

Four new eunicellin-based diterpenoids from soft coral *Cladiella conifera* (Tixier-Durivault, 1943)



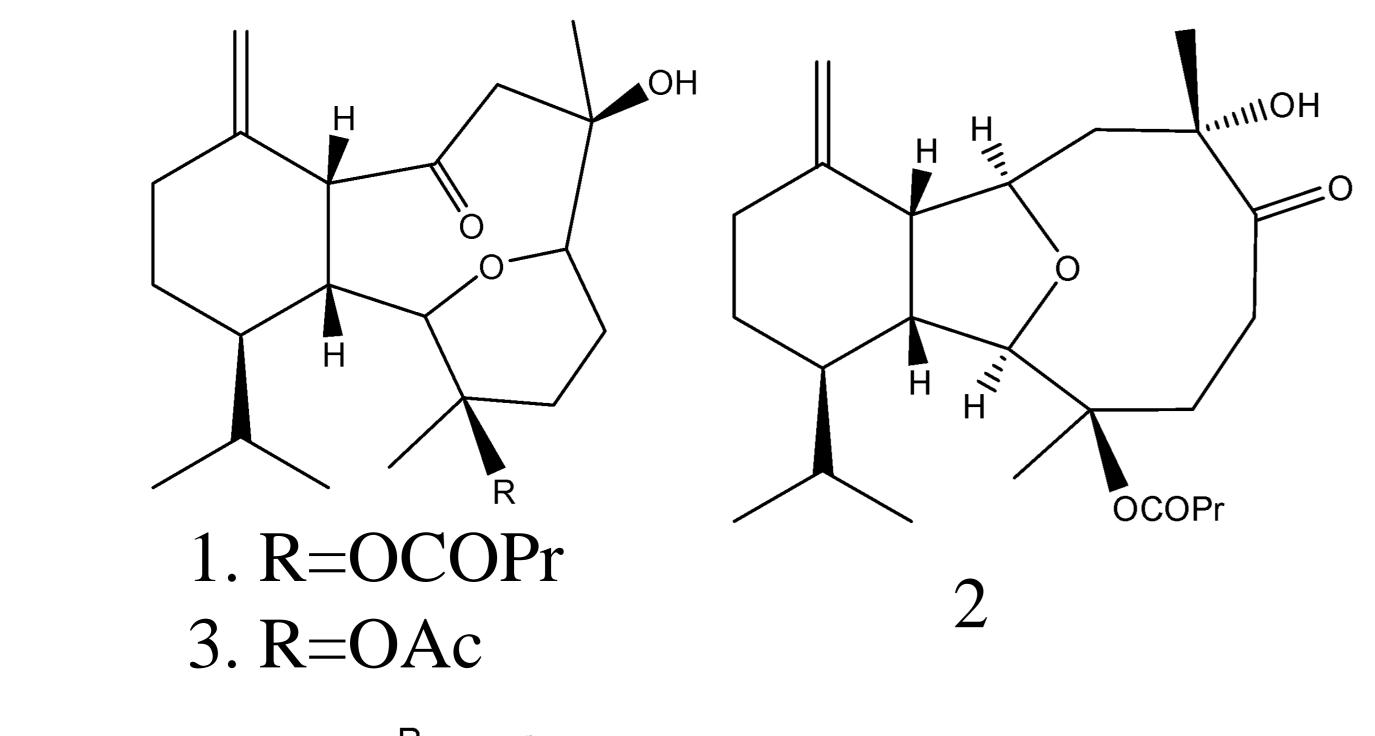
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Abstract

Eunicellin-based diterpenoid had been found to possess multiple and extensive bioactivities, which was rich in the octocoral of genus *Cladella*. Therefore, we collected *Cladella conifera* from Penghu Archipelago in Taiwan by scuba diving. It isolation of seven eunicellin type diterpenoids, including four new compounds coniferains A-D (1, 2, 6, and 7), and three known compounds, multifloralin (3), litophynol B (4), $(1R^*, 2R^*, 3R^*, 6S^*, 7S^*, 9R^*, 10R^*, 14R^*)$ -3-butanoyloxycladiell-11(17)-en-6,7-diol (5). The structures 1–5 were determined by extensive spectroscopic analysis and comparison of spectroscopic with those data release before. These compounds showed not efficient cytotoxic activity to HL-60 and HT-29 cell lines, .



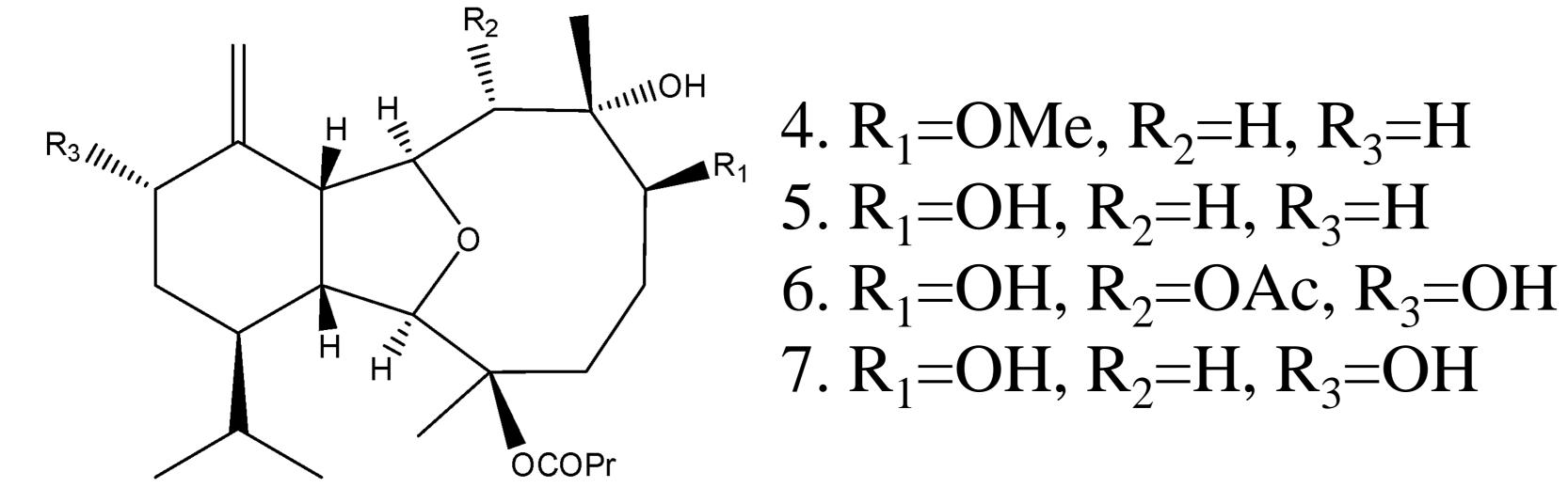


Figure 2. Specimens of *C. conifera* were collected in May 2017 by hand with SCUBA divers off the coast of Penghu Archipelago.

Figure 1. The structure of compounds

Table 1. Cytotoxic data of compounds 1–5

Effects of compounds on cell viability in HL-60 and HT-29 cells.

| | HL-60 | | HT-29 | |
|---------------------|----------------------------|------------------------------------|----------------------------|------------------------------------|
| Sample ⁻ | ^a Viability (%) | ^b IC ₅₀ (μM) | ^a Viability (%) | ^b IC ₅₀ (μM) |
| DMSO | 100.00 ± 0.58 | - | 100.00 ± 0.18 | _ |
| 1 | 100.68 ± 0.45 | > 10 | 95.34 ± 0.65 | > 10 |
| 2 | 97.85 ± 1.92 | > 10 | 103.96 ± 1.69 | > 10 |
| 3 | 89.69 ± 2.31 | > 10 | 101.50 ± 0.97 | > 10 |
| 4 | 111.64 ± 1.64 | > 10 | 95.49 ± 0.86 | > 10 |
| 5 | 99.62 ± 2.12 | > 10 | 103.79 ± 1.55 | > 10 |

^aPercentage of cell viability of 10 μ M in HL-60 (24 h) or HT-29 (72 h) cells. Resultls are expressed as mean \pm S.E.M. (n = 3).

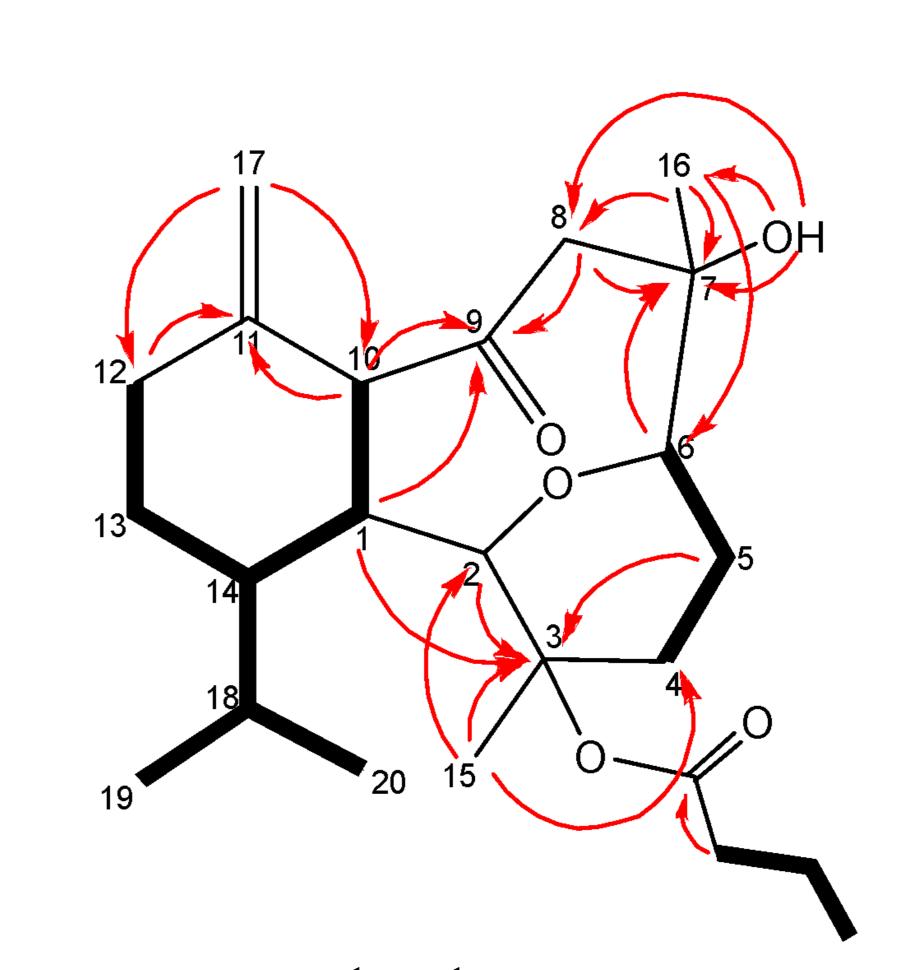


Figure 3. ¹H-¹H COSY (bold bonds) and HMBC (arrow) correlations for coniferain A

are expressed as mean ± S.E.M. (n = 3).

bConcentration necessary for 50 % inhibition (IC₅₀).