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# Chemical synthesis and self-assembly of a ladderane phospholipid

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## Supporting Information Placeholder

**ABSTRACT:** Ladderane lipids produced by anammox bacteria constitute some of the most structurally fascinating yet poorly studied molecules among biological membrane lipids. Slow growth of the producing organism and the inherent difficulty of purifying complex lipid mixtures have prohibited isolation of useful amounts of natural ladderane lipids. We have devised a highly selective total synthesis of ladderane lipid tails and a full phosphatidylcholine in order to enable biophysical studies on chemically homogeneous samples of these molecules. Additionally, we report the first proof of absolute configuration of a natural ladderane.

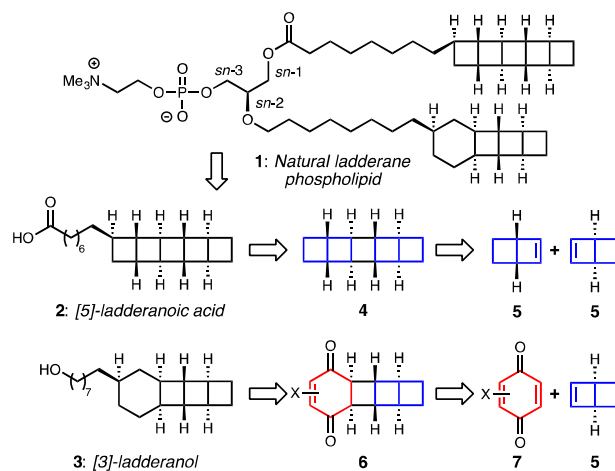
Ladderane lipids, named for the ladder-like polycyclobutane motifs in their hydrophobic tails, are produced by anaerobic ammonium oxidizing (anammox) bacteria as a significant fraction of their membrane lipids.<sup>1</sup> In a large intracellular compartment called the anammoxosome, anammox bacteria couple ammonium and nitrite to produce dinitrogen as their principle source of energy.<sup>2</sup> Experimental and computational evidence suggests that ladderane lipids form a densely packed membrane around the anammoxosome and serve to limit the transmembrane diffusion of toxic and/or valuable metabolites from anammox catabolism.<sup>3</sup> Ladderane lipids likely impart a number of previously unobserved characteristics in membrane settings, but their biophysical characterization has been hampered by a lack of access to pure material.<sup>3a</sup> An axenic culture of anammox bacteria is not available, and growth rates of enrichment cultures in laboratory reactors are slow (doubling times on the order of 1–2 weeks).<sup>4</sup> Furthermore, anammox bacteria produce an array of ladderane and non ladderane lipids, which have proven inseparable on a preparative scale.<sup>3a</sup>

In order to circumvent these difficulties, we designed a synthetic route that would allow us to access any ladderane lipid.

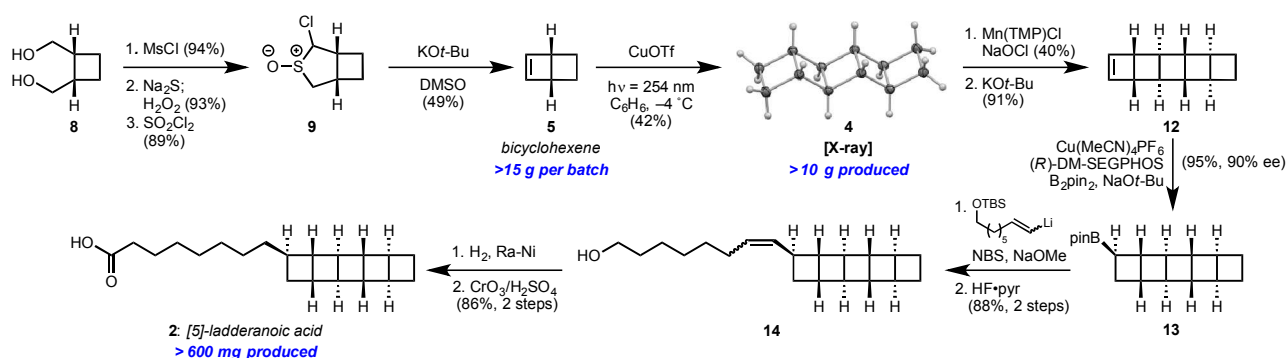
Ladderane phospholipids (e.g. **1**, Scheme 1) isolated from anammox enrichment cultures comprise a mixture containing [5]-ladderane tails, as in acid **2**, and [3]-ladderane tails, as in alcohol **3**; the nomenclature refers to the number of cyclobutane rings at the terminus of the chain.<sup>1,5</sup> Typically, a [3]-ladderane is attached to the central *sn*-2 position by an ether linkage; [5]- and [3]-ladderane lipids, as well as common straight-chain lipids, are found ether- or ester-linked at the *sn*-1 position. Polar head groups at the *sn*-3 position include phosphocholine, phosphoethanolamine, and phosphoglycerol.

Innovative synthetic efforts from Corey and Mascitti resulted in a racemic synthesis of [5]-ladderanoic acid (+/–)-**2** and an enantioselective synthesis which utilized preparative chiral HPLC to produce ca. 20 mg *ent*-**2**.<sup>6</sup>

## Scheme 1. Retrosynthetic analysis of a natural ladderane phospholipid



## Scheme 2. Synthesis of a [5]-ladderanoic acid



No synthesis of a [3]-ladderane lipid has been reported. Despite attempts towards its elucidation, the biosynthesis of ladderanes remains unsolved.<sup>7</sup>

An outline of our retrosynthetic strategy is illustrated in Scheme 1. For both **2** and **3** we planned to install the linear alkyl component at a late stage, focusing first on the polycyclic cores. We targeted the symmetric pentacyclobutane **4**, consisting of all of the cyclobutanes in the [5]-ladderane core, as a key intermediate, which we hoped to construct by dimerization of bicyclohex[2.2.0]ene **5**. For [3]-ladderanol **3**, retrosynthetic installation of two ketones and an alkene on the cyclohexane ring as **6** enabled two key disconnections: a conjugate addition reaction to install the linear alkyl chain, and an enone-olefin [2+2] photocycloaddition between a benzoquinone equivalent (**7**) and bicyclohexene **5**.

Having identified **5** as a key building block for the synthesis of both ladderane tails, we first developed a procedure to obtain this strained olefin in large quantities.<sup>8</sup> Our optimized route commences with diol **8** (Scheme 2), which is commercially available or easily accessible from the photocycloadduct of ethylene and maleic anhydride (see Supporting Information). In three steps we reached  $\alpha$ -chlorosulfoxide **9** as an inconsequential mixture of diastereomers. Treatment of **9** with excess potassium *tert*-butoxide effected an atypical sulfoxide Ramberg–Bäcklund olefination providing, after distillation, over 15 grams of **5** per batch. The use of a sulfoxide in place of the typical sulfone precursor was crucial for high yields in this transformation. The observation that the sulfoxide variant of the Ramberg–Bäcklund reaction excels in constructing strained cyclobutenes was first made by Weinges in the early 1980s, but this procedure has seen no use in synthesis.<sup>9</sup>

In order to accomplish the conversion of **5** to [5]-ladderane core **4**, we hoped to employ a procedure developed by Salomon and Kochi for the dimerization of strained, but otherwise unactivated, olefins.<sup>10</sup> Unfortunately, irradiation of THF or ether solutions of **5** with 254 nm light in the presence of catalytic copper(I) triflate (CuOTf) resulted exclusively in the formation of 1,3-cyclohexadiene (**10**) and a mixture of hexatriene isomers (**11**) (Table 1, entries 1 and 2). Encouragingly,

small amounts of **4** could be observed in reactions conducted in non-coordinating heptane (entry 3). In a key discovery, we found that the electrocyclic ring opening of **5** to **10** is mediated by CuOTf even in the absence of light, and that this undesired pathway could be slowed at temperatures below 0 °C (entries 4–6). Reactions performed in toluene – despite its high absorbance cutoff (284 nm) – delivered **5** in an improved 28% yield (entry 7). Surprisingly, the reaction proceeds optimally in benzene, which is frozen under these conditions (entries 8, 9).<sup>11</sup> The [5]-ladderane pentacyclobutane **4** can thus be obtained in 42% isolated yield. That we see more than one turnover of the CuOTf catalyst seems to indicate that there is some degree of mobility in the solvent matrix. No reaction occurs in the absence of catalyst. Despite its moderate efficiency, this transformation is highly enabling: the entire pentacyclobutane core is assembled in just five steps.

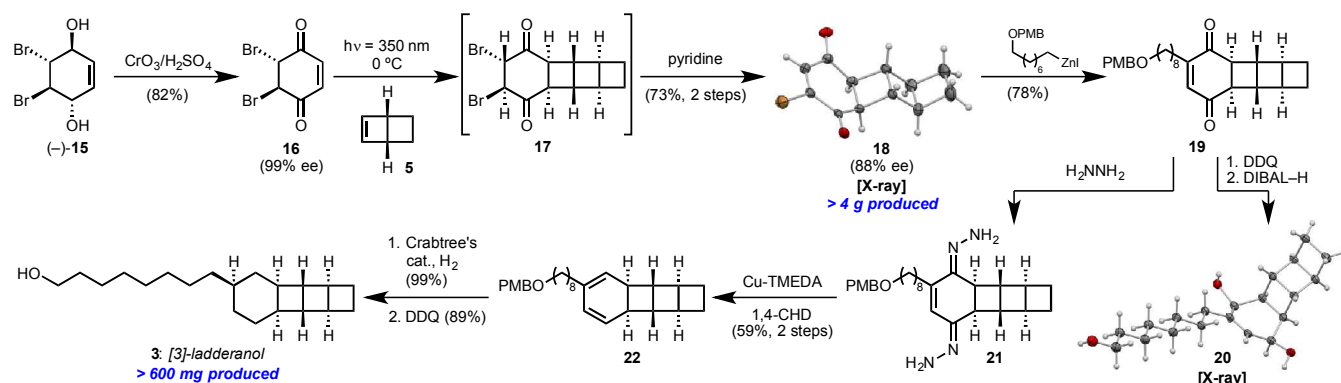
Table 1. Optimization of bicyclohexene [2+2]<sup>a</sup>

entry	solvent	temp.	h $\nu$	<b>5</b> <sup>b</sup>	<b>10</b> + <b>11</b> <sup>b</sup>	<b>4</b> <sup>b</sup>
1	THF	27 °C	254 nm	--	60%	--
2	Et <sub>2</sub> O	27 °C	254 nm	--	86%	--
3	heptane	27 °C	254 nm	--	51%	7%
4	heptane	23 °C	dark	24%	46% <sup>c</sup>	--
5	heptane	-4 °C	dark	70%	2% <sup>c</sup>	--
6	heptane	-4 °C	254 nm	43%	39%	18%
7	toluene	-4 °C	254 nm	24%	46%	28%
8	benzene	-4 °C	254 nm	21%	37%	42%

<sup>a</sup>Reactions were conducted on 1 mmol scale in 1 mL indicated solvent. <sup>b</sup>Yields calculated by comparison to <sup>1</sup>H NMR internal standard. <sup>c</sup>Only **10**.

Tetramesitylporphyrinatomanganese(III) chloride-catalyzed C–H chlorination according to Groves' procedure<sup>12</sup> was uniquely capable of functionalizing **4**, providing 40% of an intermediate chloro-ladderane

## Scheme 3. Synthesis of a [3]-ladderanol.



along with 18% recovered **4**. This is the first report of this chemistry on a cyclobutane. Subsequent elimination with potassium *tert*-butoxide delivered olefin **12** (Scheme 2). Desymmetrization of **12** was accomplished by enantioselective hydroboration with a recently disclosed copper–DM-SEGPHOS catalyst,<sup>13</sup> delivering pinacolboronic ester **13** in 95% yield and 90% ee. We employed a Zweifel olefination to install the 8-carbon alkyl chain.<sup>14</sup> Following deprotection of the primary alcohol, we obtained alkene **14** in 88% yield over two steps. Hydrogenation with Ra-Ni yielded a crystalline [5]-ladderanol intermediate (X-ray: Supporting Information). Subsequent Jones oxidation completed our synthesis of the [5]-ladderanoic acid **2**.

Our synthesis of [3]-ladderanol **3** (Scheme 3) centered on the use of chiral dibromobenzoquinone **16** both as a [2+2] partner for a high-yielding photocycloaddition with bicyclohexene **5** and as a means to relay stereochemistry into the [3]-ladderane core. Enantioenriched **16** is easily accessible from known lipase-resolved dibromodiol (–)-**15** following oxidation.<sup>15</sup> Both  $\pi$  faces of **16** are equivalent, and photocycloaddition with **5** from either side provides the same enantiomer of dibromodiketone **17**. Upon treatment with pyridine, **17** undergoes selective elimination via proton abstraction from the convex face to form vinyl bromide **18** with slight erosion of enantiopurity. Enabled by the  $C_2$ -symmetry of **16**, this two-step sequence effectively achieves an enantioselective [2+2] cycloaddition to desymmetrized **18**.

Initial attempts to install the linear alkyl chain by Negishi cross coupling were accompanied by significant loss of enantiopurity (80% ee **18** to +30% ee **19**, Table 2, entry 1). We reasoned that two pathways might be competing: (1) palladium-catalyzed cross coupling at the site of bromine substitution and (2) uncatalyzed addition of the alkylzinc iodide reagent at the unsubstituted position, followed by elimination of HBr.<sup>16</sup> Indeed, treatment of **18** with an alkylzinc iodide reagent in the absence of catalyst delivered **19** with inversion of the substitution pattern (entry 2). Alternatively, replacing the nucleophilic zinc reagent with a potassium trifluoroborate salt resulted in an enantioretentive Suzuki cross coupling (entry 3). The absolute configurations of in-

termediates **18** and **19** (as the crystalline derivative **20**, Scheme 3) were confirmed by X-ray crystallography. All that remained to elaborate **19** to a [3]-ladderane was the complete reduction of the six-membered ring. While attempting a double Wolff–Kishner reduction on bis-hydrazone **21** (Scheme 3), we discovered instead the formation of a trace amount of 1,3-cyclohexadiene **22**, arising from an apparent 2-electron oxidation and release of two molecules of dinitrogen. After extensive optimization we arrived at commercial Cu-TMEDA as an effective oxidant and 1,4-cyclohexadiene as an H atom source for this transformation. The resulting diene **22** underwent diastereoselective hydrogenation under action of Crabtree's catalyst. Finally, cleavage of the PMB ether completed our synthesis of [3]-ladderanol **3**.

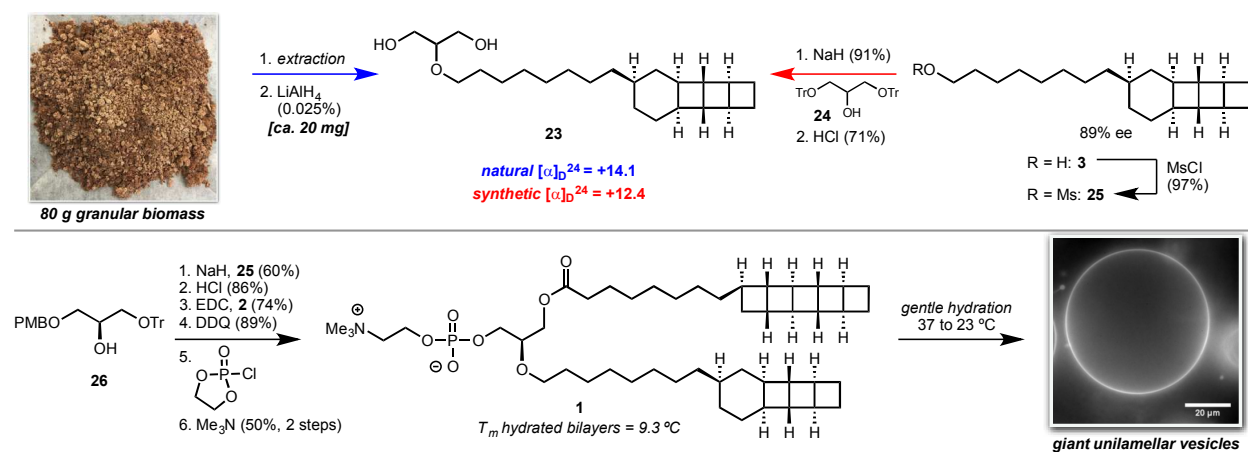
Table 2. Enantiodivergent coupling strategies<sup>a</sup>

entry	nucleophile	catalyst	yield <sup>b</sup>	ee <sup>c</sup>
1	PMBO-CH <sub>2</sub> (CH <sub>2</sub> ) <sub>6</sub> ZnI	Pd(OAc) <sub>2</sub> + XPhos	88%	+30%
2	PMBO-CH <sub>2</sub> (CH <sub>2</sub> ) <sub>6</sub> ZnI	none	64%	–80%
3	PMBO-CH <sub>2</sub> (CH <sub>2</sub> ) <sub>6</sub> BF <sub>3</sub> K	PdCl <sub>2</sub> (dppf)•CH <sub>2</sub> Cl <sub>2</sub>	68%	+80%

<sup>a</sup>Reactions conducted with 80% ee **18**. <sup>b</sup>Yields reflect isolated yields after silica chromatography. <sup>c</sup>Enantiomeric excess determined by chiral HPLC.

The glycerol stereocenter of ladderane phospholipids has been established to be (*R*), as drawn in **1**, which is consistent with other prokaryotic membrane phospholipids.<sup>17</sup> However, the absolute stereochemistry of the ladderane tails has remained unknown. With the goal of completing this assignment, we conducted a lipid extraction of biomass from an anammox enrichment culture grown over four months in a laboratory-scale bioreactor.<sup>18</sup> Cognizant that [3]-ladderane glycerol diol **23** (Scheme 4, top) is one of the most readily separable components of ladderane lipid mixtures,<sup>1,5</sup> we treated our crude extracts directly with lithium aluminum hy-

## Scheme 4. Absolute configuration of [3]-ladderane and the synthesis of a ladderane phospholipid.



dride to reduce all ester and phosphoester linkages and enrich for **23**. From 80 grams of dried biomass we obtained a sufficient amount of **23** for an optical rotation ( $[\alpha]_D^{24} = +14.1$ ). Synthetic **23** prepared via alkylation of glycerol **24** with [3]-ladderanol mesylate **25** was also found to be dextrorotatory ( $[\alpha]_D^{24} = +12.4$  at 88% ee), establishing the natural configuration of **23** as drawn.

Natural [5]-ladderane alcohol from our extracts was inseparable from the corresponding [3]-ladderane alcohol and other straight-chain lipid alcohols. To the best of our knowledge natural [5]-ladderane, first characterized as its methyl ester, has never been completely separated from contaminating [3]-ladderane methyl ester,<sup>1,5a</sup> making synthesis the only way to obtain pure [5]-ladderane lipids. We have assumed by analogy to **3** that the configuration of natural **2** is as drawn (levorotatory).

More than 20 mg of natural ladderane phosphocholine lipid **1** was readily prepared by a short sequence (Scheme 4, bottom). Alkylation of the enantiopure glycerol derivative **26** with mesylate **25** installed a [3]-ladderane at the *sn*-2 position. Removal of the trityl group and esterification with **2** installed a [5]-ladderane at the *sn*-1 position. Removal of the PMB group and installation of the phosphocholine head group completed the route to **1**.

Despite the crystallinity of multiple synthetic intermediates towards **2** and **3**, preliminary evidence indicates that pure **1** forms fluid hydrated bilayers at ambient temperature. Differential scanning calorimetry of aqueous dispersions of **1** revealed a single transition at 9.3 °C. Glass-supported bilayers, a common model for biological membranes, of **1** showed full fluorescence recovery after photobleaching, indicating free lateral mobility within the bilayer (see Supporting Information). Upon gentle hydration, **1** self-assembled into giant unilamellar vesicles (Scheme 4, bottom and Supporting Information) with uniform incorporation of a fluorescently labeled lipid (0.1% Texas Red DHPE). The discovery that pure **1** self-assembles under conditions standard for natural acyclic phosphocholine lipids suggests that conventional techniques can be used to probe

the biophysical properties of ladderane phospholipids. Efforts to validate the hypothesis that ladderanes prevent diffusion of small anammox metabolites are underway. The synthetic route presented here will allow efficient access to a range of natural and non-natural ladderane phospholipids, enabling full exploration of the structural and functional space around these fascinating molecules.

## ASSOCIATED CONTENT

### Supporting Information.

The Supporting Information is available free of charge on the ACS Publications website.

Experimental procedures, characterization, and spectral data (PDF). Cif files (PDF).

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### Notes

The authors declare no competing financial interest.

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