

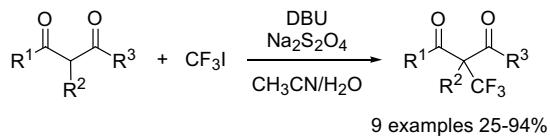
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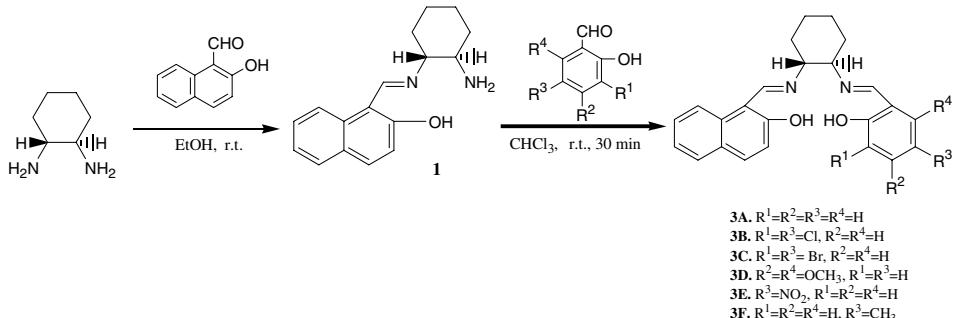
Vitaliy Petrik and Dominique Cahard*



Easy radical trifluoromethylation of a series of 1,3-dicarbonyl compounds with CF_3I . The reaction occurs in the presence of a nitrogen base and sodium dithionite in $\text{CH}_3\text{CN}-\text{H}_2\text{O}$ solution. A new access to ammonium triflinates is also reported.

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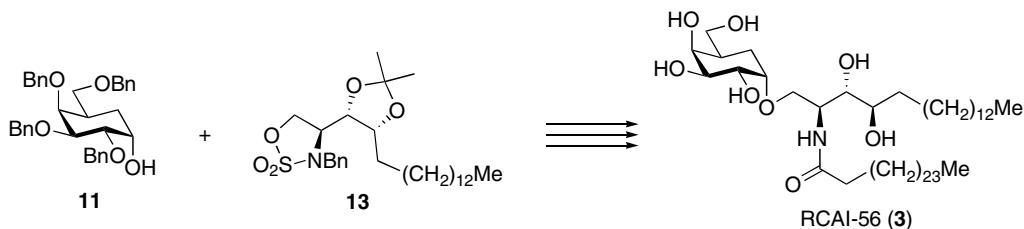
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Titration → ΔpK_a → Partition coefficient
 $(\log P)$



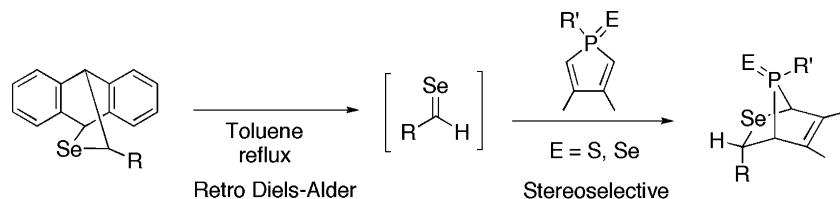
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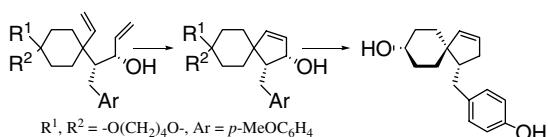
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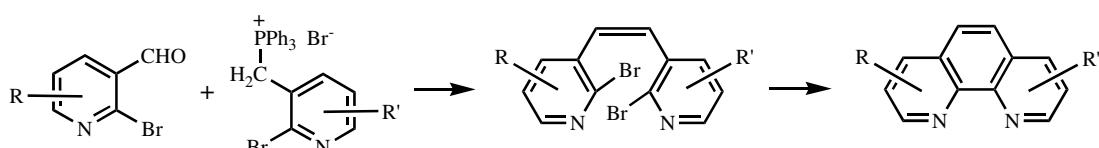
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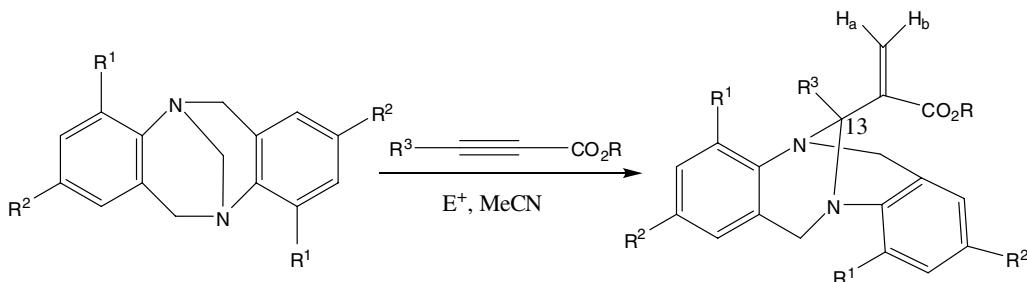
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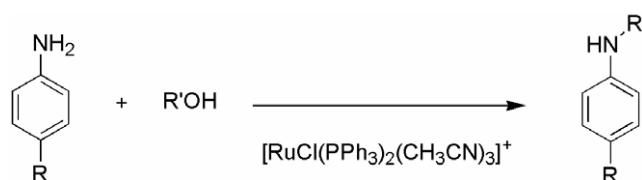
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**Selective N-monoalkylation of anilines catalyzed by a cationic ruthenium(II) compound**

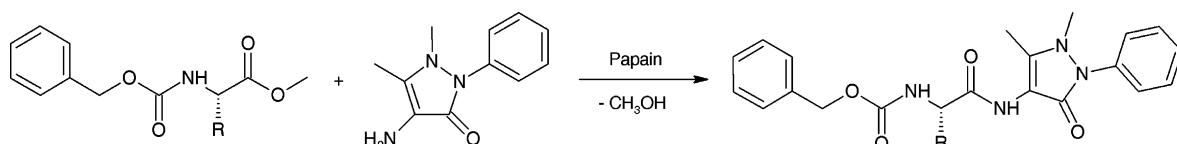
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[RuCl(PPh₃)₂(CH₃CN)₃][BPh₄] catalyzes selective monoalkylation of anilines by alcohols.**Papain-catalysed synthesis of Z-L-aminoacyl-antipyrine amides from Z-protected amino acid esters and 4-aminoantipyrine**

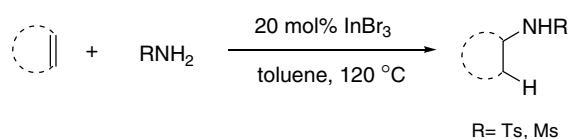
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**InBr₃ Catalyzed intermolecular hydroamination of unactivated alkenes**

Jing-Mei Huang,* Chek-Ming Wong, Feng-Xia Xu and Teck-Peng Loh*

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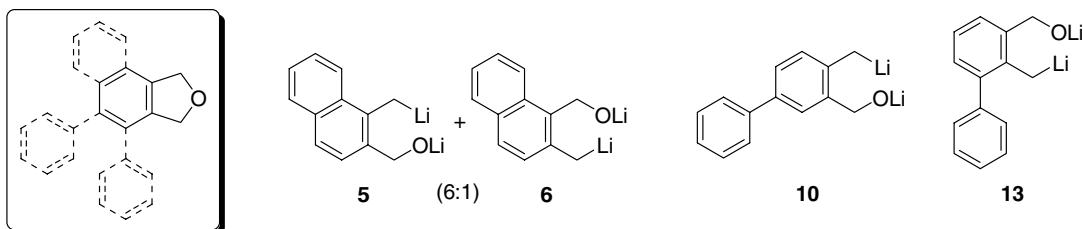


InBr₃ has been demonstrated to be a simple catalyst for the intermolecular hydroamination of unactivated alkenes to produce tosyl- and mesyl-protected amines in moderate to good yields.

Regiochemistry in the reductive opening of phthalan derivatives

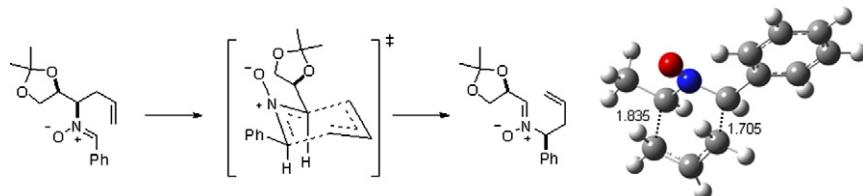
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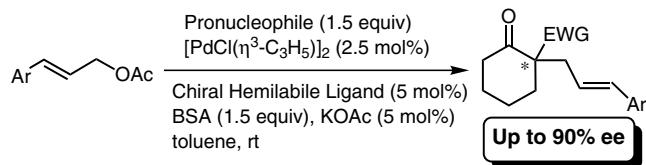
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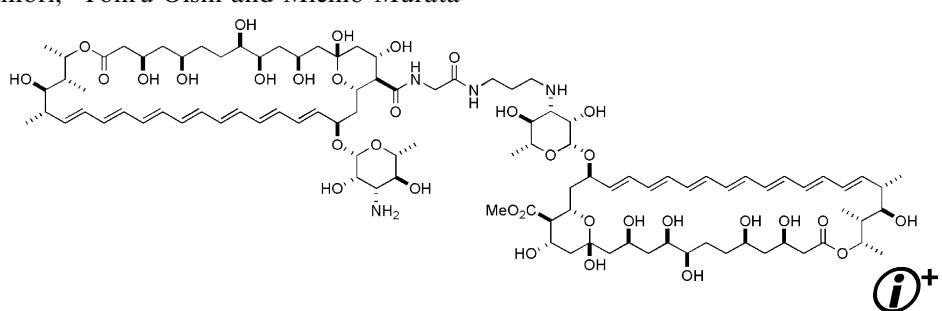
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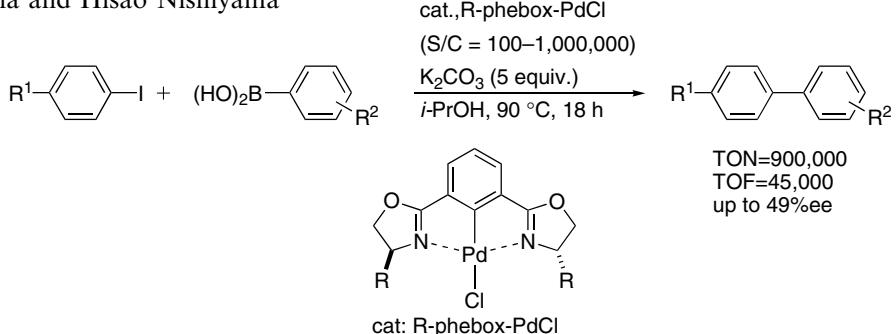
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Yuichi Umegawa, Nobuaki Matsumori,* Tohru Oishi and Michio Murata*

Based on an amphotericin B ion-channel model with close intermolecular interaction between carboxyl and amino groups, covalent dimers of AmB with potent biological activities were prepared.



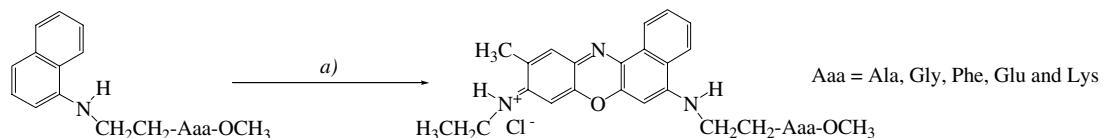
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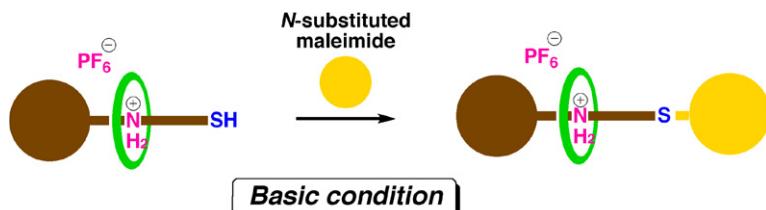
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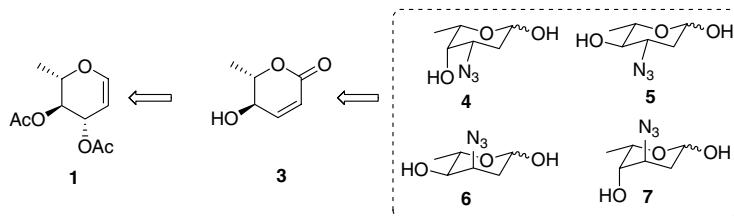
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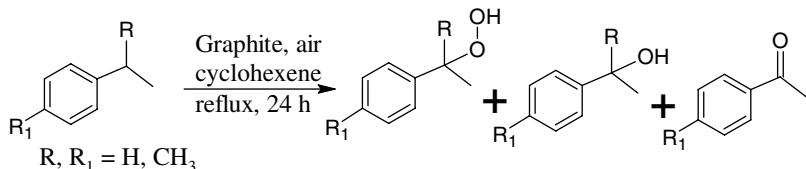
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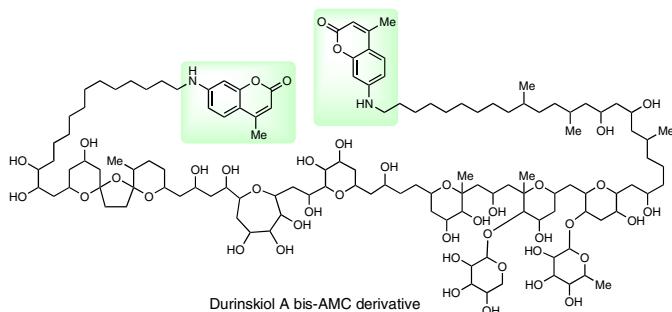
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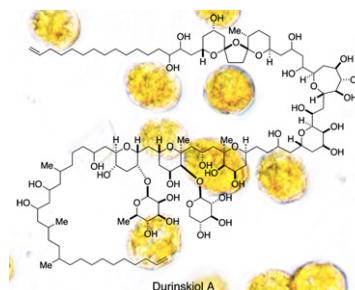
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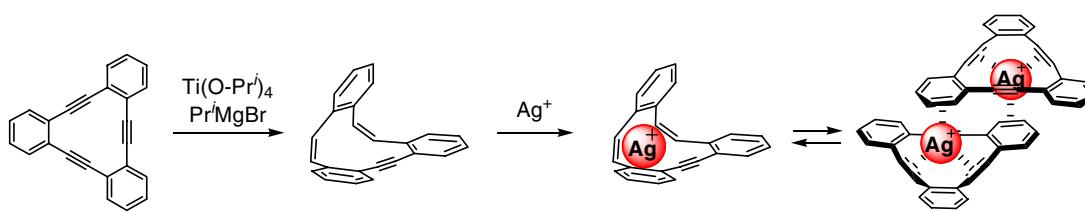
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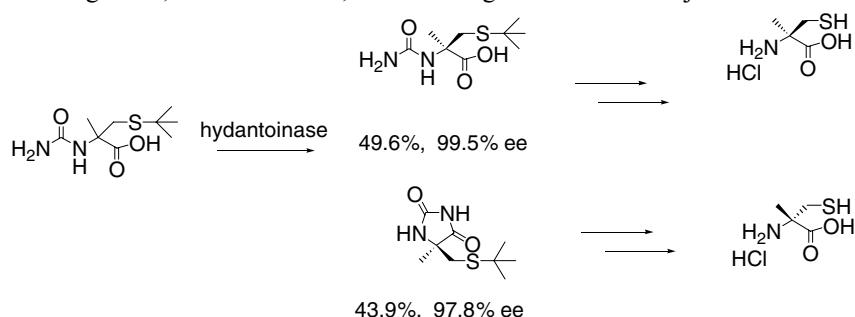
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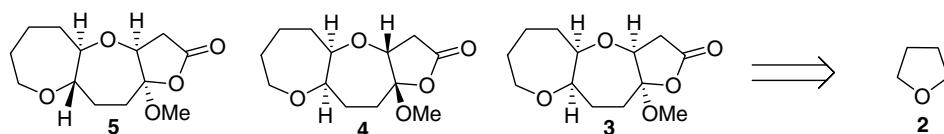
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**The furan approach to oxacycles. Part 6: From THF to fused polyoxepanes**

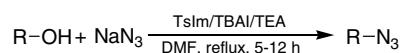
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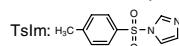
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Mohammad Navid Soltani Rad,* Somayeh Behrouz and Ali Khalafi-Nezhad

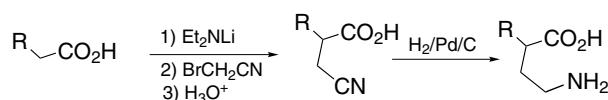


R = 1°, 2° and 3° alkyl

**A simple synthesis of γ -aminoacids**

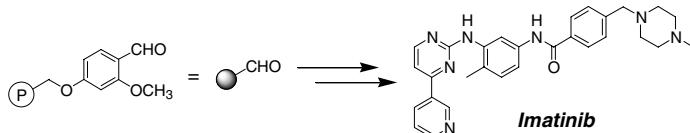
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Salvador Gil, Margarita Parra* and Pablo Rodríguez

We describe a new synthesis in two-steps of γ -aminoacids, which improves previous results.

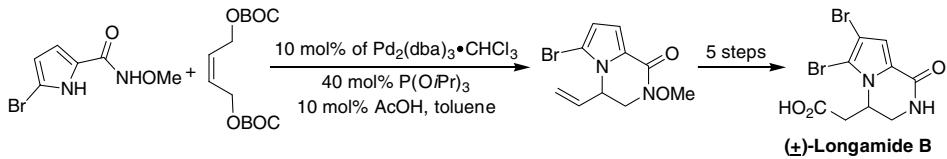
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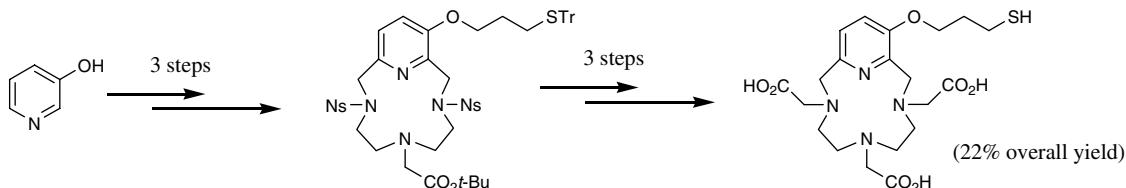
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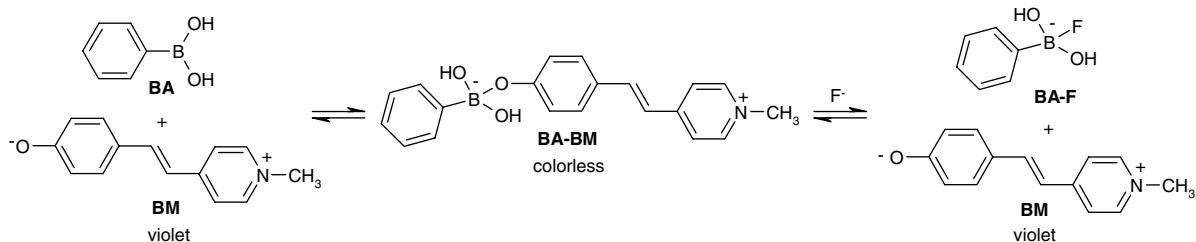
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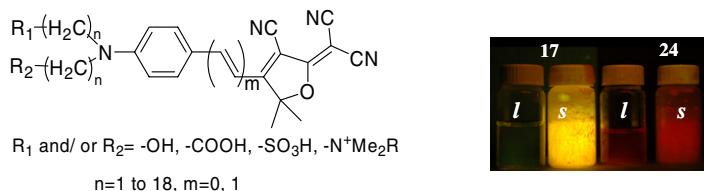
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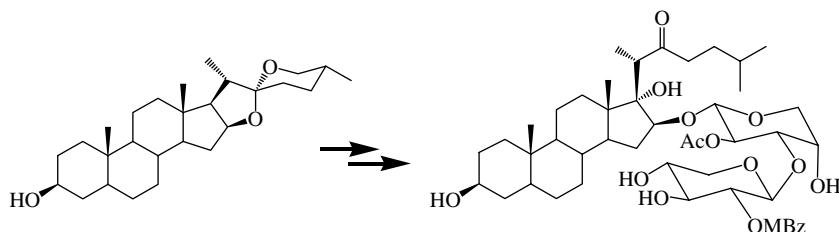
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Li-Jun Chen, Qi-Hai Xu, Hao Huang, Jing-Rong Lin and Wei-Sheng Tian*

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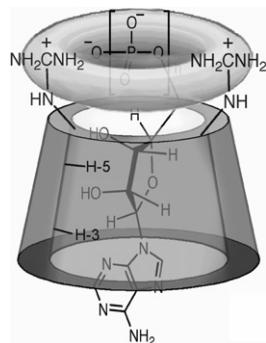


Heptakis(6-deoxy-6-guanidino)- β -cyclodextrin: an artificial model for mitochondrial ADP/ATP carrier

De-Qi Yuan,* Ayako Izuka, Makoto Fukudome, Mikhail V. Rekharsky, Yoshihisa Inoue and Kahee Fujita*

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Heptakis(6-deoxy-6-guanidino)- β -cyclodextrin, prepared by one-step reaction of heptakis(6-amino-6-deoxy)- β -cyclodextrin with 1*H*-pyrazolecarboxamidine, binds ADP/ATP very tightly.



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A long carbon-chain polyol compound with a molecular weight of 2128 mu, durinskiol A, was isolated from the cultured symbiotic dinoflagellate *Durinskia* sp. Its planar and partial relative stereostructure was elucidated based on 2D-NMR and MS/MS analysis. Durinskiol A possesses the following unique features: a 6,5,6-bis-spiroacetal ring, two terminal olefins, two sugar units, and six- and seven-membered ether rings. It caused a short body length, abnormal pigment pattern, and pericardiac and yolk-sac edema in zebrafish.

Tetrahedron Letters 2007, 48, 3423–3427 and 3429–3432.

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