx extract was dissolved in DCM and partitioned with 50% methanol (yields of 3.5 g and 0.5 g, respectively). Part of the DCM phase (2.5 g) was fractionated in silica column chromatography (silanized silica gel 60, Merck Millipore, Darmstadt, Germany) using as mobile phase a solvent gradient in decreasing order of polarity (water—250 mL, 25% methanol—250 mL, 50% methanol—250 mL, 80% methanol—250 mL, methanol—250 mL, methanol:dichloromethane (7:3)—250 mL, methanol:dichloromethane (3:7)—2 × 250 mL, dichloromethane—6 × 250 mL, and hexane—250 mL). All the fractions were monitored by analytical thin-layer chromatography (TLC; Merck Silica Gel 60 F₂₅₄, Darmstadt, Germany), visualized using iodine vapor, and pooled together according to elution similarity, resulting in five groups (F1—20 mg, F2—16 mg, F3—165 mg, F4—632 mg, and F5—1.6 g). Part of the group F5 (41 mg) was fractionated in preparative thin-layer chromatography (TLC; 60G silica gel, Merck Millipore, impregnated with 0.02% sodium fluorescein). The mobile phase was hexane:diethyl ether:acetic acid (80:20:1, *v/v/v*). The plates were visualized under UV light, and five bands were scraped off the silica gel, resulting in the subfractions S1 (3.4 mg), S2 (11 mg), S3 (3 mg), S4 (4.3 mg), and S5 (19 mg). The M extract was dissolved in 50% methanol and partitioned with DCM (yields of 59 g and 5 g, respectively). Part of the 50% methanol phase (531 mg) was fractionated by semi-preparative high-performance liquid chromatography (HPLC; Agilent 1200 series, USA) equipment using a semi-preparative column (Zorbax RX-SIL HILIC, 9.4 mm × 250 mm, 5 µm, USA), with a mobile phase of acetonitrile (solvent A) and 0.2% acetic acid (solvent B) in isocratic elution with 68% A for 20 min, and flow rate of 4 mL min^{−1}. This fractionation procedure resulted in a fraction that was rich in two MAAs (31 mg; porphyra-334 and shinorine, 70% and 30% respectively).

The chemical characterization of the active extracts and fractions was performed using analytical TLC plates, GC-MS (Gas Chromatography-Mass Spectrometry), and an analytical HPLC system. The group F5 was analyzed by analytical TLC (Merck Silica Gel 60 F₂₅₄, Darmstadt, Germany) [30]. The mobile phase was hexane:diethyl ether:acetic acid (80:20:1, *v/v/v*), and the plates were visualized using iodine vapor. The M and 80M extracts and subfractions S1 to S5 were analyzed by GC-MS and HPLC with Diode Array Detection (HPLC-DAD). For the GC-MS analysis, the samples (100 µg) were dissolved in 50 µL of pyridine and derivatized by reaction with 50 µL of BSTFA for 30 min at 80 °C. The derivatized samples were

evaluated by GC-MS (Agilent 6890N/5975, Malaysia) that was equipped with an HP-5MS column (30 m × 0.25 mm, 0.25 µm, USA) [31]. For the HPLC–DAD analysis, the samples were injected into a 1260 Infinity LC System (Agilent, Santa Clara, CA, USA). A ZORBAX Eclipse plus C18 column (150 mm × 4.6 mm, 3.5 µm, USA) was used for the analysis of the metabolites, and the mobile phase was a gradient of 0.1% acetic acid and acetonitrile [32]. The M and 80M extracts were also analyzed for MAAs through HPLC-DAD using an HILIC column (Zorbax RX-SIL, 250 mm × 4.6 mm, 5 µm, USA), and a gradient of acetonitrile:5 mM ammonium acetate (9:1, *v/v*) and acetonitrile:5 mM ammonium acetate (1:1, *v/v*) as mobile phase [33]. The MAAs were identified by comparison with isolated standards and quantified using a calibration curve of porphyra-334 ($R^2 > 0.99$).

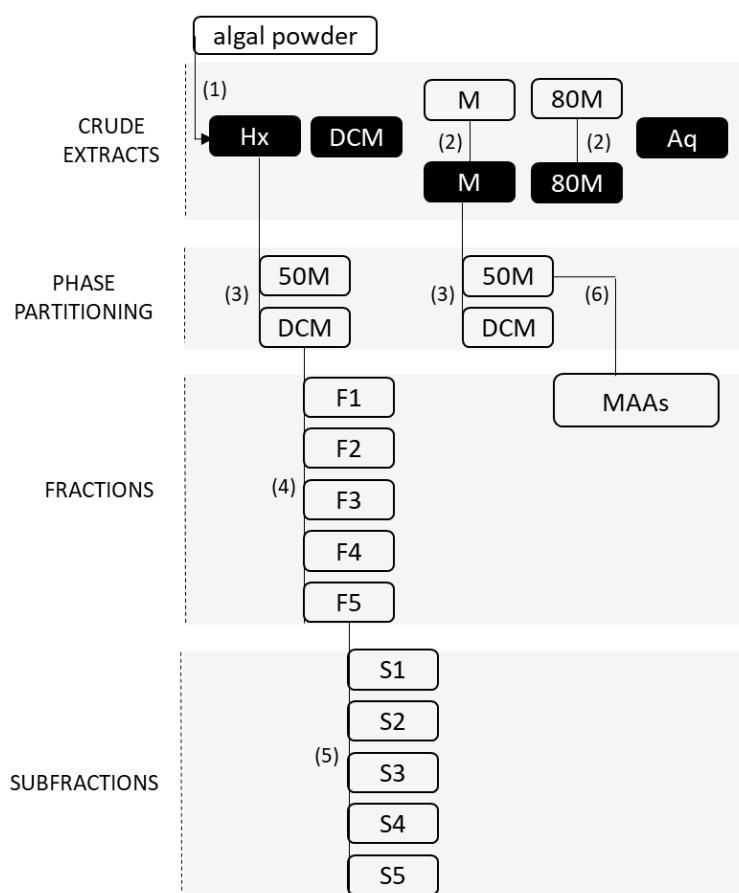


Figure 1. Scheme of the extraction and fractionation procedure depicting the crude extracts (Hx—hexane, DCM—dichloromethane, M—methanol, 80M—80% methanol, Aq—aqueous), the phase partitioning (50M—50% methanol and DCM—dichloromethane), the fractions F1 to F5, a fraction rich in two MAAs (mycosporine-like amino acids; porphyra-334 and shinorine), and the subfractions S1 to S5. 1. Extraction. 2. Desalination. 3. Partition, 4. Silica column chromatography, 5. Preparative thin layer chromatography (TLC), and 6. Semi-preparative high-performance liquid chromatography (HPLC).

2.3. Determination of the Antioxidant Capacity

All the assays were performed in 96-well clear polystyrene microplates using a microplate reader (Synergy™ H1 BioTek®, Winooski, VT, USA). All the samples (crude extracts, phase partitioning, fractions, and subfractions) were dissolved in DMSO and diluted in ultrapure water to obtain sample concentrations between 0.6 and 6 mg mL^{−1} (final concentrations in the wells between 40 and 400 µg mL^{−1}) and a DMSO concentration of 10% (*v/v*). In all the assays, a negative control (sample volume was replaced by 10% DMSO), positive control (sample volume was replaced by standard), and sample blank (sample + reagents of the assay, but without one reagent to avoid the reaction) were in-

cluded. Sample blanks are used to avoid errors from color or turbidity that may exist in the sample. Only corrected absorbances by subtracting the sample blank absorbance from the absorbance after reaction were used for calculations. There were two natural antioxidants (rutin and p-coumaric acid) and two synthetic antioxidants (Trolox and BHT—butylated hydroxytoluene) that were used as positive controls. The results were expressed in gallic acid equivalents ($\text{mg GAE} \cdot \text{g}^{-1}$ dry extract) for the Folin–Ciocalteu and metal chelating assays, Trolox equivalent ($\mu\text{mol TE} \cdot \text{g}^{-1}$ dry extract) for the FRAP assay, and protection percentages from the negative control for the β -carotene-linoleic acid assay.

2.4. Ferric Reduction Antioxidant Power (FRAP) Assay

The reducing power by the FRAP assay was conducted following the method that was described previously [34]. The FRAP solution was prepared mixing 0.3 M acetate buffer (pH 3.6), 10 mM TPTZ in 40 mM hydrochloric acid, and 20 mM ferric chloride in a ratio of 10:1:1. Briefly, a volume of 20 μL of sample (final concentration in the well: 400 $\mu\text{g mL}^{-1}$ for crude extracts and 150 $\mu\text{g mL}^{-1}$ for other samples), negative control, or standard was added to each well along with 15 μL of ultrapure water and 265 μL of FRAP solution. After incubation at 37 °C for 20 min, the absorbances were measured at 595 nm. For the sample blanks, a FRAP solution was prepared without TPTZ. A calibration curve was constructed using Trolox (final concentrations in well between 0.5 and 7 $\mu\text{g mL}^{-1}$, $R^2 > 0.99$).

2.5. Metal Chelating Ability Based on the Measurement of Iron-Ferrozine Complex

The metal chelating ability assay was performed as the method in the microplate format that was described previously [35]. Briefly, a volume of 20 μL of sample (final concentration in the well: 400 $\mu\text{g mL}^{-1}$ for crude extracts and 150 $\mu\text{g mL}^{-1}$ for other samples), negative control, or standard was added to each well, followed by the addition of 260 μL of 10% (w/v) ammonium acetate and 10 μL of 1 mM ferrous ammonium sulfate. After incubation for 5 min, a volume of 10 μL of 6.1 mM ferrozine was added to each well. The microplate was incubated for 10 min, and the absorbances were measured at 562 nm. For the sample blanks, the volume of the ferrozine solution in the well was replaced by ultrapure water. A calibration curve was constructed using gallic acid (final concentrations in the wells between 1 and 14 $\mu\text{g mL}^{-1}$, $R^2 > 0.99$).

2.6. Lipid Peroxidation Inhibition Using the β -Carotene-Linoleic Acid Assay

The assay of inhibition of β -carotene bleaching was performed with modifications from the methods that were described previously [14]. The reactive solution was prepared by mixing 16 μL of linoleic acid, 160 μL of tween 40, and 200 μL of β -carotene (2 mg mL^{-1} in dichloromethane). The solution was evaporated until the complete removal of the dichloromethane (<45 °C), and then 50 mL of ultrapure water that was bubbled with air for 30 min was added. The absorbance was adjusted to 0.90 ± 0.10 at 450 nm. For the assay, 20 μL of sample (final concentration in the well: 100 $\mu\text{g mL}^{-1}$), negative control, or standard was added to each well with 280 μL of reactive solution. The microplate was incubated at 45 °C under stirring, and the absorbance was measured at 450 nm in the time 0 min (sample blank) and 120 min.

2.7. Folin–Ciocalteu (FC) Assay

This assay was performed based on a previously described method [36]. Into each well containing 200 μL of ultrapure water, 20 μL of sample (final concentration in the well: 400 $\mu\text{g mL}^{-1}$ for crude extracts and 40 $\mu\text{g mL}^{-1}$ for fraction), negative control, or standard was added along with 20 μL of Folin–Ciocalteu reagent, and 60 μL of saturated sodium carbonate (25%; w/v). After incubation for 30 min, the absorbances were measured at 760 nm. For the sample blanks, the volume of the FC reagent in the well was replaced by ultrapure water. A calibration curve was constructed using gallic acid (final concentrations in the well between 1 and 10 $\mu\text{g mL}^{-1}$, $R^2 > 0.99$).

2.8. Interaction Indexes

Interaction indexes (Ii) were calculated based on Dettweiler et al. [9], with adaptations to use for values that are expressed in mass equivalent of the reference antioxidant or inhibition percentage. Values ≤ 0.5 indicate synergy, values between 0.5–4 indicate non-interaction, and values > 4 indicate antagonism. The index was calculated using the following formula: $(A_s \times Y_s)/A_f$, where A_s = antioxidant value of the fraction/subfraction, Y_s = yield of the fraction/subfraction, and A_f = antioxidant value of the original sample. The formula was applied to each fraction/subfraction, and the sum of all the values is the interaction index.

2.9. Statistical Analysis

The data analysis and graphing software was GraphPad 8[®]. The results were expressed as the mean $\pm$ standard deviation (SD; $n = 3$, technical triplicate). The comparisons between the means were performed by one-way ANOVA. The Tukey test was used in post hoc analysis ($p < 0.05$).

3. Results

The Aq extract showed the highest yield (34% on dry algal mass), while the Hx and DCM extracts gave the lowest yields (values $< 1\%$). The M and 80M extracts yielded 6.5% and 3%, respectively. The Hx extract was the most effective in three-antioxidant assays: lipid peroxidation inhibition (Figure 2A), FRAP (Figure 2B), and metal chelating assay (Figure 2C). However, this extract had no activity in the FC assay (Figure 2D). The DCM extract showed weak to moderate activities in the four assays (Figure 2). The M and 80M extracts were the most effective in the FC assay (Figure 2D), while the Aq extract showed lipid peroxidation inhibition (Figure 2A) and metal chelating ability (Figure 2C).

An activity-guided fractionation of the Hx extract was carried out for the identification of its active constituents (Figure 3). The fractionation procedure resulted in the fraction F5 with higher antioxidant activities than those of the crude extract (Figure 3). The level of lipid peroxidation inhibition for the fraction F5 (85% of protection) was higher than the value for p-coumaric acid—74% of protection ($100 \mu\text{g mL}^{-1}$). However, further purifications from this fraction resulted in a loss of lipid peroxidation inhibition, with values ranging from 10% to 20% of protection for all subfractionations (Figure 3A). The interaction index (Ii) among them was 0.23, suggesting that the subfractions from F5 have a strong synergistic effect. On the other hand, a synergistic effect was not observed among subfractions on FRAP and metal chelating assays (interaction index between 0.5–4). Similar activities to fraction F5 were found for subfractions S2 and S4 in the FRAP assay (Figure 3B) and for S2 in the metal chelating assay (Figure 3C).

The chemical composition of the fraction F5 includes free fatty acids, triacylglycerols, monopalmitin, chlorophyll *a* and its derivatives, cholesterol, and polar lipids (Figure 3D). The latter is also found in fractions F1 to F4 (Figure 3D). Concerning active subfractions, GC-MS analysis confirmed monopalmitin (40%) as the main peak in the subfraction S2 and cholesterol (97%) in the subfraction S4. Chlorophyll *a* and its derivatives were confirmed in the subfractions S3 and S4 using HPLC analysis.

Once the M and 80M extracts were the most active in the FC assay (Figure 2D), the search for phenolic compounds was carried out in these extracts. However, even after analysis in GC-MS (Figure S1) and HPLC (Figure 4A,B), no phenolic compounds were detected in these extracts. Carboxylic acids (e.g., citric acid), free amino acids (e.g., glutamine), heterosides (floridoside and isofloridoside), monosaccharides, and isethionic acid were characteristic components of these extracts that were detected by GC-MS (Figure S1 and Table S1), while MAAs were practically the unique class of compounds that were found in the HPLC-DAD chromatograms (Figure 4A,B). The methodology for analysis for specific MAAs allowed identification of porphyra-334 as the major MAAs of the polar extracts, followed by shinorine, palythine, asterina-330, and palythanol (Figure 4C,D).

The polar extracts contained a high amount of MAAs, with contents of 7.8% and 9.6% (dry mass of the extract), respectively, for the M and 80M extracts. Due to the remarkably high MAAs contents in the M and 80M extracts, a fraction of MAA (porphyra-334 + shinorine, 70% and 30%, respectively) was isolated and evaluated in the FC assay. This fraction ($156 \text{ mg GAE} \cdot \text{g}^{-1}$) showed a similar performance to the values that were found for well-known synthetic antioxidants (BHT = $156 \text{ mg GAE} \cdot \text{g}^{-1}$ and Trolox = $166 \text{ mg GAE} \cdot \text{g}^{-1}$) and 30 times greater than those of the original extracts ($\sim 5 \text{ mg GAE} \cdot \text{g}^{-1}$) (Figure 2D).

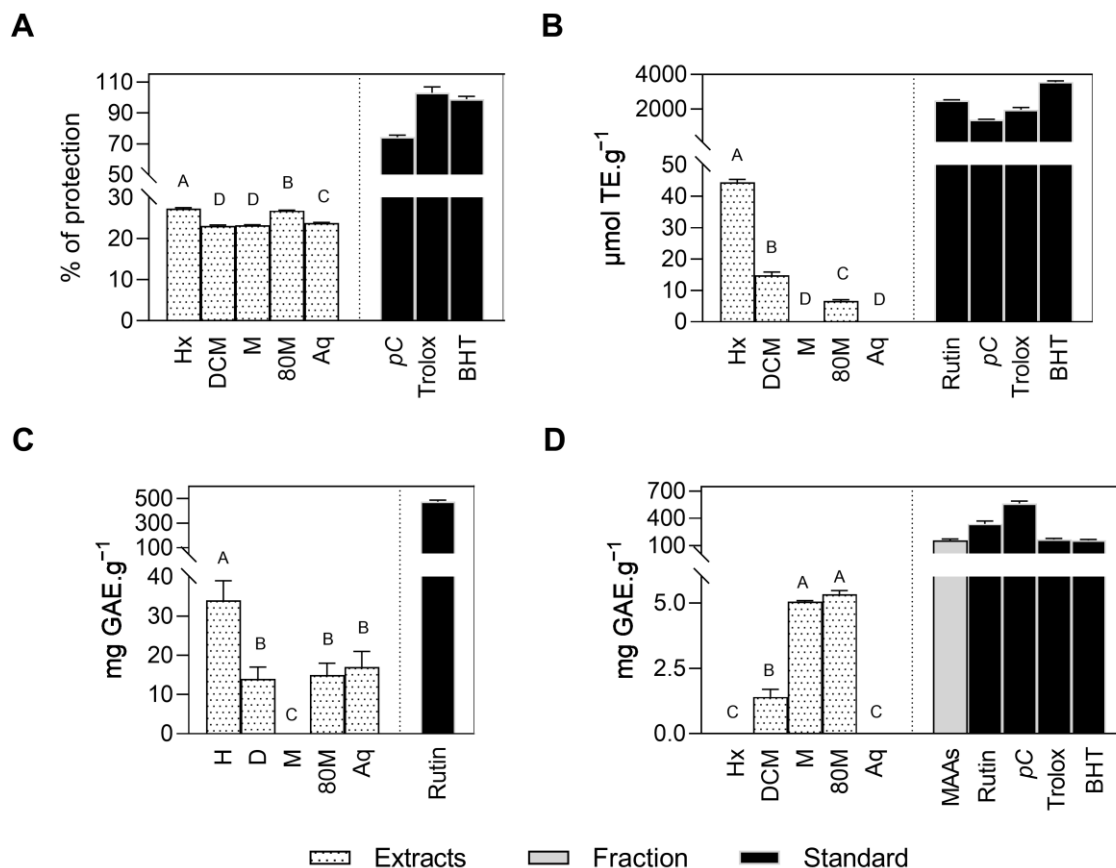


Figure 2. Antioxidant activities of crudes extracts that were obtained with increasing polarity solvents and MAAs (porphyra-334 + shinorine, 70% and 30%, respectively) from *Gracilaria domingensis* and standard references for lipid peroxidation inhibition (A), FRAP (B), ferrous chelating (C), and Folin–Ciocalteu (D) assays. The values (means $\pm$ SD; $n = 3$) were expressed in $\text{mg GAE} (\text{Gallic Acid Equivalent}) \cdot \text{g}^{-1}$ of dry extract or $\mu\text{mol TE} (\text{Trolox Equivalent}) \cdot \text{g}^{-1}$ of dry extract or % of protection. Hx—hexane, DCM—dichloromethane, M—methanol, 80M—80% methanol, A—aqueous, MAAs—mycosporine-like amino acids, BHT—butylated hydroxytoluene, and pC—p-coumaric acid.

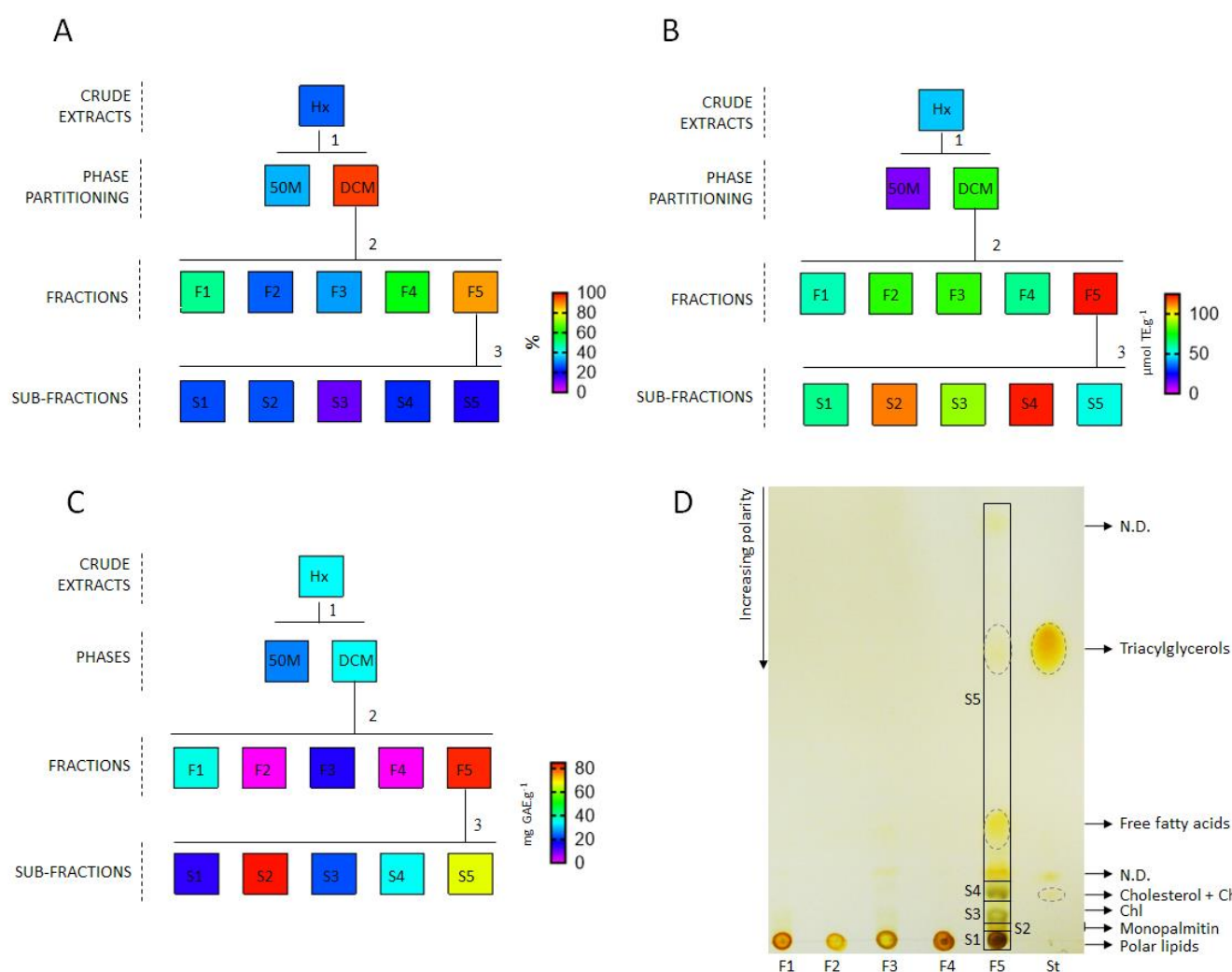


Figure 3. Variation of the antioxidant activities through color scale using the scheme of the fractionation of hexane crude extract that was guided by lipid peroxidation inhibition (**A**), FRAP (**B**), and ferrous chelating capacity (**C**). Thin-layer chromatography (TLC) (**D**) of the main lipid classes of the fractions F1 to F5, and indications of the subfractions S1 to S5 that were obtained from the fraction F5. Hx = Hexane, DCM = Dichloromethane, M = Methanol, 80M = 80% Methanol, and 50M = 50% Methanol. 1. Partition, 2. Silica column chromatography, and 3. Preparative thin layer chromatography. St = standard containing cholesterol and triacylglycerol. Chl = chlorophyll *a* and its derivatives. N.D. = not determined.

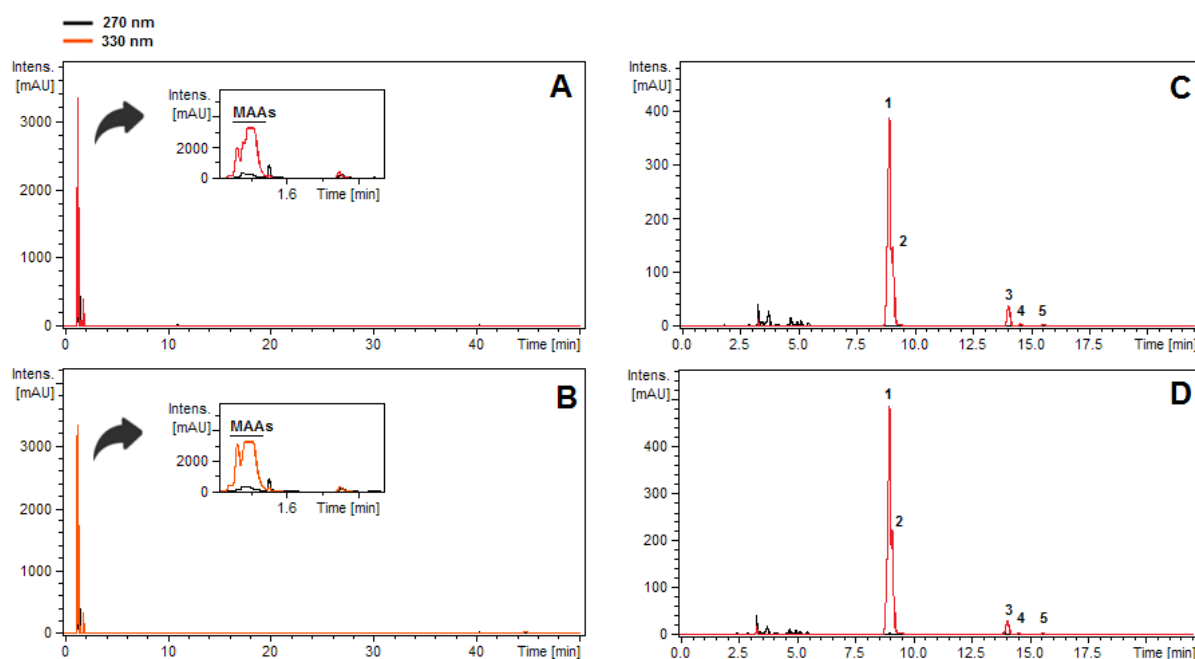


Figure 4. Global analysis of metabolites (A,B), respectively, methanol and 80% methanol extracts and specific analysis of MAAs (mycosporine-like amino acids) (C,D), respectively, methanol and 80% methanol extracts from polar extracts of *Gracilaria domingensis*. 1. Porphyra-334, 2. Shinorine, 3. Palythine, 4. Palythanol, and 5. Asterina-330.

4. Discussion

Hexane (Hx) and methanolic (M and 80M) extracts of *Gracilaria domingensis* presented the higher antioxidant activities, enabling them for further fractionation and compound identification. The synergetic effects between compounds might be responsible for the antioxidant activities that were observed for a fraction of Hx extract against lipid peroxidation inhibition. MAAs were the main components of the polar extracts. An enriched fraction with porphyra-334 and shinorine presented a Folin–Ciocalteu assay value that was 30 times greater than those of the original extracts.

Despite the low yield, the Hx extract was the most active extract, with significant activities in the three assays (FRAP, metal chelating, and lipid peroxidation inhibition). Similar results for non-polar extracts of *Gracilaria gracilis* were obtained previously with the FRAP assay [19]. The fractionation procedure from the Hx extract allowed to observe a loss in the lipid peroxidation inhibition (Figure 3A), suggesting synergistic interaction effects between the compounds. Synergistic interactions between antioxidant components are common in complex mixtures such as crude extracts [10]. Algal extracts have shown synergistic antioxidant activities when they are supplemented with isolated compounds [37] and other crude extracts [38].

In contrast to lipid peroxidation inhibition, the fractionation procedure led to the subfractions with activities that were similar to the original fraction in the FRAP and ferrous chelating assays. The main compounds from fraction F5 and its subfractions are primary metabolites with several reports of antioxidant activities. For example, chlorophyll *a* and its derivatives have iron-chelating ability [39], and monoacylglycerols are described with reducing activities in the FRAP assay [40].

In contrast to the Hx extract, the Aq extract showed the highest yield, but only weak antioxidant activities in ferrous chelating (Figure 2C) and inhibition of lipid peroxidation (Figure 2A) assays. Our previous study showed sulfated polysaccharides as the main component of this Aq extract [31]. The antioxidant properties, including lipid peroxidation inhibition and ferrous chelating activity, have already been reported for sulfated polysaccharides from *Gracilaria* [41,42].

The M and 80M extracts were the most active extracts in the FC assay. The activities for these extracts ($\sim 5 \text{ mg GAE} \cdot \text{g}^{-1}$) were higher or similar to those that were seen in another study with *Gracilaria*, such as *G. birdiae* ($\sim 1 \text{ mg GAE} \cdot \text{g}^{-1}$) [43], *G. cornea* ($\sim 1 \text{ mg GAE} \cdot \text{g}^{-1}$) [43], *Gracilaria corticata* ($\sim 2 \text{ mg GAE} \cdot \text{g}^{-1}$) [44], *G. edulis* ($\sim 4 \text{ mg GAE} \cdot \text{g}^{-1}$) [45], *G. gracilis* (3 to 6 $\text{mg GAE} \cdot \text{g}^{-1}$) [46], and *Gracilaria manilaensis* (0 to 6 $\text{mg GAE} \cdot \text{g}^{-1}$) [47]. However, surprisingly, no phenolic compounds were detected in the extracts, suggesting that the concentration of these compounds is lower than the minimum detectable amount, or they are absent in *G. domingensis*. Despite its popularity, the FC method is nonspecific since reducing non-phenolic compounds can also react with the FC reagent [6,7]. Thus, our results lead us to think that the activities using the FC assay can have a strong influence of other non-phenolic components.

Among the non-phenolic components, the M and 80M extracts have a high content of MAAs. The reducing power of MAAs was suggested by Coba et al. [48]. These authors demonstrated a powerful reducing capability of MAAs in alkaline conditions with ABTS^{•+} assay. More recently, a comparative study of the antioxidant capacity of isolated MAAs using several assays has demonstrated that these compounds have strong antioxidant activity in the FC assay in comparison to a reference antioxidant (Trolox) [33]. This is not surprising since the FC assay is performed in a high pH of ~ 10 , an ideal condition for the higher reducing potential of MAAs. Our results show that the reducing activities of a fraction of MAA were notably higher than those of the crude extracts (Figure 2D). Thus, within the studied samples, MAAs contributed to the antioxidant activities that were observed in the FC assay.

5. Conclusions

Chemical investigation of the antioxidant potential of *Gracilaria* species is rarely described in the literature. The studies that carried out chemical investigation tried to establish a relationship between antioxidant capacity and phenolic compounds through searches of these components in the extracts [11,43] or the use the Folin–Ciocalteu assay for quantification of the total phenolic content [44–47]. In the present study, we use the antioxidant activity-guided fractionation that provided us with a better understanding of the antioxidant properties of the crude extracts. The primary components were found as the most active antioxidants from non-polar extracts by the FRAP, metal chelating, and lipid peroxidation inhibition assays. On the other hand, MAAs seem to be responsible for the antioxidant potential by the FC assay. The interference of non-phenolic compounds on the FC assay should not be ignored in algal samples that are rich in MAAs, especially red algae, or other reducing components (e.g., ascorbic acid). A critical appraisal of the use of the FC assay for the total phenolic compounds determination is suggested for algae.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/phycolgy2030018/s1>, Figure S1: GC-MS chromatograms for the methanol (A) and 80% methanol (B) extracts from *Gracilaria domingensis*, showing the abundant peaks. For peak annotation, see Table S1; Table S1: Relative abundance (%) of the compounds identified in the methanol (M) and 80% methanol (80M) extracts from *Gracilaria domingensis*. Values comparable only within a same chemical class. RT—retention time, (–)—absence, and tr—trace.

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