

The Chinese University of Hong Kong Department of Chemistry Research Seminar Series

- **Speaker:** Prof. Jishan Wu Department of Chemistry National University of Singapore & Institute of Materials Research and Engineering, A*STAR
- **Title:** Open-shell Polycyclic Hydrocarbons

<< Abstract >>

I will discuss three types of polycyclic hydrocarbons with an open-shell ground states: (a) zig-zag edged nanographenes such as periacenes;^[1] (b) Z-shaped quinoidal hydrocarbons such as zethrenes,^[2] and (c) extended *p*-quinodimethanes.^[3] The discussion will include their challenging synthesis, their characterizations of the ground-state structures and their unique optical, electronic and magnetic properties.^[4]



References

[1] (a) Yao, J. et al., Chem. Eur. J. 2009, 15, 9299; (b) Li, J. et al., Chem. Sci. 2012, 3, 846.

[2] (a) Sun, Z. et al., J. Am. Chem. Soc. 2011, 133, 11896. (b) Li, Y. et al., J. Am. Chem. Soc. 2012, 134, 14913.

(c) Z. Sun et al., J. Am. Chem. Soc. 2013, 135, 18299. (d) Zeng, W. et al., Chem. Eur. J. 2013, 19, 16841.

[3] (a) Zeng, Z. et al., J. Am. Chem. Soc. 2012, 134, 14513. (b) Zeng, Z. et al., J. Am. Chem. Soc. 2013, 135, 6363. (c) Zeng, Z. et al., Angew. Chem. Int. Ed. 2013, 52, 8561.

[4] See our recent review articles: (a) Sun, Z. et al., Chem. Soc. Rev. 2012, 41, 7857. (b) Sun, Z. et al., Chem. Asian J. 2013, 8, 2894.

Date: January 9, 2014 (Thursday)

Time: 4:00 p.m.

Venue: LG23, Science Centre

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< Abstract >

Carbon – carbon (C - C) bonds constitute the major framework of organic molecules and carbon – hydrogen (C - H)bonds are abundant in their peripheries. Such nonpolar –bonds are thermodynamically stable and kinetically inert in general. Selective activation of those ubiquitous bonds offers a straightforward method to construct and functionalize organic skeletons. Herein is presented ring expansion from orthocyclophanes to metacyclophanes by means of sequential actions of light and a metal catalyst. Formally, non-strained C–H and C–C bonds are cleaved and exchanged without any atoms or groups eliminated. Of note is that the entire transformation is energetically uphill. The involvement of photocyclization renders it possible to drive the thermodynamically disfavored transformation forward. Furthermore, the present work has established stereospecificity in the transfer of central chirality on a tertiary carbon to planar chirality of the resulting substituted metacyclophane.





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The Chinese University of Hong Kong Department of Chemistry

Research Seminar Series

- **Speaker:** Prof. William R. Roush The Scripps Research Institute Jupiter, Florida U.S.A.
- Title:Synthesis of Chiral Allylboron and Enolborane
Reagents via Allene and Acrylamide
Hydroboration Reactions and Applications to
the Synthesis of Natural Products
- **Date:** January 29, 2014 (Wednesday)
- **Time:** 2:30 p.m.

Venue: Room 703 Mong Man Wai Building



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Contact Person: Prof. Henry N.C. Wong