

1 Supplementary Materials

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3 **Table S1.** Secondary metabolites of the genus *Litophyton* from 1975 to July, 2023.

No.	Name	Class	Species	Locality	Bioassays	Ref.
1	(-)-bicyclogermacrene	bicyclogermacrene sesquiterpene	<i>Litophyton arboreum</i>	near Bali, Indonesia	antiproliferative activities against the L-929 and K-562 cell lines (GI ₅₀ 186±6 and 200±8 μM, respectively), cytotoxic effect against the HeLa cell line (CC ₅₀ 182±8 μM)	[24]
2	(2E,6E)-3-isopropyl-6-methyl-10-oxoundeca-2,6-dienal	sec-germacrane sesquiterpene	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 (IC ₅₀ 3.7 and 2.2 mg/mL, respectively), non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
3	alismol	guaiane sesquiterpene	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	inhibitory activity against HIV-1 protease receptor (IC ₅₀ 7.20±0.7 μM), cytotoxic activities against the HeLa and Vero cell lines (IC ₅₀ 30 and 49 μM, respectively)	[25]
			<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic effects against the HepG2, MDA and A549 cancer cell lines (IC ₅₀ 4.52, 7.02 and 9.23 μg/mL, respectively)	[26]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antibacterial activities against <i>Escherichia coli</i> ATCC 10536, <i>Pseudomonas aeruginosa</i> NTCC 6750, <i>Bacillus cereus</i> ATCC 9634, <i>Bacillus subtilis</i> ATCC6633, <i>Staphylococcus aureus</i> ATCC5141 (MIC 9.2, 10.4, 4.3, 5.0, and 9.2 μg/mL, respectively), antifungal activities against <i>Candida albicans</i> and <i>Aspergillus niger</i> (MIC 10.1 and 7.2 μg/mL, respectively), cytotoxic effects against MCF-7, HCT-116 and HepG2 cell lines (IC ₅₀ 44.52±0.5, 31.45±1.19, and 10.71±1.18 μM, respectively)	[27]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	a –	[17]
4	10-O-methyl alismoxide	guaiane sesquiterpene	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic activities against the HeLa, Vero and U937 cell lines (IC ₅₀ 38±0.7, 49.8±0.5 and 50±0.23 μM, respectively)	[25]

No.	Name	Class	Species	Locality	Bioassays	Ref.
5	alismoxide	guaiane sesquiterpene	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	inactive against the HeLa, Vero and U937 cell lines (all IC ₅₀ >100 μM)	[25]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell line (all IC ₅₀ >100 μmol/mL)	[28]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	a –	[17]
6	alismorientol B	guaiane sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antibacterial activities against <i>Escherichia coli</i> ATCC 10536, <i>Pseudomonas aeruginosa</i> NTCC 6750, <i>Bacillus cereus</i> ATCC 9634, <i>Bacillus subtilis</i> ATCC6633, <i>Staphylococcus aureus</i> ATCC5141 (MIC 3.4, 7.7, 1.3, 6.8, and 8.3 μg/mL, respectively), antifungal activities against <i>Candida albicans</i> and <i>Aspergillus niger</i> (MIC 6.0 and 8.4 μg/mL, respectively), cytotoxic effects against MCF-7, HCT-116 and HepG2 cell lines (IC ₅₀ 4.32±0.13, 27.86±1.28 and 12.27±1.09 μM, respectively)	[27]
7	litoarbolide A	guaiane sesquiterpene	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
8	4α,7β,10α-trihydroxyguai-5-ene	guaiane sesquiterpene	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
9	leptocladol B	guaiane sesquiterpene	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	anti-malarial activity against chloroquine-resistant <i>Plasmodium falciparum</i> W2 (IC ₅₀ 4.3 mg/mL), no anti-malarial effect against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration	[17]

No.	Name	Class	Species	Locality	Bioassays	Ref.
					of 4.76 mg/mL	
10	nephthetraol	guaiane sesquiterpene	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	anti-malarial activity against chloroquine-resistant <i>Plasmodium falciparum</i> W2 (IC ₅₀ 3.2 mg/mL), no anti-malarial effect against hloroquine-sensitive <i>Plasmodium falciparum</i> D6 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
11	litopharbol	pseudoguaiane sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antibacterial activities against <i>Escherichia coli</i> ATCC 10536, <i>Pseudomonas aeruginosa</i> NTCC 6750, <i>Bacillus cereus</i> ATCC 9634, <i>Bacillus subtilis</i> ATCC6633, <i>Staphylococcus aureus</i> ATCC5141 (MIC 9.0, 3.6, 1.8, 5.8, and 9.6 µg/mL, respectively), antifungal activities against <i>Candida albicans</i> and <i>Aspergillus niger</i> (MIC 12.5 and 12.9 µg/mL, respectively), cytotoxic effects against MCF-7, HCT-116 and HepG2 cell lines (IC ₅₀ 9.42±0.17, 26.21±2.249 and 38.92±1.12 µM, respectively)	[27]
12	litopharbdiol	pseudoguaiane sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a –	[18]
13	3α,6α-epidioxyhimachal-1-ene	himachalene sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabi (Red Sea)	^b antiproliferative effects toward MCF-7, HCT116 and HepG2 cell lines	[29]
14	5β,8β-epidioxy-11-hydroxy-6-eudesmene	eudesmane sesquiterpene	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	cytotoxic effect against A549 (IC ₅₀ 67.3±9.9 µmol/mL), inactive against the MCF-7 and HepG2 cell line (both IC ₅₀ >100 µmol/mL)	[28]
15	chabrolidione B	seco-eudesmane sesquiterpene	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell line (all IC ₅₀ >100 µmol/mL)	[28]
16	teuhetenone A	tri-nor-eudesmane sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antibacterial activities against <i>Escherichia coli</i> ATCC 10536, <i>Pseudomonas aeruginosa</i> NTCC 6750, <i>Bacillus cereus</i> ATCC 9634, <i>Bacillus subtilis</i> ATCC6633, <i>Staphylococcus aureus</i> ATCC5141 (MIC 1.9, 2.8, 5.4, 4.3, and 10.9 µg/mL, respectively), antifungal activities against <i>Candida albicans</i>	[27]

No.	Name	Class	Species	Locality	Bioassays	Ref.
					and <i>Aspergillus niger</i> (MIC 4.1 and 7.4 µg/mL, respectively), cytotoxic effect against and HepG2 cell line (IC ₅₀ 39.23±3.21 µM), non-toxic to the MCF-7 and HCT-116 cell lines (both IC ₅₀ >100 µM)	
17	calamusin I	tri-nor-eudesmane sesquiterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antibacterial activities against <i>Escherichia coli</i> ATCC 10536, <i>Pseudomonas aeruginosa</i> NTCC 6750, <i>Bacillus cereus</i> ATCC 9634, <i>Bacillus subtilis</i> ATCC6633, <i>Staphylococcus aureus</i> ATCC5141 (MIC 3.2, 1.2, 6.1, 7.8, and 7.4 µg/mL, respectively), antifungal activities against <i>Candida albicans</i> and <i>Aspergillus niger</i> (MIC 3.2 and 5.6 µg/mL, respectively), cytotoxic effects against MCF-7 and HepG2 cell lines (IC ₅₀ 6.43±0.23 and 13.46±1.16 µM, respectively), non-toxic to the HCT-116 cell line (IC ₅₀ >100 µM)	[27]
18	7-oxo-tri-nor-eudesm-5-en-4β-ol	tri-nor-eudesmane sesquiterpene	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell line (all IC ₅₀ >100 µmol/mL)	[28]
19	11,12-dihydroxy-6,10-eremophiladiene	eremophilane sesquiterpene	<i>Litophyton nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	a –	[23]
20	linardosinene B	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	cytotoxic effect against the THP-1 cell line (IC ₅₀ 59.49 µM)	[12]
21	linardosinene C	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	cytotoxic activities against the SNU-398 and HT-29 cell lines (IC ₅₀ 24.3 and 44.7 µM, respectively), inactive against the Capan-1 and A549 cell lines (both IC ₅₀ >50 µM)	[12]
22	linardosinene D	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inhibitory effect against BRD4 at a concentration of 10 µM with inhibitory rate 16.3%	[30]
23	linardosinene E	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inhibitory effect against BRD4 at a concentration of 10 µM with inhibitory rate 15.8%	[30]

No.	Name	Class	Species	Locality	Bioassays	Ref.
24	linardosinene F	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inhibitory effect against BRD4 at a concentration of 10 μ M with inhibitory rate 17.6%	[30]
25	linardosinene G	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inhibitory effect against BRD4 at a concentration of 10 μ M with inhibitory rate 18.1%	[30]
26	linardosinene I	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inhibitory activity against PTP1B (IC_{50} 10.67 \pm 1.81 μ g/mL), cytotoxic activities against the HT-29, Capan-1 and SNU-398 cell lines (IC_{50} 35.48 \pm 11.22, 42.55 \pm 2.51, and 25.17 \pm 11.13 μ M, respectively)	[23]
27	emnal-1(10)-ene-7 β ,12 α -diol	nardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inactive against PTP1B (IC_{50} >20 μ g/mL) and HT-29, Capan-1 and SNU-398 cell lines (all IC_{50} >50 μ M)	[23]
28	paralemnolin J	nardosinane sesquiterpenes	<i>Litophyton setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC_{50} >20 μ M)	[15]
29	(1S,8S,8aS)-1-[(E)-2'-acetoxyl-1'-methylethenyl]-8,8a-dimethyl-3,4,6,7,8,8a-hexahydronaphthalen-2(1H)-one	nardosinane sesquiterpenes	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC_{50} >20 μ M)	[15]
30	linardosinene A	nornardosinane sesquiterpenes	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inactive against the THP-1 cell line and the PTP1B, BRD4, HDAC1 and HDAC6 protein kinases	[12]
31	linardosinene H	eremophilane-nardosinane bis-sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inactive against PTP1B (IC_{50} >20 μ g/mL) and HT-29, Capan-1 and SNU-398 cell lines (all IC_{50} >50 μ M)	[23]
32	lineolemnene A	neolemnane sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	cytotoxic activity against the SNU-398 cell line (IC_{50} 44.4 μ M), inactive against the HT-29, Capan-1 and A549 cell lines (all IC_{50} >50 μ M)	[12]
33	lineolemnene B	neolemnane sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	cytotoxic activity against the SNU-398 cell line (IC_{50} 27.6 μ M), inactive against the HT-29, Capan-1 and A549 cell lines (all IC_{50} >50 μ M)	[12]

No.	Name	Class	Species	Locality	Bioassays	Ref.
34	lineolemnene C	neolemnane sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inactive against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >50 μM)	[12]
35	4-acetoxy-2,8-neolemnadien-5-one	neolemnane sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	a ₋	[12]
			<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 μM)	[15]
36	paralemnolin E	neolemnane sesquiterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 μM)	[15]
37	lineolemnene D	seconeolemnane sesquiterpene	<i>L. nigrum</i>	Xisha Islands, Hainan, China (South China Sea)	inactive against the THP-1 cell line and the PTP1B, BRD4, HDAC1 and HDAC6 protein kinases	[12]
38	nephtenol	cembrane diterpene	<i>Litophyton viridis</i>	Serwaru, Leti Island, Indonesia	a ₋	[19]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a ₋	[31]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a ₋	[18]
39	2-hydroxynephtenol	cembrane diterpene	<i>L. viridis</i>	Serwaru, Leti Island, Indonesia	a ₋	[19]
40	(3 <i>E</i> ,11 <i>E</i>)-cembra-3,8(19),11,15-tetraene-7α-ol	cembrane diterpene	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a ₋	[31]
41	sarcophytol M	cembrane diterpene	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	inhibitory activity against HIV-1 protease receptor (IC ₅₀ 15.7±0.10 μM), cytotoxic activities against the HeLa, Vero and U937 cell lines (IC ₅₀ 27.5±0.2, 22±0.2 and 31.7±3.2 μM, respectively)	[25]
42	11-acetoxy-15,17-dihydroxy-2,12-epoxy-(3 <i>E</i> ,7 <i>E</i>)-1-cembra-3,7-diene	cembrane diterpene	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	antiproliferative effects toward MCF-7, HCT116 and HepG2 cell lines (IC ₅₀ 19.1±0.032, 22.0±0.092, 24.0±0.032 μM, respectively)	[29]

No.	Name	Class	Species	Locality	Bioassays	Ref.
43	11 β ,12 β -epoxypukalide	cembrane diterpene	<i>L. arboreum</i>	near Bali, Indonesia	antiproliferative activities against the L-929 and K-5629 cell lines (both GI ₅₀ >129 μ M, respectively), cytotoxic effect against the HeLa cell line (CC ₅₀ 115 \pm 6 μ M)	[24]
44	litophynin A	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	insect growth inhibitory against the silkworm <i>Bombyx mori</i> L. (ED ₅₀ 12 ppm)	[32]
45	litophynin B	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	insect growth inhibitory against the silkworm <i>Bombyx mori</i> L. (ED ₅₀ 2.7 ppm)	[32]
46	litophynin C	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	insect growth inhibitory against the second instar larvae of the silkworm <i>Bombyx mori</i> L. (ED ₅₀ 25 ppm)	[33]
47	litophynin D	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	significant brine shrimp lethality (LD ₅₀ 0.9 ppm)	[34]
48	litophynin E	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	^a –	[34]
			<i>Litophyton</i> sp.	Nango-cho, Miyazaki Prefecture, Japan	^b positive in a hemolytic reaction test	[37]
49	litophynin F	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	inactive against the second instar larvae of the silkworm <i>Bombyx mori</i> L. at 100 ppm)	[35]
			<i>Litophyton viscudium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 18 μ M)	[16]
50	litophynin G	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	insect growth inhibitory against the second instar larvae of the silkworm <i>Bombyx mori</i> L. (ED ₅₀ 42 ppm)	[35]
51	litophynin H	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	inactive against the second instar larvae of the silkworm <i>Bombyx mori</i> L. at 100 ppm	[35]
			<i>Litophyton</i> sp.	Nango-cho, Miyazaki Prefecture, Japan	^b positive in a hemolytic reaction test	[37]
52	litophynin I	eunicellane diterpene	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	significant molluscicidal activity against the muricid gastropod <i>Drupella fragum</i> with 100% mortality within 24 hours at 30 ppm, repellent activity against the gastropod when impregnated on filterpaper by 45 μ g/cm ² .	[36]
53	litophynin J	eunicellane	<i>Litophyton</i> sp.	Sukumo Bay, Kochi	significant molluscicidal activity against the muricid	[36]

No.	Name	Class	Species	Locality	Bioassays	Ref.
		diterpene		Prefecture, Japan	gastropod <i>Drupella fragum</i> with 100% mortality within 24 hours at 30 ppm, repellent activity against the gastropod when impregnated on filterpaper by 45 µg/cm ² .	
54	litophynol A	eunicellane diterpene	<i>Litophyton</i> sp.	Nango-cho, Miyazaki Prefecture, Japan	^b positive in a hemolytic reaction test	[37]
55	litophynol B	eunicellane diterpene	<i>Litophyton</i> sp.	Nango-cho, Miyazaki Prefecture, Japan	^b positive in a hemolytic reaction test	[37]
56	litophynin I monoacetate	eunicellane diterpene	<i>Litophyton</i> sp.	Nango-cho, Miyazaki Prefecture, Japan	^b positive in a hemolytic reaction test	[37]
57	^c 6-oxo litophynin H	eunicellane diterpene	<i>L. viscidium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 20 µM)	[16]
58	^c 6-oxo litophynin H 12-acetate	eunicellane diterpene	<i>L. viscidium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 20 µM)	[16]
59	^c 6-oxo litophynol A	eunicellane diterpene	<i>L. viscidium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 5.7 µM)	[16]
60	^c 6- <i>epi</i> litophynol A	eunicellane diterpene	<i>L. viscidium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 4.2 µM)	[16]
61	^c 6-methyl litophynin E	eunicellane diterpene	<i>L. viscidium</i>	Otsuki Town, Kochi Prefecture, Japan	cytotoxic activity against the HL-60 cell line (IC ₅₀ 50 µM)	[16]
62	lemnabourside	serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 µM)	[15]
63	biflora-4,9,15-triene	serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 µM)	[15]
64	litosetoenin A	5,9-cyclized serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 µM)	[15]
65	litosetoenin B	5,9-cyclized serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 µM)	[15]
66	litosetoenin C	5,9-cyclized	<i>L. setoensis</i>	Singaraja, Bali Island,	weak cytotoxic activities against the HT-29, Capan-1, A549,	[15]

No.	Name	Class	Species	Locality	Bioassays	Ref.
		serrulatane diterpene		Indonesia	and SNU-398 cell lines (all IC ₅₀ >20 μM)	
67	litosetoenin D	5,9-cyclized serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 μM)	[15]
68	litosetoenin E	5,9-cyclized serrulatane diterpene	<i>L. setoensis</i>	Singaraja, Bali Island, Indonesia	weak cytotoxic activities against the HT-29, Capan-1, A549, and SNU-398 cell lines (all IC ₅₀ >20 μM)	[15]
69	all- <i>trans</i> -peridinin	tetraterpene	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	antiproliferative activities against the HUVEC and K-562 cell lines (GI ₅₀ 48.4±1.2 and 53.8±3.9 μM, respectively), cytotoxicity against the HeLa cell line (IC ₅₀ 51.9±0.7 μM)	[31]
70	4α-methyl-3β,8β-dihydroxy-5α-ergost-24(28)-en-23-one	4α-methylated steroid	<i>L. viridis</i>	Lesser Sunda Islands, Indonesia	a –	[40]
71	4α,24-dimethyl-5α-cholest-24(28)-en-3β,8β,18-triol	4α-methylated steroid	<i>Litophyton mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activities against the K562 and A549 cell lines (IC ₅₀ 8.9±0.9 and 25.7±1.5 μM, respectively)	[41]
72	(22 <i>E</i> ,24 <i>R</i>)-4α,24-dimethyl-5α-cholest-22-en-3β,8β,18-triol	4α-methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	inactive against the K562 and A549 cell lines (both IC ₅₀ >50 μM)	[41]
73	(22 <i>E</i>)-4α,24-dimethyl-5α-cholesta-2,24(28)-dien-3β,8β,18-triol	4α-methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	inactive against the K562 and A549 cell lines (both IC ₅₀ >50 μM)	[41]
74	nebrosteroid M	4α-methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activity against the A549 cell line (IC ₅₀ 20.4±1.1 μM), inactive against the K562 cell line (IC ₅₀ >50 μM)	[41]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	cytotoxic activity against the A549 and MCF-7 cell line (IC ₅₀ 36.9±2.9 and 55.3±4.9 μmol/mL, respectively), inactive against the HepG2 cell line (IC ₅₀ >100 μmol/mL)	[28]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]

No.	Name	Class	Species	Locality	Bioassays	Ref.
75	(22E,24R)-4 α ,24-dimethyl-5 α -cholest-22-en-3 β ,8 β ,11 β -triol	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activities against the K562 and A549 cell lines (IC ₅₀ 7.7 \pm 0.8 and 20.8 \pm 1.2 μ M, respectively)	[41]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
76	nebrosteroid D	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activities against the K562 and A549 cell lines (IC ₅₀ 6.0 \pm 0.5 and 22.1 \pm 1.4 μ M, respectively)	[41]
77	nebrosteroid A	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activity against the K562 cell line (IC ₅₀ 5.8 \pm 0.8 μ M)	[41]
78	23 ξ -acetoxy-4 α ,24-dimethyl-5 α -cholest-24(28)-en-3 β ,8 β ,11 β -triol	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	cytotoxic activity against the K562 cell line (IC ₅₀ 5.6 \pm 1.2 μ M)	[41]
79	(22E,24R)-4 α ,23,24-trimethyl-5 α -cholest-22-en-3 β ,8 β ,11 β -triol	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	inactive against the K562 and A549 cell lines (both IC ₅₀ >50 μ M)	[41]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial activity against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 and chloroquine-resistant <i>Plasmodium falciparum</i> W2 at the concentration of 4.76 mg/mL, non-toxic to the Vero cell line at the concentration of 4.76 mg/mL	[17]
80	(22Z)-4 α ,24 ξ -dimethyl-5 α -cholest-22-en-3 β ,8 β ,11 β -triol	4 α -methylated steroid	<i>L. mollis</i>	Hurghada, Egypt (Red Sea)	a-	[41]
81	4 α ,24-dimethyl-cholest-22Z-en-3 β -ol	4 α -methylated steroid	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a-	[45]
82	24-methylenecholest-5-en-3 β ,7 β ,19-triol	steroid	<i>L. viridis</i>	Serwaru, Leti Island, Indonesia	a-	[39]
			<i>L. arboreum</i>	Xisha Islands, Hainan, China (South China Sea)	a-	[48]
			<i>L. arboreum</i>	Sharm El-Sheikh, Egypt	cytotoxic activities against the HeLa, Vero and U937 cell	[25]

No.	Name	Class	Species	Locality	Bioassays	Ref.
				(Red Sea)	lines (IC ₅₀ 8±0.5, 11.4±0.04 and 16.4±1.25 µM, respectively)	
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a –	[45]
			<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic effects against the HepG2, MDA and A549 cancer cell lines (IC ₅₀ 8.5, 5.5 and 9.3 µg/mL, respectively)	[26]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a –	[18]
83	7-acetoxy-24-methylenecholest-5-en-3β,7β,19-triol	steroid	<i>L. viridis</i>	Serwaru, Leti Island, Indonesia	a –	[39]
			<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	inhibitory activity against HIV-1 protease receptor (IC ₅₀ 4.85±0.18 µM), cytotoxic activities against the HeLa, Vero and U937 cell lines (IC ₅₀ 5.3±0.60, 31.3±14.03 and 10.6±0.12 µM, respectively)	[25]
			<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic effects against the HepG2, MDA and A549 cancer cell lines (IC ₅₀ 6.07, 6.3 and 6.2 µg/mL, respectively)	[26]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	a –	[17]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	^b antiproliferative effects toward MCF-7, HCT116 and HepG2 cell lines	[29]
84	litosterol	steroid	<i>L. viridis</i>	Ishiqaki Isoland, Okinawa, Japan	a –	[47]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a –	[45]
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	a –	[17]
85	5,6-epoxylitosterol	steroid	<i>L. viridis</i>	Ishiqaki Isoland, Okinawa, Japan	antileukemic activity against the P388 cell line (IC ₅₀ 0.5 µg/mL)	[47]
86	(24E)-24-ethyl-5α-cholesta-8,24(28)-diene-3β,12β,19-triol	steroid	<i>L. arboreum</i>	Xisha Islands, Hainan, China (South China Sea)	a –	[48]

No.	Name	Class	Species	Locality	Bioassays	Ref.
87	24-methylcholesta-5,24(28)-diene-3- β -ol	steroid	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic activities against the HeLa and Vero cell lines (IC ₅₀ 48 \pm 8.7 and 100 \pm 1.2 μ M, respectively), inactive against the U937 cell line (IC ₅₀ >100 μ M)	[25]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a –	[45]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a –	[18]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	cytotoxic activities against the A549, MCF-7 and HepG2 cell line (IC ₅₀ 28.5 \pm 4.4, 70.0 \pm 2.3 and 77.6 \pm 3.3 μ mol/mL, respectively)	[28]
88	3 β ,7 β -dihydroxy-24-methylenecholesterol	steroid	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	anti-malarial activity against chloroquine-resistant <i>Plasmodium falciparum</i> W2 (IC ₅₀ 4.0 mg/mL), no anti-malarial effect against chloroquine-sensitive <i>Plasmodium falciparum</i> D6 at the concentration of 4.76 mg/mL	[17]
89	chabrolosteroid I	steroid	<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	no anti-malarial effect against chloroquine-resistant <i>Plasmodium falciparum</i> W2 and chloroquine-sensitive <i>Plasmodium falciparum</i> D6 at the concentration of 4.76 mg/mL	[17]
90	13,14- <i>seco</i> -22-norergosta-4,24(28)-dien-19-hydroperoxide-3-one	steroid	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	^b antiproliferative effects toward MCF-7, HCT116 and HepG2 cell lines	[29]
91	sarcsteroid F	steroid	<i>Litophyton</i> sp.	Egyptian Red Sea	a –	[21]
92	24-methylenecholestane-1 α ,3 β ,5 α ,6 β ,11 α -pentol-11-monoacetate	steroid	<i>Litophyton</i> sp.	Egyptian Red Sea	a –	[21]
93	erythro- <i>N</i> -dodecanoyl-docosasphingina-(4 <i>E</i> ,8 <i>E</i>)-dienine	ceramide	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic activity against the HeLa cell line (IC ₅₀ 38.17 \pm 0.7 μ M), inactive against the Vero and U937 cell lines (both IC ₅₀ >100 μ M)	[25]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	^b antiproliferative effects toward MCF-7, HCT116 and HepG2 cell lines	[29]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia	a –	[27]

No.	Name	Class	Species	Locality	Bioassays	Ref.
				(Red Sea)		
			<i>L. arboreum</i>	Neweba, Egypt (Red Sea)	a –	[17]
94	erythro- <i>N</i> -palmityl-octadecasphing-4(<i>E</i>),8(<i>E</i>)-dienin	ceramide	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	a –	[45]
			<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a –	[27]
95	thymine	nucleotide	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell lines (all IC ₅₀ >100 µg/mL), inactive against against <i>Leishmania major</i> (IC ₅₀ >100 µg/mL)	[49]
96	thymidine	nucleotide	<i>L. arboreum</i>	Jeddah, Saudi Arabia (Red Sea)	a –	[27]
97	uracil	nucleotide	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell lines (all IC ₅₀ >100 µg/mL), inactive against against <i>Leishmania major</i> (IC ₅₀ >100 µg/mL)	[49]
98	uridine	nucleotide	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell lines (all IC ₅₀ >100 µg/mL), inactive against against <i>Leishmania major</i> (IC ₅₀ >100 µg/mL)	[49]
99	PGB ₂ methyl ester	prostaglandin	<i>L. arboreum</i>	gulf of Aqaba, Eilat, Israel (Red Sea)	a –	[50]
100	litophytolide A	γ-lactone	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	a –	[51]
101	litophytolide B	γ-lactone	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	a –	[51]
102	^c (<i>R</i>)-5-((<i>S</i> ,2 <i>Z</i> ,4 <i>E</i>)-1-hydroxydeca-2,4-dien-1-yl)dihydrofuran-2(3 <i>H</i>)-one	γ-lactone	<i>L. arboreum</i>	gulf of Aqaba, Eilat, Israel (Red Sea)	toxic to brine shrimp <i>Artemia salina</i> (CC ₅₀ 15.3 µg/mL), antibacterial activities against <i>Staphylococcus aureus</i> (diameters of inhibition zone 7.8 mm) and <i>Bacillus subtilis</i> (diameters of inhibition zone 5.6 mm), inactive against <i>Escherichia coli</i> and <i>Saccharomyces cerevisiae</i>	[50]
103	^c	γ-lactone	<i>L. arboreum</i>	gulf of Aqaba, Eilat,	toxic to brine shrimp <i>Artemia salina</i> (CC ₅₀ 21.4 µg/mL),	[50]

No.	Name	Class	Species	Locality	Bioassays	Ref.
	(R)-5-((S,4E,8E)-1-hydroxy-4,8-dimethyltrideca-4,8-dien-1-yl)-5-methylidihydrofuran-2(3H)-one			Israel (Red Sea)	antibacterial activities against <i>Staphylococcus aureus</i> (diameters of inhibition zone 18.6 mm) and <i>Bacillus subtilis</i> (diameters of inhibition zone 14.7 mm), inactive against <i>Escherichia coli</i> and <i>Saccharomyces cerevisiae</i>	
104	methyl (5Z,8Z,11Z,14Z,17Z)-5,8,11,14,17-icosapentaenoate	fatty acid	<i>Litophyton</i> sp.	Sukumo Bay, Kochi Prefecture, Japan	a-	[51]
105	arachidonic acid	fatty acid	<i>L. arboreum</i>	gulf of Aqaba, Eilat, Israel (Red Sea)	a-	[50]
106	eicosapentaenoic acid	fatty acid	<i>L. arboreum</i>	gulf of Aqaba, Eilat, Israel (Red Sea)	a-	[50]
107	docosahexaenoic acid	fatty acid	<i>L. arboreum</i>	gulf of Aqaba, Eilat, Israel (Red Sea)	a-	[50]
108	chimyl alcohol	glycerol ether	<i>L. arboreum</i>	Sharm El-Sheikh, Egypt (Red Sea)	cytotoxic activity against the HeLa and Vero cell line (IC ₅₀ 23.35±5.8 and 60±1.14 μM, respectively), inactive against the U937 cell lines (both IC ₅₀ >100 μM), inhibitory activity against HIV-1 protease receptor (IC ₅₀ 26.6±2.6 μM)	[25]
			<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell lines (all IC ₅₀ >100 μmol/mL)	[28]
109	batyl alcohol	glycerol ether	<i>L. arboreum</i>	Hurghada, Egypt (Red Sea)	inactive against the A549, MCF-7 and HepG2 cell lines (all IC ₅₀ >100 μmol/mL)	[28]

4 ^a The sign ‘-’ indicated no bioassay was recorded in the work. ^b No specific data was provided in the paper. ^c Compound was named based on its structure.