New Anti-Inflammatory Cembranes from the Cultured Soft Coral *Nephthea columnaris*

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Abstract: Two new cembranes, columnariols A (1) and B (2), were isolated from the cultured soft coral *Nephthea columnaris*. The structures of cembranes 1 and 2 were elucidated by spectroscopic methods. In the anti-inflammatory effects test, cembranes 1
and 2 were found to significantly inhibit the accumulation of the pro-inflammatory iNOS and COX-2 protein of the lipopolysaccharide (LPS)-stimulated RAW264.7 macrophage cells. Compound 1 exhibited moderate cytotoxicity toward LNCaP cells with an IC$_{50}$ value of 9.80 $\mu$g/mL.

**Keywords:** Nephthea columnaris; cembrane; octocoral; antiinflammatory; iNOS; COX-2; cytotoxicity

1. Introduction

Among the natural products isolated from soft corals, the cembranes are major representative compounds. Previous investigations of soft corals belonging to the genus *Nephthea* (family Nephtheidae), collected off the waters of Taiwan, have yielded several interesting cembranes [1,2]. In our continuing studies, a sample originally collected off the coast of Southern Taiwan, identified as *Nephthea columnaris* (Studer, 1895), yielded two new cembranes, columnariols A (1) and B (2) (Scheme 1). In this paper, we deal with the isolation, structure determination and bioactivity of cembranes 1 and 2.

![Nephthea columnaris](image)

**Scheme 1.** The soft coral *Nephthea columnaris* and the structures of columnariols A (1) and B (2).

2. Results and Discussion

Columnariol A (1) was isolated as a colorless oil and the molecular formula for this compound was determined to be C$_{20}$H$_{34}$O$_{3}$ (four unsaturations) using HRESIMS (C$_{20}$H$_{34}$O$_{3}$ + Na, m/z 345.23989, calcld 345.24002). Comparison of the $^{13}$C-NMR and DEPT data with the molecular formula indicated there must be two exchangeable protons, which required the presence of two hydroxyl groups. This deduction was supported by a broad absorption in the IR spectrum at 3419 cm$^{-1}$. The $^{13}$C-NMR data for 1 confirmed the presence of 20 carbon signals (Table 1), characterized by DEPT spectrum as five methyls, six sp$^{3}$ methylenes, three sp$^{3}$ methines (including two oxymethines), two oxygenated sp$^{3}$ quaternary carbons, two sp$^{2}$ methines and two sp$^{2}$ quaternary carbons. Based on the $^{1}$H and $^{13}$C-NMR spectra (Table 1), 1 was determined to contain two isolated methyl-bearing trisubstituted double bonds ($\delta$H 5.48, 1H, d, $J$ = 10.8 Hz, H-3; 5.02, 1 H, dd, $J$ = 10.8, 4.4 Hz, H-7; 1.69, 3 H, s, H$_{3}$-18; 1.60, 3 H, s, H$_{3}$-19; $\delta$C 126.4, CH-3; 139.1, C-4; 123.8, CH-7; 136.2, C-8; 15.3, CH$_{3}$-18; 17.3, CH$_{3}$-19). Thus, from the reported data, the proposed skeleton of 1 was suggested to be a cembrane with two rings.
Table 1. $^1$H (400 MHz, CDCl$_3$) and $^{13}$C (100 MHz, CDCl$_3$) NMR data, $^1$H–$^1$H COSY and HMBC correlations for cembrane 1.

| Position | $\delta$H ($J$ in Hz) | $\delta$C, Multiple | $^1$H–$^1$H COSY | HMBC |
|----------|------------------------|----------------------|------------------|-------|
| 1        | 1.79 m                 | 49.9, CH             | H-2, H$_2$-14    | C-2, -3 |
| 2        | 4.65 dd (10.8, 4.4)    | 70.1, CH             | H-1, H-3         | C-1, -3, -4, -12, -14 |
| 3        | 5.48 d (10.8)          | 126.4, CH            | H-2, H$_3$-18    | C-2, -5, -18 |
| 5        | 2.20 ddd (12.4, 3.6, 3.6); 1.98 m | 40.0, CH$_2$ | H-2, H-6 | C-3, -4, -6, -7, -18 |
| 6        | 2.31 ddd (14.0, 9.6, 3.6); 2.15 m | 25.1, CH$_2$ | H$_2$-5, H-7 | C-4, -7, -8 |
| 7        | 5.02 dd (10.8, 4.4)    | 123.8, CH            | H$_2$-6, H$_3$-19 | C-9, -19 |
| 8        |                        | 136.2, C             |                  |       |
| 9        | 2.16 m; 1.99 m         | 35.3, CH$_2$         | H$_2$-10         | C-7, -8, -10, -11, -19 |
| 10       | 1.82 m; 1.29 m         | 31.6, CH$_2$         | H$_2$-9, H-11    | C-8, -9, -11, -12 |
| 11       | 3.84 d (9.2)           | 72.4, CH             | H$_2$-10         | C-9, -12, -20 |
| 13       | 2.46 ddd (14.0, 3.6, 3.6); 1.39 m | 33.7, CH$_2$ | H$_2$-14 | C-1, -11, -12, -14, -20 |
| 14       | 1.75 m; 1.66 m         | 16.0, CH$_2$         | H-1, H$_2$-13    | C-1, -2, -12, -13 |
| 15       |                        | 71.5, C              |                  |       |
| 16       | 1.17 s                 | 28.3, CH$_3$         | C-1, -15, -17    |       |
| 17       | 1.12 s                 | 27.8, CH$_3$         | C-1, -15, -16    |       |
| 18       | 1.69 s                 | 15.3, CH$_3$         | H-3              | C-3, -4, -5 |
| 19       | 1.60 s                 | 17.3, CH$_3$         | H-7              | C-7, -8, -9 |
| 20       | 1.04 s                 | 24.0, CH$_3$         |                  | C-11, -12, -13 |

From the $^1$H–$^1$H COSY spectrum of 1 (Table 1), it was possible to differentiate among the separate spin systems of H-3/H-2/H-1/H$_2$-14/H$_2$-13, H$_2$-5/H$_2$-6/H-7 and H$_2$-9/H$_2$-10/H-11. These data, together with the HMBC correlations between H-1/C-2, -3; H-2/C-1, -3, -4, -14; H-3/C-2, -5; H$_2$-5/C-3, -4, -6, -7; H$_2$-6/C-4, -7, -8; H-7/C-9; H$_2$-9/C-7, -8, -10, -11; H$_2$-10/C-8, -9, -11, -12; H-11/C-9, -12; H$_2$-13/C-1, -11, -12, -14; and H$_2$-14/C-1, -2, -12, -13, observed in an HMBC experiment, established the connectivity from C-1 to C-14 in a 14-membered ring. The vinyl methyls attached at C-4 and C-8 were confirmed by the HMBC correlations between H-3, H$_2$-5/C-18; H$_3$-18/C-3, -4, -5; and H-7, H$_2$-9/C-19; H$_3$-19/C-7, -8, -9, and were further supported by the allylic couplings between H-3/H$_2$-18 and H-7/H$_3$-19. An isopropyl carbinol group at C-1 was elucidated by the HMBC correlations between H$_3$-16/C-1, -15, -17 and H$_3$-17/C-1, -15, -16. The HMBC correlations between H-2 ($\delta$H 4.65) and C-12, an oxygenated quaternary carbon at $\delta$C 75.3; and H$_3$-20/C-11, -12, -13, suggested an oxygen atom had to be positioned between C-2 and C-12 to form an ether bridge. Thus, the remaining hydroxyl group was positioned at C-11, an oxygen-bearing methine.

As those of the cembranoid derivatives isolated from Formosan soft coral [3], the relative configuration of compound 1 was deduced from NOESY correlations and Chem3D Ultra 9.0 (Figure 1). If H-1 assumes a $\beta$-orientation and this proton correlated with H-2, but not with H-3, and a large coupling constant was recorded between H-2 and H-3 ($J = 10.8$ Hz), indicated that the dihedral angle between H-2 and H-3 is approximately 180° and the configurations of both chiral carbons C-1 and C-2 were assigned as $R^*$. Furthermore, H-11 correlated with H-3, but not with H$_3$-20, indicating that the chiral carbons C-11 and C-12 can be assigned as $S^*$ and $R^*$, respectively. No correlation was found between
H-3/H3-18 and H-7/H3-19, indicating that C-3/4 and C-7/8 carbon-carbon double bonds had an E-configuration, by modeling analysis.

**Figure 1.** Key NOESY correlations of compound 1.

Cembrane 2 (columnariol B), obtained as a colorless oil, showed an [M + Na]+ signal at m/z 345.23977 in the HRESIMS, suggesting the molecular formula C20H34O3 (calcd C20H34O3 + Na, 345.24002), with four units of unsaturation. The IR absorption of 2 at 3355 cm⁻¹ indicated the presence of hydroxyl functionality. ¹H and ¹³C-NMR spectral data (Table 2) showed that the structure of 2 contained an isopropyl carbinol, two isolated methyl-bearing trisubstituted olefins, an exocyclic olefin, an aliphatic methine carbon, two oxymethine carbons and six methylene carbons. These data suggested that 2 possessed a cembrane skeleton with functionalities of an isopropyl carbinol, two isolated methyl-bearing trisubstituted double bonds, an exocyclic double bond and two hydroxy groups. ¹H–¹H couplings in the COSY spectrum of 2 enabled identification of the C-3/-2/-1/-14/-13, C-5/-6/-7, C-9/-10/-11, C-3/-18 (by allylic coupling) and C-11/C-20 (by allylic coupling) units (Table 2), which were assembled with the assistance of an HMBC experiment. In the HMBC experiment of 2, the isopropyl carbinol group at C-1 was confirmed by correlations between H3-16/C-1, -15, -17 and H3-17/C-1, -15, -16. The vinyl methyl groups attached at C-4 and C-8 were confirmed by correlations between H-3/C-18; H3-18/ C-3, -4, -5; and H-7, H2-9/C-19, H3-19/C-7, -8, -9, respectively. An exocyclic carbon-carbon double bond at C-12 was confirmed by the correlations between H2-13/C-20 and H2-20/C-11, -12, -13. Thus, the remaining hydroxyl groups had to be positioned at C-2 and C-11, the oxymethines at δC 71.7 and 70.8. Based on the above findings, the planar structure of 2 was established. The 2D NMR correlations observed fully supported the locations of functional groups (Table 2).

In the NOESY experiment of 2 (Figure 2), H-1 correlated with H-3, but not with H-2, and large coupling constants were recorded between H-1/H-2 (J = 10.0 Hz) and H-2/H-3 (J = 9.2 Hz) and indicating that the dihedral angles between H-1/H-2 and H-2/H-3 are approximately 180° and the configurations of both chiral carbons C-1 and C-2 were assigned as R*. H-11 correlated with H3-18, but not with H2-20 confirming the S*-configuration of C-11 by modeling analysis. Furthermore, no correlation was found between H-3/H3-18 and H-7/H3-19, indicating that C-3/4 and C-7/8 carbon-carbon double bonds had an E-configuration.
Table 2. $^1$H (400 MHz, CDCl$_3$) and $^{13}$C (100 MHz, CDCl$_3$) NMR data, $^1$H–$^1$H COSY and HMBC correlations for cembrane 2.

| Position | $\delta_H$ (J in Hz) | $\delta_C$, Multiple | $^1$H–$^1$H COSY | HMBC |
|----------|----------------------|----------------------|-------------------|-------|
| 1        | 1.54 m               | 51.9, CH             | H-2, H$_2$-14     |       |
| 2        | 4.43 dd (10.0, 9.2)  | 71.7, CH             | H-1, H-3          | C-4   |
| 3        | 5.21 d (9.2)         | 128.3, CH            | H-2, H$_3$-18     | C-5, -18 |
| 4        |                      | 140.0, C             |                   |       |
| 5        | 2.26 m; 2.07 m       | 40.0, CH$_2$         | H$_2$-6           | C-3, -4, -7 |
| 6        | 2.26 m; 2.17 m       | 24.5, CH$_2$         | H$_2$-5, H-7      | C-4, -5, -7, -8 |
| 7        | 5.02 dd (6.8, 6.8)   | 124.5, CH            | H$_2$-6, H$_3$-19 | C-9, -19 |
| 8        |                      | 134.6, C             |                   |       |
| 9        | 2.18 m; 2.03 m       | 33.9, CH$_2$         | H$_2$-10          | C-7, -8, -11, -19 |
| 10       | 1.75 m; 1.53 m       | 32.4, CH$_2$         | H$_2$-9           | C-11  |
| 11       | 4.08 dd (7.2, 4.0)   | 70.8, CH             | H$_2$-10, H$_2$-20a | C-12  |
| 12       |                      | 152.0, C             |                   |       |
| 13       | 2.17–2.03 m          | 34.9, CH$_2$         | H$_2$-14, H$_2$-20 | C-1, -11, -12, -14, -20 |
| 14       | 1.18 m; 1.07 m       | 27.4, CH$_2$         | H-1, H$_2$-13     | C-1, -2, -12, -13, -15 |
| 15       |                      | 75.0, C              |                   |       |
| 16       | 1.29 s               | 30.3, CH$_3$         |                   | C-1, -15, -17 |
| 17       | 1.31 s               | 24.2, CH$_3$         |                   | C-1, -15, -16 |
| 18       | 1.73 s               | 15.7, CH$_3$         | H-3               | C-3, -4, -5 |
| 19       | 1.58 s               | 16.2, CH$_3$         | H-7               | C-7, -8, -9 |
| 20a/b    | 5.11 s; 4.85 s       | 109.6, CH$_2$        | H$_2$-11, H$_2$-13 | C-11, -12, -13 |

Figure 2. NOESY correlations of compound 2.

In the *in vitro* anti-inflammatory activity test, the upregulation of the pro-inflammatory iNOS (inducible nitric oxide synthase) and COX-2 (cyclooxygenase-2) protein expression of LPS (lipopolysaccharide)-stimulated RAW264.7 macrophage cells was evaluated using immunoblot analysis. At a concentration of 50 $\mu$M, compounds 1 and 2 were found to significantly reduce the levels of iNOS and COX-2, respectively, relative to the control cells stimulated with LPS only (Figure 3). To evaluate the cytotoxic effects of compounds 1 and 2 on the viability of RAW264.7 macrophage cells, we used the alamar blue assay. Both compounds 1 and 2 (20 and 50 $\mu$M) did not significantly affect the viability of macrophage cells 16h after treatment. Thus, compounds 1 and 2 might be promising as anti-inflammatory agents, as they do not exhibit cytotoxicity to RAW264.7 macrophage cells.
Figure 3. Effects of compounds 1 and 2 on pro-inflammatory inducible nitric oxide synthase (iNOS) and cyclooxygenase-2 (COX-2) protein expression in the lipopolysaccharide (LPS)-stimulated murine macrophage cell line, RAW264.7. (A) Relative density of iNOS immunoblot; (B) relative density of COX-2 immunoblot. The relative intensity of the LPS-stimulated group was taken to be 100%. Band intensities were quantified by densitometry and are indicated as the percent change relative to that of the LPS-stimulated group. Compounds 1, 2 and dexamethasone (Dex) significantly inhibited LPS-induced iNOS protein expression in macrophages. The experiment was repeated three times (* p < 0.05, significantly different from the LPS-stimulated group).

Table 3. Cytotoxic Data of Cembranes 1 and 2.

| Compounds | Cell Lines IC50 (μg/mL) |
|-----------|-------------------------|
|           | MOLT-4 | SUP-T1 | DLD-1 | LNCap | MCF7 |
| 1         | NA     | NA     | NA    | 9.80  | NA   |
| 2         | NA     | NA     | NA    | NA    | NA   |
| Doxorubicin* | 0.001  | 0.19   | 0.08  | 2.68  | 1.39 |

*Doxorubicin was used as a positive control and NA = not active at 20 μg/mL for 72 h.
3. Experimental Section

3.1. General Experimental Procedures

Optical rotation values were measured with a Jasco P-1010 digital polarimeter (Japan Spectroscopic Corporation, Tokyo, Japan). IR spectra were obtained on a Varian Digilab FTS 1000 FT-IR spectrophotometer (Varian Inc., Palo Alto, CA, USA); absorptions are reported in cm$^{-1}$. NMR spectra were recorded on a Varian Mercury Plus 400 NMR spectrometer (Varian Inc., Palo Alto, CA, USA) using the residual solvent (CDCl$_3$, $\delta_H$ 7.26 ppm for $^1$H NMR and $\delta_C$ 77.1 ppm for $^{13}$C NMR) as the internal standard. Coupling constants ($J$) are given in Hz. ESIMS and HRESIMS were recorded using a Bruker 7 Tesla solariX FTMS system (Bruker, Bremen, Germany). Column chromatography was performed on silica gel (230–400 mesh, Merck, Darmstadt, Germany). TLC was carried out on precoated Kieselgel 60 F$_{254}$ (0.25 mm, Merck, Darmstadt, Germany); spots were visualized by spraying with 10% H$_2$SO$_4$ solution followed by heating. Normal phase HPLC (NP-HPLC) was performed using a system comprised of a Hitachi L-7110 pump (Hitachi Ltd., Tokyo, Japan), a Hitachi L-7455 photodiode array detector (Hitachi Ltd., Tokyo, Japan) and a Rheodyne 7725 injection port (Rheodyne LLC, Rohnert Park, CA, USA). A normal phase column (Supelco Ascentis® Si Cat #: 581515-U, 25 cm $\times$ 21.2 mm, 5 $\mu$m, Sigma-Aldrich, St. Louis, MO, USA) was used for HPLC. The reversed phase HPLC (RP-HPLC) was performed using a system comprised of a Hitachi L-2130 pump (Hitachi Ltd., Tokyo, Japan), a Hitachi L-2455 photodiode array detector (Hitachi Ltd., Tokyo, Japan), a Rheodyne 7725 injection port (Rheodyne LLC, Rohnert Park, CA, USA). A reversed phase column (Supelco Ascentis® Si Cat #: 581343-U, 25 cm $\times$ 10.0 mm, 5 $\mu$m, Sigma-Aldrich, St. Louis, MO, USA) was used for RP-HPLC.

3.2. Animal Material

Specimens of the octocoral *Nephthea columnaris* (Studer, 1895) were collected originally by hand using SCUBA equipment off the coast of the Southern Taiwan, and transplanted to five 0.6-ton cultivating tanks equipped with a flow-through sea water system in February 2012. The cultured octocorals for this research work were collected from the tanks in May 2013. Prof. Chang-Feng Dai, Institute of Oceanography, National Taiwan, identified the soft coral. Living reference specimens are being maintained in the authors’ marine organism cultivating tank and a voucher specimen (NMMBA-TWSC-12005) was deposited in the National Museum of Marine Biology and Aquarium, Taiwan.

3.3. Extraction and Isolation

Sliced bodies of *Nephthea columnaris* (wet weight 800.0 g, dry weight 76.6 g) were extracted with a mixture of MeOH and DCM (1:1) (1.6 L $\times$ 5). The extract was partitioned between EtOAc and H$_2$O. The EtOAc layer (7.4 g) was separated on silica gel and eluted using $n$-hexane/EtOAc (stepwise, 100:1–pure EtOAc) to yield 17 fractions A–Q. Fraction J was chromatographed on NP-HPLC using a mixture of $n$-hexane and acetone (2:1) to afford 14 fractions J1–J14. Fraction J3 was separated by NP-HPLC using a mixture of $n$-hexane and acetone (2:1) as the mobile phase to yield 6 fractions J3A–J3F. Fraction J3B was purified by RP-HPLC using a mixture of acetonitrile and water (1:1) as the
mobile phase to afford 1 (2.3 mg, \( t_R = 22 \text{ min} \)). Fraction J5 was purified by RP-HPLC using a mixture of MeOH and water (3:2) as the mobile phase to afford 2 (2.0 mg, \( t_R = 22 \text{ min} \)).

Columnariol A (1): Colorless oil; \( [\alpha]_D^{23} + 90 \) (c 0.8, MeOH); IR (neat) \( \nu_{\text{max}} \) 3419 cm\(^{-1}\); \(^1\)H (400 MHz, CDCl\(_3\)) and \(^{13}\)C (100 MHz, CDCl\(_3\)) NMR data, see Table 1; ESIMS: \( m/z \) 345 [M + Na]\(^+\); HRESIMS: \( m/z \) 345.23989 (calcd for C\(_{20}\)H\(_{34}\)O\(_3\) + Na, 345.24002).

Columnariol B (2): Colorless oil; \( [\alpha]_D^{23} - 53 \) (c 0.7, MeOH); IR (neat) \( \nu_{\text{max}} \) 3355 cm\(^{-1}\); \(^1\)H (400 MHz, CDCl\(_3\)) and \(^{13}\)C (100 MHz, CDCl\(_3\)) NMR data, see Table 2; ESIMS: \( m/z \) 345 [M + Na]\(^+\); HRESIMS: \( m/z \) 345.23977 (calcd for C\(_{20}\)H\(_{34}\)O\(_3\) + Na, 345.24002).

3.4. In Vitro Anti-Inflammatory Assay

According to our previous and other studies for the in vitro anti-inflammatory activity assay, we used LPS induced RAW murine macrophage cell line which was purchased from American Type Culture Collection (ATCC, Manassas, VA, USA) [4–7]. The in vitro anti-inflammatory activity of Compounds 1 and 2 was measured by examining the inhibition of lipopolysaccharide (LPS)-induced upregulation of pro-inflammatory iNOS (inducible nitric oxide synthase) and COX-2 (cyclooxygenase-2) protein expression in macrophage cells using Western blotting analysis [7–9]. Briefly, inflammation in macrophages was induced by incubating them for 16 h in a medium containing only LPS (10 ng/mL) without compounds. For the anti-inflammatory activity assay, Compounds 1, 2 and dexamethasone (10 \( \mu \)M) were added to the cells 10 min before the LPS challenge. The cells were then for western blot analysis. The immunoreactivity data were calculated with respect to the average optical density of the corresponding LPS-stimulated group. RAW264.7 macrophage cells viability was determined after treatment with alamar blue (Invitrogen, Carlsbad, CA, USA), the tetrazolium dye that is reduced by living cells to fluorescent products. This assay is similar in principle to the cell viability assay using 3-(4,5-dimethyldiazol-2-yl)-2,5-diphenyltetrazolium bromide and has been validated as an accurate measure of the survival of RAW264.7 macrophage cells [10,11]. For statistical analysis, the data were analyzed by a one-way analysis of variance (ANOVA), followed by the Student-Newman-Keuls post hoc test for multiple comparisons. A significant difference was defined as a \( p \)-value of <0.05.

3.5. MTT Antiproliferative Assay

Based on our previous and other studies from the cytotoxicity, there are different tumor cell lines (include MOLT-4, SUP-T1, DLD-1, LNCaP and MCF7 cells) which were obtained from ATCC [12–16]. Cells were maintained in RPMI 1640 medium supplemented with 10% fetal calf serum, 2 mM glutamine and antibiotics (100 units/mL penicillin and 100 \( \mu \)g/mL streptomycin) at 37 °C in a humidified atmosphere of 5% CO\(_2\). Cells were seeded at 4 \( \times \) 10\(^4\) per well in 96-well culture plates before treatment with different concentrations of the tested compounds. In cell culture experiments, compounds 1 and 2 were dissolved in 100% dimethyl sulfoxide (DMSO) (clear). The final concentration of DMSO in the culture medium was 0.02% and the compounds were made concentrations of 1.25, 2.5, 5, 10 and 20 \( \mu \)g/\( \mu \)L prior to the experiments. After treatment for 72 h, the cytotoxicity of the tested compounds was determined using a MTT cell proliferation assay (thiazolyl blue tetrazolium bromide, Sigma-M2128). The MTT is reduced by the mitochondrial dehydrogenases of viable cells to a purple formazan product. The MTT-formazan product was dissolved in DMSO. Light absorbance values
(OD = OD\text{570} − OD\text{620}) were recorded at wavelengths of 570 and 620 nm using an ELISA reader (Anthos Labtec Instrument, Salzburg, Austria) to calculate the concentration that caused 50% inhibition (IC\text{50}), i.e., the cell concentration at which the light absorbance value of the experiment group was half that of the control group. These results were expressed a percentage of the control ± SD established from \text{n} = 4 wells per one experiment from three separate experiments.

4. Conclusions

Octocorals have been well-recognized as an important source of potential bioactive marine natural products. However, because of the octocorals are claimed to be threatened species and most of the compounds from octocorals are difficult to obtain by chemical methods at this stage, bioactive substances from cultured marine invertebrates will play an important role in this field. Our studies on \textit{Nephthea columnaris} for the extraction of additional natural substances, have led to the isolation of two new cembranes, columnnariols A (1) and B (2). Compounds 1 and 2 are potentially anti-inflammatory and may become lead compounds in future marine anti-inflammation drug development [17,18]. To the best of our knowledge, this is the first time to study the natural products from \textit{N. columnaris}. These results suggest that continuing investigation of novel secondary metabolites together with the potentially useful bioactivities from this marine organism are worthwhile for future drug development. The octocoral \textit{Nephthea columnaris} will be transplanted to culturing tanks located in the National Museum of Marine Biology and Aquarium, Taiwan, for extraction of additional natural products to establish a stable supply of bioactive material [19].

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Author Contributions

Yang-Chang Wu and Ping-Jyun Sung designed the whole experiment and contributed to manuscript preparation. Ting-Hsi Hsiao and Chun-Sung Sung researched data and wrote the manuscript. Yu-Hsuan Lan, Yi-Chen Wang, Mei-Chin Lu and Zhi-Hong Wen analyzed the data and performed data acquisition.

Conflicts of Interest

The authors declare no conflict of interest.
References

1. Duh, C.-Y.; Wang, S.-K.; Weng, Y.-L.; Chiang, M.Y.; Dai, C.-F. Cytotoxic terpenoids from the Formosan soft coral *Nephthea brassica*. *J. Nat. Prod.* 1999, 62, 1518–1521.

2. Duh, C.-Y.; Wang, S.-K.; Weng, Y.-L. Brassicolene, a novel cytotoxic diterpenoid from the Formosan soft coral *Nephthea brassica*. *Tetrahedron Lett.* 2000, 41, 1401–1403.

3. Wang, S.-K.; Hsieh, M.-K.; Duh, C.-Y. New diterpenoids from soft coral *Sarcophyton ehrenbergii*. *Mar. Drugs* 2013, 11, 4318–4327.

4. Ho, F.M.; Lai, C.-C.; Huang, L.-J.; Kuo, T.-C.; Chao, C.-M.; Lin, W.-W. The anti-inflammatory carbazole, LCY-2-CHO, inhibits lipopolysaccharide-induced inflammatory mediator expression through inhibition of the p38 mitogen-activated protein kinase signaling pathway in macrophages. *Br. J. Pharmacol.* 2004, 141, 1037–1047.

5. Park, J.S.; Woo, M.S.; Kim, S.Y.; Kim, W.K.; Kim, H.S. Repression of interferon-gamma-induced inducible nitric oxide synthase (iNOS) gene expression in microglia by sodium butyrate is mediated through specific inhibition of ERK signaling pathways. *J. Neuroimmunol.* 2005, 168, 56–64.

6. Lin, Y.-F.; Kuo, C.-Y.; Wen, Z.-H.; Lin, Y.-Y.; Wang, W.-H.; Su, J.-H.; Sheu, J.-H.; Sung, P.-J. Flexibilisquinone, a new anti-inflammatory quinone from the cultured soft coral *Sinularia flexibilis*. *Molecules* 2013, 18, 8160–8167.

7. Huang, S.-Y.; Chen, N.-F.; Chen, W.-F.; Hung, H.-C.; Lee, H.-P.; Lin, Y.-Y.; Wang, H.-M.; Sung, P.-J.; Sheu, J.-H.; Wen, Z.-H. Sinularin from indigenous soft coral attenuates nociceptive responses and spinal neuroinflammation in carrageenan-induced inflammatory rat model. *Mar. Drugs* 2012, 10, 1899–1919.

8. Jean, Y.-H.; Chen, W.-F.; Sung, C.-S.; Duh, C.-Y.; Huang, S.-Y.; Lin, C.-S.; Tai, M.-H.; Tzeng, S.-F.; Wen, Z.-H. Capnellene, a natural marine compound derived from soft coral, attenuates chronic constriction injury-induced neuropathic pain in rats. *Br. J. Pharmacol.* 2009, 158, 713–725.

9. Jean, Y.-H.; Chen, W.-F.; Duh, C.-Y.; Huang, S.-Y.; Hsu, C.-H.; Lin, C.-S.; Sung, C.-S.; Chen, I.-M.; Wen, Z.-H. Inducible nitric oxide synthase and cyclooxygenase-2 participate in anti-inflammatory and analgesic effects of the natural marine compound lemnalol from Formosan soft coral *Lemnalia cervicorni*. *Eur. J. Pharmacol.* 2008, 578, 323–331.

10. Chen, L.-C.; Lin, Y.-Y.; Jean, Y.-H.; Lu, Y.; Chen, W.-F.; Yang, N.-S.; Wang, H.-M.D.; Jang, I.-Y.; Chen, I.-M.; Su, J.-H.; et al. Anti-inflammatory and analgesic effects of the marine-derived compound comaparvin isolated from the crinoid *Comanthus bennetti*. *Molecules* 2014, 19, 14667–14686.

11. Oliveira, T.; Figueiredo, C.A.; Brito, C.; Stavroullakis, A.; da Silva Velozo, E.; Nogueira-Filho, G.; Prakki, A. Effect of *Allium cepa* L. on lipopolysaccharide-stimulated osteoclast precursor cell viability, count, and morphology using 4′,6-diamidino-2-phenylindole-staining. *Int. J. Cell Biol.* 2014, 2014, doi:10.1155/2014/535789.

12. Shih, F.-Y.; Chen, T.-H.; Lu, M.-C.; Chen, W.-F.; Wen, Z.-H.; Kuo, Y.-H.; Sung, P.-J. Cladieunicellins K and L, new eunicellin-based diterpenoids from an octocoral *Cladiella* sp. *Int. J. Mol. Sci.* 2013, 14, 21781–21789.
13. Chen, K.-H.; Dai, C.-F.; Lu, M.-C.; Li, J.-J.; Chen, J.-J.; Chang, Y.-C.; Su, Y.-D.; Wang, W.-H.; Sung, P.-J. Secondary metabolites from the soft coral *Sinularia arborea*. *Mar. Drugs* **2013**, *11*, 3372–3380.

14. Alley, M.C.; Scudiero, D.A.; Monks, A.; Hursey, M.L.; Czerwinski, M.J.; Fine, D.L.; Abbott, B.J.; Mayo, J.G.; Shoemaker, R.H.; Boyd, M.R. Feasibility of drug screening with panels of human tumor cell lines using a microculture tetrazolium assay. *Cancer Res.* **1988**, *48*, 589–601.

15. Scudiero, D.A.; Shoemaker, R.H.; Paull, K.D.; Monks, A.; Tierney, S.; Nofziger, T.H.; Currens, M.J.; Seniff, D.; Boyd, M.R. Evaluation of a soluble tetrazolium/formazan assay for cell growth and drug sensitivity in culture using human and other tumor cell lines. *Cancer Res.* **1988**, *48*, 4827–4833.

16. Lu, M.-C.; Hwang, S.-L.; Chang, F.-R.; Chen, Y.-H.; Chang, T.-T.; Hung, C.-S.; Wang, C.-L.; Chu, Y.-H.; Pan, S.-H.; Wu, Y.-C. Immunostimulatory effect of *Antrodia camphorata* extract on functional maturation of dendritic cells. *Food Chem.* **2009**, *113*, 1049–1057.

17. Wei, W.-C.; Sung, P.-J.; Duh, C.-Y.; Chen, B.-W.; Sheu, J.-H.; Yang, N.-S. Anti-Inflammatory activities of natural products isolated from soft corals of Taiwan between 2008 and 2012. *Mar. Drugs* **2013**, *11*, 4083–4126.

18. Senthilkumar, K.; Kim, S.-K. Marine invertebrate natural products for anti-Inflammatory and chronic diseases. *Evid. Based Complement. Altern. Med.* **2013**, *2013*, doi:10.1155/2013/572859.

19. Leal, M.C.; Calado, R.; Sheridan, C.; Alimonti, A.; Osinga, R. Coral aquaculture to support drug discovery. *Trends Biotechnol.* **2013**, *31*, 555–561.

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