



Antioxidant Activities of Phycocyanin: A Bioactive Compound from *Spirulina platensis*

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Authors' contributions

This work was carried out in collaboration among all authors. Author GAG designed the study, wrote the protocol and wrote the manuscript. Author SMES performed the laboratory research work, wrote the literature survey and managed the analyses of the study. Author MSH performed the statistical analysis, managed the literature searches and wrote the first draft of the manuscript. All authors read and approved the final manuscript.

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ABSTRACT

Background and Objectives: The cyanobacterium *Spirulina* also called blue green algae is a class of gram negative bacteria which possesses wide range of bioactive colored components as Phycocyanin, carotenoids and chlorophyll. *Spirulina* is one of the microalgae containing nutrients that have been used as a functional food in addition to therapeutic and pharmaceutical applications. This study aimed to evaluate the biochemical composition of *Spirulina platensis* biomass and its ethanolic and aqueous extracts, as well as, evaluate the antioxidant activities of the biomass, ethanolic, aqueous and the purified Phycocyanin.

Materials and Methods: The chemical compositions of *Spirulina platensis* were determined, as well as the antioxidant activity of extracts, Phycocyanin, Phycocyanopeptide and Phycocyanobilin using (DPPH) radical-scavenging activity.

Results: Results show that biomass has higher total proteins (49.72±0.508%), total carbohydrates (10.3±0.330%), moisture content (7.5±0.685%), lipids (7.2±0.105%) and Minerals (6.9±0.130%). In

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biomass, the total phenols ($51.20 \pm 0.25 \mu\text{g/mL}$) and flavonoids ($97.73 \pm 1.858 \mu\text{g/mL}$) were high compared to the ethanolic (49.48 ± 0.130 and $69.07 \pm 1.814 \mu\text{g/mL}$) and aqueous (15.27 ± 0.639 and $4.67 \pm 0.611 \mu\text{g/mL}$) extracts respectively. In the phenolic compounds, pyrogallol was identified as the major compound in biomass and aqueous extract (63.85 and 12.33%) respectively, E-vanillic acid in ethanolic (18.20 %), whereas, hesperidin (3.517 and 1.639%) were major flavonoids found in aqueous and ethanolic extracts respectively. The DPPH scavenging activity was found higher in ethanolic extract compared to aqueous while in bioactive, the order of antioxidant activity was Phycocyanin > Phycocyanobilin > Phycocyanopeptide.

Conclusions: The study data regarding to *Spirulina* nutritional value, makes *Spirulina* an excellent choice when formulating diets and combating malnutrition. Furthermore, it is a strong antioxidant and could be used as alternative treatments as anticancer, antidiabetic, and anti-inflammatory agent.

Keywords: *Spirulina platensis*; antioxidants; chemical composition; phycococynin.

1. INTRODUCTION

Spirulina, a microscopic, filamentous, dried biomass of *Arthrospira plantensis*, is an oxygenic photosynthetic cyanobacterium found in fresh and marine waters worldwide [1]. It is consumed by both humans and animals as a food supplement [2,3]. *Spirulina* has been labeled as a superfood because of its richness in carbohydrates, proteins, sterols, polyunsaturated fatty acids and minerals such as selenium, calcium, chromium, iron, manganese, zinc and magnesium [4]. It is also one of the natural source of provitamin A and vitamin B₁₂, E, C, phenolic acids, linoleic acid, and xanthophylls phytopigments [5]. Multiple reported activities of *spirulina* such as anti-inflammatory, anticancer, cholesterol-lowering and antiviral, are attracting the attention of scientists for using *spirulina* as a nutraceutical and pharmaceutical sources [6-8]. Phycocyanin (PC), an extracted pigment from *Spirulina platensis*, has been widely used as a natural blue colour for multi-purposes [9]. In addition, PC being natural antioxidant is useful for treatment of many diseases such as Alzheimer's, Parkinsons and Huntingtons diseases [10]. Phycocyanobilin is one of the two compounds extracted from PC, it has free radical scavenger and thus protecting the living cells from oxidative stress [11,12]. The current study aimed to evaluate the biochemical components of *Spirulina* alga biomass and determine the antioxidant activities of its ethanolic and aqueous extract, as well as, the antioxidant activity of Phycocyanin, the *Spirulina* bioactive compound.

2. MATERIALS AND METHODS

2.1 Microalgae Preparation

Spirulina (Arthrospira) platensis strain was cultured at National Research Centre (NRC),

Unit algae technology, Giza, Egypt. The cultures were allowed to grow for eighteen days in shaking incubator with a light intensity of $50 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ and 12:12 light/dark regime, at $30 \pm 1^\circ\text{C}$, pH 9.0 [13].

2.2 Drying of Fresh Biomass of *Spirulina platensis*

The biomass of *spirulina platensis* was collected by filtration through a nylon cloth, then the adhered salts were removed by washing with distilled water and stored at -20°C [14]. The harvested fresh biomass was subjected to different methods of drying including drum drying, spray drying, sun drying and oven drying [15].

2.3 Preparation of Ethanolic and Aqueous Extracts of *Spirulina platensis*

2.3.1 Ethanolic extract

Ten grams of *Spirulina platensis* dried powder were extracted by adding 100 ml of 80% ethanol and kept overnight on a rotary shaker at room temperature. The mixture was filtered and centrifuged at 6,000 rpm for 15 min., the supernatant collected and stored for further investigations [14].

2.3.2 Aqueous extract

Ten grams of dried *Spirulina platensis* powder (by spray drying which give the best method of drying) were extracted in 100 mL distilled water kept on a rotary shaker for 2 h and then kept at 4°C for 24 h according to the method of Anwer et al. [14].

2.4 Chemical Composition of *Spirulina platensis*

2.4.1 Determination of total soluble proteins

The concentration of the total soluble protein was determined in the spirulina biomass according to Bradford [16] using bovine serum albumin as a standard.

2.4.2 Determination of free amino acids

The free amino acids were extracted by mixing 5 g of *Spirulina platensis* powder with 50 ml ethanol 80% at 70°C and refluxing for 1 h. This method was repeated three times, the extracts were collected, filtered and the supernatant was evaporated at 55°C. The dried film was dissolved in 10 ml of isopropanol (10%) and the total volume was completed to 50 ml. The isopropyl extracts were kept at -20°C until further analysis [17].

2.4.3 Determination of total carbohydrates

Total carbohydrates were determined according to the method of A.O.A.C [18]. Approximately 0.1 g of sample dried by air and hydrolyzed by 1 N HCl in boiling water bath for 6 h. The resultant solution was filtered, neutralized and the final volume was made up to 100 ml with distilled water. Total reducing sugars was measured calorimetrically in 1ml of sample by alkaline potassium ferricyanide reagent at 420 nm using UV-VIS spectrophotometer.

2.4.4 Determination of moisture content

Moisture content was measured by drying approximately 2 g of sample in a ventilated oven at 70°C for 24 h, then at 105°C for 3 h. Samples cooled over CaCl₂ in glass desiccator, the weighed samples were heated many times at the same temperature until constant weight was achieved and moisture content was calculated according to A.O.A.C [18].

2.5 Determination of Pigments

Chlorophyll-a and carotenoids were extracted and estimated according to A.O. A.C [18]. 0.5 g samples (fresh alga biomass and both extracts) were homogenized in a mortar with 85% acetone in the presence of washed dried sand and a little amount of CaCO₃ (0.1 g) in order to neutralize organic acids in the homogenate of the fresh alga biomass. The homogenate was then filtered through sintered glass funnel. The residue

washed several times with acetone until the filtrate became colorless. The optical density of this extract was determined using a spectrophotometer (UV-Vis spectrophotometer UV 9100 B, LabTech) at 662, 644 nm for (Chl. a, and Chl. b, respectively) and at 440.5 nm for carotenoids. The extraction and purification of PC from *Spirulina platensis* was followed by earlier method mentioned in our previous study [19].

2.6 Determination of Total Flavonoids

Total flavonoid content of spirulina biomass and both extracts were determined by the aluminum chloride colorimetric assay as described by Marinova et al. [20]. One ml of each sample was added to 10 ml volumetric flask containing 4 ml of distilled water. 0.3 ml 5% NaNO₂ was added then 0.3 ml 10% AlCl₃ after 5 min. At the 6th min, 2 ml of 1M NaOH was added and total volume was completed to 10 ml with dH₂O. The solution was well mixed and the absorbance was reading at 510 nm. The concentration of total flavonoids was calculated using quercetin standard curve.

2.7 Determination of Total Phenolic Compounds

Total soluble phenolics of spirulina biomass and both extracts were determined using the method described by Shahidi and Naczki [21]. A known weight of each sample was immersed in 10 ml ethanol 80% and kept in a dark bottle for 24 h at 0°C. The samples were re-extracted 3 times then the clarified extract was completed to 50 ml with ethanol 80%. Then 1 ml of the extract was mixed with 0.5 ml Folin-Denis reagent in a test tube and thoroughly shaken. After 3 min, 1 ml of saturated Na₂CO₃ (33 g anhydrous salt were dissolved in 100 ml dH₂O at 70-80°C, then cooled and kept overnight at room temperature then filtered. The absorbance was measured at 725 nm using spectrophotometer (UV-Vis spectrophotometer UV 9100 B, LabTech). The concentration of total soluble phenols was calculated using gallic acid standard curve. The total phenols concentration was expressed as µg equivalents of gallic acid per g DW of the sample.

2.8 Identification and Quantification of Phenolic Compounds of *Spirulina platensis* Biomass and Both Extracts by HPLC

The Phenolic compounds in *Spirulina* biomass and both extract were identified by HPLC [22].

5 g of sample were homogenized with methanol and centrifuged at 10000 rpm for 10 min. The supernatant was filtered through 0.2 µm Millipore filter membrane, 3 ml of filtrate was collected in a vial for HPLC injection. The stationary phase is octadecylsilyl (reversed phase) and the gradient separation was carried out with ethanol and acetonitrile as a mobile phase at flow rate of 1ml/min. The standard of phenolic compounds purchased from sigma were dissolved in a mobile phase and injected into HPLC. Retention time and peak area were used to calculate the concentration of phenolic compounds by using the data analysis of Hewlett Packard software.

2.9 Antioxidant Activity of Extracts, Phycocyanin, Phycocyanopeptide and Phycocyanobilin

The antioxidant activity of the ethanolic and aquatic extracts of *spirulina platensis* and its bioactive compounds phycocyanin, phycocyanopeptide and phycocyanobilin were determined based on 1,1-Diphenyl-2-Picrylhydrazyl (DPPH) radical-scavenging activity [23]. In the DPPH assay, 1 ml of sample was mixed with 0.5 ml of 50 µM DPPH in ethanol and kept in dark for 30 min. The mixture absorbance was measured at 517 nm., and vitamin C standard (5–30 µg/ml) was used as positive controls. The radical scavenging activity was

determined based on percentage of inhibition by using the following formula:

$$\% \text{ inhibition} = [(A \text{ control} - A \text{ sample}) / A \text{ control}] \times 100.$$

3. RESULTS AND DISCUSSION

3.1 Major Chemical Components of *Spirulina platensis* Biomass and Its Both Extracts

The chemical composition of *Spirulina platensis* biomass results show, *Spirulina platensis* have the following chemical composition: high protein content, 49.72 (% w/w), total lipids 7.2% (w/w), total carbohydrates 10.3% (w/w), moisture 7.5% (w/w), Ash 9.86%, crude fiber 8.52% and minerals 6.9 % (w/w) (Fig. 1). These results were supported by El-Baz et al. [24] who found the protein content of *Spirulina platensis* was (35–65%) while, Wong and Chan [25] reported *spirulina* has 10% (w/w) of minerals content, in addition, Aly and Gad [26] found the *Spirulina platensis* biomass contain fat and total carbohydrates contents were 8 and 10%, respectively. Different studies reported that the *Spirulina platensis* is a rich protein source, essential amino acids, polyunsaturated fatty acids γ-linolenic acid (GLA), sterols and some more vital elements like magnesium, calcium, zinc, iron, manganese, selenium and chromium [26,27].

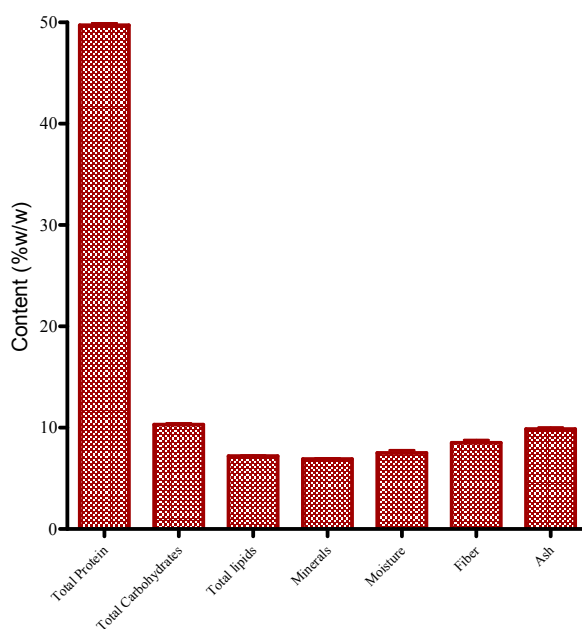


Fig. 1. Major chemical composition of *Spirulina platensis*
Data presented as the means of three replicates ± SD

The obtained data show the ethanolic extract of *Spirulina platensis* contained higher concentrations of carotenoids, chlorophyll-a, total phenols and flavonoids and lower concentration of PC, free amino acids, total proteins and carbohydrates (Table 1). On the other hand, the aqueous extract of *Spirulina platensis* contained higher contents of phycocyanin, free amino acids, total proteins and carbohydrates, while it contained lower concentration of carotenoids, chlorophyll-a, total phenols and flavonoids. These results show the main constituent of biomass alga and its water extract contain high percentage of total proteins (49.72 and 48.01% w/w), total carbohydrate (10.3 and 9.4%), total free amino acids (4.71 and 4.68%) respectively compared to ethanolic extract (Table 1).

Multiple studies show that the phytopigments such as carotenoids, chlorophyll-a and phycocyanin play an important role as antioxidant and anticancer agents [28,29]. The results indicated the *Spirulina platensis* have the highest content of carotenoids and chlorophyll-a (4.69 and 25.59 mg/g dw) followed by the ethanolic extract (4.25 and 25.43 mg/g dw). On the other hand, the concentration of PC in *Spirulina platensis* (0.335 mg/mL) was higher than aqueous extract (0.185 mg/mL) followed by ethanolic extract (0.007 mg/mL).

The total phenols in *Spirulina* alga (51.20 µg/mL) were higher than ethanolic extract (49.48 µg/mL), while water extract show the lowest total phenols content (15.26 µg/mL). *Spirulina* significantly contained the highest content of total flavonoids (97.73 µg/mL), followed by its ethanolic extract (69.07 µg/mL) and the lowest content was observed in its water extract (4.67 µg/mL).

3.2 Identification and Quantification of Phenolic Compounds of *Spirulina platensis* by HPLC

Spirulina platensis and its ethanolic and aqueous extracts were subjected to HPLC analysis to investigate the types of phenolic compounds responsible for possible antioxidant activities. In this investigation, thirty-three phenolic compounds were identified and quantified in *Spirulina platensis* and its extracts by comparing its HPLC chromatograms with the standard compounds based on the retention time. The identified phenols were pyrogallol, gallic, chlorogenic, caffeine, vanillic, p-Coumaric and other phenols. Flavonoids were naringin, hesperidin, rutin, quercetrin, naringenin,

catechin, hesperidin and other flavonoids (Tables 2 and 3). The results show low amounts of phenol and flavonoid compounds in water extract of alga compared with ethanolic extract. Pyrogallol was identified as the major phenol compound in *Spirulina platensis* (638.50 mg/100 g) and its water extract (12.33 mg/100 g), while, E-Vanillic was the major compounds in ethanolic extract (18.20 mg/100 g). Also, water extract contained high amount of Ellagic and not detected in ethanolic extract and the major flavonoid compound in *Spirulina platensis* biomass was Hesperidin (9.013 mg/100 g). Also, *Spirulina* exhibited highest amount of naringin (4.743 mg/100 g) followed by ethanolic extract (0.652 mg/100 g) and water extract (0.359 mg/100 g). The quercetin content was 0.922 mg/100 g in *Spirulina* biomass followed by alcoholic extract (0.278 mg/100 g), it was not detected in the water extract. Generally, HPLC analysis show the content of phenolic compounds of *Spirulina platensis* and its extracts have the following order, *Spirulina platensis* biomass > ethanolic extract > water extract.

These results were supported by the studies of Syarina et al. [30] on *S. platensis* aqueous extract. They recorded that cinnamic acid, naringenin and kaempferol have identified in *S. platensis* aqueous extract by LC-MS analysis. Also, El-Baky et al. [31] showed the presence of large number of phenolic acids and few numbers of flavonoids which includes gallic, cinnamic, chlorogenic, p-OH-benzoic, ferulic acids, vanillic, quimic and caffeic in *Spirulina platensis*.

Colla et al. [32] mentioned that phenolic compounds in *Spirulina* such as organic acids (chlorogenic, synaptic, quimic, salicylic, trans-cinnamic and caffeic) can act as antioxidants synergistically or individually. Also, Miranda et al. [33] identified the main phenolic compounds in *Spirulina*, which include trans-cinnamic, chlorogenic, salicylic, synaptic and caffeic acids. In addition, the antioxidant and pharmacological activities of phenols and flavonoids in *Spirulina* were reported by many studies, such as cinnamic acid and their derivatives inhibit diabetic complications and moderately induce corneal epithelial wound healing by inhibiting protein tyrosine phosphatases (PTP)-1β gene [34]. Also, Naringenin has inhibitory effect on the pro-inflammatory cytokine (IL-6, IL-1β, IL-8 and TNF-α) response stimulated by lipopolysaccharide in both whole blood and macrophages [35]. Kaempferol has been reported to possess antioxidant and anti-

Table 1. Major chemical components of *Spirulina alga* and its water and ethanolic extracts

	Phytopigments			Phenolic compounds ($\mu\text{g/mL}$)	
	chlorophyll-a (mg/g DW)	Carotenoids (mg/g DW)	Phycococynin (mg/ml)	Total phenols	Total flavonoid
<i>Spirulina platensis</i>	25.59 ^a ±0.025	4.69 ^a ±0.048	0.335 ^a ±0.00	51.20 ^a ±0.025	97.73 ^a ±1.858
Ethanolic Extract	0.652 ^c ±0.017	0.50 ^c ±0.007	0.185 ^b ±0.00	15.27 ^c ±0.639	4.67 ^c ±0.611
Aqueous Extract	25.43 ^b ±0.021	4.25 ^b ±0.104	0.007 ^c ±0.00	49.48 ^b ±0.130	69.07 ^b ±1.814

Each value represents the means \pm SD. Different letters refer to significant differences ($P \leq 0.05$)

Table 2. Types, retention time (min) and the content of phenols (mg/100 g) identified by HPLC analysis of *Spirulina platensis* biomass and its aqueous and ethanolic extracts

	Compound	<i>Spirulina platensis</i>		Aqueous extract		Ethanolic extract	
		(mg/100 g)	min	(mg/100 g)	min	(mg/100 g)	min
1	Pyrogallol	638.5	6.875	12.33	6.841	8.50	6.896
2	Gallic	4.64	7.011	1.63	7.003	0.36	7.065
3	4-Amino-benzoic	0.32	8.246	0.05	8.342	0.02	8.319
4	Protocatchuic	1.26	8.694	0.27	8.509	0.78	8.467
5	Catechol	7.62	8.636	2.55	8.656	0.49	8.612
6	P-OH- benzoic	1.90	9.136	0.13	9.048	0.18	9.027
7	Caffeine	0.75	9.762	0.15	9.655	0.21	9.653
8	Chlorogenic	3.06	9.859	0.52	9.840	0.94	9.807
9	Vanillic	2.62	9.875	1.31	9.880	0.57	9.907
10	Caffeic	5.97	10.324	0.28	10.246	0.43	10.102
11	P-Coumaric	0.28	10.900	0.11	10.349	0.33	10.322
12	Ferulic	0.39	11.657	0.11	11.586	0.21	11.601
13	Iso-Ferulic	0.22	11.938	0.04	11.820	0.09	11.839
14	E-Vanillic	31.53	12.107	4.47	12.209	18.20	12.196
15	Alpha-Coumaric	1.26	12.524	0.18	12.280	0.24	12.292
16	Benzoic	6.82	13.268	0.34	13.241	1.06	13.283
17	Ellagic	19.83	13.488	2.59	13.393	ND	ND
18	3,4,5-methoxy-cinnamic	1.40	14.183	0.53	14.181	2.91	14.210
19	Rosmarinic	0.18	14.282	0.04	14.380	0.03	14.197
20	Coumarin	1.08	14.552	0.11	14.385	0.27	14.373
21	Cinnamic	0.12	15.285	0.02	15.262	0.04	15.238
22	salicylic	3.89	16.258	0.82	16.480	1.01	16.370

Table 3. Quantity of flavonoids (mg/100 g) identified by HPLC analysis of *Spirulina platensis* biomass and its extracts

	Compound	<i>Spirulina platensis</i>		Aqueous extract		Ethanolic extract	
		(mg/100 g)	min	(mg/100 g)	min	(mg/100 g)	min
1	Catechein	3.77	8.246	0.86	8.425	0.47	8.469
2	Epicatechein	1.68	10.324	0.09	9.597	0.46	9.663
3	Naringin	4.74	12.340	0.36	12.087	0.65	12.239
4	Hespirdin	9.01	12.438	3.52	12.514	1.64	12.538
5	Rutin	1.23	12.575	0.12	12.780	1.15	12.806
6	Quercetrin	2.26	13.482	0.09	13.680	0.65	13.709
7	Quercetin	0.92	14.887	ND	ND	0.28	14.655
8	Naringenin	0.89	15.034	0.14	15.192	0.15	15.196
9	Hespirtin	5.16	15.324	0.34	15.634	0.41	15.431
10	Kampferol	1.20	16.243	0.04	15.927	0.26	16.224
11	Apigenin	0.54	16.600	0.01	16.641	0.02	16.683

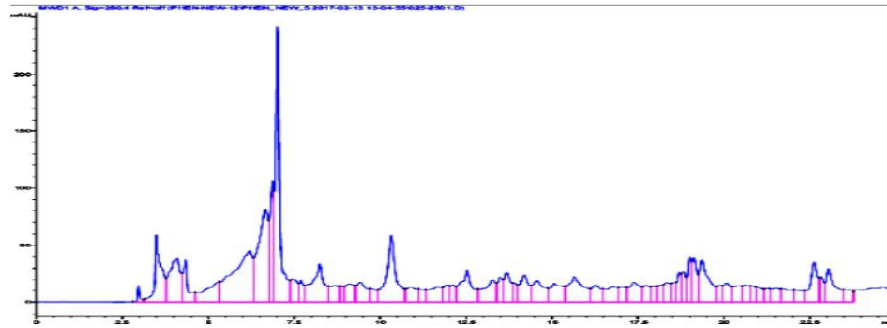


Fig. 2. HPLC-UV chromatogram of phenolic compounds of *Spirulina platensis* alga

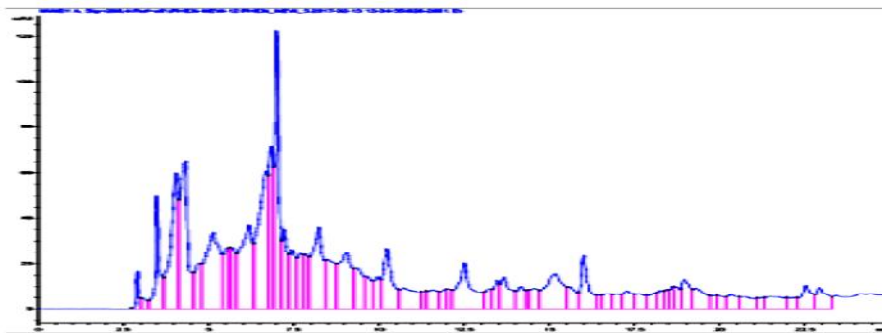


Fig. 3. HPLC -UV chromatogram of phenolic compounds of aqueous extract of *Spirulina platensis*

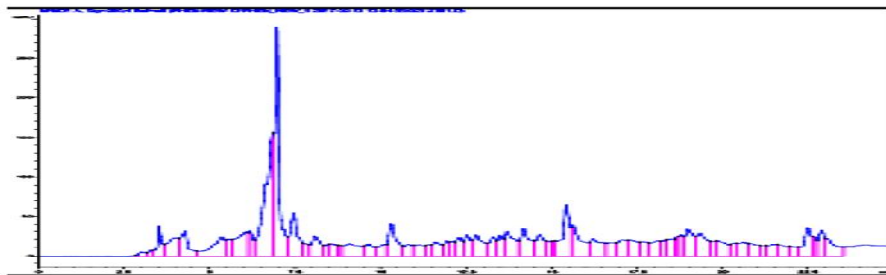


Fig. 4. HPLC -UV chromatogram of phenolic compounds of ethanolic extract of *Spirulina platensis*

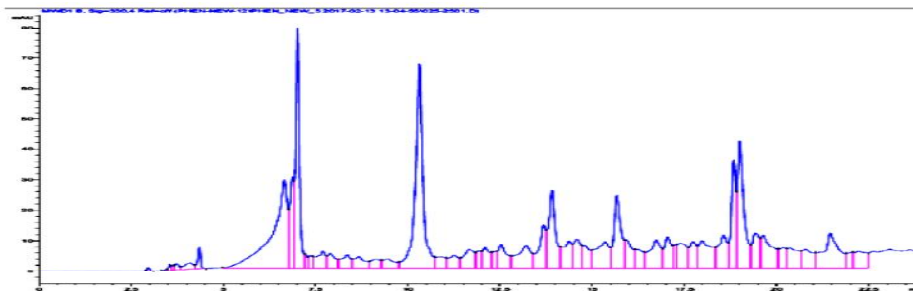


Fig. 5. HPLC -UV chromatogram of flavonoids of *Spirulina platensis*

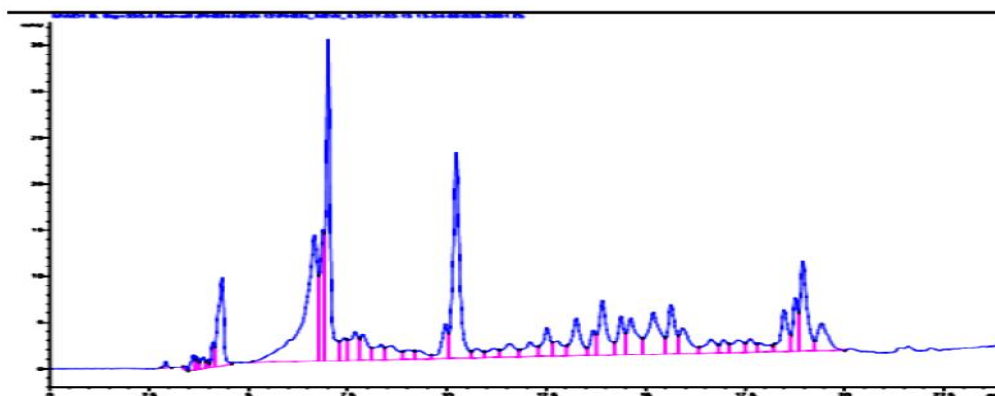


Fig. 6. HPLC -UV chromatogram of flavonoids of aqueous extract of *Spirulina platensis*

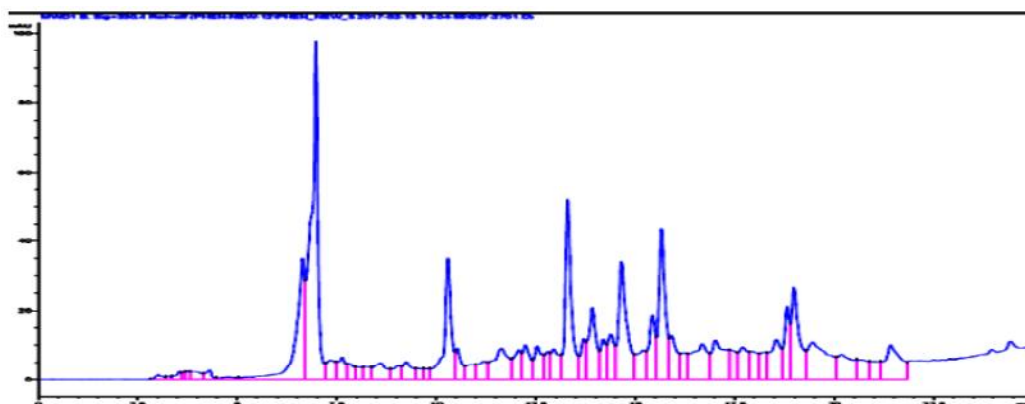


Fig. 7. HPLC -UV chromatogram of flavonoids of ethanolic extract of *Spirulina platensis*

inflammatory activity [36]. Chlorogenic acid (CGA) has many biological activities as antibacterial, antioxidant, anticarcinogenic, hypoglycemic and hypolipidemic [37-39].

3.3 Antioxidant Activities

3.3.1 DPPH assay of *Spirulina platensis* ethanolic and aqueous extracts

The free radical scavenging ability of aqueous and ethanolic of *Spirulina platensis* biomass was assessed on the basis of the radical scavenging effect on the 1,1-diphenyl-2-picryl-hydrazil (DPPH) free radical. The antioxidant activity increased with increasing the concentration of sample solution [40]. The highest antioxidant activity in term of % inhibition was detected in ethanolic extract followed by vitamin C and water extract. The obtained results show that the ethanolic extract of *Spirulina platensis* has higher scavenging activity (96.33%) than water extract (73.58%) at 30 µg/ml, while the activity in biomass was (90.60%) (Fig. 8).

Generally, it could be concluded that ethanolic extract possessed the highest antioxidant activity comparing with aqueous extract of *Spirulina platensis* due to the presence of high content of antioxidant phenols or flavonoids in the ethanolic extract. The ethanolic extract of *Spirulina maxima* possessed a great ability of scavenging DPPH radical activity compared to positive synthetic antioxidants butylated hydroxytoluene (BHT) and butylated hydroxyanisole (BHA) depending on phenolic contents in extract [41,42].

The carotenoids in *Spirulina platensis* act as antioxidant and it has the ability to quench singlet oxygen and scavenge of the free radicals, also, α -tocopherol act as antioxidant compound in brown algae [43].

3.3.2 DPPH assay of phycocyanin and its two bioactive components

Both of the phycocyanopeptide and phycocyanobilin had high radical scavenging

activity in DPPH assay. The antioxidant activity in term of % inhibition was detected in phycocyanopeptide and phycocyanobilin were (93.59%) and (95.48%) respectively compared to phycocyanin (97.32%). The antioxidant capacity of phycocyanin and its bioactive components (phycocyanopeptide and phycocyanobilin) was showed the following descending order, phycocyanopeptide < phycocyanobilin < phycocyanin and the

increased activity as their concentrations were increased. Based on the DPPH assay, the antioxidant activity of phycocyanin, phycocyanobilin and phycocyanopeptide at 5 µg/mL was about 46.71, 42.03 and 40.00%, respectively (Fig. 9). These results are in agreement with Estrada et al. [44] who observed that an increase in the amount of phycocyanin caused an increase in the antioxidant activity.

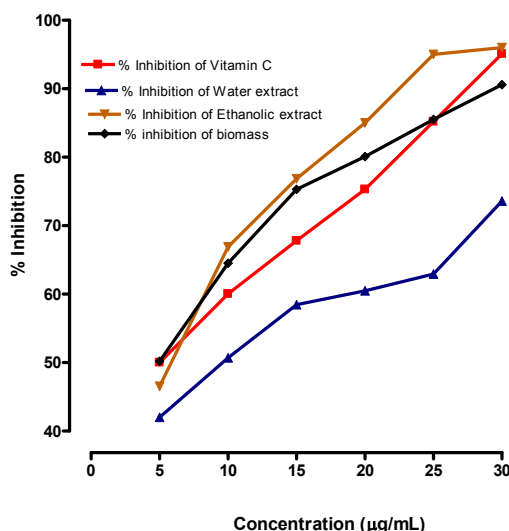


Fig. 8. DPPH free radical scavenging activity of biomass, aqueous and ethanolic extracts of *Spirulina platensis* compared with vitamin C

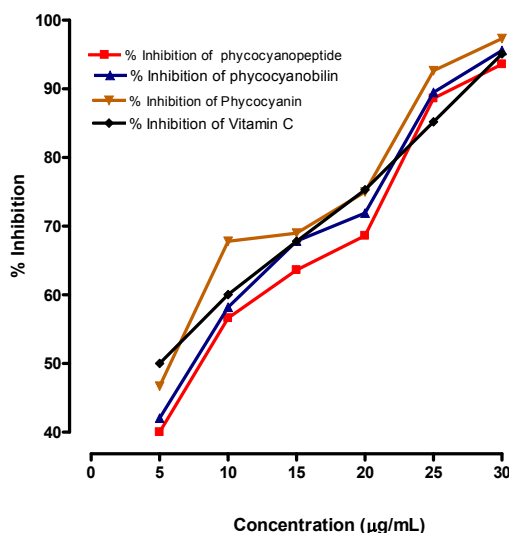


Fig. 9. Antioxidant activity of phycocyanin and its bioactive components (phycocyanopeptide and phycocyanobilin) as function of concentration

Phycocyanin is used as a food coloring and has a great antioxidant [45,46]. It is the major water-soluble antioxidant constituent in *Spirulina*; its activity was about 20 times more efficient than vitamin C [47]. The covalently-linked tetrapyrrole chromatophore phycocyanobilin is suggested to be involved in the scavenging activity of phycocyanin [48]. Also, Cherdkiatikul & Suwanwong [49] mentioned that the two components of phycocyanin (phycocyanopeptide and phycocyanobilin) were possessed strong antioxidant activity. For instance, selenium-containing phycocyanin from *Spirulina* has been shown to have strong alkoxyl, superoxide and hydrogen peroxide radical scavenging activities [50]. In addition to, the selenium possessed antioxidant effect and plays an important role in protecting against oxidative stress [51]. Also, Cherdkiatikul and Suwanwong [49] found that the apophycocyanin possessed strong antioxidant. The strong antioxidant activity is related to selenium content. Selenium binds to phycobiliproteins, increasing their antioxidant properties [52]. The antioxidant activity of free phycocyanobilin was higher than phycocyanobilin bound in phycocyanin (PC) when denaturated or trypsin digestion of PC [53].

4. CONCLUSION

Total biochemical were found higher in *Spirulina* biomass whereas its ethanolic extract contained the highest total phenols and flavonoids while total proteins and phycocyanins were found higher in the aqueous extract of *Spirulina*. The antioxidant activity of phycocyanin was found highest followed by ethanolic extract whereas aqueous extract was found least potent.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

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