



**International Journal of Biology, Pharmacy  
and Allied Sciences (IJBPAS)**

*'A Bridge Between Laboratory and Reader'*

[www.ijbpas.com](http://www.ijbpas.com)

---

---

## MICROSCOPIC EVALUATION AND PHYSIOCHEMICAL ANALYSIS OF ALGAE *ARTHROSPIRA PLATENSIS*

RANJITHA S<sup>2</sup>, THIRMUAL M<sup>1\*</sup>, MONISHA ANANDAN<sup>2</sup>, DIVYA R<sup>2</sup>

1: Department of Pharmacognosy, SRM College of Pharmacy, SRM Institute of Science and Technology, Kattankulathur, India –603203

2: Faculty of pharmacy, Dr.M.G.R Educational and Research Institute, Velappanchavadi, Chennai

\*Corresponding Author: Dr. Thirumal M: E Mail: [thirumal\\_sel@yahoo.co.in](mailto:thirumal_sel@yahoo.co.in)

Received 24<sup>th</sup> Sept. 2023; Revised 25<sup>th</sup> Oct. 2023; Accepted 9<sup>th</sup> Jan. 2023; Available online 1<sup>st</sup> Nov. 2024

<https://doi.org/10.31032/IJBPAS/2024/13.11.8235>

### ABSTRACT

**Objective:** To evaluate the microscopic analysis and physicochemical examination of algae, *Arthrospira platensis*

**Methods:** Macroscopical and microscopical examinations were done on dried samples. Plant material underwent an initial phytochemical analysis. There were additionally standardisations using additional WHO-recommended metrics.

**Results:** Powder microscopy studies on algae of *Arthrospira platensis* species has been documented as per standard procedures. The proportion of foreign substances, the swelling index, the ash values, the loss on drying, the extractive values, and other physicochemical parameters were also determined. Initial phytochemical analysis found alkaloid, steroids, saponins, flavonoids, phenolic compounds, glycosides, and tannin in the sample.

**Conclusions:** *Arthrospira platensis* microscopic and physicochemical examination is used for establishing standards for sample verification, quality, and the utmost.

**Keywords:** *Arthrospira platensis*, powder microscopy, physicochemical parameters, phytochemical analysis, algae

## INTRODUCTION

Blue-green algae (cyanobacteria) of the genus *Arthrospira*, including *A. platensis*, *A. maxima*, and *A. fusiformis*, which are naturally floating in seawater, are used to make the well-known dietary supplement spirulina. In reservoirs with alkaline water, spirulina typically occurs and is commercially produced in subtropical and tropical parts of North America, Asia, and Central Africa. Spirulina is well-liked by consumers and used as a dietary supplement because of its high nutrient contents and many therapeutic properties [1, 2]. The photosynthetic cyanobacterium known as *Arthrospira* (*Spirulina* sp.) has lately acquired popularity as a dietary ingredient and nutritional supplement due to its high protein content of over 60% and other purported health advantages [3]. Many ailments, including diabetes, obesity, arthritis, anemia, cardiovascular conditions, allergies, tumors, and cancer can reportedly be treated with *Arthrospira*. Moreover, it has antioxidant, anti-inflammatory, and anti-virus properties [4]. Spirulina (*Arthrospira*) is one of the best sources of protein. It contains 60–70% protein [5]. Blue-green microalgae *Spirulina platensis*, which has been widely used as food and feed additives in agriculture, the food industry, pharmaceuticals, perfume manufacturing, medicine, and science, are one of the trends in biotechnology [6]. Research has shown

that consuming spirulina for four weeks lowers human serum cholesterol levels by 4.5% [7] and body weight is greatly reduced. Tumor necrosis factor is activated in macrophages by spirulina extract, suggesting a potential mechanism for tumor elimination [8]. A study indicated that spirulina could treat allergic rhinitis [9]. A variety of allergic reactions, including asthma, atopic dermatitis, and allergic rhinitis, are protected against by a number of chemicals made from marine organisms, including cyanobacteria. [10] Because antibacterial and antimycotic compounds have been identified from *Spirulina platensis* through medicinal research, spirulina has potent antiviral activity. These findings demonstrate excellent in vitro HIV-1 restraint in human lymphocyte lines and human monocytes [11]. Spirulina has been researched as a stimulant of animal cell growth and as an alginate-based residual water treatment [12]. Spirulina supplementation has also been shown to reduce the incidence of anemia that occurs during pregnancy and breastfeeding. In their research, Kapoor and Mehta [13]. In the presence of acidic polysaccharides from *A. platensis*, macrophages produced a significant amount of tumor necrosis factor (TNF) [14]. Infected Balb/C mice were used to assess *S. platensis* extract's immunostimulatory effects [15]. Thus, in

this study, we describe an effort to standardise *A. platensis* through microscopic assessment and physiochemical investigation.

## MATERIALS AND METHOD

### Chemicals

All of the chemicals employed in the study, including phloroglucinol, glycerin, hydrochloric acid, chloral hydrate, potassium hydroxide, were of analytical grade.

### Plant material

*Arthrospira platensis*'s fresh algae were procured from the area of Auroville Pondicherry, India. Prof. P. Jayaraman, Ph.D., Director of the Plant Anatomy and Research Centre (PARC) in Tambaram, confirmed and authenticated the algae.

### Macroscopic and microscopic analysis

The plant's macroscopy and microscopy were investigated using the Brain et al. approach [16]. For powder microscopic, A small dried piece of the sample was soaked for 4 h in distilled water. A drop of 50% glycerol was used to mount the algal sample suspended in water on a tiny slide. A Zeiss Axiolab-5 Trinocular microscope and a Zeiss Axiocam-208 colour digital camera were used to observe the characters in bright field and phase contrast I and II. Diagnostic characteristics were photographed and recorded as photomicrographs. A scale bar served as the indicator for the magnifications [17, 18].

### Physiochemical analysis

The official method suggested and in compliance with the WHO recommendations on quality control methods for medicinal plant material were used to assess the physiochemical Ash and extractive values [19-21].

### Preliminary phytochemical screening

The usual approach was used to conduct preliminary screening. Kokate [22].

## RESULT AND DISCUSSION:

### Macroscopic characteristics

The algae sample is dark green with a characteristic odor and saline mucilaginous taste. It forms the gel with water.

### Microscopical characteristics

#### Powder microscopy

The fine powder was dyed with phloroglucinol and concentrated HCl, as well as mounted in glycerin. When examined under a microscope, it revealed helical filament bodies and solitary cells with a cylindrical form (**Figure 1, 2**).

#### Macroscopical evaluation:

*Spirulina platensis*, an air-dried coarse powdered algae, had its ash values (total ash, water-soluble ash, acid insoluble ash, and sulphated ash) and extractive values (water soluble extractive value, alcohol soluble (extractive values), as well as moisture content) assessed (loss on drying). All of the values have been determined and listed in **Table 1**.

### Preliminary phytochemical Evaluation:

Initial phytochemical analysis found alkaloid, steroids, saponins, flavonoids, phenolic substances, glycosides, and tannin in the sample (Table 2).

### Physiochemical parameters

The physiochemical parameters, including ash values, drying losses, swelling index, and the percentage of foreign matter, were determined and are displayed in Table 3.



Figure: 1 *Arthrospira paltensis*



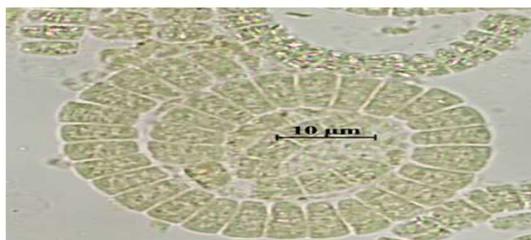
Single cells and filamentous body under a bright field light



Single cells and filamentous body under a phase contrast I



Single cells and filamentous body under a phase contrast II

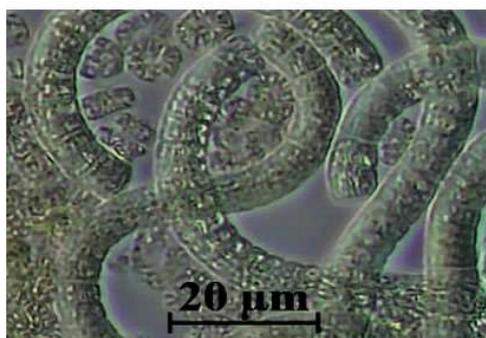


Single cells and filamentous body under a bright field light in 100X



Single cells and filamentous body under a phase contrast II enlarged view

Single cells under a phase contrast II enlarged view



Filamentous body under a phase contrast II enlarged view

Figure 2: Powder microscopy of *Arthrospira platensis*

Table 1: Extractive value of Crude drugs

S. No.	solvent	Extractive values (%w/w)
1.	Petroleum ether	2.23
2.	Chloroform	2.51
3.	Ethyl acetate	2.15
4.	Ethanol	15.4
5.	Water	5.4

Table 2: Preliminary phytochemical of crude drug

S. No.	Constituents	Report
1.	Alkaloids	+
2.	Proteins	+
3.	Carbohydrates and amino acids	+
4.	Steroids	+
5.	Saponins	+
6.	Flavonoids	+
7.	Phenolic compounds	+
8.	Glycosides	+
9.	Tannins	+

Table 3: Physicochemical Parameters

S. No.	PHYSICOCHEMICAL PARAMETERS	VALUE(%W/W)
1.	Foreign matter	0.23
2.	Loss on drying	11.5
3.	Total ash Value	9
4.	Acid-insoluble ash	1.8
5.	Water soluble ash	1.9
6.	Sulphated Ash	5

## CONCLUSION

Although there are advanced research techniques available today for evaluating plant medications, One of the simplest and most affordable ways to start identifying the correct source materials is still using the microscopic approach. In the current study, *A. platensis* leaf physiochemical analysis and microscope evaluation were performed. *Arthrospira platensis* (Family: oscillatoriaceae). The extractive values can be used to assess the chemical components of a crude medication and to estimate which components are soluble in a given solvent. Alkaloids, steroids, saponins, flavonoids, phenolic compounds, glycosides, and tannin are present, according to preliminary phytochemical investigation. The information from the initial phytochemical test will be useful in figuring out how innovative the drug truly. A drug's ash values provide information about any earthy or inorganic components as well as other contaminants that may be present. The percentages of total ash, acid-insoluble ash, and water-soluble ash are calculated. The identification of used-up or tampered pharmaceuticals is the main application of extractive values. In conclusion, the goal of the current effort was to establish criteria that could determine the legitimacy of this plant's medical value. To support and authenticate the drug, physiochemical

benchmarks and microscopic analysis may be helpful.

## ACKNOWLEDGMENT

The authors has no conflict of interest

## REFERENCE

- [1] Czerwonka A, Kaławaj K, Sławińska-Brych A, Lemieszek MK, Bartnik M, Wojtanowski KK, Zdzisińska B, Rzeski W. Anticancer effect of the water extract of a commercial *Spirulina* (*Arthrospira platensis*) product on the human lung cancer A549 cell line. *Biomedicine & Pharmacotherapy*. 2018 Oct 1;106:292-302.
- [2] S. Hosseini, S. Shahbazizadeh, K. Khosravi-Darani, M. Mozafari, *Spirulina paltensis*: food and function, *Curr. Nutr. Food Sci.* 9 (2013) 189–193.
- [3] Pan-utai W, Iamtham S. Extraction, purification and antioxidant activity of phycobiliprotein from *Arthrospira platensis*. *Process Biochemistry*. 2019 Jul 1; 82: 189-98.
- [4] S. Papadaki, K. Kyriakopoulou, I. Tzovenis, M. Krokida, Environmental impact of phycocyanin recovery from *Spirulina platensis* cyanobacterium, *Innov. Food Sci. Emerg. Technol.* 44 (2017) 217–223. doi:10.1016/J.IFSET.2017.02.014

- [5] Y. Ishimi, F. Sugiyama, J. Ezaki, M. Fujioka, and J. Wu, "Effects of spirulina, a blue-green alga, on bone metabolism in ovariectomized rats and hindlimb-unloaded mice," *Bioscience, Biotechnology and Biochemistry*, vol. 70, no. 2, pp. 363–368, 2006
- [6] Basirath Roof, Ayflegul Zeker and Knur. The Growth of *Spirulina platensis* in Different Culture Systems under Greenhouse Condition. *Turkey Journal of Biology*, 31, 2006, 47-52.
- [7] Henrikson SJ. Principles of Microbe and Cell Cultivation. Blackwell Scientific Publications, Oxford, London, 1994.
- [8] Shklar and Schwartz. Semicontinuous cultivation of the cyanobacterium *Spirulina platensis* in a closed photobioreactor. *Brazilian Journal of Chemical Engineering*, 23(1), 1988, 23-28.
- [9] C. Cingi, M. Conk-Dalay, H. Cakli, and C. Bal, "The effects of spirulina on allergic rhinitis," *European Archives of Oto-RhinoLaryngology*, vol. 265, no. 10, pp.1219–1223, 2008.
- [10] T.-S. Vo, D.-H. Ngo, and S.-K. Kim, "Marine algae as a potential pharmaceutical source for anti-allergic therapeutics," *Process Biochemistry*, vol. 47, no. 3, pp. 386–394, 2012.
- [11] Bhat V.B., Madyastha M., scavenging of Peroxynitrite by phycocyanin and phycocyanobilin from *Spirulina platensis*: protection against oxidative damage to DNA: *Biochemical and Biophysical Research Communications*. 2001;285:262-266.
- [12] Saranraj D Stella, G Usharani and S Sivasakthi. Effective recycling of Lignite Fly Ash for the laboratory cultivation of Blue Green Algae – *Spirulina platensis*. *International Journal of Microbiology Research*, 4(3), 2013, 219 - 226.
- [13] R. Kapoor and U. Mehta, "Supplementary effect of spirulina on hematological status of rats during pregnancy and lactation," *Plant Foods for Human Nutrition*, vol. 52, no. 4, pp. 315–324, 1998.
- [14] M. L. Parages, R. M. Rico, R. T. Abdala-D'iaz, M. Chabrigon, T. G. Sotiroudis, and C. Jimenez, "Acidic polysaccharides of *Arthrospira (Spirulina) platensis* induce the synthesis of TNF-  $\alpha$  in RAW macrophages," *Journal of Applied Phycology*, vol. 24, pp. 1537–1546, 2012.
- [15] M. Soltani, A.-R. Khosravi, F. Asadi, and H. Shokri, "Evaluation

- of protective efficacy of *Spirulina platensis* in Balb/C mice with candidiasis,” *Journal of Medical Mycology*, vol. 22, no. 4, pp. 329–334, 2012.
- [16] Brain K R, Turner T D. The practical evaluation of phytopharmaceuticals. Bristol: Wright-Sciencetechnica; 1975, p. 4-9.
- [17] Alejandra Guasto and Wojciech Waliszewski, *Arthrospira platensis*. Monograph Colegio Bolivar 2018-201. Available from: <https://www.colegiobolivar.edu.co/garden/wp-content/uploads/2019/06/Alejandra-Guasto-Arthrospira-Platensis.pdf>.
- [18] Edis Koru (2012). *Earth Food Spirulina (Arthrospira): Production and Quality Standarts, Food Additive*, Prof. Yehia El-Samragy (Ed.), ISBN: 978-953-51-0067-6, InTech, Available from: <http://www.intechopen.com/books/food-additive/earth-food-spirulina-arthrospira-production-and-qualitystandards>
- [19] WHO. Quality control methods for medicinal plant material. Geneva: Organisation Mondiale De La Sante; 1992, p. 22-34.
- [20] Ministry of Health and Welfare. *Indian pharmacopeia*. 4th ed. New Delhi: Ministry of Health and Welfare, Controller of Publications; 1996, p. A53-A54.
- [21] Khandelwal KR. *Practical pharmacognosy*. 18th ed. Pune: Nirali Publication; 2007.
- [22] Torey A, Sasidharan S, Yeng C, Latha LY. Standardization of *Cassia spectabilis* with respect to authenticity, assay and chemical constituent analysis. *Molecules* 2010; 15: 3411-3420.