

LONG-CHAIN ALIPHATIC WAX ESTERS ISOLATED FROM THE SPONGE *Chalinula saudensis* (DEMOSPONGIA) ALONG THE JEDDAH COAST OF THE RED SEA

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ABSTRACT

The sponge *Chalinula saudensis*, which occurs along the Jeddah coast, has only recently been isolated and identified. In this study, the total crude organic matter of the sponge was extracted by solvents. The total crude extract was further separated by partitioning it with hexane and water, then with water and chloroform, and finally with water and t-butanol. The chloroform layer was subjected to separation by preparative layer chromatography on silica. One fraction contained four long-chain fatty acid esters, C₂₈H₅₆O₂, C₃₀H₆₀O₂, C₃₂H₆₂O₂ and C₃₆H₇₀O₂. The second ester, C₃₀H₆₀O₂, has been identified in the fire corals *Millepora dichotoma* and *Millepora platyphylla*. The others have not previously been reported from marine organisms; however similar long-chain esters with different long aliphatic chains and with different molecular weights have been identified from other marine organisms. These compounds are normally waxy and their presence in *Chalinula saudensis* plays a vital role in the biosynthetic pathways. They also act as insulators against seasonal variations.

RESUMO

A esponja *Chalinula saudensis* ocorre ao longo da costa de Jeddah, Arabia Saudita, mas apenas recentemente foi isolada e identificada. No presente estudo a matéria orgânica total da esponja foi extraída por solventes e o extrato foi separado por partição sucessiva através do emprego de hexano e água, clorofórmio e água e finalmente t-butanol e água. A camada contendo clorofórmio foi então separada por cromatografia em sílica. Os resultados mostraram a presença de ésteres de quatro ácidos graxos de cadeia longa (C₂₈H₅₆O₂, C₃₀H₆₀O₂, C₃₂H₆₂O₂ e C₃₆H₇₀O₂), sendo que o segundo deles foi também identificado nos corais *Millepora dichotoma* e *Millepora platyphylla*. Não se tem evidência da presença dos demais compostos em outros organismos marinhos, embora haja relatos para ésteres semelhantes de cadeia longa, mas contendo diferentes cadeias alifáticas e diferentes pesos moleculares. Os compostos isolados em *C. saudensis* são geralmente ceras e sua presença na esponja tem importância não só nas rotas de biosíntese, mas servem como isolantes nas variações sazonais adversas.

Descriptors: *Chalinula saudensis*, Fatty acid esters, Jeddah coast.
Descritores: *Chalinula saudensis*, Ácidos graxos, Costa de Jeddah.

INTRODUCTION

Chalinula saudensis is a rare sponge. It was identified for the first time in the Red Sea along the Jeddah coast. The taxonomic identification was undertaken jointly by French and Saudi scientists (VACELET et al., 2001). It belongs to the class Demospongia, order Haploscleroidea, and family Chalinidae. This is a rare sponge in the sense that it has been found in very small quantities and has its full blooming season during the months of November-

December. Because it is a new species, very little work has been done on it. We sought, therefore, to identify some of its chemical constituents. This marine organism is contained within with a thick, round, irregular crust. It is attached to the substratum by a short thick peduncle with oscules located on its upper surface. The colony is 30 cm in maximum extension and is found at a depth of 20 – 30 m, close to the black coral (*Antipathes* sp). It is vivid blue in life and has a soft resilient body (Fig. 5).

MATERIAL AND METHODS

The sponge *Chalinula saudensis* was brought from near the mouth Obhur Creek on the Jeddah coast. It was removed by hand from a hard substrate, of a mainly black dead coral skeleton, by the diver (Al-SOFYANI). There were variations in the temperature, pH and salinity of the sea water in the various seasons. During the summer the temperature of the water column was 27.0°C, pH was 8.27 and the salinity was 39.83‰, whereas in winter the temperature of the water column was 25.5°C, pH was 8.02 and the salinity was 39.48‰. The specimen was taken to the laboratory immediately and washed with sea water; it was then dried and subjected to solvent extraction (methanol: chloroform: water, 2:4:1, v/v). It was stirred at room temperature for 24 hours and then filtered. The residue was subjected to a second solvent extraction and the two extracts were combined and evaporated to dryness under vacuum. It was observed that 71 gm of the wet organism yielded 253 mg of crude extract. The crude extract was partitioned with hexane: water (1:1, v/v), shaken vigorously and then separated. The water layer was again taken up in an equal volume of chloroform and the water layer was separated. This water layer was finally partitioned with t-butanol and the two layers were separated. The chloroform layer was treated with anhydrous sodium sulfate to remove traces of moisture. It was then evaporated to dryness under vacuum. The gummy material was dissolved in a little chloroform and applied to a silica preparative-layer chromatography plate (1 mm thickness). It was developed in a solvent system (methanol:chloroform, 5:95 ; v/v). Spots with an R_f value of 0.6 were eluted from the plate, dissolved in chloroform, filtered, and evaporated. It was further purified in a small column containing neutral alumina (grade III, 6" × 0.5") and finally evaporated. Attempts made to crystallize it were unsuccessful.

Samples were analyzed by GC-MS, on a Shimadzu QP-2010. The instrument was programmed as follows: injector temperature was 250°C, the injection mode was split, and the initial column temperature was 50°C (held for one min); it was increased at a rate of 10°C/min until a final temperature of 300°C was reached. It was held at that temperature for 5 min. The column was TDX-1, with a stationary phase thickness of 0.25 µm, a diameter of 0.25 mm, and a length of 30 m. The carrier gas was helium; the pressure was maintained at 69.4 kPa. The total flow rate was 13.9 mL/min and the column flow

rate was 1.22 mL/min., linear velocity was 40 cm/sec and the split ratio was 10. The ion source temperature was 250°C and the solvent cut time 4 min. Data collection was obtained on a full scan mode (40-600 µ).

RESULTS AND DISCUSSION

The term wax esters indicates long chain carboxylic acids with long chain alcohols. They are present in many marine species (SARGENT, 1978). Such long chain esters have frequently been found in organisms that experience short periods of food supply followed by long periods of food shortage. They play an important role in cuticular water proofing, buoyancy and energy metabolism (BEITTENMILLER, 1996). The presence of wax esters has also been reported from marine particulate matter. A mixture of saturated and unsaturated wax esters has been detected over a range of $C_{28} - C_{42}$ (WAKEMAN; FREW, 1982).

Individual saturated wax esters have been identified in different marine organisms. For example, hexadecanyl hexadecanoate (C_{32}) has been identified in the coral *Sinularia micrclavate* from China (ZHUIJIN et al., 1990) while hexadecanyl octadecanoate (C_{34}) has been reported in the soft coral *Sarcophyton sp* from China (ROSENBERG et al., 1982). Bandar et al., (1987) detected saturated C_{36} wax esters in a marine organism from Sri Lanka. Various mass spectrometric methods have been established to identify waxy esters (AASEN et al., 1971). In these studies, it was concluded that each peak is composed of a number of wax esters that have the same total number of carbons but that differ in alkyl and acyl moieties. The purpose of our study was to characterize the wax esters in this new species of sponge. Mass spectrometry indicated the presence of four long-chain fatty acid esters (Figs 1-4) in the chloroform fraction (purified by preparative layer chromatography). These were identified as tetradecanyl tetradecanoate ($C_{28}H_{56}O_2$), tetradecanyl hexadecanoate ($C_{30}H_{62}$), tetradecanyl 9-octadecenoate ($C_{32}H_{60}O_2$) and 9-heptadecanyl nonadecanoate, $C_{36}H_{70}O_2$. Thin Layer chromatography on silica plates was observed with solvent system (hexane:ether:acetic acid, 20:80:1, v/v); the plate was exposed to iodine vapors. The appearance of the spots indicated these compounds to be the major components (approximately 40% of the total spots), however no attempt was made to quantify them.

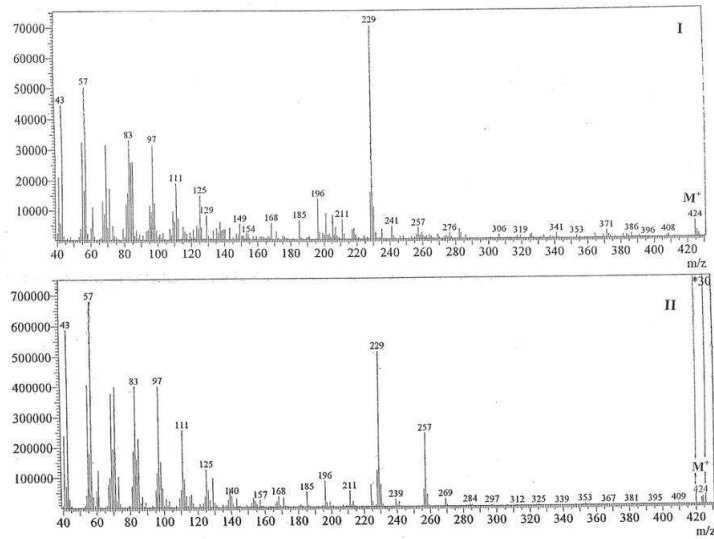


Fig. 1. Mass-spectrum of teradecanyl tetradecanoate (I) $C_{28}H_{56}O_2$ ($M/Z = 424$) isolated from a sponge along coast of Jeddah. The CG-MS indicates the presence of one compound 14/14. The second GC-MS (II) indicates a mixtures of isomers containing chains of 14/14 and 12/16 carbon atoms.

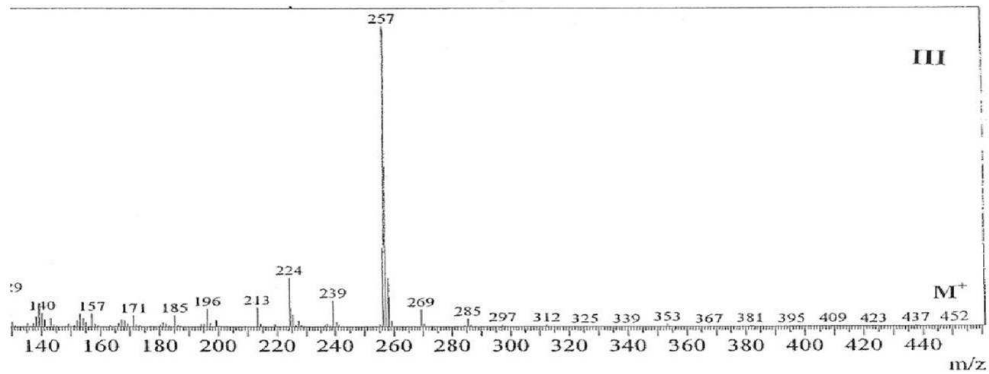


Fig. 2. Mass spectrum of tetradecanyl hexadecanoate (III) $C_{30}H_{60}O_2$ ($m/z = 452$) from a sponge indicates the presence of one compound with 14/16 carbon chain.

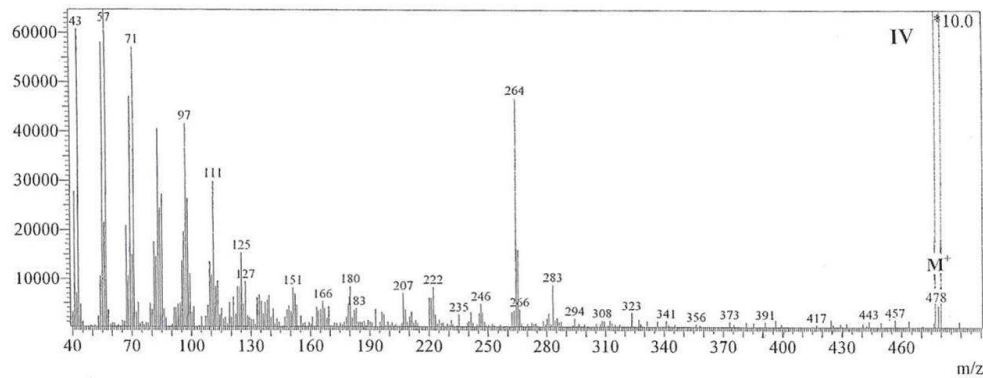


Fig. 3. Mass spectrum of tetradecanyl 9-octadecenoate (IV) $C_{32}H_{64}O_2$ ($m/z = 478$) from a sponge indicates the presence of 16/18 chain.

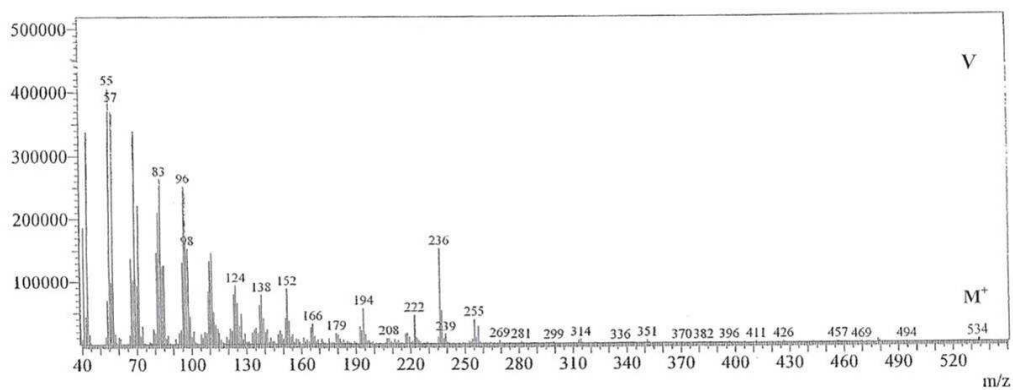


Fig. 4. Mass spectrum of 9-heptadecenyl nonadecanoate (V) $C_{32}H_{70}O_2$ ($m/z=534$) from a sponge indicates the presence of 17/19 chain.

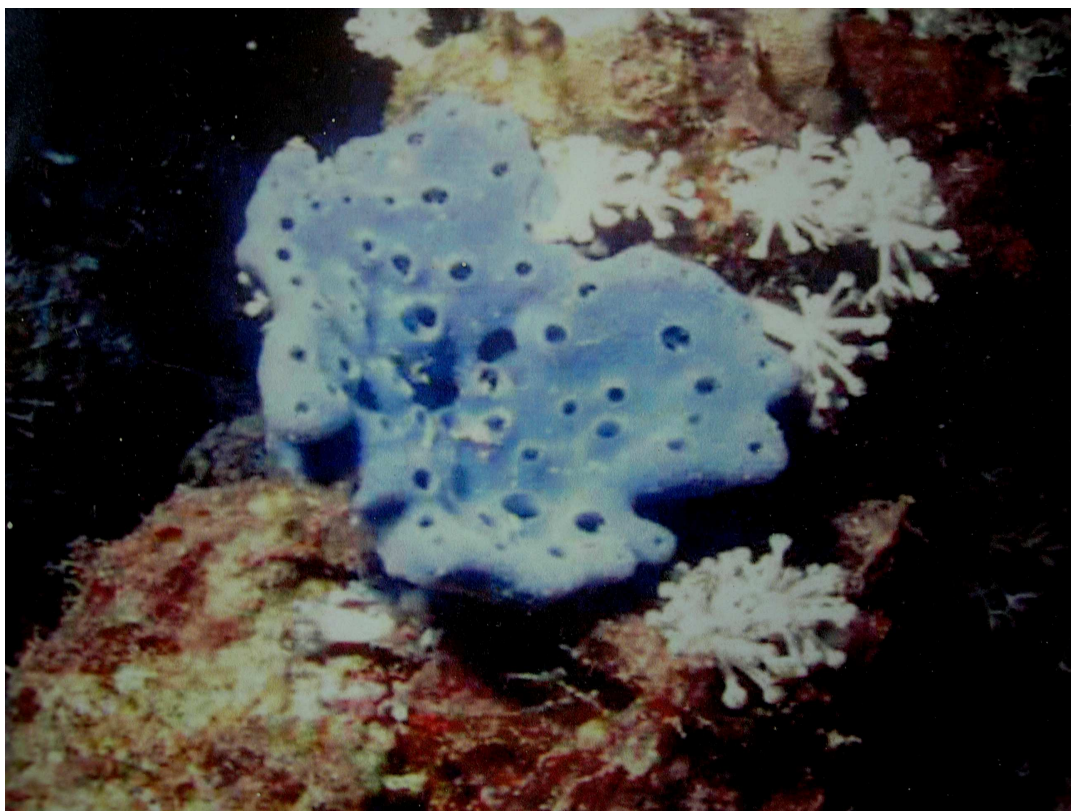


Fig. 5. *Chalinula saudensis* from the coast of Jeddah. The blue color is distinctly visible in the rocky bottom.

In the present study the GC quadruple was carefully examined to calculate the diagnostic ions, R' being the alkyl group in the alcohol moiety and R being the acyl group in the acid moiety. The relative

intensities of $[R'^{-1}]$, CO_2R' , $CH_2=C(OH)-OR'$, $CH_2=C^+(OH)R'$ were determined (Table 1). Similarly, the intensities of R , the acyl group in the acid moiety were determined; it was calculated for $[RCO_2H-$

Propyl]⁺, [RCO]⁺, [RCO₂H]⁺ and [RCOOH₂]⁺ (Table 2). The ion [RCO₂H]⁺ could be formed by a MacLafferty rearrangement. It is also possible that the alkoxy fission with transfer of one hydrogen atom from position 2 is responsible for the formation of this ion; RCO₂H⁺. RCO⁺ and RCO₂⁺ ions are formed by the normal cleavage of the bonds adjacent to the carbonyl group. From these two tables it can be calculated that two isomers were possible for tetradecanyl tetradecanoate (C₂₈H₅₆O₂); these were in the ratios of 14:0/14:0 and 12:0/16:0, as is evident from the m/z 229 and 257 respectively, while the second compound (C₂₈H₅₆O₂) existed in 14:0/16:0 form. The third compound (C₃₀H₆₀O₂) showed the existence of one double bond at position 9 of the alcohol moiety, 14:1/16:0, giving one of the fragments [R'-H] at 236. The molecular ion m/z 534 was not

revealed in the GC-MS; however, comparison with the NIST library indicated 96% resemblance. Tetradecanyl 9-octadecanoate has previously been identified in the fire corals *Millepora dichotoma* and *Millepora platyphylla* (Al-LIHAIBI et al., 2002). It would be interesting to compare and contrast the results of other groups of workers (WAKEMAN; FREW, 1982) who isolated slightly different long chain fatty acid esters from particulate matter and stated that each of them existed in isomeric form. Reyhage and Stenhagen (1963) compared C₃₂H₆₂O₂ with the one that was obtained synthetically. Compounds (II), (III) and (IV) were also isolated by Wakeman and Frew (1982) from biological and geochemical samples; our GC-MS seems to be identical with the one described by them.

Table 1. Intensities of characteristic ions containing the alcohol moiety in mass spectra of saturated and unsaturated wax esters of *Chalinula saudensis* from Red Sea coast.

No.	Molecular Formula	M ⁺		SI	R.T	Fraction	Wax		Ester		[R'-1]		CO ₂ R ⁺		CH ₂ =C ⁺ Mass		OH ⁺ OR ⁺ Int.		CH ₂ =C ⁺ Mass		OH ₂ ⁺ OR ⁺	
		Mass	Int.				Alcohol	Acid	Mass	Int.	Mass	Int.	Mass	Int.	Mass	Int.	Mass	Int.				
I	C ₂₈ H ₅₆ O ₂	424	7.76	89	26.22	CS04	14	14	196	19.31	241	6.23	256	2.29	257	5.45						
II	C ₂₈ H ₅₆ O ₂	424	0.12	91	27.74	CS04	14	14	196	12.64	241	2.36	256	8.28	257	35.42						
							12	16	168	5.13	211	7.73	228	17.67	229	75.38						
III	C ₃₀ H ₆₀ O ₂	452	0.07	90	29.77	CS04	14	16	196	6.31	241	0.66	256	26.50	257	100						
IV	C ₃₂ H ₆₂ O ₂	478	0.62	91	29.48	CS04	14	18 ^c	196	3.29	241	3.27	256	0.80	257	0.93						
V	C ₃₆ H ₇₀ O ₂	534	---b	92	29.57	CS05	17 ^d	19	236	44.78	281	1.15	298	0.12	299	0.81						

a-Studied previously as a mixture of 14/16 and 12/18 (Al-LIHAIBI et al. 1999).

b-Molecular weight according to NIST library.

c-Double bond present in the acyl moiety.

d-Double bond present in the alcohol moiety.

Table 2. Intensities of characteristic ions containing the acyl moiety in mass spectra of saturated and unsaturated wax esters of *Chalinula saudensis* from Red Sea coast.

No.	Molecular Formula	M ⁺		SI	R.T	Fraction	Wax		Ester		RCO ₂ H ⁺ - Propyl		RCO ⁺		RCO ₂ H ⁺		RCO ₂ H ₂ ⁺	
		Mass	Int.				Alcohol	Acid	Mass	Int.	Mass	Int.	Mass	Int.	Mass	Int.		
I	C ₂₈ H ₅₆ O ₂	424	7.76	89	26.22	CS04	14	14	185	9.18	211	9.62	228	22.35	229	100		
II	C ₂₈ H ₅₆ O ₂	424	0.12	91	27.74	CS04	14	14	185	7.30	211	7.73	228	17.67	229	75.38		
							12	16	213	2.55	239	3.61	256	8.28	257	35.42		
III	C ₃₀ H ₆₀ O ₂ a	452	0.07	90	29.77	CS04	14	16	213	6.61	239	8.97	256	26.50	257	100		
IV	C ₃₂ H ₆₂ O ₂	478	0.62	91	29.48	CS04	14	18 ^c	239	0.80	264	50.32	282	0.57	283	9.38		
V	C ₃₆ H ₇₀ O ₂	534	---b	92	29.57	CS05	17 ^d	19	255	9.31	281	1.15	298	0.12	299	0.81		

a-Studied previously as a mixture of 14/16 and 12/18 (WAKEMAN and FREW, 1982).

b-Molecular weight according to NIST library.

c-Double bond present in the acyl moiety.

d-Double bond present in the alcohol moiety.

CONCLUSION

Chalinula saudensis is a sponge from the Jeddah coast that has been isolated and characterized taxonomically only recently. Four long-chain aliphatic wax esters from the sponge, C₂₈H₅₆O₂, C₃₀H₆₀O₂, C₃₂H₆₂O₂ and C₃₆H₇₀O₂, were isolated and identified for the first time. Each ester was composed of a total of from 28 to 36 carbon atoms. Only one of the esters, C₃₀H₆₀O₂, has been reported previously from a marine species (the fire coral) although similar esters have

been reported from a wide variety of marine organisms and marine particulate matter. The compound C₂₈H₅₆O₂ occurred in two isomeric forms but each of the remaining compounds occurred as a single isomer. In marine organisms, wax esters probably function as an energy reserve (REINHART; VLEET, 1985). Further work will be needed to determine the precise location of these esters within the sponge and to characterize the biochemical pathways by which these compounds are produced.

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(Manuscript received 04 January 2009; revised 14 April 2010; accepted 04 November 2010)