

GTR ORGANIC SEMINAR

The Versatile Chemistry of N-Heterocycle- stabilized Iodanes – From Synthesis to Halogen Bond Catalysis



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4:30 pm ~ 6:00 pm
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The Venture Hall, 3F, VBL



Host: Kazuaki Ishihara
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The Versatile Chemistry of *N*-Heterocycle-stabilized Iodanes – From Synthesis to Halogen Bond Catalysis

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The covalent attachment of donor-substituents is an essential feature in iodane chemistry. Intramolecular donor-iodine interactions stabilize the hypervalent iodine atom by forming (pseudo)cyclic structures that have increased stability and solubility compared to their unsubstituted derivatives. Most of the well-established donor-substituted iodanes are based on tethered oxygen ligands, particularly carboxylic acids, amides and ethers, to form benziodoxol(on)es.^[1] Despite their great success as versatile ligands in transition metal chemistry, *N*-heterocycles have been treated nevercal.

We systematically investigate the chemistry of *N*-heterocycle-stabilized iodanes (NHIs) and will share our latest results regarding their synthesis, stability and structural aspects.^[2] Based on these findings, the great potential of *N*-heterocycle-substituted iodoarenes (NHAIs) in enantioselective oxidation catalysis and NHI-based cyclic iodonium salts in XB-catalysis will be discussed intensively.^[3,4] Finally, the first experimental structure elucidation of an *in situ* generated chiral iodane based on vibrational circular dichroism (VCD) and DFT will be presented.^[5]



References

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- [5] T. P. Golub, A. H. Abazid, B. J. Nachtsheim, C. Merten, *Angew. Chem.* **2022**, DOI: <https://doi.org/10.1002/anie.202204624>.