

SYNTHESIS OF AMPHIMEDINE, A NEW FUSED AROMATIC
ALKALOID FROM A PACIFIC SPONGE, *Amphimedon* sp.

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Abstract —Synthesis of the cytotoxic fused pentacyclic
aromatic alkaloid, amphimedine **1**, is described.

Although marine organisms have been a rich source of structurally diverse natural products, relatively few alkaloids have been isolated from marine sources.¹

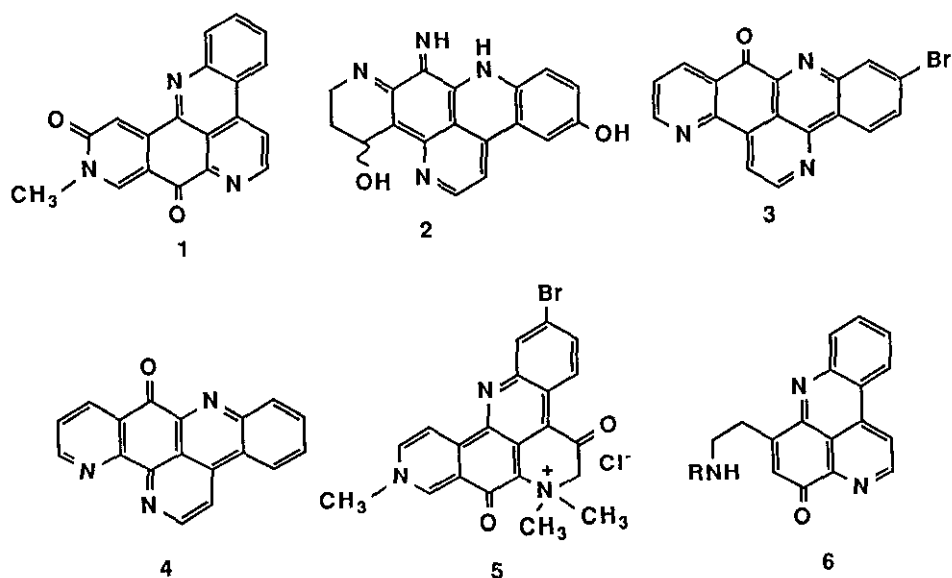
In 1983 Schmitz and co-workers reported the isolation of a novel cytotoxic fused aromatic alkaloid, named amphimedine **1**, from an *Amphimedon* sp. of sponge found near Guam island.² The structure **1** was assigned on the basis of an extensive long-range heterocorrelation and carbon-carbon correlation analyses.

To date, the structurally related fused aromatic alkaloids include calliactine **2**,³ 2-bromo-leptoclinidinone **3**,⁴ ascididemin **4**,⁵ petrosamine **5**,⁶ and cystodytins A, B, and C **6**.⁷

Their highly fused structures have proven not only to be challenging structural elucidation problems and but also to be challenging targets for synthesis.

Although a few synthetic studies⁸ toward them have been reported recently, no total synthesis has yet been accomplished. In this paper, we report the first synthesis of amphimedine **1**.

Our starting material was the *o*-nitrobenzoylacetanilide **7** [mp 93.5-95°C; ms m/z 344 (M^+)], which was prepared in quantitative yield by heating 2,5-dimethoxyaniline with ethyl *o*-nitrobenzoylacetate in toluene and a slight amount of pyridine at 140°C for 6 h. Cyclization of **7** in 80% H₂SO₄ at 75°C for 30 min gave in 53% yield the quinolone **8** [mp 208-209°C; ms m/z 326 (M^+)], which was converted to the 2-chloroquinoline **9** [mp 225-226°C; ms m/z 346 (M^++2)] in 66% yield using PCl₅/POCl₃ at 70°C for 45 min. Oxidative demethylation of **9** with ceric ammonium nitrate (CAN)⁹ in aqueous CH₃CN at 0°C for 15 min afforded the quinolinequinone **10** [mp 188-190°C(decomp); ms m/z 316 (M^++2)] in 77% yield.

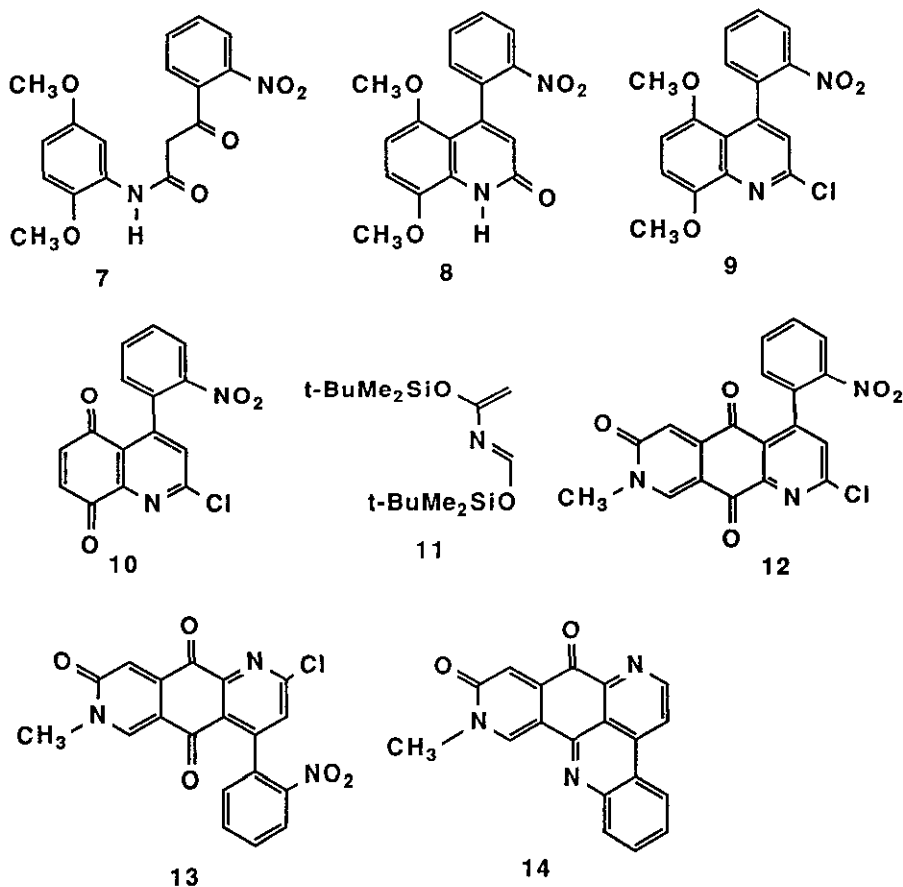


The crucial step, formation of an isoquinolone ring, was performed according to the procedure of Ghosez and co-workers.¹⁰ The Diels-Alder reaction of **10** with 2-aza-1,3-bis(tert-butyldimethylsilyloxy)-1,3-butadiene **11**¹⁰ in CHCl_3 at 35°C for 8 h gave a mixture of crude adducts after acidic workup; which was methylated with $\text{CH}_3\text{I}/\text{K}_2\text{CO}_3/\text{tris}[2-(2\text{-methoxyethoxy})\text{ethyl}]\text{amine}¹¹ in DMF at room temperature for 1 h to provide two N-methylisoquinolones [**12** (7%) and **13** (8%)] after silica gel chromatography [**12**: mp $286\text{--}288^\circ\text{C}$; ms m/z 397 (M^++2 , 4), 395 (M^+ , 11), 351 (34), 349 (100); $^1\text{H-nmr}$ (400MHz, CDCl_3) δ 3.73 (3H, s), 7.03 (1H, s), 7.26 (1H, dd, $J = 1.2, 7.3$ Hz), 7.48 (1H, s), 7.72 (1H, ddd, $J = 1.2, 7.6, 8.2$ Hz), 7.80 (1H, ddd, $J = 1.2, 7.3, 7.6$ Hz), 8.38 (1H, dd, $J = 1.2, 8.2$ Hz), 8.67 (1H, s). **13**: mp $> 300^\circ\text{C}$; ms m/z 397 (M^++2 , 1), 395 (M^+ , 3), 351 (37), 349 (100); $^1\text{H-nmr}$ (400MHz, CDCl_3) δ : 3.64 (3H, s), 7.29 (1H, dd, $J = 1.2, 7.3$ Hz), 7.35 (1H, s), 7.51 (1H, s), 7.70 (1H, ddd, $J = 1.2, 7.6, 8.2$ Hz), 7.79 (1H, ddd, $J = 1.2, 7.3, 7.6$ Hz), 8.32 (1H, s), 8.35 (1H, dd, $J = 1.2, 8.2$ Hz)].$

Finally, catalytic hydrogenation of **12** with 10% Pd-C/ Et_3N in MeOH at room temperature for 20 h afforded in 13% yield amphimedine **1** [mp $> 300^\circ\text{C}$; high-resolution ms Calcd for $\text{C}_{19}\text{H}_{11}\text{N}_3\text{O}_2$: 313.0852, Found 313.0857; uv (EtOH): λ_{max} nm (e) 235 (38,000), 281 (10,000), 340 (7,000); ms

m/z 313 (M^+ , 100), 298 (41); ir (KBr): $\nu_{\max}$ 1680, 1640 cm^{-1} ; $^1\text{H-nmr}$ (400MHz, 2:1 $\text{CF}_3\text{COOD}:\text{CDCl}_3$) δ 4.09 (3H, s), 8.21 (1H, t like), 8.38 (1H, t like), 8.49 (1H, s), 8.69 (1H, d, $J = 8.1$ Hz), 8.97 (1H, d, $J = 8.1$ Hz), 9.19 (1H, s), 9.32 (1H, d, $J = 6.5$ Hz), 9.51 (1H, d, $J = 6.5$ Hz); $^{13}\text{C-nmr}$ (100.4 MHz, 2:1 $\text{CF}_3\text{COOD}:\text{CDCl}_3$) δ 40.03, 114.36, 115.08, 119.16, 120.85, 125.38, 125.58, 133.28, 133.65, 137.72, 139.64, 139.94, 144.41, 145.62, 146.47, 147.76, 148.11, 166.34, 173.58]. Synthetic amphimedine had spectral properties (^1H -, ^{13}C -nmr, HRms, uv) and HPLC mobility identical with those of a natural specimen.

In a similar manner the regio isomer **13** afforded an isomer **14** of amphimedine in 11% yield [mp $> 300^\circ\text{C}$; ms m/z 313 (M^+ , 100), 298 (69), 285 (23); $^1\text{H-nmr}$ (400MHz, 2:1 $\text{CF}_3\text{COOD}:\text{CDCl}_3$)



δ 4.10 (3H, s), 7.92 (1H, s), 8.15 (1H, t like), 8.34 (1H, t like), 8.53 (1H, d, $J = 8.5$ Hz), 8.91 (1H, d, $J = 8.1$ Hz), 9.45 (2H, s like), 9.68 (1H, s)].

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