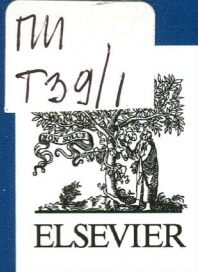


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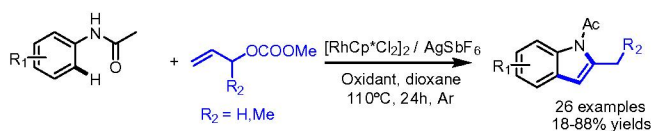
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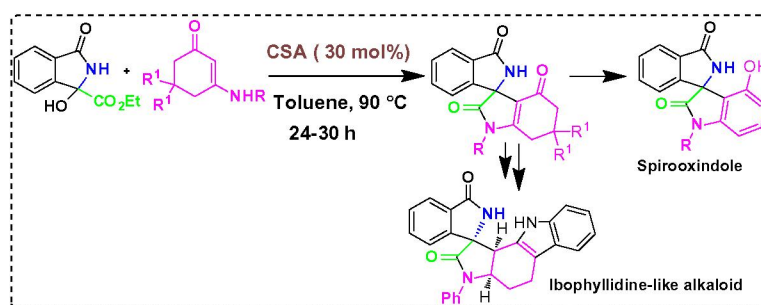
A practical Rh-catalyzed reaction was developed to achieve 2-alkyl-substituted indole synthesis. The reaction can tolerate a variety of synthetically important functional groups (*aryl-Br*, *heterocycle*, *amino acid*). The indole products can also be transformed into other important skeletons. Two bioactive compounds, that is, indomethacin and pravadoline were prepared using the new method.



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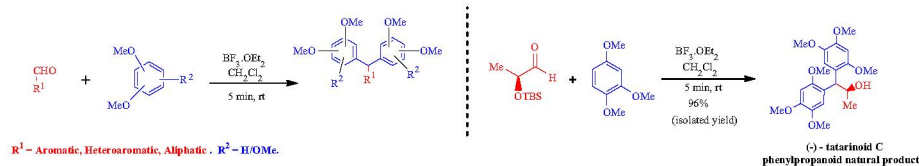
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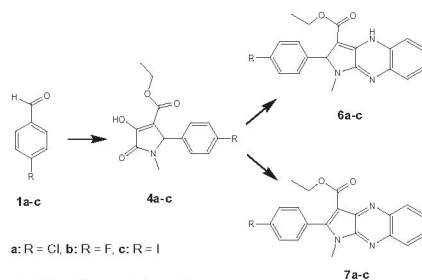
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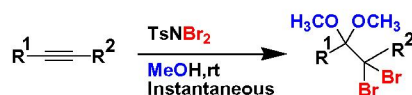
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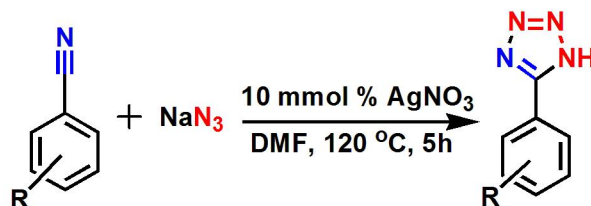
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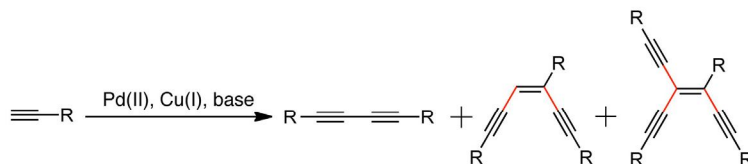
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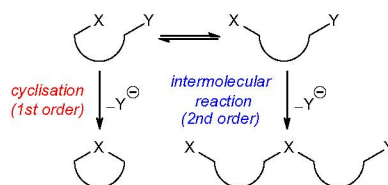
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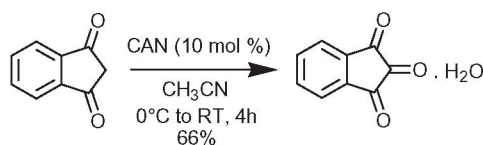
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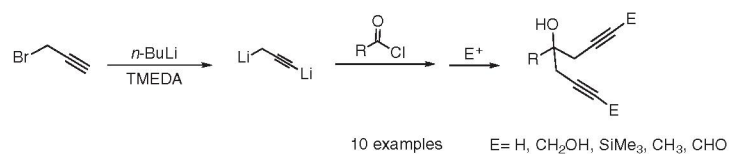
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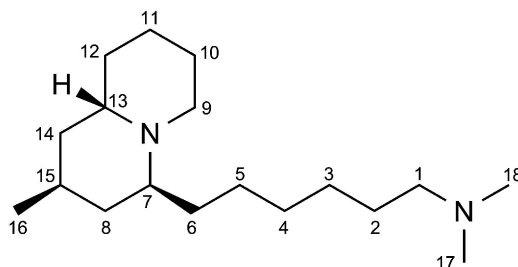
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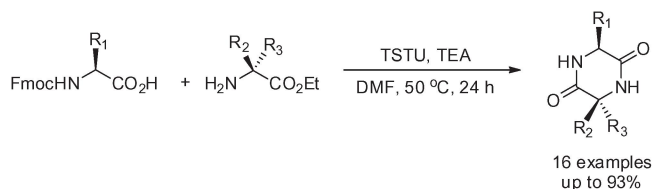
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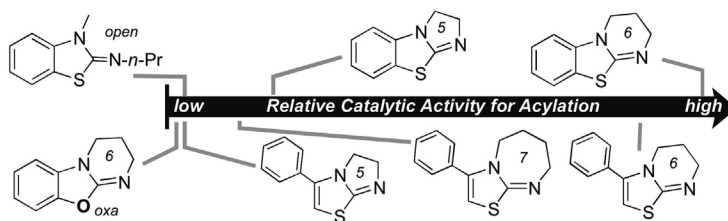
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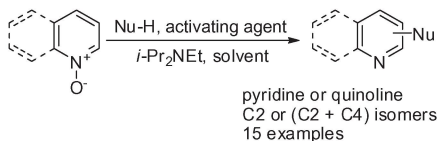
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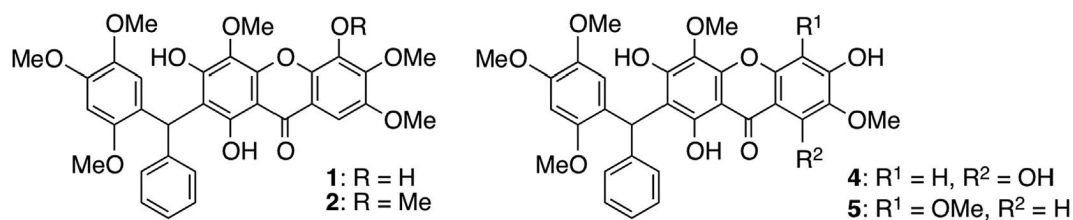
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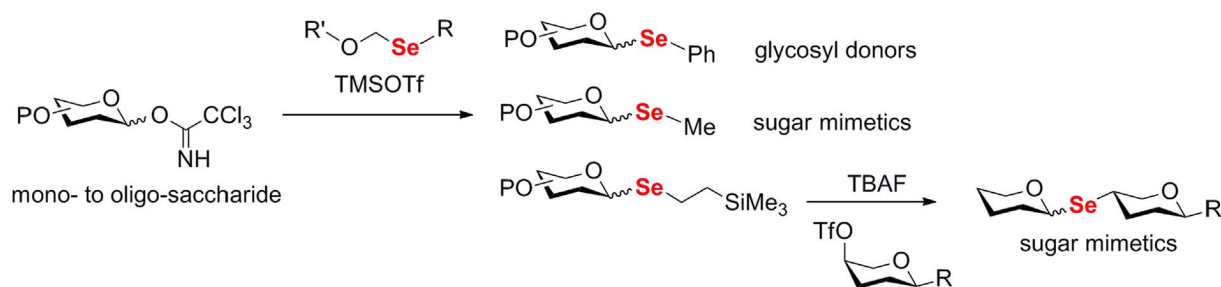
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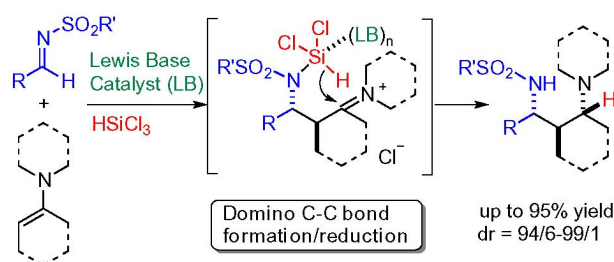
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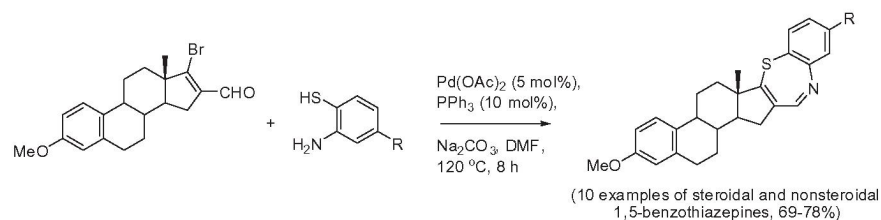
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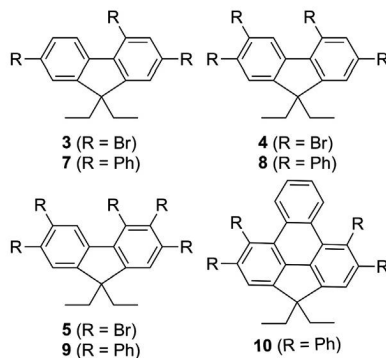
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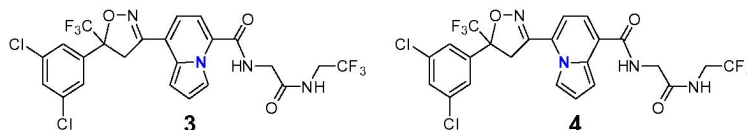
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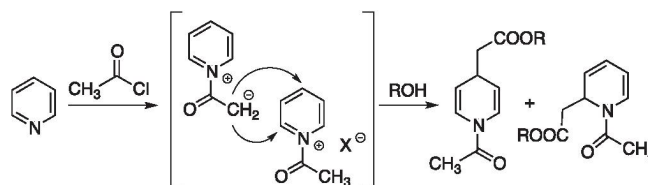
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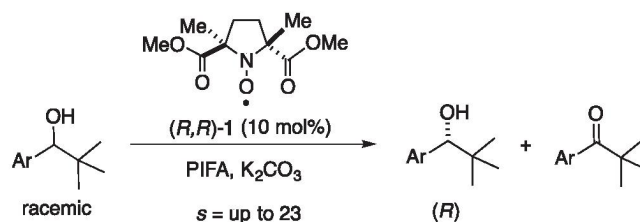
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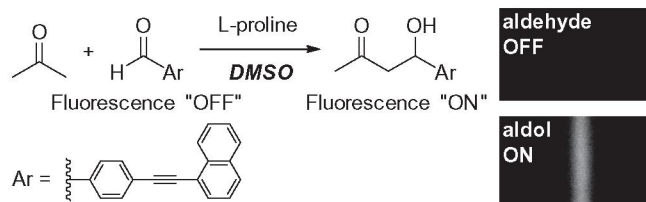
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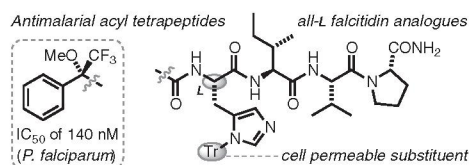
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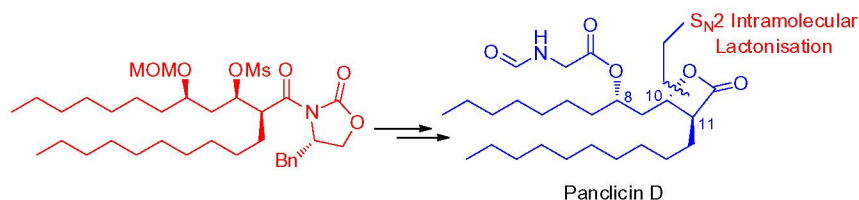
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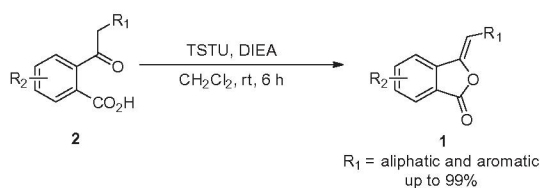
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CORRIGENDA


Corrigendum to “An effective and regioselective bromination of 1,4,5,8-naphthalenetetracarboxylic dianhydride using tribromoisocyanuric acid” [Tetrahedron Lett. 54 (2013) 6314–6318] p 1959

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Corrigendum to “Synthesis of imidazo[1,2-*a*]pyridine in the presence of iodine–water catalytic system” p 1960
[Tetrahedron Lett. 55 (2014) 1159–1163]

I. R. Siddiqui*, Pragati Rai, Rahila, Anushree Srivastava, Shayna Shamim

*Corresponding author

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