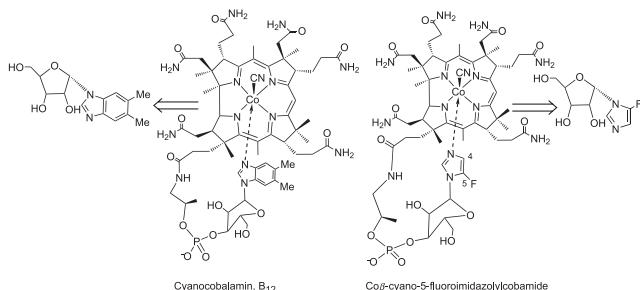


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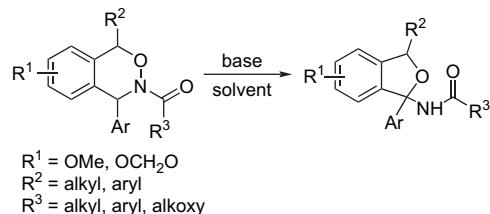


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Novel rearrangement of 1*H*-2,3-benzoxazines to cyclic N-acyl hemiaminals: application to the synthesis of 1-arylnaphthalene skeletal congeners pp. 39–44

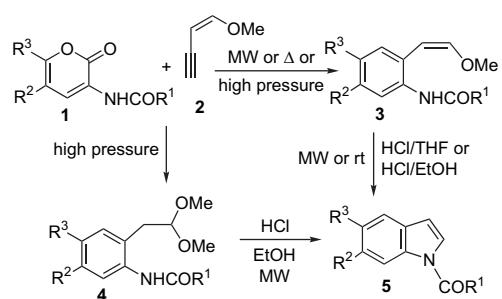
Xiu Fang Zheng, Xiao Lei Wang, Jun Biao Chang*, Kang Zhao*

The rearrangement of 1*H*-2,3-benzoxazine derivatives has been investigated. The reaction affords cyclic hemiaminal derivatives for their conversion to the corresponding 1-arylisobenzofurans, which can be trapped by various dienophiles to afford skeletal congeners of 1-arylnaphthalene lignans.



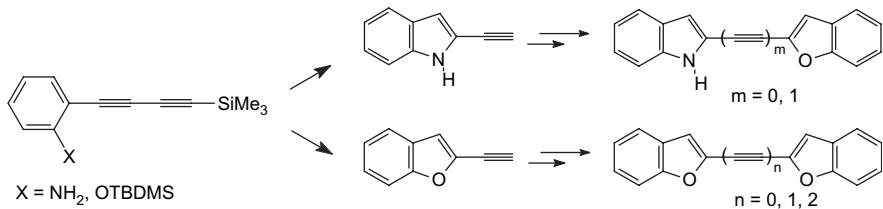
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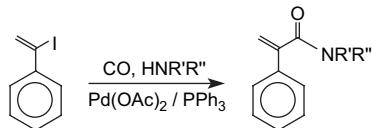
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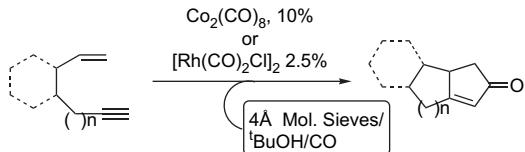
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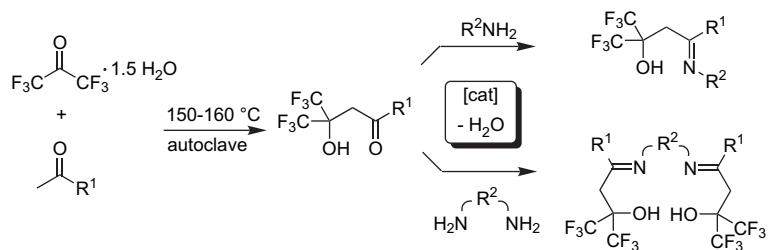
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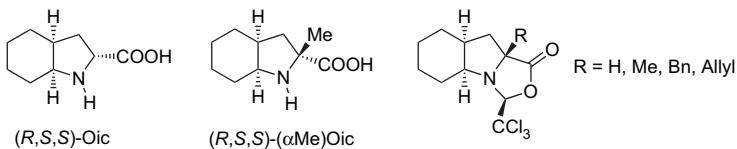
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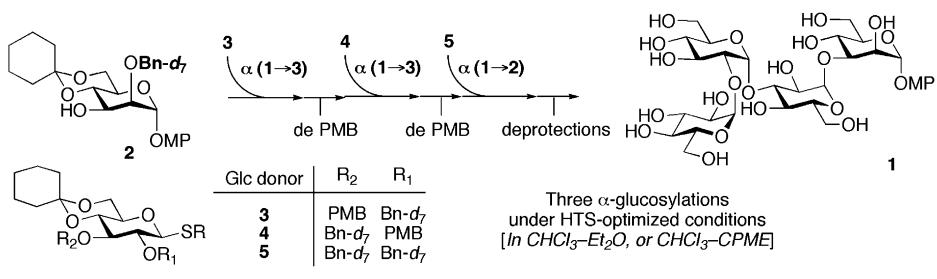
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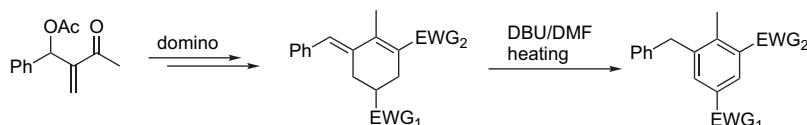
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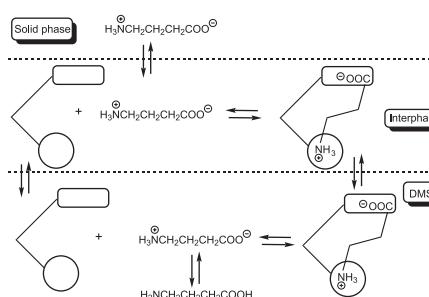


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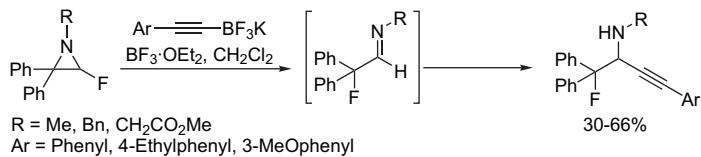
Ditopic receptors as efficient solid–liquid extractants for ω -amino acids are studied. In DMSO, both free non-zwitterionic amino acid along with its zwitterionic complexed form are present. Thus, the ratio between amino acid extracted and ligand can be higher than 1, being the highest ratio for 4-aminobutanoic acid.



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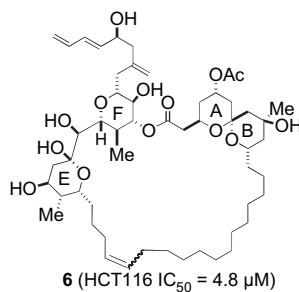
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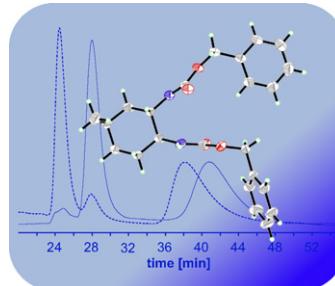
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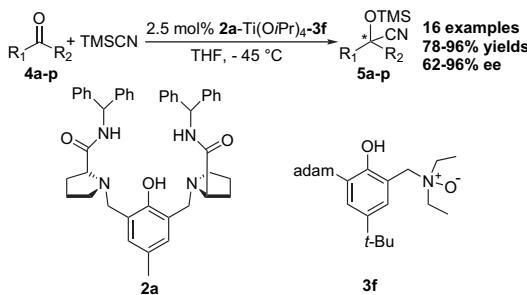
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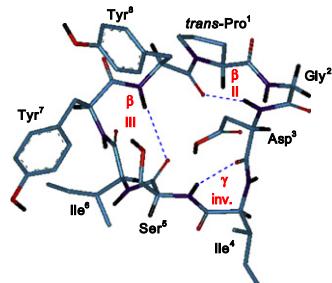
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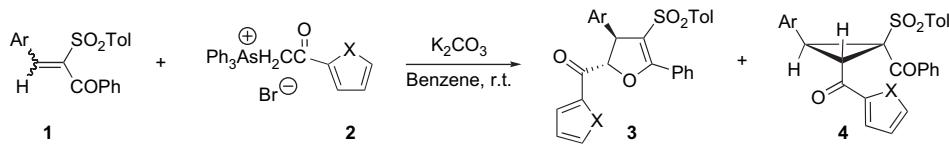
Alassane Wélé, Claudine Mayer, Quentin Dermigny, Yanjun Zhang, Alain Blond, Bernard Bodo*



Cycloreticulin A: *cyclo*(Pro¹-Gly²-Asp³-Ile⁴-Ser⁵-Ile⁶-Tyr⁷-Tyr⁸). Cycloreticulin B: *cyclo*(Pro¹-Mso²-Tyr³-Gly⁴-Thr⁵-Val⁶-Ala⁷-Val⁸).

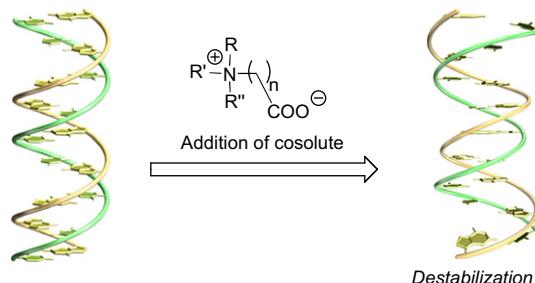
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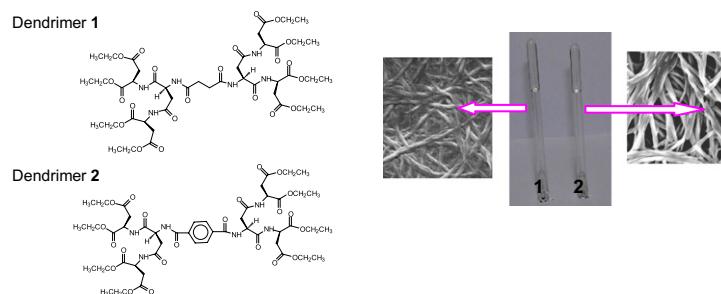
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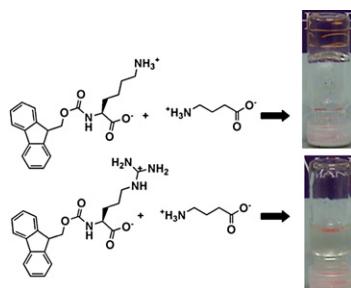
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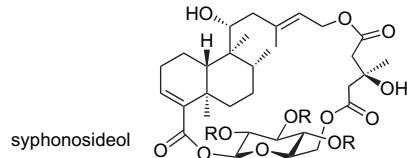
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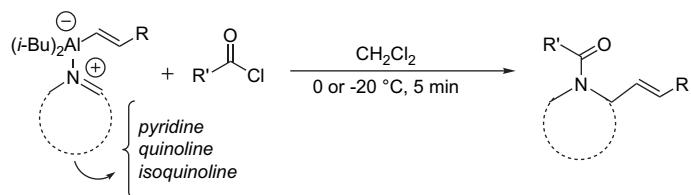
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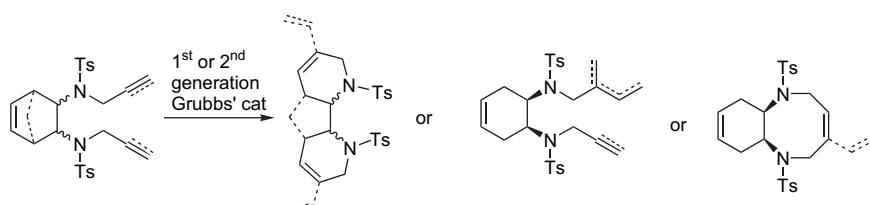
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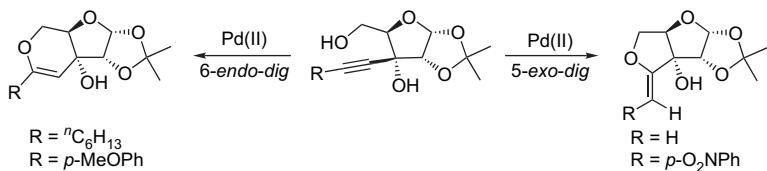
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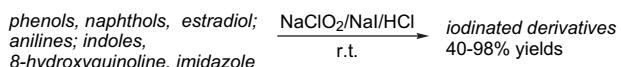
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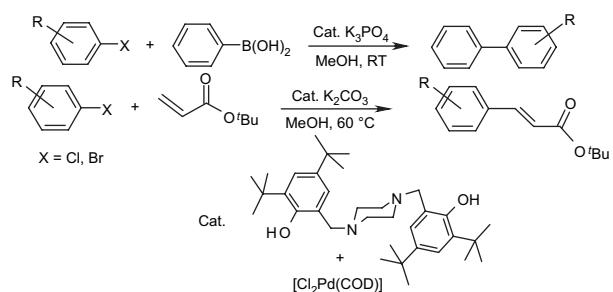
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An inexpensive and highly stable ligand 1,4-bis(2-hydroxy-3,5-di-*tert*-butylbenzyl)piperazine for Mizoroki–Heck and room temperature Suzuki–Miyaura cross-coupling reactions

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Sasmita Mohanty, D. Suresh, Maravanji S. Balakrishna*, Joel. T. Mague

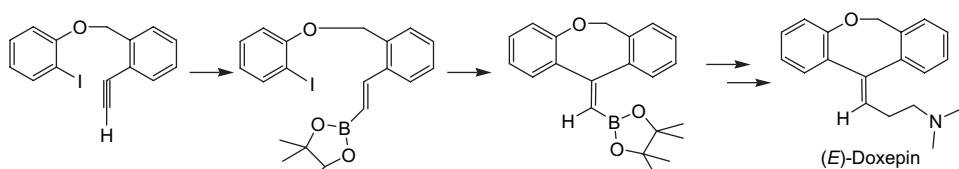


The bulky ligand 1,4-bis(2-hydroxy-3,5-di-*tert*-butylbenzyl)piperazine with $[\text{Cl}_2\text{Pd}(\text{COD})]$ is an efficient catalyst for Suzuki and Heck coupling reactions.

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Cuihua Xue, Shi-Hao Kung, Jian-Zhong Wu, Fen-Tair Luo*



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 [†] Supplementary data available via ScienceDirect



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