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Preliminary phytochemical studies on some selected seaweeds from Gulf of Mannar, India

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ABSTRACT

Objective: To explore the phytochemical constituents of *Ulva reticulata* (*U. reticulata*) and *Sargassum wightii* (*S. wightii*). **Methods:** The preliminary phytochemical screening was performed by Harborne method. **Results:** The results of the phytochemical screening revealed the presence of steroids, phenolic groups, saponins, tannin, flavonoids, carbohydrates, coumarins, and xantoproteins in the extracts of *U. reticulata* and steroids, phenolic groups, saponins, tannin, flavonoids, carbohydrates, carboxylic acid, coumarins, and xantoproteins were detected in the extracts of *S. wightii*. **Conclusions:** The solvent extracts of *U. reticulata* and *S. wightii* show a number of metabolites presence, further work will emphasize the isolation and characterization of active principles responsible for bio-efficacy and bioactivity.

1. Introduction

Seaweeds are primitive on-flowering plants without true root stem and leaves. They include one of the commercially important marine renewable prosperity. Seaweeds have been used as food stuff in the Asia diet for centuries as it contains carotenoids, dietary fibres, proteins, essential fatty acids, vitamins and minerals. Fresh and dry seaweeds are extensively consumed by people especially living in the coastal areas. From the literature, it is observed that the edible seaweeds contain a significant amount of the protein, vitamins and minerals, which are essential nutrition for human^[1]. Bio-stimulant properties of seaweeds are explored for use in agriculture and the antimicrobial activities for the development of novel antibiotics. Seaweeds have some valuable medicinal components such as antibiotics, laxatives, anticoagulants, anti-ulcer products and suspending agents in radiological preparations. Seaweeds have recently received significant attention for their potential as natural antioxidants. Most of the compounds of marine algae show anti-bacterial

activities. Many metabolites isolated from marine algae have bioactive efforts^[2–4]. Among different compounds with functional properties, antioxidants are the most widely studied. Oxidative stress is an important factor in the pathological genesis, from cancer to cardiovascular and degenerative disease. To date, there are quite a lot of reports on antibacterial activity of solvent extracts from marine algae. However, there are very few reports pertaining to antifungal activity of crude solvent extracts from the seaweeds representing Phaeophyceae and Rhodophyceae^[5]. Seaweeds have been considered as potential source of marine medicinals including antimicrobial, cancer therapies^[6] hypocholesterolemic and anthelmintic substances. Many scientists also reported antimicrobial activities of marine algae^[7–12]. Several compounds from the ocean show pharmacologica activities and bioactive compounds, primarily for treating deadly diseases like cancer, acquired immuno deficiency syndrome, arthritis *etc.*, while some compounds have been used to treat inflammation *etc.* Historically seaweeds provide essential economic, environmental, aesthetic, and cultural benefits to humanity^[6]. For centuries, many of the seaweed secondary metabolites (SSM) have been used for traditional medicines due to their therapeutic potentials^[13]. Recent studies have shown that marine algae are tremendous source of

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structurally novel and diverse array of marine secondary metabolites^[11,12]. Marine algae are continuously exposed to many biotic and abiotic pressures which influence the organism's physiology, and in turn leads to the production of multifunctional natural secondary metabolites. So far, more than 2400 SSM are described and many of the SSM are natural blueprints for the development of new drugs^[14,15]. Several of these compounds are constitutive, existing in biologically active forms in healthy seaweeds. The major secondary metabolites produced by seaweeds are halogenated compounds^[16] displaying antibacterial, antifungal, antiviral, antifouling and anti-feedent properties. Although thousands of bioactive compounds have been discovered, the need for novel therapeutic compounds is still urgent in concern of number of new diseases and resistant strains of microorganisms. Few reports are available on the biopotential and biochemical studies on the seaweeds from Gulf of Mannar and peninsular coast of India^[17]. With this knowledge the present study was aimed to develop standard method for the extraction of *Ulva reticulata* (*U. reticulata*) and *Sargassum wightii* (*S. wightii*) against the selected pathogenic microorganisms.

2. Materials and methods

2.1. Collection of samples

The samples of *U. reticulata* and *S. wightii* were collected by handpicking at Rasthacaud coastal waters (Gulf of Mannar Coast, Lat N 08008'308'' E77032'80''). The collected samples were cleaned well with seawater to remove all the extraneous matter such as epiphytes, sand particles, pebbles and shells and brought to the laboratory in plastic bags. The samples were then thoroughly washed with tap water followed by distilled water. For drying, washed seaweeds were blotted on the blotting paper and spread out room temperature in shade. Shade dried samples were grounded into fine powder using tissue blender. The powdered samples were then stored in refrigerator for further use.

Table 1. Phytochemical screening of different solvents extract of *U. reticulata*.

Phytochemical	Acetone	Benzene	Chloroform	Ethanol	Petroleum ether	H ₂ O
Alkaloids	–	–	–	–	–	–
Phenol	+	+++	+	+	++	+
Flavonoids	–	+	+	–	++	+++
Saponins	–	+++	+++	–	++	–
Protein	–	–	–	–	–	–
Quinone	–	–	–	–	–	–
Steroids	++	+	–	+	–	–
Tannin	+	–	+	+	+	+++
Xanthoprotein	++	–	–	–	–	–
Carboxylic acid	–	–	–	–	–	–
Coumarins	–	–	+++	–	+	+
Carbohydrates	–	–	+++	–	–	–

2.2. Preparation of extracts

The powdered samples (2 g) and packed in Soxhlet apparatus and extracted with ethanol, acetone, petroleum ether, chloroform, benzene and water for 8 h. The crude extracts were weighed and deep frozen (–20°C) until tested. The preliminary phytochemical screening was performed by Harborne method^[18].

3. Results

3.1. *U. reticulata*

By preliminary phytochemical screening of twelve different chemical compounds (steroids, alkaloids, phenolic groups, saponins, tannin, flavonoids, anthraquinone, carbohydrates, carboxylic acid, coumarins, proteins and xantoproteins) were tested in six different extracts. Thus out of (6×12 = 72) tests for the presence or absence of the above compounds, only 26 gave positive results and the remaining 46 gave negative results. The 26 positive results showed the presence of steroids, phenolic groups, saponins, tannin, flavonoids, carbohydrates, coumarins, and xantoproteins. Alkaloids, proteins, quinone and carboxylic acid did not show any positive result for their presence in any of the six extracts tested.

Phenolic group, tannins showed the maximum presence in five different extracts followed by flavonoids in 4 extracts, steroids, saponins and coumarins in 3 different extracts. Among the six different extracts, chloroform extract showed the presence of maximum number (6) of compounds. Next to that, petroleum extracts showed 5 compounds. Acetone, water and benzene extracts showed 4 compounds each and ethanol extracts showed only three compounds (Table 1).

3.2. *S. wightii*

The preliminary phytochemical studies on acetone,

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