

Analysis of the Chemical Composition of macro Alga from area of Syrian coast

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□ ABSTRACT □

Two spaces of algae p. capillacea and Ulvafasciata were collected from Jablah coast south of Lattakia coast.

The results are revealing that all study samples contain a significant amount of hydrocarbons and carboxyl Acids and alcohols compounds.

The hydrocarbons were more preferential in p. capillacea but ketones and alcohols were more preferential in Ulvafasciata, and aldehydes was preferential too in this space especially the compound 16- Heptadecanal . 1-(+)-ascorbic acid 2- 6 dihexadecanoate is the preferential in the two spaces as carboxylic acid derivatives, and cholesterol is the specific compound in alcohols of Ulvafasciata.

Key words: Algae , hydrocarbons, carboxyl Acids, alcohols, GC/MS

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تحليل التركيب الكيميائي لبعض طحالب الماكرو من مناطق في الشاطئ السوري

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□ ملخص □

تم دراسة التركيب الكيميائي لنوعين من الطحالب هما *p. capillacea* و *Ulvafasciata* اللذين تم جمعهما من شاطئ مدينة جبلة جنوب اللاذقية، لقد أشارت النتائج إلى أن العينات المدروسة تحتوي على تراكيز مهمة من الفحوم الهيدروجينية و الأحماض الكربوكسيلية و الكحولات حيث كانت الفحوم الهيدروجينية مميزة عند النوع *p. capillacea* بينما الكيتونات و الكحولات كانت مميزة عند النوع *Ulvafasciata* كذلك الأمر بالنسبة للأدهيدات أيضاً في هذا النوع وبصورة خاصة المركب 16- Heptadecanal .

كان المركب *2- 6dihexadecanoate- ascorbic acid(+)-1* المركب المميز في كلا النوعين من مشتقات

الأحماض الكربوكسيلية و *cholesterol* كان المركب المميز من الكحولات في النوع *Ulvafasciata*.

الكلمات المفتاحية: الطحالب، الأحماض الكربوكسيلية، الكحولات، GC/MS

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Introduction

Recent trends in drug research from natural sources suggest that algae are a promising group to furnish novel biochemically active substances (Bazeset al., 2006; Chew et al., 2007; Mayer et al., 2007) because the marine algae are one of the most important organisms for producing antibiotic because of their broad spectrum of biological activities such as antimicrobial (Bouhlalet al. 2010, Chihebet al. 2009) antiviral (Bouhlalet al. 2010, Bouhlalet al. 2011, Kim and Karadeniz 2011, Bansemiret al., 2006), antifungal (de Felicio et al., 2010, Bhadury et al. 2004), antioxidant activities (Devi et al., 2011), and anticancer (Kim et al., 2011).

Therefore, research activities concerning the investigating of products of these organism, not only for a better understanding of nature and its economic importance (Myahob, 1991, Myahobet al., 1992), but also to discover metabolites of possible use for human in different fields of interest.

The antimicrobial potential of macro algae from Syrian coast remains unexplored, Many chemically compounds of marine algae with antimicrobial activity have been isolated and have new pharmaceuticals such as phenols compounds, sterols, acidic compounds, terpenes, heterocyclic carbons etc. (Bhacuniet al. 2005; Li et al., 2007; Bouhlalet al., 2011; Priyadharshiniet al., 2011). The present study was undertaken to analyze and detect the compounds of extracts of two species of marine benthic algae in Syrian coast.

Material and Methods:

Two species of algae *P. capillacea* and *Ulva fasciata* were collected from Jablah coast south of Lattakia coast between 1 and 4 m of depth and were rinsed with sterile seawater to remove all extraneous matter and dried in shadow and on air ambient until completely dry (Lima-Filho et al., 2002; Ibtissamet al., 2009).

Chemical extraction and analysis

The dried seaweeds were crushed by an electric grinder and the obtained powder was then stored at -18 °C until the extraction step. The powder of dried seaweeds (50g) (desired solvent [hexane, chloroform, methanol and petroleum ether] in cold maceration method for 48 hours using aspirated bottle and the extracts were concentrated under vacuum using a rotary evaporator. The residues were then diluted in 2 ml of pure dichloromethane and analysis by GC/MS the analysis was performed with a Hewlett-Packard Gas Chromatograph 6890 series linked to a Hewlett-Packard 7683B series mass spectrometer system equipped with a HP-5 capillary column (30m x 0.25mm, film thickness 0.25µm) the temperature was programmed from 50°C to 280°C at a rate of 5°C/min the ion source was set at 250°C and ionization voltage at 70 eV. Helium was used as carrier gas. The GC/MS identification was based on the interpretation of the spectral fragmentation followed by comparison of the spectra obtained with those of authentic samples. Searches in HP Mass spectral library NIST.

Wiley were also applied

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