

Supplementary Materials

New Naphtho- γ -Pyrones Isolated from Marine-Derived Fungus *Penicillium* sp. HK1-22 and Their Antimicrobial Activities

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Figure S21. HPLC chromatogram of the crude extract from the fungus *Penicillium* sp. HK1-22.

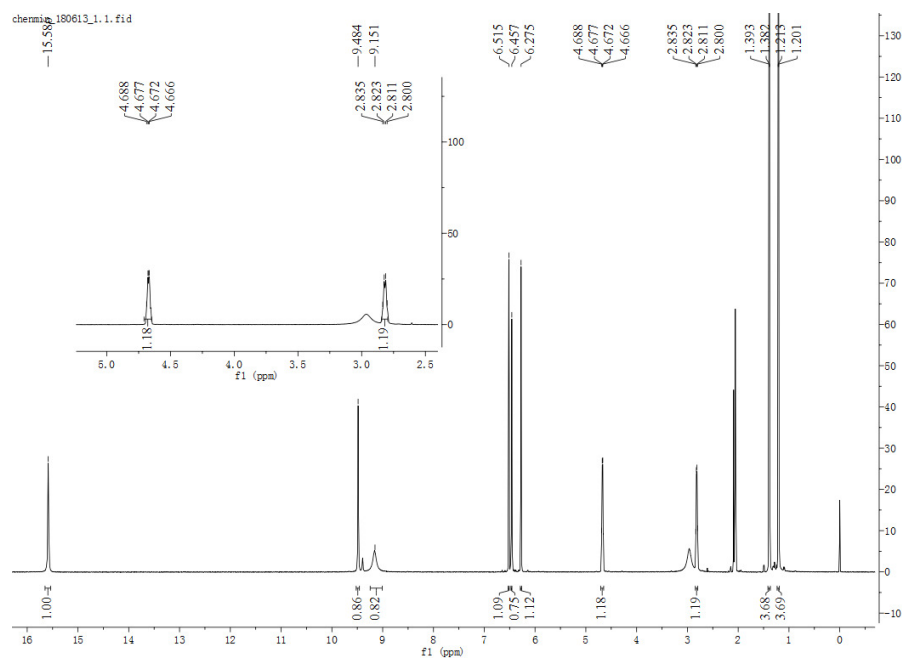


Figure S1. ^1H NMR spectrum of compound **1** in acetone- d_6 (600 MHz).

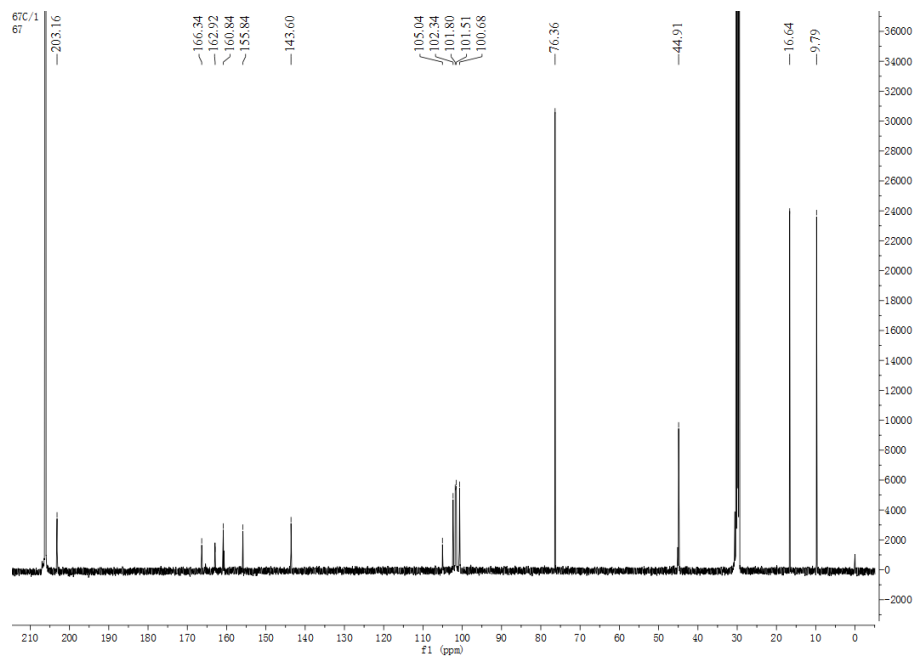


Figure S2. ^{13}C NMR spectrum of compound **1** in acetone- d_6 (150 MHz).

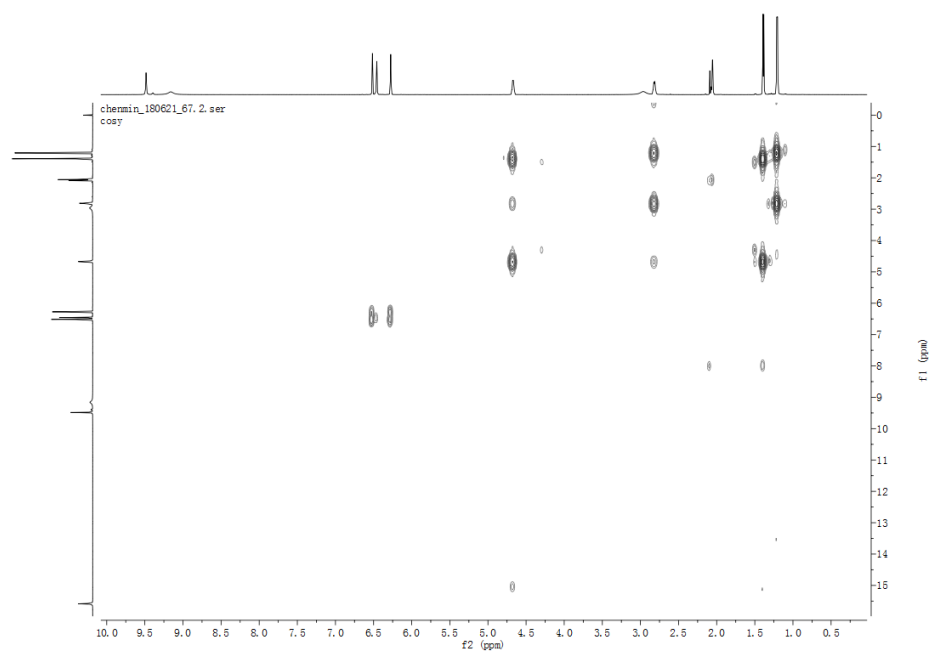


Figure S3. ^1H - ^1H COSY spectrum of compound **1** in acetone- d_6 (600 MHz).

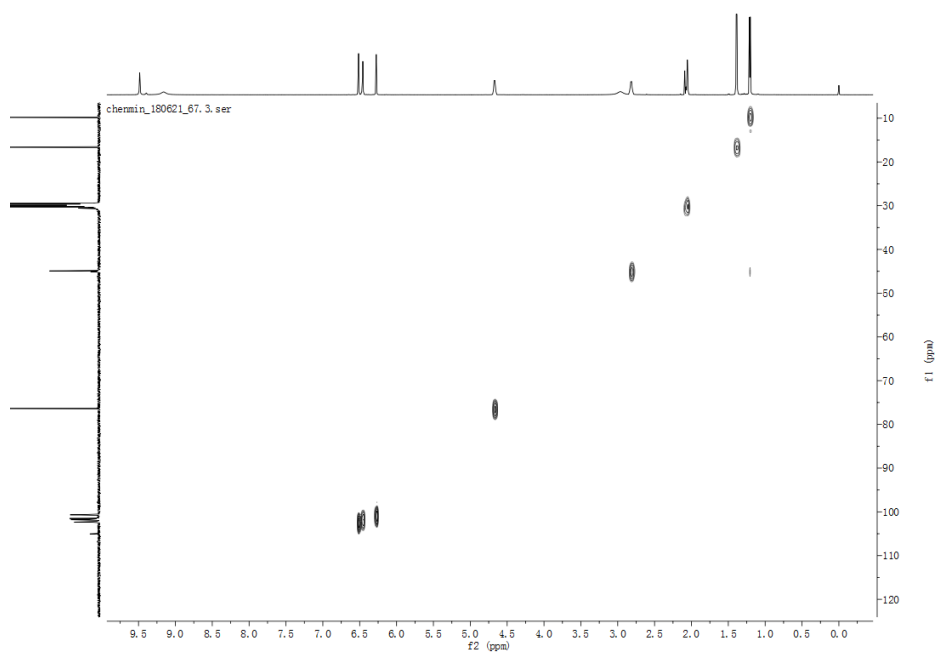


Figure S4. HSQC spectrum of compound **1** in acetone- d_6 (600 MHz).

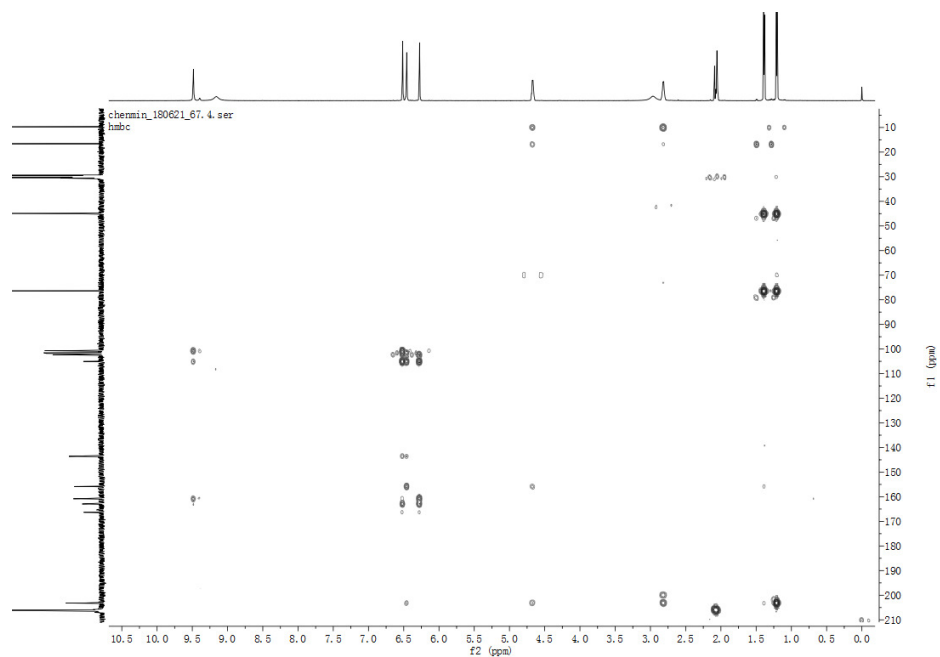
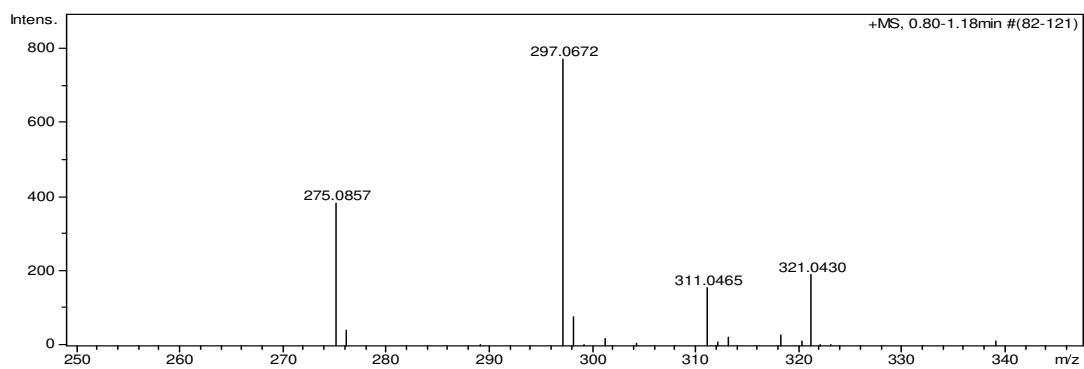


Figure S5. HMBC spectrum of compound **1** in acetone-*d*₆ (600 MHz).



HRESIMS *m/z* 275.0857 (calcd for C₁₅H₁₅O₅, 275.0860),

297.0733 (calcd for C₁₅H₁₄NaO₅, 275.0860)

Figure S6. HRESIMS spectrum (positive ion mode) of compound **1**.

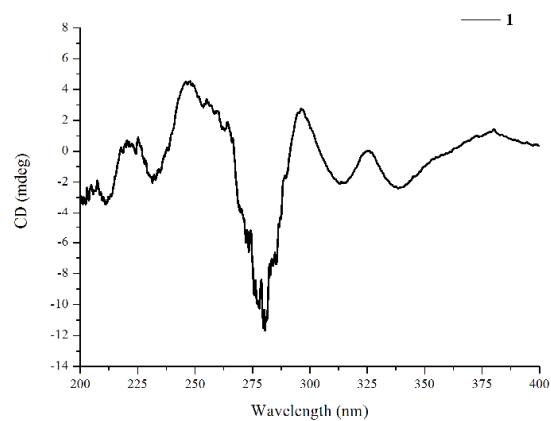


Figure S7. ECD spectrum of compound 1.

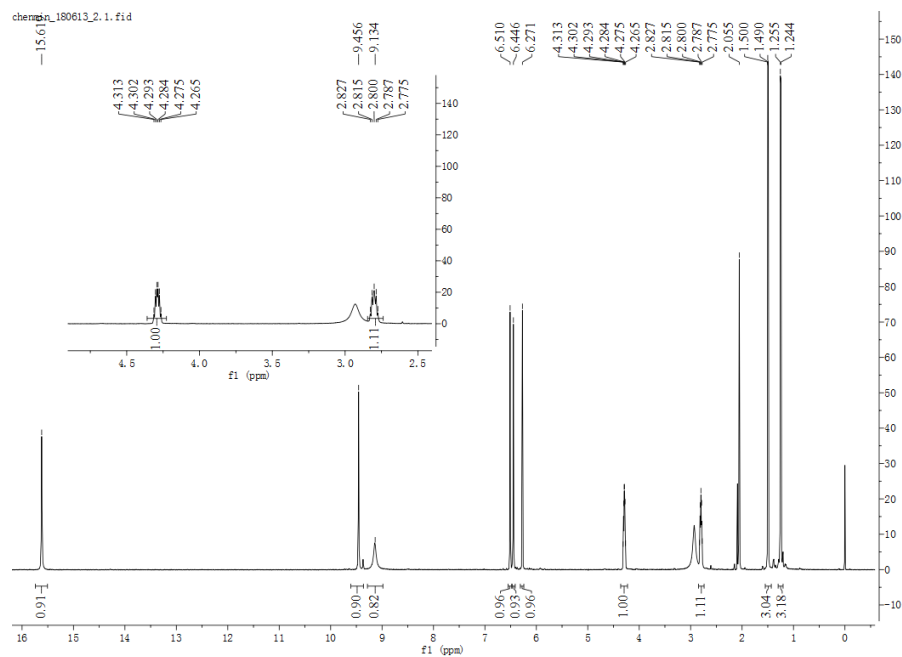


Figure S8. ^1H NMR spectrum of compound 2 in acetone- d_6 (600 MHz).

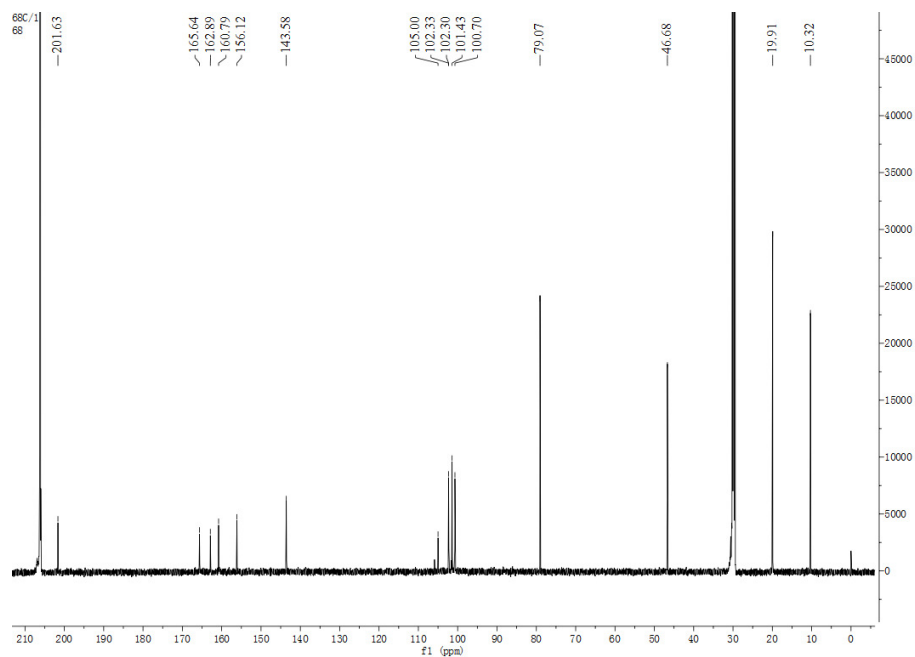


Figure S9. ¹³C NMR spectrum of compound 2 in acetone-*d*₆ (150 MHz).

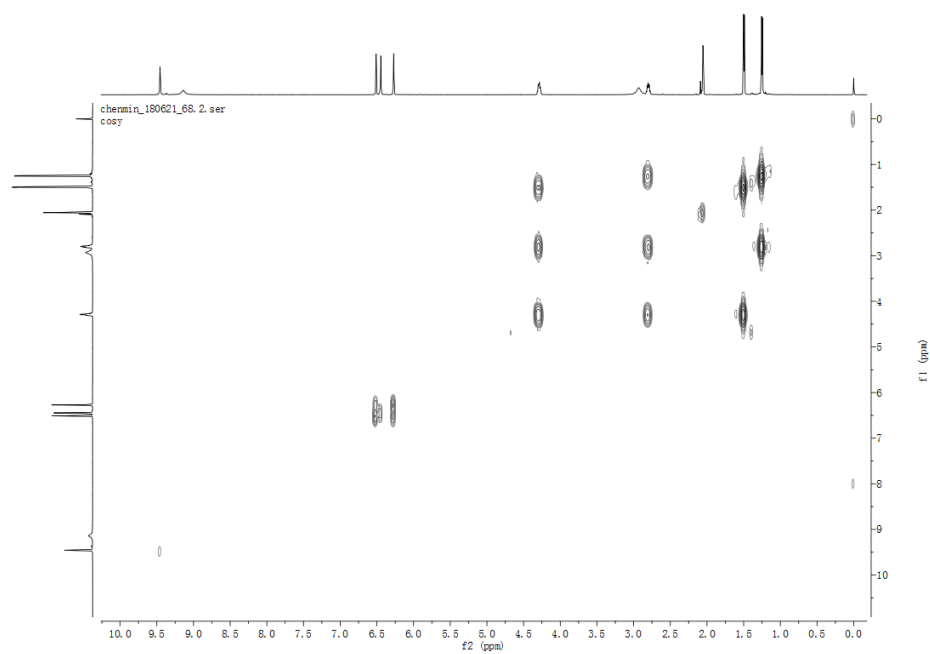


Figure S10. ¹H-¹H COSY spectrum of compound 2 in acetone-*d*₆ (600 MHz).

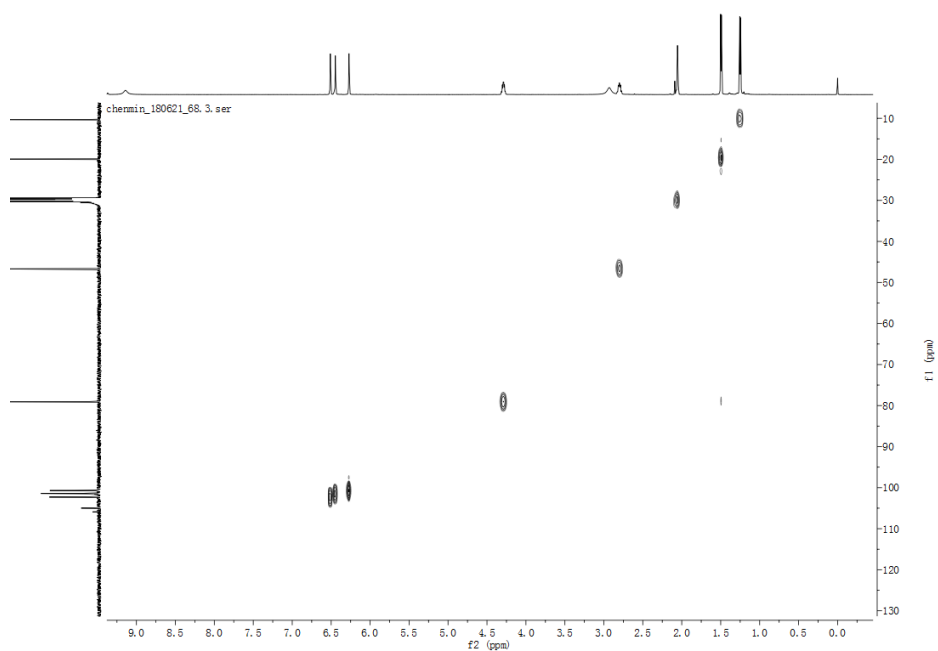


Figure S11. HSQC spectrum of compound **2** in acetone-*d*₆ (600 MHz).

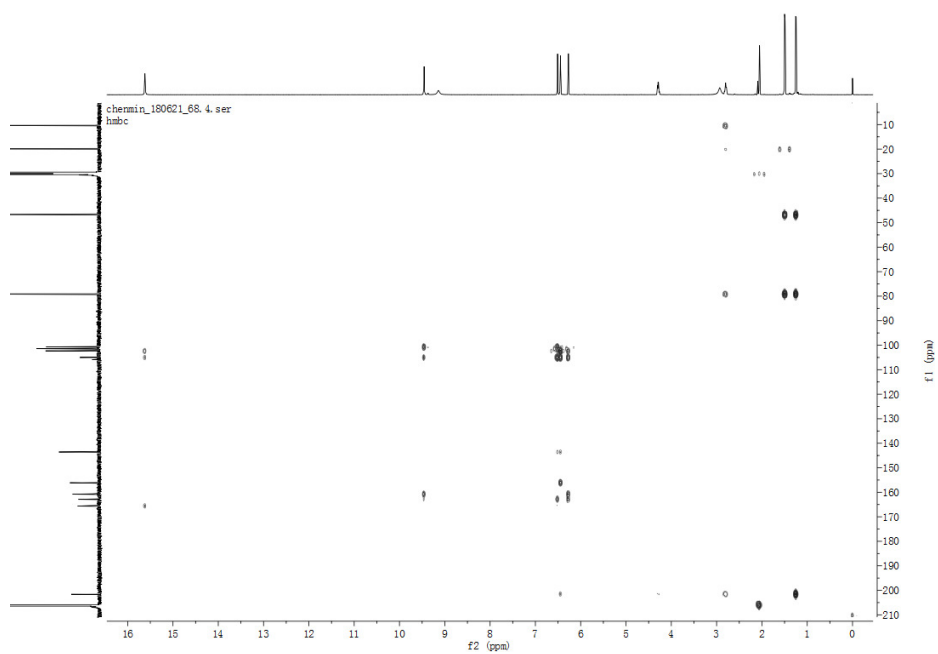
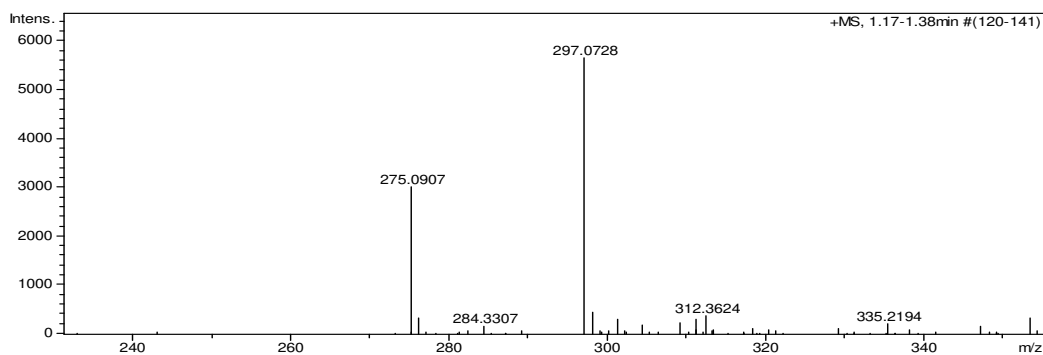


Figure S12. HMBC spectrum of compound **2** in acetone-*d*₆ (600 MHz).



HRESIMS m/z 275.0907 (calcd for $C_{15}H_{15}O_5$, 275.0914),

297.0728 (calcd for $C_{15}H_{14}NaO_5$, 297.0733)

Figure S13. HRESIMS spectrum (positive ion mode) of compound 2.

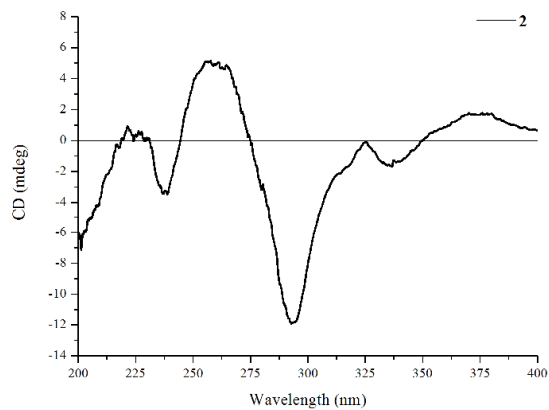


Figure S14. ECD spectrum of compound 2.

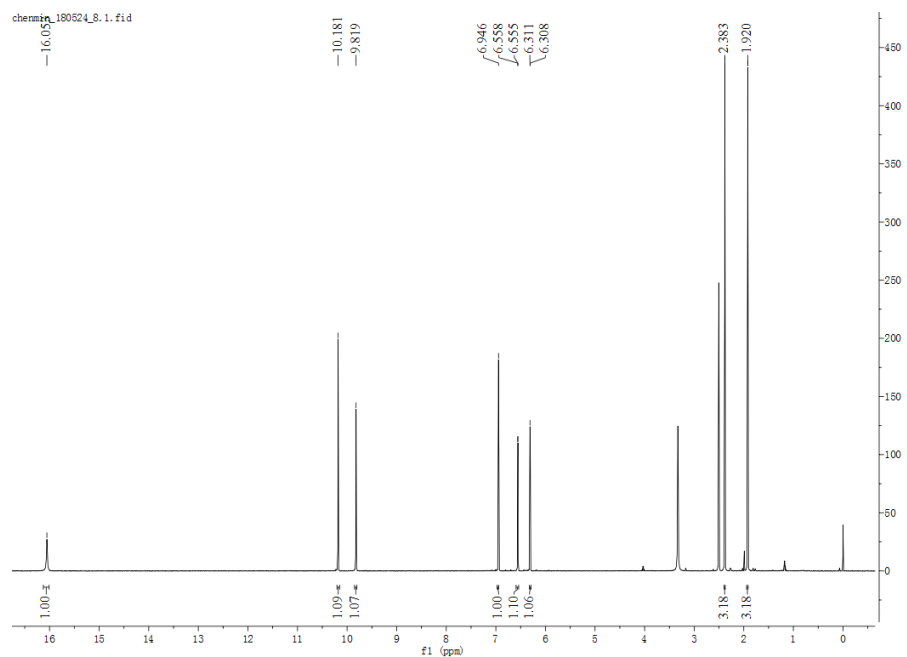


Figure S15. ^1H NMR spectrum of compound **3** in acetone- d_6 (600 MHz).

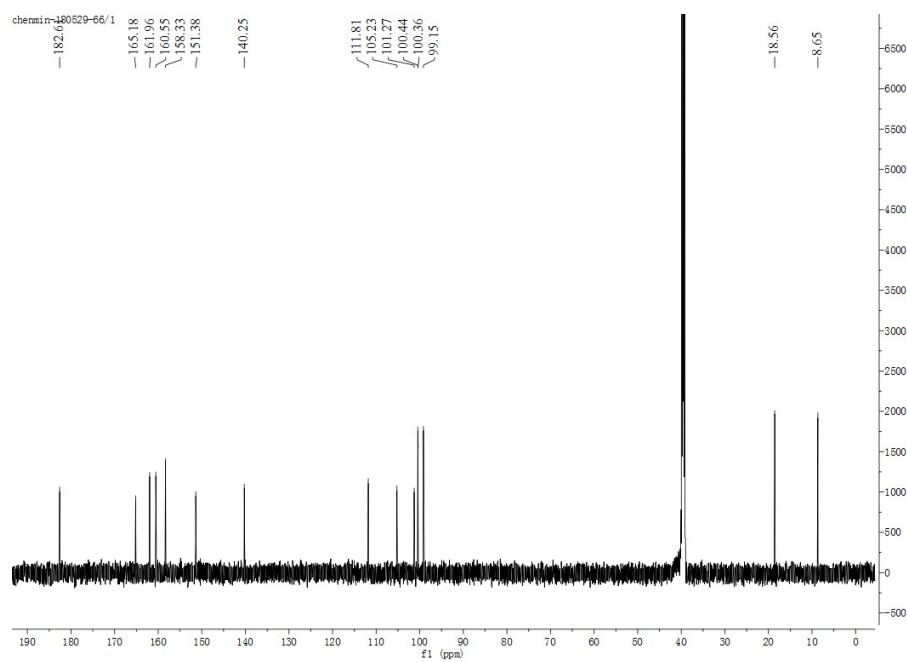


Figure S16. ^{13}C NMR spectrum of compound **3** in acetone- d_6 (150 MHz).

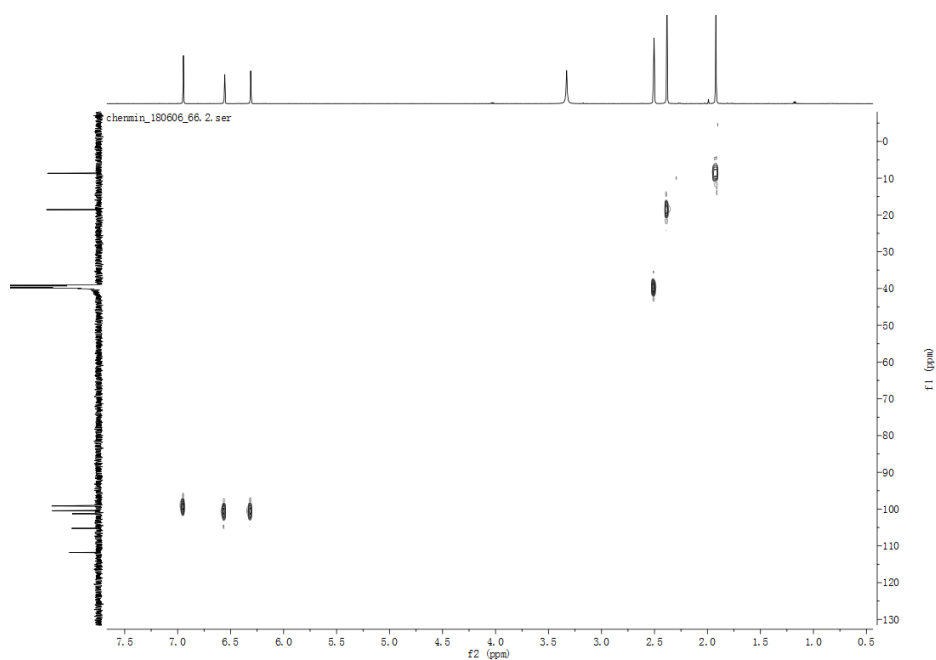


Figure S17. HSQC spectrum of compound **3** in acetone-*d*₆ (600 MHz).

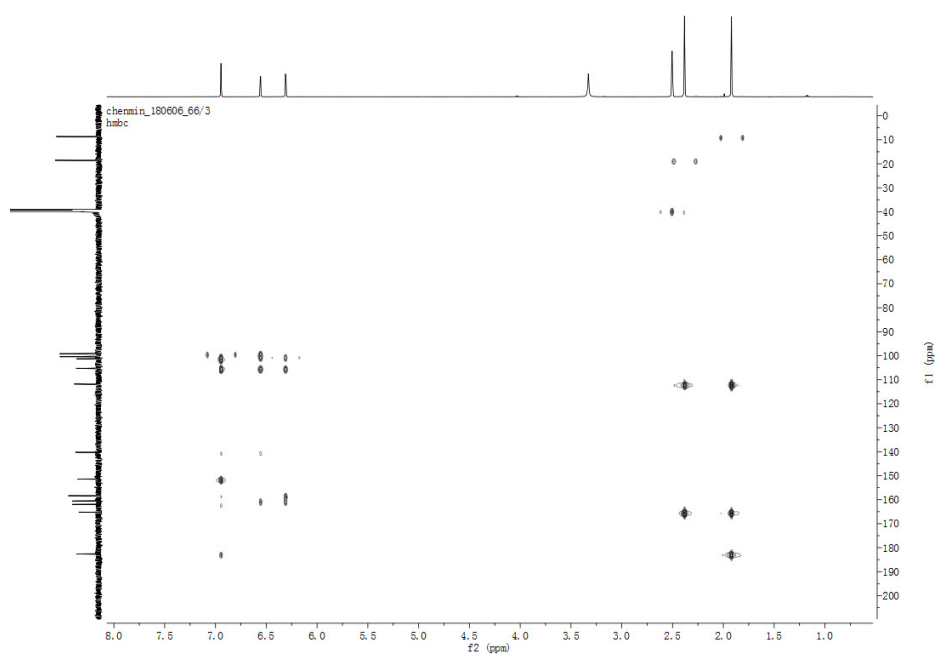
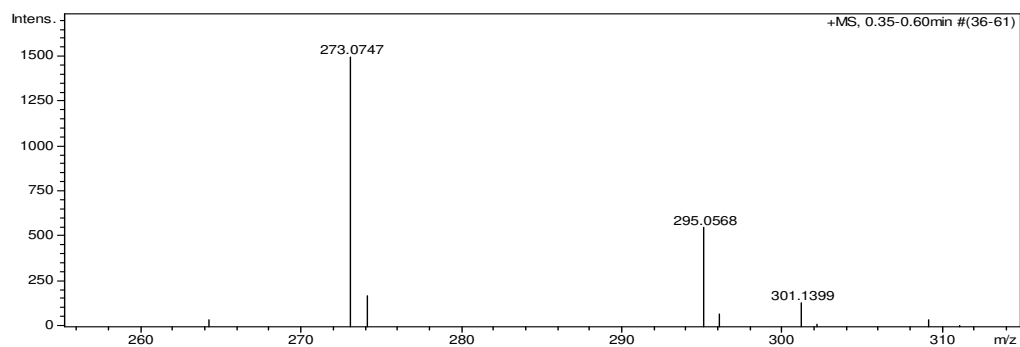


Figure S18. HMBC spectrum of compound **3** in acetone-*d*₆ (600 MHz).



HRESIMS m/z 273.0747 (calcd for $C_{15}H_{13}O_5$, 273.0757),

295.0568 (calcd for $C_{15}H_{12}NaO_5$, 295.0577)

Figure S19. HRESIMS spectrum (positive ion mode) of compound **3**.

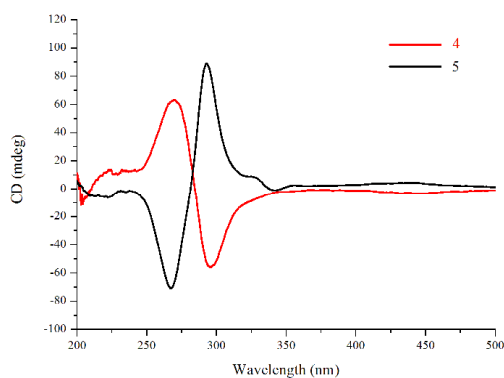


Figure S20. ECD spectrum of compounds **4** and **5**.

The 1H NMR, ESIMS and ECD data of **4** and **5**.

Isochaetochromin B₁ (4): yellow amorphous powder; ECD (0.30 mM, MeOH) λ_{max} ($\Delta\epsilon$) 295 (−55.52), 270 (+62.20) nm; 1H NMR (acetone- d_6 , 600 MHz) δ_H 6.71 (1H, s, H-10), 6.55 (1H, s, H-7), 6.43 (1H, s, H-7'), 6.13 (1H, s, H-10'), 4.71 (1H, dq, J = 3.0, 5.4 Hz, H-2), 4.23 (1H, dq, J = 13.2, 5.4 Hz, H-2'), 2.85 (1H, dq, J = 3.0, 6.6 Hz, H-3), 2.77 (1H, dq, J = 13.2, 6.6 Hz, H-3'), 1.41 (3H, d, J = 5.4 Hz, CH₃-11), 1.41 (3H, d, J = 5.4 Hz, CH₃-11'), 1.22 (3H, d, J = 6.6 Hz, CH₃-12), 1.22 (3H, d, J = 6.6 Hz, CH₃-12'). ESIMS m/z 547 $[M + H]^+$.

Isochaetochromin B₂ (5): yellow amorphous powder; ECD (0.30 mM, MeOH) λ_{max} ($\Delta\epsilon$) 294 (+88.07), 267 (−70.79) nm; ¹H NMR (acetone-*d*₆, 600 MHz) δ_{H} 6.71 (1H, s, H-10), 6.56 (1H, s, H-7), 6.44 (1H, s, H-7'), 6.13 (1H, s, H-10'), 4.72 (1H, dq, *J* = 3.0, 5.4 Hz, H-2), 4.24 (1H, dq, *J* = 11.4, 5.4 Hz, H-2'), 2.85 (1H, dq, *J* = 3.0, 6.6 Hz, H-3), 2.77 (1H, dq, *J* = 11.4, 6.6 Hz, H-3'), 1.41 (3H, d, *J* = 5.4 Hz, CH₃-11), 1.41 (3H, d, *J* = 5.4 Hz, CH₃-11'), 1.23 (3H, d, *J* = 6.6 Hz, CH₃-12), 1.23 (3H, d, *J* = 6.6 Hz, CH₃-12'). ESIMS *m/z* 547 [M + H]⁺.

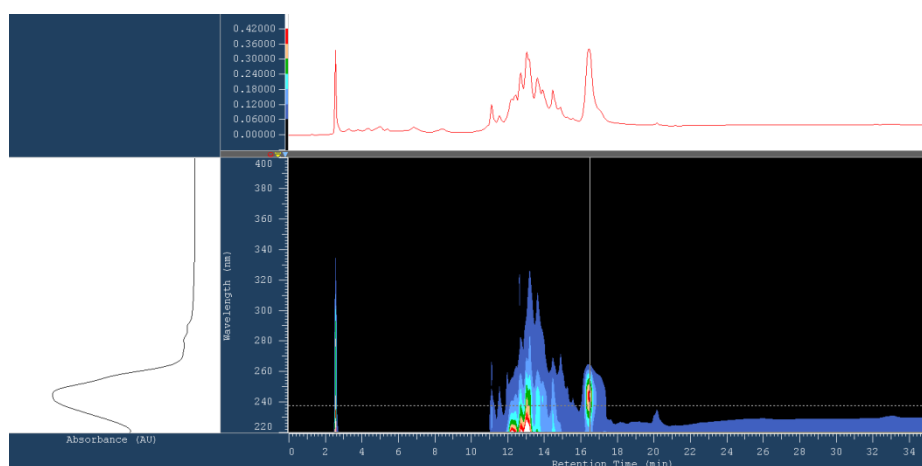


Figure S21. HPLC chromatogram of the crude extract from the fungus *Penicillium* sp. HK1-22.

HPLC was performed on a Hitachi L-2000 system using an analytical C₁₈ (Apollo, 5 μ m, 250 mm $\times$ 4.5mm) column coupled with a 2455 UV detector. The conditions of the HPLC analysis: solvents: A, water; B, MeOH. Linear gradient: 0 min, 20% B; 5 min, 20% B; 10 min, 80% B; 15 min 100% B; 60 min 100%. Temperature: 30 $^{\circ}$ C. Flow rate: 1 mL/min. UV detection at λ = 220 nm.