New 5-LOX and 15-LOX Inhibitors Isolated from Red Sea Derived Brown Alga *Sarragassum Cinnerum*: An In Silico-Supported Study

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Abstract

Sargassum is a brown algal genus inhabiting tropical region. Metabolomic profiling of *Sarragassum cinnerum* "Sargassaceae", dereplicated eleven compounds **1-11**, further phytochemical investigation afforded two new aryl cresol **12-13**, along with eight known compounds **14-21**. Both new metabolites along with **19** showed moderate *in vitro* antiproliferative activity against HEPG2, MCF7, and CAC02. Molecular targets of the bioactive compounds using a pharmacophore-based virtual screening, predicts 5-LOX and 15-LOX as the most probable target linked to their observed antiproliferative activity. The validation step revealed **12** and **13** inhibited 5-LOX more prudentially than 15-LOX, while **19** showed a convergent inhibitory activity toward both enzymes. Further in-depth *in silico* analysis revealed the molecular interactions inside both enzymes active sites and explained the varying inhibitory activity for **12**, and **13** toward 5-LOX and 15-LOX. Taken together, unique metabolites in *S. cinnerum* had potential anticancer activity supported with *in-silico* investigations to facility drug discovery and development processes.

Introduction

Worldwide, the macro-algal genus *Sargassum* C. Agardh (1820) includes over 537 species, as well as 426 infra-specific names in the database¹. At present, 361 of the species names have been flagged as accepted taxonomically based on the record literature under the species name¹. *Sargassum* is a cosmopolitan brown algal genus, inhabiting temperate subtropical and tropical marine environments, which is identified by non-filamentous thallus with a holdfast that arms to form many central axes². They have specific leaves, receptacles, and the vesicles which are located on the axes near the leaves hold the algal structure upright when submerged³.

Sargassum species are nutritious and valuable source of bioactive compounds like vitamins, carotenoids, dietary fibers, proteins, and minerals⁴. Additionally, many biologically active compounds such as terpenoids, flavonoids, sterols, sulfated polysaccharides, polyphenols, sargaquinoic acids, sargachromenol, pheophytine were separated from different *Sargassum* species⁴. These isolated compounds exhibit distinct biological activities like analgesic, anti-inflammatory, antioxidant, neuroprotective, anti-microbial, anti-tumor, fibrinolytic, immune-modulatory, anti-coagulant, hepatoprotective, anti-viral activity. Therefore, *Sargassum* species have considerable potential to be utilized in pharmaceutical and nutraceutical areas^{4,5}.

According to literature, eicosanoids was formed from arachidonic acid (AA) oxidation cascade; which have been linked to pathogenesis for number of human diseases including cancer. Nowadays; there is enough evidence supporting their part in tumorigenesis and metastases⁶⁻⁹.

Although most consideration has focused on prostaglandins (PGs) and other cyclooxygenase (COX) derived metabolites. There is growing evidence suggests that lipoxygenase (LOXs) catalyzed products, such as leukotrienes (LTs), have also profound biological effects on the progression of human cancers⁶⁻⁹.

LOXs are a family of non-heme iron-containing enzymes; that catalyze the oxygenation of polyunsaturated fatty acids⁹. Several later records on the relation between LOXs and cancer development support a critical role for 5-lipoxygenase (5-LOX) and 15-lipoxygenase (15-LOX) during the initial stages of prostate, colorectal and pancreatic carcinogenesis^{6,8}. Consequently; using LOXs inhibitors has a vital effect on reducing the growth of these tumor cells^{6,8}.

In the present study, metabolomic profiling and phytochemical investigation of *S. cinnerum* were investigated using liquid chromatography high resolution mass spectrometry (LC-HRESIMS), chromatographic, and spectroscopic techniques. The antiproliferative activity of the isolated compounds was *in vitro* tested on breast Michigan Cancer Foundation-7 (MCF-7), hepatic G2 (HepG2), and colorectal adenocarcinoma-2 (CAC02) cancer cell lines. Since LOXs have a role in viability of tumor cells^{6,8}, several isolated compounds were assayed for their 5-LOX and 15-LOX inhibitory activities. Docking and dynamic studies were conducted to determine the interactions of these compounds inside the enzymes active sites.

Results And Discussion

Chemical Dereplication of *Sarragassum cinnerum*

Metabolomic profiling of *S. cinnerum* crude extract, dereplicated eleven compounds, using LC-HRESIMS. The identified metabolites **1-11** belonged to different chemical classes, including tetrahydrofuran, hydroquinone, plastoquinone, sterols, meroditerpenoids, and sulfoglycolipid (Supplementary Table S1, Figures S1-2, Figure1)

Analyzing of *S. cinnerum* crude extract, several hits were proposed (Supplementary Table S1, Figure S1-2, Figure1). The molecular ion mass peaks at *m/z* 215.1283, and 277.2162 [M-H]⁺, for the predicted molecular formulas C₁₁H₂₀O₄ and C₁₈H₃₀O₂ gave hits of the tetrahydrofuran antibacterial, and bisnorditerpenes type cytotoxicity against P388 cells (5R,7S,8S)-Communiol A **1**, and Hedaol A **2**, respectively, that were previously isolated from *Sargassum* spp.^{10,11}. The mass ion peaks at *m/z* 307.2624, and 343.2276 corresponding to the suggested molecular formulas C₂₀H₃₄O₂, and C₂₂H₃₀O₃ [M+H]⁺ fit a fatty acid, and hydroquinone anti-inflammatory derivative compound Arachidonic acid **3**, and Sargachromanol A **4**, that was previously isolated from *Sargassum pallidum*, and *Sargassum siliquastrum*, respectively^{12,13}. The ion mass peaks at *m/z* 395.2950, 425.3420, 427.3576, and 487.3060 [M+H]⁺ for the predicted molecular formulas C₂₇H₃₈O₂, C₂₉H₄₄O₂, C₂₉H₄₆O₂, and C₂₉H₄₂O₆ gave hits of the antiviral plastoquinones 2-Geranylgeranyl-6-methylbenzoquinone **5**, which was isolated from *Sargassum micracanthum*¹⁴, the anticancer steroidal nucleus of 24-Ethylcholesta-4,24(28)-dien-3,6-dione **6**, Saringosterone **7**, which were isolated from *Sargassum carpophyllum*, and *Sargassum asperfolium*, respectively^{15,16}, and the antioxidant meroditerpenoids of Nahocol A **8**, which were isolated from *Sargassum siliquastrum*¹⁷. Two major ion peaks with the *m/z* values of 445.3682 and 459.2749 [M+H]⁺ with molecular formulas C₂₉H₄₈O₃ and C₂₇H₃₈O₆ were detected and dereplicated as 24xi-Hydroperoxy-24-vinylcholesterol **9**, and Sargathunbergol A **10**, respectively, which were isolated earlier from *Sargassum carpophyllum*, and *Sargassum thunbergii*, respectively^{15,18}.

In addition, the mass ion peaks at m/z 553.2681 [M-H]⁺, for the predicted molecular formula C₂₅H₄₆O₁₁S was dereplicated sulfoglycolipid derivative 1-*O*-(11-Hexadecenyl)-3-*O*-(6'-sulfo- α -D-quinovopyranosyl) glycerol **11**, which was previously detected in *Sargassum hemiphyllum* (Figure1)¹⁹.

Phytochemical investigation of *Sarragassum cinnerum*

Based on the physicochemical and chromatographic properties, the spectral analyses from UV, ¹H, and DEPT-Q NMR, as well as comparisons with the literature and some authentic samples, the crude methanolic extract of *S. cinnerum* afforded the new aryl cresol **12–13**, along with the known *O*-Cresol **14**²⁰, *m*-Cresol **15**²¹. Additionally, arachidonic acid **16**²², eicosenoic acid **17**²², 1-*O*-arachidonyl-glycerol **18**²³, 1-*O*- arachidonyl-3-*O*-(α -D-glucopyranosyl) glycerol **19**²³, 7- β -methyl androsterol **20**²⁴, and 1-deoxy- β -D-psicosofuranose **21**²⁵, were identified (Figure 2). All characterized compounds **14**, and **15** were isolated herein for the first time from the genus *Sarragassum* (Figures S3-28, Figure 2).

Analysis of the HRESIMS, 1D and 2D NMR data of compounds **12–13** suggested a possible plastoquinones core scaffold¹¹. The HRESIMS data for compound **12** showed an adduct pseudo-molecular ion peak at m/z 314.2607 [M + H]⁺ (calc. for C₂₂H₃₄O, 314.2604), suggesting 7 degrees of unsaturation. The ¹H and DEPT-Q ¹³C NMR data (Table 1, Figures S3–S4), along with the Heteronuclear Single Quantum Correlation Experiment (HSQC) data (Figure S5), suggested six characteristic resonances appeared: three aromatic methine groups at δ_H 6.68 (1H, s) δ_C 91.1, δ_H 5.18 (1H, m) δ_C 116.0, δ_H 6.98 (1H, d, J = 8.0) δ_C 123.6, δ_H 7.13 (1H,d, J = 8.0) δ_C 123.1, three quaternary carbons at δ_C 153.8, 140.6, and 134.5, and one methyl group at δ_H 1.34 (1H, s) δ_C 29.8, suggesting the characteristic core structure for a tri-substituted benzene unit¹¹.

Table 1. DEPT-Q (400 MHz) and ¹H (100 MHz) NMR data of compounds **12, 13** in DMSO-*d*₆; Carbon multiplicities were determined by the DEPT-Q experiments

Position	12		13	
	δ_C	δ_H (J in Hz)	δ_C	δ_H (J in Hz)
1	153.8, qC		153.8, qC	
2	123.1 , CH	7.13, <i>d</i> (8.0)	123.1 , CH	7.13, <i>d</i> (8.0)
3	123.6 , CH	6.98, <i>d</i> (8.0)	123.6 , CH	6.98, <i>d</i> (8.0)
4	134.5 , qC		140.6 , qC	
5	140.6, qC		116.0 , CH	6.68, <i>s</i>
6	116.0 , CH	6.68, <i>s</i>	134.5 , qC	
7	29.8, CH ₃	1.34, <i>s</i>	31.9, CH ₃	1.23, <i>s</i>
1'	33.4 , CH ₂	2.26, <i>m</i>	33.7 , CH ₂	2.26, <i>m</i>
2'	20.3, CH ₂	2.03, overlapped	20.5, CH ₂	2.03, overlapped
3'	27.1, CH ₂	2.01, overlapped	27.1, CH ₂	2.01, overlapped
4'	127.9 , CH	5.31-5.35, <i>m</i>	127.9 , CH	5.31-5.35, <i>m</i>
5'	128.8 , CH	5.31-5.35, <i>m</i>	128.8 , CH	5.31-5.35, <i>m</i>
6'	25.5, CH ₂	2.78, overlapped	25.5, CH ₂	2.78, overlapped
7'	128.0 , CH	5.31-5.35, <i>m</i>	128.1 , CH	5.31-5.35, <i>m</i>
8`	128.3 , CH	5.31-5.35, <i>m</i>	128.4 , CH	5.31-5.35, <i>m</i>
9`	25.6, CH ₂	2.78, overlapped	25.6, CH ₂	2.78, overlapped
10`	128.2 , CH	5.31-5.35, <i>m</i>	128.3 , CH	5.31-5.35, <i>m</i>
11`	129.4, CH	5.31-5.35, <i>m</i>	129.4, CH	5.31-5.35, <i>m</i>
12`	24.9 , CH ₂	1.52, overlapped	24.9 , CH ₂	1.52, overlapped
13`	28.9 , CH ₂	1.24, overlapped	28.9 , CH ₂	1.24, overlapped
14`	22.4 , CH ₂	1.25, overlapped	22.4 , CH ₂	1.25, overlapped
15`	14.5 ,CH ₃	0.89, <i>t</i> (6.6)	14.3 ,CH ₃	0.85, <i>t</i> (6.6)

qC, quaternary, CH, methine, CH₂, methylene, CH₃, methyl carbons

NMR data also showed eight aliphatic methylene groups at δ_H 1.20-2.8 δ_C 20.5-33.7 (Table 1), three olefinic methine groups at δ_H 5.31-5.35 (6H, m) δ_C 127.9-129.4. These signals are suggestive characteristics for 4,7,11-pentadecenyl moiety, where the Heteronuclear Multiple-Bond Correlation (HMBC) experiment

of **12** (Figure 3) confirmed the position of the three olefinic methine groups at 4,7,11 of the alkene side chain. Moreover, HMBC experiment showed the 3J -HMBC correlation of the proton H-1' δ_H 2.26 (δ_C 33.4) with the quaternary carbonyl carbon C-4' (δ_C 134.5). Accordingly, compound **12** was identified as 4-(1-(4,7,11-pentadecenyl)-*o*-cresol.

The molecular formula of compound **13** was identical to that of **12** based on HRESIMS ($C_{22}H_{34}O$). The 1H and ^{13}C NMR data was also very close to those of compound **12** for the 4,7,11-pentadecenyl moiety, but differed in the resonated chemical shifts of the aromatic attached methyl group of the core tri-substituted benzene unit (Table 1). Comparing the DEPT-Q ^{13}C NMR data of compound **13** with those of **12** showed a downfield shifting of carbons C-7 ($\Delta\delta_C +2.1$), compared with those of compound **12** (Table 1). This suggested a positional difference of the location of the aromatic attached methyl group in the tri-substituted benzene unit *versus* **12** (Table 1, Figure 2, Figures S8–S12). The assignment of the location of the aromatic attached methyl group in **13** was aided by the HMBC experiment. A 3J -HMBC correlation (Figure 4) of compound **13** proton H-7 δ_H 1.23 (δ_C 31.9) with the quaternary carbonyl carbon C-4 (δ_C 140.6) and a 4J -HMBC correlation of the proton H-7 δ_H 1.23 (δ_C 31.9) with the methylene carbon C-1' (δ_C 33.7) confirmed the meta-location of aromatic attached methyl group at the cresol moiety. Accordingly, compound **13** was identified as 4-(1-(4,7,11-pentadecenyl)-*m*-cresol.

Antiproliferative activity of the isolated metabolites

The isolated compounds **12–21** were *in vitro* screened for their ability as antiproliferative against hepatic, breast, and colorectal carcinoma cell lines (HEPG2, MCF7, and CACO2, respectively). Results showed that compounds **12**, **13**, and **19** were able to inhibit the growth of all tested cell lines moderately with IC_{50} values ranged from 11.2 ± 0.6 to $21.6 \pm 1.3 \mu M$ (Table 2).

Table 2. *In vitro* antiproliferative activity of the isolated compounds **12–21** expressed as $IC_{50} \pm$ (SSEM) μM

IC_{50} (μM)			
Code	HEPG2	MCF7	CACO2
12	14.5 ± 0.8	17.6 ± 0.9	18.2 ± 0.7
13	13.1 ± 1.1	12.7 ± 1.3	11.2 ± 0.6
14	> 50	> 50	> 50
15	> 50	> 50	> 50
16	> 50	> 50	> 50
17	> 50	> 50	> 50
18	> 50	> 50	> 50
19	18.5 ± 1.4	21.6 ± 1.3	15.7 ± 0.9
20	> 50	> 50	> 50
21	> 50	> 50	> 50
Doxorubicin	4.2 ± 0.3	3.8 ± 0.2	3.4 ± 0.1

IC_{50} value of compounds against each cancer cell line, which was defined as the concentration (μM) that caused a 50% inhibition of cell growth *in vitro*, data were expressed as mean $\pm$ SEM ($n = 3$). One-way analysis of variance (ANOVA) followed by Dunnett's test using PASW Statistics® version 18 (Quarry Bay, Hong Kong) was applied. GraphPad Prism software version 6 (La Jolla, CA, USA) was used for statistical calculations. * statistically significant at $p < 0.05$. Doxorubicin a positive control

Virtual screening-based target identification

Characterization of the biological target for a certain molecule is a true challenge, however, the continuous development of *in silico* tools including molecular modeling and virtual screening has significantly improve the success rate of finding suitable molecular targets. Many online target identification platforms are currently available, and their search protocols are either structural based or ligand-based. PharmMapper is one of these online platforms that can screen and suggest the most probable protein targets of a query molecule based on its pharmacophore model²⁶. The basic principle of pharmacophore-based screening is that the binding of certain molecules with their protein targets is mainly determined by key pharmacophore maps (i.e. spatial arrangement of structural features). Thus, molecules that their shapes are able to fit with these pharmacophore maps have the highest probability to bind the same protein target. Consequently, PharmMapper was used to propose a proper protein target for compounds **12**, **13** and **19**. 5-LOX and 15-LOX were found to be the top-scoring hits for these metabolites. In addition, they have a direct link to tumorigenesis^{8,27–30}. Consequently, they were selected for further *in vitro* and *in silico* validations.

LOX inhibition assay

To validate the preliminary virtual screening prediction, compounds **12**, **13** and **19** were assayed for their 5-LOX and 15-LOX inhibitory activities. Interestingly the three compounds achieved potent enzyme inhibition toward 5-LOX (IC_{50} 1.3 ± 0.1 to $2.1 \pm 0.4 \mu M$, Table 3). However, their activity against 15-LOX was weaker, particularly compounds **12** and **13** (IC_{50} 25.3 ± 0.4 and $23.6 \pm 0.3 \mu M$, respectively) that were more selective for 5-LOX (Table 3).

Table 3. Docking scores, binding free energies, K_i and IC_{50} values of compounds 12, 13, and 19 together with the co-crystallized inhibitors NDGA and AA

Compound	ΔG_{Vina}^*		ΔG_{FEP}^{**}		ΔG_{KDEEP}^{***}		$K_i^{\#}$		$IC_{50}^{\#}$	
	5-LOX	15-LOX	5-LOX	15-LOX	5-LOX	15-LOX	5-LOX	15-LOX	5-LOX	15-LOX
12	-9.3	-5.1	-8.1	-4.4	-7.7	-4.6	0.9 ± 0.1	17.4 ± 0.2	1.6 ± 0.3	25.3 ± 0.4
13	-8.9	-5.5	-8.0	-4.7	-7.5	-4.5	0.7 ± 0.2	14.3 ± 0.4	1.3 ± 0.1	23.6 ± 0.3
19	-9.1	-7.7	-7.9	-7.1	-7.6	-7.2	1.4 ± 0.2	4.2 ± 0.1	2.1 ± 0.4	6.7 ± 0.3
NDGA ^{##}	-7.2	-6.9	-7.0	-6.5	-6.8	-6.5	6.9 ± 0.1	6.1 ± 0.2	8.8 ± 0.3	-9.5 ± 0.5
AA ^{##}	-7.6	-7.0	-6.2	-6.4	-7.1	-6.3	-	-	-	-

Lipoxygenase (LOX), nordihydroguaiaretic acid (NDGA), arachidonic acid (AA)

*Vina docking scores calculated in kcal/mol

**MDS-derived binding free energies calculated in kcal/mol by FEP method

***Neural networking-derived binding free energies calculated in kcal/mol by KDEEP software

[#]*In vitro* inhibition constant (K_i) and inhibition concentration 50 (IC_{50}) expressed as μM

^{##}The reported co-crystallized ligands

Moreover, they showed inhibitory constants (K_i) ranged from 0.7 ± 0.2 to $17.4 \pm 0.2 \mu M$ (Table 3), and these values were most agree with the competitive inhibition of both enzymes³¹.

The results of enzyme inhibition assay were also correlated with those of the antiproliferative ones for HepG2 and MCF-7 and CACO2. Surprisingly, overexpression of 5-LOX has been reported in breast, liver and colorectal cancers²⁷⁻²⁹. Also 15-LOX has been reported to be over expressed in a number of tumors like prostate and breast cancers. Hence, these enzymes can be considered promising targets for cancer therapy.

Molecular Docking and dynamic simulation

5-LOX has a hydrophobic active site⁹, that harbor a catalytic iron (Fe^{+2}), and such hydrophobicity is essential to allow efficient binding with the hydrophobic substrate, arachidonic acid (AA) (Figure 2)⁹. Compounds **12**, **13**, and **19** have extended unsaturated hydrophobic side chains that resemble AA (Figure 5).

Molecular docking experiments revealed that these compounds could bind with the 5-LOX's active site efficiently with binding scores ranged from -8.9 to -9.3 kcal/mol (Figure 5), and their bindings were even better than the co-crystallized ligands (Table 3). Additionally, the phenolic moiety of both compounds was involved in H-bonding with HIS-600 similarly to the co-crystallized redox-type inhibitor, nordihydroguaiaretic acid (NDGA) (Figure 5).

Compounds 12 and 13's hydrophobic side chains were able to adapt themselves inside the hydrophobic U-shaped active site, where they took convergent orientations, but slightly different from that of AA (Figure 5). LEU-368, ILE-406, LEU-414, and LEU-607 were the main amino acid residues involved in the hydrophobic interactions with their side chains, while PHE-359, TRP-599, and PRO-569 interacted with their aromatic moieties. The binding mode of compound 19 was quite different, where its polar carbohydrate head interacted with LYS-409, GLN-413, and ILE-673 through four strong hydrogen bonds ($< 2.5\text{\AA}$), while its hydrophobic tail interacted with LEU-368, LEU-414, TRP-599, and LEU-607 (Figure 5).

Subsequent Molecular Dynamic Simulation (MDS) experiments (50ns) revealed that the three compounds 12, 13, and 19 were able to stabilize the enzyme's active site.

Compounds 12 and 13's positions remained to change over the first 32.4 ns (RMSD $\sim 3.4\text{\AA}$). Afterward, they became stable till the end of the simulation (average RMSD values of 2.67 and $2.59\text{\AA}$, respectively), where their extended hydrocarbon chains became more relaxed and straight (Figure 6). The H-bonds between their phenolic group and HIS-600 remained unchanged throughout the MDS. Starting from 22.6ns, GLN-363's side chain became also involved in H-bonding with the phenolic group of both compounds (Figure 6). Additionally, compound 12's tail remained imbedded inside a hydrophobic pocket consists of the side chains of TRP-147, PHE-151, LEU-368, LEU-373, and LEU-414, while compound 13's tail, settled inside another hydrophobic pocket consists of TRP-147, LEU-414, ILE-415, and VAL-433 (Figure 6).

Similarly, the hydrophobic part of compound 19 was compacted at the beginning of MDS, and gradually become more extended till stabilization at 25.4ns (RMSD = $2.75\text{\AA}$), where PHE-359, PRO-569, and TRP-599 became involved in hydrophobic interactions with the molecule's tail. Furthermore, the side chain of LYS-409 became involved in an additional H-bonding with the molecule's hydrophilic carbohydrate part (Figure 6). Further binding free energy calculations (ΔG_{FEP} and ΔG_{KDEEP}) revealed that compounds 12, 13, and 19 got higher binding free energy values than that of the co-crystallized ligands (Figure 7), and were in good accordance with the *in vitro* enzyme inhibition (Figure 7).

On the other hand, docking scores of the three compounds against 15-LOX were significantly lower, particularly for compounds 12 and 13 (Figure 7) that showed only hydrophobic interactions (with PHE-184, TYR-185, PHE-365, LEU-374, LEU-415, LEU-419, LEU-420, VAL-603, LEU-609, and LEU-610) inside the enzyme's active site. Upon MDS (Figure 7), both compounds showed unstable positioning inside 15-LOX active site, where the surrounding hydrophobic amino acid residues were able to keep them in position till ~ 23.4ns (RMSD ~ 3.5 Å). Afterward, their position inside the active site began to change dramatically and their RMSDs reached about 7.1 Å and remained to fluctuate till the end of MDS with gross averages of 8.1 and 7.5 Å, respectively over the 50ns of simulation. This obvious instability of compounds 11 and 12 reflected on their binding free energies and in vitro inhibitory activities that were significantly lower in comparison with 5-LOX (Figure 7).

Such instability was not the case with compound 19, where the hydrophilic carbohydrate moiety was able to form a network of H-bonds (7 H-bonds) with TYR-185, GLN-425, ARG-429, and ASP-602. Furthermore, these hydrophilic interactions remained unchanged over the course of MDS, and thus compound 20's RMSD was at equilibrium (~2.6 Å) till the end of MDS. Such structural and dynamic information can explain the convergent inhibitory activity of compound 19 against both 5-LOX and 15-LOX (Figure 7), and the moderate selective inhibition of 5-LOX by compounds 12 and 13 over 15-LOX.

Materials And Methods

Algae material

The marine algae *S. cinnerum* was collected during January 2020 along the shore of the Red Sea in Hurghada, Egypt. The samples were collected in sterilized polyethylene bags, and kept in an icebox, for transportation to the laboratory. Samples were washed thoroughly with sterile distilled water to remove any associated debris. A voucher specimen (2020-BuPD 55) was deposited at the Department of Pharmacognosy, Faculty of Pharmacy, Beni-Suef University, Egypt.

Chemicals and reagents

The solvents used in this work included *n*-hexane (*n*-hex., boiling point b.p. 60–80 °C), dichloromethane (DCM), ethyl acetate (EtOAc), *n*-butanol (*n*-but.), and methanol (MeOH) were purchased from El-Nasr Company for Pharmaceuticals and Chemicals (Egypt). High Performance Liquid Chromatography (HPLC) and deuterated solvents used for chromatographic and spectroscopic analyses were purchased from Sigma-Aldrich (Saint Louis, Missouri, USA), including HPLC-methanol, HPLC-water, HPLC-acetonitrile, deuterium oxide (D₂O), methanol (CD₃OD), and dimethyl sulfoxide (DMSO-*d*₆). Column chromatography (CC) was performed using silica gel 60 (63–200 µm, E. Merck, Sigma-Aldrich), and Sephadex LH-20 (0.25–0.1 mm, GE Healthcare, Sigma-Aldrich), while silica gel GF254 for Thin-layer chromatography (TLC) (El-Nasr Company for Pharmaceuticals and Chemicals, Egypt) was employed for vacuum liquid chromatography (VLC). Thin-layer chromatography (TLC) was carried out using pre-coated silica gel 60 GF254 plates (E. Merck, Darmstadt, Germany; 20 × 20 cm, 0.25 mm in thickness). Spots were visualized by spraying with *para*-anisaldehyde (PAA) reagent (85 : 5 : 10 : 0.5 absolute EtOH : sulfuric acid : G.A.A. : *para*-anisaldehyde), followed by heating at 110 °C³². For the biological study, doxorubicin (Sigma-Aldrich, Germany) was used as a positive control while the HEPG2, MCF7, and CACO2 cancer cell lines were obtained from the American Type Culture Collection (ATCC, Rockville, MD, USA; HPACC, Salisbury, UK) and were routinely sub-cultured twice per week.

Spectral analyses

Proton ¹H and Distortionless Enhancement by Polarization Transfer-Q (DEPT-Q) ¹³C NMR spectra were recorded at 400 and 100 MHz, respectively. Tetramethylsilane (TMS) was used as an internal standard in deuterium oxide (D₂O), methanol (CD₃OD), and dimethyl sulfoxide (DMSO-*d*₆), using the residual solvent peak ($\delta_{\text{H}} = 4.78$), ($\delta_{\text{H}} = 3.34, 4.78$ and $\delta_{\text{C}} = 49.9$) and ($\delta_{\text{H}} = 2.50$ and $\delta_{\text{C}} = 39.5$) as references, respectively. Measurements were performed on a Bruker Advance III 400 MHz with BBFO Smart Probe and a Bruker 400 MHz EON Nitrogen-Free Magnet (Bruker AG, Billerica, MA, USA). Carbon multiplicities were determined using a DEPT-Q experiment. The ultraviolet radiation (UV) spectrum in methanol was obtained using a Shimadzu UV 2401PC spectrophotometer (Shimadzu Corporation – UV-2401PC/UV-2501PC, Kyoto, Japan). Infrared (IR) spectra were measured using a Jasco FTIR 300E infrared spectrophotometer. HRESIMS data were obtained using an Acquity Ultra Performance Liquid Chromatography system coupled to a Synapt G2 HDMS quadrupole time-of-flight hybrid mass spectrometer (Waters, Milford, MA, USA). HPLC chromatographic separations were conducted using an Agilent 1260 Infinity preparative pump (G1361A), Agilent 1260 diode array detector VL (G1315 D), Agilent 1260 Infinity Thermostand column compartment (G1361 A), Agilent 1260 Infinity preparative autosampler (G2260A) and a YMC-Pack ODS-A A-324 column (i.d. 10 × 300 mm, YMC, Kyoto, Japan).

Extraction and fractionation of algae material

Sarragassum cinnerum (0.5 kg) were collected and air-dried in the shade for one month. After drying, the brown algae were finely powdered using an OC-60B/60B grinding machine (60–120 mesh, Henan, Mainland China). The finely powdered algae was extracted by maceration using 70% methanol (3 L, 3×, seven days each) at room temperature, and concentrated under vacuum at 45 °C using a rotary evaporator (Buchi Rotavapor R-300, Cole-Parmer, Vernon Hills, IL, USA) to afford 75 g crude extract. The dry extract was suspended in 100 mL distilled water (H₂O), and successively portioned with solvents of different polarities (*n*-Hex., DCM, EtOAc, and *n*-but.). The organic phase in each step separately evaporated under reduced pressure to afford the corresponding fractions I (8.0 g), II (1.5 g), III (1.5 g) and IV (3.0 g), respectively, while the remaining mother liquor was then concentrated down to give the aqueous fraction (V). All resulting fractions were kept at 4 °C for biological and phytochemical investigations.

Metabolomic Analysis Procedure

The crude methanolic extract from *S. cinnerum* was prepared at 1 mg/mL for mass spectrometry analysis. The recovered methanolic extract was subjected to metabolic analysis using LC-HRESIMS according to Abdelmohsen et al. 2014³³. An Acquity Ultra Performance Liquid Chromatography system connected to a

Synapt G2 HDMS quadrupole time-of-flight hybrid mass spectrometer (Waters, Milford, USA) was used. Positive and negative ESI ionization modes were utilized to carry out the high-resolution mass spectrometry coupled with a spray voltage at 4.5 kV, the capillary temperature at 320 °C, and mass range from m/z 150–1500. The MS dataset was processed and data were extracted using MZmine 2.20 based on the established parameters²². Mass ion peaks were detected and accompanied by chromatogram builder and chromatogram deconvolution. The local minimum search algorithm was addressed and isotopes were also distinguished via the isotopic peaks of grouper. Missing peaks were displayed using the gap-filling peak finder. An adduct search along with a complex search was carried out. The processed data set was next subjected to molecular formula prediction and peak identification. The positive and negative ionization mode data sets from the respective extract were dereplicated against the DNP (Dictionary of Natural Products) databases.

Isolation and purification of compounds

Fraction I (8 g) was subjected to normal VLC fractionation using silica gel GF₂₅₄ (column 6 × 30 cm, 50 g). Elution was performed using *n*-hex. : EtOAc gradient mixtures in order of increasing polarities (0, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 60, 80 and 100%, 500ml each). The effluents from the column were collected in fractions (100 mL each); and each collected fraction was concentrated and monitored by TLC using the system *n*-hex. : EtOAc 8 : 2 and PAA reagent. Similar fractions were grouped and concentrated under reduced pressure to provide three sub-fractions (I₁–I₃). Subfraction II₂ (3.0 g) was further fractionated on silica gel 60 (100 × 1 cm, 50 g). Elution was performed using *n*-hex. : EtOAc gradient mixtures in the order of increasing polarities (0, 1, 2, 3,4,5,6,7,8,9 and 10%, 1 L each, FR 3 mL min⁻¹), to afford four sub-subfractions (II₂₋₁–II₂₋₄). Sub-subfraction II₂₋₁ (50 mg) was further fractionated on silica gel 60 (100 × 1 cm, 20 g). Elution was performed using *n*-hex. : EtOAc isocratic mixture (1%, 500 mL, FR 3 mL min⁻¹) to afford compound **17** (20 mg). Sub-subfractions II₂₋₂, and II₂₋₄ (70, 30 mg each) was further fractionated on C-18 RP-HPLC using H₂O–CH₃CN (10–60%, 30 min, 5 mL/min) to afford compound **12** (20 mg), **13** (10 mg), **14** (10 mg), **15** (7 mg). Sub-subfraction II₂₋₃ (100 mg) was further fractionated on silica gel 60 (100 × 1 cm, 20 g). Elution was performed using *n*-hex. : EtOAc isocratic mixture (5%, 500 mL, FR 3 mL min⁻¹) to afford compound **16** (50 mg). Finally, subfraction II₃ was further fractionated on silica gel 60 (100 × 1 cm, 20 g). Elution was performed using *n*-hex. : EtOAc isocratic mixture (1%, 500 mL, FR 3 mL min⁻¹) to afford compound **20** (30 mg). Fraction II (1.5 g) was subjected to normal VLC fractionation on a silica gel (column 6 × 30 cm, 50 g). Elution was performed using DCM : MeOH gradient mixtures in the order of increasing polarities (0, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 60, 80 and 100%, 1 L each). The effluents were collected in fractions (100 mL each); each fraction was concentrated and monitored by TLC using the system DCM : MeOH 9.5 : 0.5 and PAA reagent. Similar fractions were grouped and concentrated under reduced pressure to provide two sub-fractions (II₁–II₂), which were further purified on a Sephadex LH₂₀ column (0.25–0.1 mm, 100 × 0.5 cm, 100 gm) which eluted with MeOH to afford compound **18** (16 mg), and **19** (6 mg), separately.

Crystallization of fractions IV was performed separately using CH₂Cl₂, and afforded compounds **21** (2 g).

4-(1-(4,7,11-pentadecenyl)-*o*-cresol (12): Yellow oil; [UV (MeOH) $\lambda_{\max}$ (log_e) 225 (5.5), 260 (6.0), 300 (4.5) nm; IR $\nu_{\max}$ (KBr) 3429, 3100, 3000, 1680, 1600, 1475, 1450, 1300, 835, 601 cm⁻¹; NMR data; see Table 1; HRESIMS m/z 314.2607 [M + H]⁺ (calc. for C₂₂H₃₄O, 314.2604).

4-(1-(4,7,11-pentadecenyl)-*m*-cresol (13): Yellow oil; UV (MeOH) $\lambda_{\max}$ (log_e) 225 (5.5), 260 (6.0), 300 (4.5) nm; IR $\nu_{\max}$ (KBr) 3429, 3100, 3000, 1680, 1600, 1475, 1450, 1300, 835, 601 cm⁻¹; NMR data; see Table 1; HRESIMS m/z 314.2609 [M + H]⁺ (calc. for C₂₂H₃₄O, 314.2604).

Antiproliferative assay

The antiproliferative activity of the isolated compounds **12–21** was measured by the sulphorhodamine B (SRB) assay as described by Skehan et al. 1990³⁴, and Vichai and Kirtikara 2006³⁵, on breast (MCF-7), liver (HepG2) and colorectal (CACO2) cancer cell lines. Cells were seeded in 96-well microtiter plates at initial concentration of 3×10³cell/well in 150µL, fresh medium and left for 24h to attach to the plates. Different concentrations 0, 5, 12.5, 25, 50µg/mL of the respective compound were added. The plates were incubated for 48h. The cells were fixed with 50µL cold trichloroacetic acid (10% final concentration) for 1h at 4°C. The plates were washed with distilled water (automatic washer Tecan, Germany) and stained with 50µL 0.4% SRB dissolved in 1% acetic acid for 30 min., at room temperature. Then they were washed with 1% acetic acid and air-dried. The dye was solubilized with 100 µL/well of 10M trisbase (pH 10.5). The optical density of each well was measured spectrophotometrically at 570nm using an ELISA microplate reader (Sunrise Tecan reader, Germany). Doxorubicin was used as a positive control. The mean background absorbance's was automatically subtracted and mean values of each drug concentration were calculated. The experiment was repeated three times, and then the IC₅₀ values were calculated.

Lipoxygenase (LOXs) inhibition assay

The ability of the isolated compounds **12–13**, **19** to inhibit 5-LOX and 15-LOX enzymes (IC₅₀ and K_i values, µM) was determined using human recombinant enzyme assay kits (catalogue no 60402 and 10011263, Cayman Chemical, Ann Arbor, MI, USA) following manufacturer's specifications³⁶. Stock solutions were freshly prepared before use and buffer solution (0.1M Tris HCl, PH, 7.4) was used. 10µL of each compound were prepared, dissolved in the least amount of DMSO and diluted with the stock solution to be in concentrations of (0.001, 0.1, 1, 5, 10µM) in a final volume of 210mL. The kinetic parameters for both 5-LOX and 15-LOX were determined by measuring the increase in absorbance at 238nm in an Agilent 8453 Diode Array Spectrophotometer (Agilent Technologies, Santa Clara, CA, USA). Substrate concentration was ranged from 5 to 50µM. Substrate concentrations (5, 10, 20, 30, 40, 50µM) were monitored in triplicate for each sample³⁷. Doxorubicin was used as a positive control.

Docking study

The crystal structures of both 5-LOX and 15-LOX (PDB: 6N2W and 4NRE) were used for the docking analysis using Autodock Vina docking machine³⁸. The co-crystallized ligands nordihydroguaiaretic acid (NDGA) and AA were used to determine the binding sites. The ligand to binding site shape matching root mean

square (RMSD) threshold was set to 2.0 Å. The interaction energies were determined using the Charmm Force Field (v.1.02) with 10.0 Å as a non-bonded cutoff distance and distance dependent dielectric. Then, 5.0 Å was set as an energy grid extending from the binding site³⁹. The tested compounds were energy minimized inside the selected binding pocket. The editing and visualization of the generated binding poses were performed using Pymol software⁴⁰.

Molecular Dynamic Simulation

Molecular dynamic simulations (MDS) for ligand enzyme complexes were performed according to the previous protocol⁴¹, using the Nanoscale Molecular Dynamics (NAMD) 2.6 software⁴², applying the CHARMM27 force field⁴³. Hydrogen atoms were added to the protein structures using the psfgen plugin included in the Visual Molecular Dynamic (VMD) 1.9 software⁴⁴. Afterward, the whole systems were solvated using TIP3P water particles and 0.15M NaCl. The energy of the generated systems was firstly minimized and gradually heated to 300K and equilibrated for 200/seconds. Subsequently, the MDS was continued for 20ns, and the trajectory was stored every 0.1ns and further analyzed with the VMD 1.9 software. The MDS output were sampled every 0.1ns to evaluate the conformational changes of the entire system to analyze the root mean square deviation (RMSD) and root mean square fluctuation (RMSF). The topologies and parameters of the tested compounds were prepared using the VMD Force Field Toolkit (ffTK), and the online software Ligand Reader and Modeler (<http://www.charmm-gui.org/?doc=input/ligandrm>)⁴⁵. MDS-derived binding free energies (ΔG) were calculated using the free energy perturbation (FEP) method through the web-based software Absolute Ligand Binder along with MDS using NAMD software^{45,46}. Moreover, ΔG was calculated using another web-based software utilizing neural networking in its calculations, namely KDEEP (<https://www.playmolecule.org/Kdeep/>)⁴⁷.

Statistical analysis

All *in vitro* experiments were performed in triplicate. Pooled data were presented as the mean $\pm$ standard error of the mean (SEM) of at least three independent experiments. The differences among various treatment groups were determined by ANOVA, followed by Dunnett's test using PASW Statistics® version 18 (Quarry Bay, Hong Kong). A difference of $p < 0.05$ was considered statistically significant compared with a vehicle-treated control group and shown by a *symbol. The IC₅₀ values were determined using a nonlinear regression curve fitting analysis using GraphPad Prism software version 6 (La Jolla, CA, USA).

Conclusions

In the present study, phytochemical investigation of the brown algae *S. cinnerum* with the guidance of LC-HRESIMS dereplication, afforded two new phenolic derivatives (compounds **12** and **13**) together with further nine known ones. These two new compounds along with compound **19** exhibited moderate *in vitro* antiproliferative activity against hepatic, breast, and colorectal carcinoma cell lines (HEPG2, MCF7, and CACO2, respectively) with IC₅₀ values ranged from 11.2 $\pm$ 0.6 to 21.6 $\pm$ 1.3 μ M. A subsequent comprehensive pharmacophore-based virtual screening was conducted to putatively figure out the molecular targets of these active compounds. Both 5-LOX and 15-LOX were found to be the most probable hits. *In vitro* testing results supported the preliminary virtual screenings outcomes, where both new compounds **12** and **13** showed higher selective inhibition toward 5-LOX over 15-LOX, while compound **19** was able to inhibit both enzymes. Additional kinetic investigations showed that the three compounds exhibit competitive inhibition of both enzymes. To get a deeper insight into the mode of interaction inside the active site of each enzyme, we conducted a series of *in silico* experiments (docking, MDS, and binding free energy calculations) that putatively explained the lower inhibitory activity of compounds **12** and **13** against 15-LOX in comparison with 5-LOX to the lack of strong H-bond interactions inside the 15-LOX's active site. In turn, both compounds exhibited significant instability and lower binding free energy that translated eventually into weaker activity against 15-LOX. The present study shows the potential of marine natural products in providing unique metabolites with potent biological activities and highlighted the power of *in silico* investigations in the facilitating of drug discovery and development processes.

Declarations

Supplementary materials: The NMR data are available as supplementary data.

Author contributions: Conceptualization: URA, AHE, AMS; methodology: AHE, URA, SIA, AMS; software: AHE, URA, AMS, SIA, OMH, AM; formal analysis: AHE, URA, SIA, AMS, MA, EMM, OMH; investigation: AHE, URA, SIA, AMS, HS, KAY; resources: AHE, URA, SIA, AMS, ASA; data curation: AHE, URA, SIA, AMS, AMS; writing—original draft: AHE, AMS, URA; writing—review and editing: URA, AHE, SIA, AMS, HS, AM, MA, MMA, EMM, OMH, KAY, ASA, MA, AMS; supervision: URA, SIA; project administration: SIA; funding acquisition: SIA. All authors have read and agreed to the published version of the manuscript.

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Figures

Figure 1

Dereplicated metabolites from LC-HRESIMS analysis of *Sarragassum cinnerum*

Figure 2

Structures of compounds isolated from *Sarragassum cinnerum* 12-21 together with 5-LOX and 15-LOX's co-crystalized ligands AA 16 and NDGA

Figure 3

Selected HMBC (->) correlations of compound 12

Figure 4

Selected HMBC (->) correlations of compound 13

Figure 5

Binding modes of compounds 12, 13, and 19 inside 5-LOX's active site, A, C, and E: Their binding modes upon docking, B, D, and F: Their binding modes over 50 ns MDS. G and H: Binding modes of the co-crystallized ligands AA and NDGA

Figure 6

Binding modes of compounds 12, 13, and 19 inside 15-LOX's active site, A, C, and E: Their binding modes upon docking. B, D, and F: Their binding modes over 50 ns MDS. G and H: Binding mode of the co-crystallized ligands AA

Figure 7

RMSDs of compounds 12, 13 and 19 inside 5-LOX and 15-LOX's active sites over 50 ns of MDS

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