# **Division of Synthetic Chemistry** – Structural Organic Chemistry –

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## **Scope of Research**

Fundamental studies are being conducted for the creation of new functional  $\pi$ -systems with novel structures and properties, and for evaluation of their application as organic semi-conducting materials for photovoltaic and electroluminescent

devices. The major subjects are: 1) organochemical transformation of fullerenes  $C_{60}$  and  $C_{70}$ , specifically organic synthesis of endohedral fullerenes by the technique of molecular surgery; 2) generation of ionic fullerene species and their application for the synthesis of functional material; 3) synthesis of new carbon-rich materials by the use of transition metal complex; and 4) creation of new functional  $\pi$ -materials with unique photoelectric properties.

#### **KEYWORDS**

π-Conjugated Systems Functional Materials Organic Solar Cells Endohedral Fullerenes Perovskite-Based Solar Cells



## **Selected Publications**

Futagoishi, T.; Murata, M.; Wakamiya, A.; Murata, Y., Trapping N<sub>2</sub> and CO<sub>2</sub> on the Sub-Nano Scale in the Confined Internal Spaces of Open-Cage  $C_{60}$  Derivatives: Isolation and Structural Characterization of the Host–Guest Complexes, *Angew. Chem. Int. Ed.*, **54**, 14791-14794 (2015). Nishimura, H.; Eliseeva, M. N.; Wakamiya, A.; Scott, L. T., 1,3,5,7-Tetra(Bpin)azulene by Exhaustive Direct Borylation of Azulene and 5,7-Di(Bpin)azulene by Selective Subsequent Deborylation, *Synlett*, **26**, 1578-1580 (2015).

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Wang, S.; Yang, D.-T.; Lu, J.; Shimogawa, H.; Gong, S.; Wang, X.; Mellerup, S. K.; Wakamiya, A.; Chang, Y.-L.; Yang, C.; Lu, Z.-H., In Situ Solid-State Generation of (BN)<sub>2</sub>-Pyrenes and Electroluminescent Devices, *Angew. Chem. Int. Ed.*, **54**, 15074-15078 (2015).

Chaolumen; Murata, M.; Sugano, Y.; Wakamiya, A.; Murata, Y., Electron Deficient Tetrabenzo-Fused Pyracylene and Conversions into Curved and Planar π-Systems with Distinct Emission Behaviors, *Angew. Chem. Int. Ed.*, **54**, 9308-9312 (2015).

## Trapping $N_2$ and $CO_2$ on the Sub-nano Scale in the Confined Internal Spaces of Open-cage $C_{60}$ Derivatives: Isolation and Structural Characterization of the Host–Guest Complexes

We have previously reported the synthesis of open-cage  $C_{60}$  **1** with a 17-membered-ring opening and one sulfur atom on the rim. It was found that **1** was able to encapsulate molecular nitrogen and carbon dioxide after its exposure to high pressures of N<sub>2</sub> and CO<sub>2</sub> gas. A subsequent reduction of one of the four carbonyl groups on the rim of the opening induced a contraction of the opening ( $\rightarrow$ **2**) and trapped the guest molecules inside. The molecular structures of N<sub>2</sub>@**2** and CO<sub>2</sub>@**2** were determined by single-crystal X-ray diffraction analyses, which revealed a short N $\equiv$ N triple bond for the encapsulated N<sub>2</sub> and an non-symmetric molecular structure for the encapsulated molecule of CO<sub>2</sub>.



Figure 1. The structures of open-cage  $C_{60}$  derivative 1 and host–guest complexes  $N_2@2$  and  $CO_2@2$ .

## Hole-transporting Materials with a Two-dimensionally Expanded $\pi$ -System around an Azulene Core for Efficient Perovskite Solar Cells

Two-dimensionally expanded  $\pi$ -systems, consisting of partially oxygen-bridged triarylamine skeletons that are connected to an azulene (1–3) or biphenyl core (4), were



**Figure 2.** The structures of 1-4 with two-dimensionally expanded  $\pi$ -systems.

synthesized. When tetra-substituted azulene **1** was used as a hole-transporting material (HTM) in perovskite solar cells, the observed performance (power conversion efficiency = 16.5%) was found to be superior to that of the current HTM standard Spiro-OMeTAD. Based on a comparison of the optoelectronic and electrochemical properties of **1**–**4** and Spiro-OMeTAD, we were able to elucidate the factors that are required for HTMs to act efficiently in perovskite solar cells.

## Electron Deficient Tetrabenzo-fused Pyracylene and Conversions into Curved and Planar $\pi$ -Systems with Distinct Emission Behaviors

Polycyclic aromatic compounds containing fully unsaturated five-membered ring(s) have been intensively studied because of their unique properties, which include high electron affinity and reactivity. We demonstrated an efficient route for the synthesis of electron-deficient tetrabenzo-fused pyracylene 1 by using intramolecular oxidative C-H coupling. Compound 1 was shown to possess high electron affinity and to undergo addition reactions with *n*-butyllithium or benzyne. These reactions led to either a 1,4-addition compound 2 or triptycene-type adduct 3 with a curved or planar  $\pi$ -system, respectively. These compounds showed distinct emissions in the solid state with high quantum yields. X-ray diffraction analyses demonstrated that the 2 forms dimers with effective  $\pi$ -overlap, which can cause a significant red-shift in the fluorescence by the stabilization of excimer.



Figure 3. Conversions of electron deficient TBP 1 into curved and planar  $\pi$ -systems 2 and 3 having distinct emission behaviors.